The bioinorganic chemistry of copper-containing systems: From Type-3 systems pertinent to Alzheimer's disease to mononuclear hydrolysis involved in biological development

Giordano Faustini Zimmerer Da Silva

University of South Florida

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The Bioinorganic Chemistry Of Copper-Containing Systems: From Type-3 Systems
Pertinent To Alzheimer’s Disease To Mononuclear Hydrolysis Involved In Biological Development

by

Giordano Faustini Zimmerer Da Silva

A dissertation submitted in partial fulfillment of the requirements for the degree of Doctor of Philosophy Department of Chemistry College of Arts and Sciences University of South Florida

Co-Major Professor: Li-June Ming, Ph.D.
Co-Major Professor: Brian T. Livingston, Ph.D.
Steven H. Grossman, Ph.D.
Randy L. Larsen, Ph.D.

Date of Approval: May 9, 2007

Keywords: kinetics, reactive oxygen species, metallopeptide, amyloid, enzymology, sea urchin, metalloprotease

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NOTE TO THE READER

Note to Reader: The original of this document contains color that is necessary for understanding the data. The original dissertation is on file at the USF library in Tampa, FL.
DEDICATION

Although the graduate school experience is highly a personal journey, it would be foolish to believe that I was solely responsible for my accomplishments. I then would like to dedicate this work to all of those who have played a major role in this difficult yet rewarding part of my life. First my grandmother Ermelinda who is the sole reason why I am in the US, my parents Mauricio and Marta for sacrificing the presence of their son, and my sister Marcella for being sibling, friend and parent through part of my formative years. Second I must dedicate this work to the Kelley family for being my parents, brother, and sisters; Moms, Pops, John, Tere, and Abbie, I would not be here today without your love and support. Last and not least I must also dedicate this work to Rachel for being a friend, always in my heart, through difficult and joyous times, for her love and support without which I would have never remained in the US. I truly hope that the rest of my career will make the efforts of all of those herein mentioned worthwhile.
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<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Description</th>
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<tbody>
<tr>
<td>AD</td>
<td>Alzheimer’s disease</td>
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<tr>
<td>Aβ</td>
<td>Amyloid-β</td>
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<tr>
<td>CuO</td>
<td>Copper-oxygen intermediate</td>
</tr>
<tr>
<td>Hey</td>
<td>Hemocyanin</td>
</tr>
<tr>
<td>Tyr</td>
<td>Tyrosinase</td>
</tr>
<tr>
<td>ROS</td>
<td>Reactive Oxygen Species</td>
</tr>
<tr>
<td>kDa</td>
<td>kilo Daltons</td>
</tr>
<tr>
<td>APP</td>
<td>Amyloid Precursor Protein</td>
</tr>
<tr>
<td>ADP</td>
<td>Adenosine 5’ diphosphate</td>
</tr>
<tr>
<td>ATP</td>
<td>Adenosine 5’ triphosphate</td>
</tr>
<tr>
<td>ET</td>
<td>Electron Transfer</td>
</tr>
<tr>
<td>DTT</td>
<td>Dithiothreitol</td>
</tr>
<tr>
<td>EDTA</td>
<td>Ethylenediaminetetraacetic acid</td>
</tr>
<tr>
<td>NADP⁺</td>
<td>Nicotinamide adenine dinucleotide phosphate, oxidized form</td>
</tr>
<tr>
<td>NADPH</td>
<td>Nicotinamide adenine dinucleotide phosphate, reduced form</td>
</tr>
<tr>
<td>NAD⁺</td>
<td>Nicotinamide adenine dinucleotide, oxidized form</td>
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</tbody>
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NADH: Nicotinamide adenine dinucleotide, reduced form
PAGE: Polyacrylamide gel electrophoresis
SDS: Sodium dodecyl sulfate
Tris: Tris(hydroxymethyl)aminomethane
DNA: Deoxyribonucleic acid
mRNA: Messenger ribonucleic acid
cDNA: Complementary DNA
COX: Cytochrome oxidase activity
CSF: Cerebral spinal fluid
mtDNA: Mitochondrial DNA
dsDNA: Double stranded deoxyribonucleic acid
SOD: Superoxide dismutase
MnSOD: Manganese dependent superoxide dismutase
CuZnSOD: Copper-Zinc dependent superoxide dismutase
PHF: Paired helical fragments
CNS: Central nervous system
$^1$H NMR: Proton nuclear magnetic resonance
CW-EPR: Continuous wave electron paramagnetic resonance
ESEEM: Electron spin echo envelope modulation
RAGE: Receptor for advanced glycation endproducts
ERAB: Endoplasmic reticulum amyloid- β binding protein
PBN: $N$-tert-butyl-a-phenylnitrone
PBS: Phosphate buffered saline
MNP: 2-methyl-2-nitrosopropane
PrP: Prion protein
DOPA: 3,4-dihydroxyphenylalanine
XAS: X-ray absorption spectroscopy
EXAFS: Extended X-ray absorption fine structure
DβM: Dopamine β hydroxylating monooxygenase
PHM: Peptidylglycine α-hydroxylating monooxygenase
PPh3: Triphenylphosphine
THB: 1,2,3-trihydroxybenzene
DMSO: Dimethyl sulfoxide
HEPES: N-(2-hydroxyethyl)-piperazine-N'-2-ethanesulfonic acid
MES: 2-Morpholinoethanesulfonic acid
TAPS: N-Tris(hydroxymethyl)methyl-3-aminopropanesulfonic acid
CAPS: N-Cyclohexyl-3-aminopropanesulfonic acid
MBTH: 3-methyl-2-benzothiazolinone hydrazone
d6DMSO: Deuterated dimethyl sulfoxide
UV-Vis: Ultraviolet-visible electronic spectroscopy
LMCT: Ligand to metal charge transfer transition
MALDI-TOF: Matrix assisted laser desorption mass spectrometry time of flight
KIE: Kinetic isotope effect
DTC: Di-tertbutyl catechol

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DCC: Dichlorocatechol
CMC: Critical micelle concentration
GSH: Reduced glutathione
NQI: Nuclear quadrupole interactions
BP10: Blastula Protease 10
CuBP10: Copper derivative of Blastula protease 10
ZnBP10: Zinc derivative of Blastula protease 10
BMP-1: Bone morphogenetic protein 1
EGF: Epidermal growth factor
ECM: Extracellular matrix
suBMP: Sea urchin bone morphogenetic protein 1
TGF-β: Transformation growth factor β
CUB: Complement-like domains
Ni-NTA: Nickel nitrilotriacetic acid
BAPNA: N-benzoyl-arginine-p-nitroanilide
IPTG: Isopropyl-β-thiogalactopyranoside
PMSF: Phenylmethyl sulfonyl fluoride
PCR: Polymerase chain reaction
OD: Optical density
CD: Circular dichroism
BCA: Bicinchoninic acid
APS: Ammonium persulfate
TEMED: N,N,N',N'-Tetramethylethylenediamine

OP: 1,10-phenanthroline

Arg-NHOH: Arginine hydroxamate

MM3: Molecular mechanics protocol
The Bioinorganic Chemistry of Copper-Containing Systems: from Type-3 Systems Pertinent to Alzheimer’s Disease to Mononuclear Hydrolysis Involved in Biological Development

Giordano F.Z. da Silva

ABSTRACT

Although transition metals are essential for life, misregulation of redox-active metal uptake, delivery, storage, and excretion has been linked with a series of neurodegenerative disorders. Alzheimer’s disease (AD) is considered an epidemic and is the most widespread of all forms of dementia. Copper ions found in large concentrations localized in amyloid-β plaques in the brain of AD patients have been linked with the generation of reactive oxygen species which are suspected to be the culprits leading to neuronal cell death. Herein a series of mechanistic and spectroscopic studies elucidate the chemistry about the metal-centered oxidation of biomolecules, including catecholamine neurotransmitters and some analogues by copper-complexes of amyloid-β peptide.

Transition metals can also be useful tools for characterization of metalloproteins due to their unique chemical and spectroscopic features. Herein a series of studies of the native Zn$^{2+}$ and Cu$^{2+}$-derivative of recombinant Blastula Protease 10 (BP10) from the sea urchin Paracentrotus lividus are presented in order to elucidate its catalytic mechanism, with the use of enzymology, metal substitution, and electronic absorption spectroscopy.
CHAPTER I. COPPER-DIOXYGEN IN METABOLISM AND CATALYSIS

I. BACKGROUND

The interaction of transition metals with oxygen (O₂, or dioxygen) is of paramount importance in biological systems. The metal-O₂ interaction is ubiquitous in metabolic pathways of aerobic organisms on the earth’s crust and it is fundamental to homeostasis by serving as catalysts in redox reactions as well as for the binding and delivery of O₂ in aerobes.¹ For these purposes, iron and copper are the predominant, albeit not the only, transition metals utilized in metalloproteins and metalloenzymes. Classic examples of iron-containing O₂ transport metalloproteins include hemoglobin,² myoglobin,² and hemerythrin,³ while redox-active non-heme iron metalloenzymes involved in O₂ metabolism include methane monooxygenase,⁴ and protocatechuate 3,4 dioxygenase.⁵ Copper is also well represented in the area of O₂-binding with hemocyanin⁶ being the classic example, while catalytic copper-centers include tyrosinase,⁶ catechol oxidase,⁷ and galactose oxidase.⁸
Of particular interest in the recent years has been the nature of copper-O$_2$ centered chemistry in metalloenzymes. Several research groups have pursued spectroscopic characterization, crystallography, and biomimetic studies of metalloproteins that contain copper and can interact with O$_2$ through direct binding or for activation of O$_2$ in chemical reactions like hydroxylation and oxygenation. These copper metalloenzymes are collectively termed Type-3 copper proteins attributed to the magnetically coupled di-copper center.$^6$ Although the topic in this review is primarily from a health/disease or a more medicinal perspective, the lessons learned from the investigation of the chemistry of Type-3 copper proteins have some overlap with the area of neurodegeneration involving activated or reduced forms of O$_2$. The focus of this chapter will be the parallels of what is known about the chemistry of Type-3 copper oxidases and how it may relate to Alzheimer’s disease.

The ability of binuclear copper centers in proteins to bind O$_2$ for transport and to use copper-O$_2$ (CuO$_2$) as the catalytic center for O$_2$ activation has provided a wealth of understanding in both experimental and theoretical chemistry. Understanding the interaction of O$_2$ with copper has been an intense area of study for bioinorganic chemists, both toward the native molecules and in biomimetic systems.$^9,10$ Research in this area has utilized the spectroscopic properties of the Type-3 copper proteins, as well as crystal structures when available, to serve as templates toward the synthesis of small model systems that allow the interactions of copper with O$_2$. Typically, the short-lived CuO$_2$ intermediates cannot be well resolved in the native systems either due to influence of the size of proteins in spectroscopic techniques that rely in magnetic resonance or because
crystal structures cannot offer a fast enough process in order to catch the transient species responsible for catalysis. Even in small biomimetic compounds, the unstable formation of CuO₂ intermediates has forced the determination of the structure of reactive intermediates at very low temperatures.¹¹ In fact, a significant effort has been made toward a systematic approach to synthesize biomimetic systems to approximate the chemical and spectroscopic properties of Type-3 copper proteins. The choice of ligands, bridging ligands, solvent systems, and temperature can play a significant role in the elucidation of both redox-activity of model systems and their electronic properties that give rise to the now typical spectroscopic fingerprints of Type-3 oxidases.

In Type-3 metalloprotein biomimetic systems, the coordination environment around the metal ion is typically constituted of strong donor ligands such as the nitrogen atoms of pyridines that can mimic the histidine residues which have been proposed to assist the stabilization of metal-oxygen intermediates for electrophilic substitutions and 2-electron oxidation chemistry. The stabilization of highly electrophilic/nucleophilic CuO₂ intermediates is also assisted by a large hydrophobic environment. However, several factors can influence the reactivity of these compounds; in essence, any alteration on solvent, and coordinated (and bridging) ligands will change the reactivity of these biomimetic compounds.¹² The now classic example of biomimetic O₂-binding copper complexes that mimic the spectroscopic and chemical properties of their natural templates can be traced back to the efforts of Nobumasa Kitajima, Kenneth D. Karlin, and Edward I. Solomon from the late 1980’s to today.¹³-¹⁶
Although seemingly unrelated, the knowledge gained from investigating copper-O\textsubscript{2} interactions in metalloproteins and enzymes and their biomimetic analogues, may yield significant insight into the possible mechanism for metal-centered generation of reactive oxygen species (ROS). These ROS are generated at a metal center and may either diffuse away and cause damage to redox-sensitive biomolecules or remain bound to the metal and itself be an efficient redox catalyst. The common ground between these two areas of research then can be elucidated as the copper-O\textsubscript{2} interaction. From the detailed spectroscopy of biomimetic compounds and their protein templates, we have a library of diagnostic patterns to elucidate the nature of interactions of redox-active metals and O\textsubscript{2}. From the detailed mechanistic investigation of the chemistry catalyzed by the metalloproteins and their model compounds we are now well aware of what reactivity can be expected from different conformations of reactive CuO\textsubscript{2} intermediates. In recent years, a darker side of copper-O\textsubscript{2} chemistry has surfaced in biological studies. As research in the areas of cancer, apoptosis, diabetes, heart disease, and neurodegenerative disorders becomes more focused and detailed, the metal-centered generation of ROS has become a common theme. In fact, much of the focus has been placed on the nature of ROS rather than the interaction of the redox-active metal center with the ROS. These ROS are the radical and reduced forms of oxygen that although present under normal metabolic conditions, could potentially damage oxidizable moieties in their environment if not properly regulated due to the much higher oxidation potential of ROS compared to O\textsubscript{2}. ROS alone are able to damage every major type of biological molecule, including membranes (by peroxidation of lipids),\textsuperscript{17} proteins (by oxidation of certain residues and
nitration by nitric oxide), DNA (by destruction of the structural stability of the double helix as ribose is oxidized), and smaller molecules such as neurotransmitters. According to recent studies, OH•, O₂•⁻, and H₂O₂ have been assigned as the culprits in several types of neurological disorders, with a large number of research groups focusing on the role of ROS in the etiology of Alzheimer’s disease (AD). However, as chemists further investigate the pathway of ROS generation and the fate of such molecules, it becomes more evident that a redox-active metal is necessary in order for the damaging and often lethal chemical imbalance that is caused by ROS (E° = -300 mV for the one electron reduction of O₂) to achieve its full neurodegenerative potential; in other words it is the metallo-ROS interaction that appears to be the culprit in oxidative stress that leads to neurodegeneration, not ROS alone. Recent observations demonstrate that although oxygen radicals and oxidation agents like H₂O₂ cannot achieve high enough concentration in order to be the sole culprits in the neuropathology of AD due to inherent ability of organisms to cope with this oxidative stress, the presence of a metallo-ROS can form a very potent redox catalyst.

Alzheimer’s disease is one of several neurodegenerative disorders affecting a large percentage of the population. An estimated 3 million individuals in the United States suffer from the slow, yet aggregate symptoms of AD. The number is expected to double by the year 2030. The progressive nature of AD that has even been called an epidemic, has also dire effects on the families of those diagnosed with the disease. The AD patients are “memory timers”, bringing recognition of events and people around them slowly to extinction with each passing day. With the gradual, yet always progressive and
degenerative nature of AD, families are placed under considerable stress while caring for their loved ones and coping with the loss of their existence in the minds of those afflicted by AD. To that end, numerous scientific studies, amounting to tens of thousands, have been published, reflecting the interest in research funding and efforts toward understanding the neuropathology of this disease.

Largely, AD neuropathology concerns the interaction of a 40-42 amino acid peptide splice fragment from the ubiquitous apolipoprotein (or amyloid precursor protein, APP) with a number of intracellular components. These amyloid-β (Aβ) peptides are generated by the cleavage of APP by α, β, and γ secretases. Aβ in the form of insoluble plaques contains up to mM amounts of Zn$^{2+}$, Cu$^{2+}$, and Fe$^{3+}$ in the neocortical region of the brain. However, the cause/effect connection of the metallo-Aβ plaques with AD is still under debate. Aβ has been the elusive culprit in AD studies with results ranging from free-radical generation to disruption of mitochondrial cell membrane potential, all hallmarks of oxidative stress. Formation of Aβ plaques in the presence of metals has dominated a large percentage of investigations in correlating the physiology of AD with the metal chelating event (Aβ serving as the ligand) and generation of ROS. The subsequent observations of chemical events once metallo-Aβ is formed have offered numerous hypotheses into the oxidation stress proposed as one of the main causes of AD. Herein we present a compendium of observations that bring together efforts in the area of metal-O$_2$ chemistry from a catalysis perspective to better understand what role O$_2$ and its reduced forms may have in disease. Relating the neuropathology of AD with possible
metal-centered pathways is a recent area of focus that help clarify the homeostatic generation of ROS and the possible effects of non-homeostasis ROS generation.

II. METABOLIC GENERATION OF ROS

Electron donors in the tricarboxylic acid cycle in mitochondria (i.e. NADH and succinate) to O$_2$ are responsible for the generation of ATP for upkeep of cellular processes. Electron transfer (ET) processes rely on a series of molecular complexes to ensure that the transduction of oxidative energy and that the use of proton energy in ATP synthesis is carried out efficiently. The complexes that are functionally connected to mitochondrial energy transduction include: Complex I (NADH:ubiquinone oxido-reductase), Complex II (succinate:ubiquinone oxido-reductase), Complex III (ubiquinol:ferricytochrome c oxido-reductase), Complex IV(ferrocytochrome c:oxygen oxidoreductase), and Complex V (ATP-synthase).$^{1,27}$

In the inner mitochondrial membrane, within Complexes I, III, and IV the energy transduced from ET is conserved by coupled proton translocation that is responsible for the generation of a membrane electrochemical potential of protons used in ATP synthesis. The whole ET system is reversible and an electron flow can be generated against the current. However, cytochrome $aa_3$ in Complex IV and O$_2$, or the final step in ET is irreversible, shifting the equilibrium in the system toward ATP synthesis. Cytochrome $aa_3$ retains all the partially reduced oxygen intermediates bound to its active sites until the O$_2$ itself is completely reduced to water. However, through auto-oxidation
that can affect their reduced forms, other elements in the mitochondrial ET chain like ubiquinones and the cytochrome $b$ family could transfer the electrons directly to $O_2$, but do not retain the partially reduced $O_2$ intermediates in their active sites until the $O_2$ is completely reduced to water. Because $O_2$ accepts only one electron at a time, the superoxide radical ($O_2^{•–}$) is released. In the cytochrome $b$ family, it is noteworthy that cytochrome $b_{566}$ is closely involved with the processes of energy transduction in Complex III, wavering continually between a very low potential state of approximately $–30 \text{ mV}$ and a very high one of approximately $245 \text{ mV}$. The low potential of cytochrome $b_{566}$ could play a primary role in the formation of mitochondrial $O_2^{•–}$ because of an increase in its redox potential inhibiting the univalent transfer of electrons to oxygen. Therefore, electron leakage from the cytochrome $b_{566}$ would appear to be a real possibility which can thus bring about a continuous release of oxygen free radicals. During aging, the increasing amounts of these radicals that manage to escape the local defense mechanisms (e.g. scavengers, electron-trapping agents, etc.) may lead to multiple changes in the chemical and physical state of the membranes. As a matter of fact, superoxide generation$^{28,29}$ is significantly greater in the brain mitochondria of aged rats rather than young rats. This seems to be related to the fact that, with the exception of cytochrome $aa_3$ in Complex IV, the metabolic levels of the electron carriers does not undergo significant change with aging. However, the decrease in either the amount of cytochrome $aa_3$ or its catalytic cytochrome oxidase activity (COA)$^{30-34}$ in synaptic mitochondria from some cerebral regions (frontal cortex, parieto-temporal cortex, hippocampus, cerebellum, etc.)$^{35,36}$ may account for the finding that stoichiometric calculations show aging is
related to an increase in the percentage of ubiquinones and cytochrome \textit{b} family. Although this increase is not dramatic, relatively it does explain how electrons can "escape" the ET sequence from electron donors to \textit{O}_2. The \text{K}_m for cytochrome \textit{c} is constant in aged rats but the \text{V}_\text{max} decreases, suggesting that the COA activity is not related to the functional integrity of mitochondria accompanying senescence.\textsuperscript{35} Moreover, the COA activity drops significantly less in cortical synaptic mitochondria from old rats fed a hypocaloric diet,\textsuperscript{37} and declines in the homogenate of old rat cerebral cortex\textsuperscript{38} and in insect mitochondria.\textsuperscript{39}

Electrons can leak from the energy-transduction sequences even in young animals, indicating that the formation of superoxide radicals could be associated with the normal process of mitochondrial respiration.\textsuperscript{40} The production of these radicals causes cell damage because of the dismutase reaction in which \textit{H}_2\textit{O}_2 is formed and which, with the involvement of low-molecular-weight iron and copper complexes, leads to the highly dangerous hydroxyl radical. The catalytic activity of Complex IV (COA activity) is low in three cortical areas but not in putamen and hippocampus of AD patients compared with age-matched controls.\textsuperscript{41,42} There is no correlation between the changes in COA and those of other marker mitochondrial enzymes, such as glutamate dehydrogenase and citrate synthase, leading to the conclusion that the decrease in COA activity is not related to the loss of mitochondria.\textsuperscript{41} The decrease in COA activity in cortical areas and in the hippocampus from AD patients\textsuperscript{43,44} suggests a primary defect of Complex IV, resulting in more \textit{O}_2•⁻ released. A specific decrease in COA in the platelets of AD patients has also been reported\textsuperscript{45} but not confirmed.\textsuperscript{46} COA is heterogeneously distributed in the cerebral
spinal fluid (CSF). Expression of mRNA for COA in normal human and monkey brain is high in those regions which are most vulnerable to AD pathology and is particularly reduced in the same regions from AD patients. In the mid-temporal gyrus, but not in the primary motor cortex, of AD patients there is a 50-65% decrease in the mRNA levels of the mitochondrial DNA (mtDNA)-encoded COA subunits I and III. However, the mitochondrial-encoded 12S ribosomal RNA (a mitochondrial transcript) does not change, suggesting that the observed reduction of COA I and III mRNA is not due to loss of mitochondria but to a specific alteration of transcriptional regulation. A behavioral study in rats treated with the selective sodium azide COA inhibitor showed significant inhibition of a low-threshold form of hippocampal long-term potentiation and impaired spatial learning. This finding supports the theory that Complex IV alteration is somehow involved in the pathogenesis of AD and also raises the possibility of developing an animal model reflecting this aspect of AD provided it is specific.

III. O$_2^{-}$ AND SOD

Superoxide radicals are described as having considerable reactivity, short half-life, and limited diffusion through membranes. However, the latter property has been questioned, as these radicals can appear in the brain extracellular space. O$_2^{-}$ has a dual effect where it may help protect against infectious microorganisms, but they can also be harmful as it participates in the formation of the very reactive hydroxyl radicals (OH•). Moreover, they can inactivate a number of useful enzymes, such as antioxidat
enzymes (catalase\textsuperscript{55} and glutathione peroxidase\textsuperscript{56}), enzymes involved in neurotransmission (glutamine synthase\textsuperscript{57}), in signal transduction (adenylate cyclase\textsuperscript{58}), and in energy transduction (creatine phosphokinase\textsuperscript{59} and NADH dehydrogenase and ATPase\textsuperscript{60}).

In AD brain and fibroblasts there is evidence of partial uncoupling of mitochondrial oxidation and phosphorylation\textsuperscript{61,62}. Apart from the neurodegeneration induced by the impairment of energy metabolism, these oxidative abnormalities could contribute to the accumulation of cytoskeletal material. Addition of an uncoupler to cultured fibroblasts from normal subjects causes the appearance of epitopes recognized by antibodies to paired helical filaments (PHF) and Alz-50 monoclonal antibodies\textsuperscript{61}, thus reproducing a pattern that is characteristic of fibroblasts from AD patients\textsuperscript{63}.

When O\textsubscript{2} accepts an electron from a reducing agent, the primary product is O\textsubscript{2}\textbullet\textsuperscript{−} that, in aqueous environments, is in equilibrium with its protonated form (\textbullet\textsubscript{O}O\textsubscript{2}H). When O\textsubscript{2}\textbullet\textsuperscript{−} and \textbullet\textsubscript{O}O\textsubscript{2}H approach equal molar concentrations, spontaneous dismutation occurs, and H\textsubscript{2}O\textsubscript{2} plus \textsuperscript{1}O\textsubscript{2} (singlet oxygen)\textsuperscript{64} are generated.

\[
2 \text{O}_2\textbullet^- + 2H^+ \leftrightarrow \text{H}_2\text{O}_2 + \text{O}_2 \text{ (or } \text{^1O}_2) \quad \text{Scheme (I)}
\]

In scheme 1, O\textsubscript{2}\textbullet\textsuperscript{−} can be converted into H\textsubscript{2}O\textsubscript{2} catalyzed by superoxide dismutase (SOD) that is present in varying concentrations in neural cells. Thus, the conversion removes O\textsubscript{2}\textbullet\textsuperscript{−} and prevents its direct toxic action as well as its interaction with metal ions to increase the production of hydroxyl radicals (Scheme 9 below).
The rate constant for SOD-catalyzed dismutation is approximately four orders of magnitude greater than that for the spontaneous dismutation of O$_2^{•−}$ at physiological pH. For SOD protection/activity to work properly, it is absolutely vital for other enzymes (e.g., catalase, glutathione peroxidase, etc.) to convert H$_2$O$_2$ immediately into H$_2$O, thus preventing the transformation of H$_2$O$_2$ by metal complexes into the highly toxic OH• (Scheme 9 below). In this last case, the intervention of SOD may be dangerous for neuronal cells in spite of what we expect as a beneficial biological scavenging of a highly reactive radical.

The predominant two types of SOD are Mn-dependent and Cu-Zn dependent. The manganese-dependent (Mn-SOD) is located in the mitochondria, where it interacts with the O$_2^{•−}$ leaking from the ET chain. The copper- and zinc-dependent (Cu,Zn-SOD) is located in the neural cytosol where it carries out a more general catalytic function. Auto-oxidizable electron carriers located on the internal mitochondrial membrane can generate O$_2^{•−}$ which is enzymatically dismutated to H$_2$O$_2$. However, a few reactions catalyzed by some enzymes (e.g., monoamine oxidase and L-aminoacid oxidase) can produce H$_2$O$_2$ directly. Thus, H$_2$O$_2$ may be generated either as a direct product or from each of the various sources of O$_2^{•−}$ by auto-oxidation of a variety of low-molecular weight molecules, as byproducts of various enzyme catalyses such as between xanthine and xanthine oxidase, and by the mitochondrial ET system.

The hydroxyl radical is one of the most potent reactive metabolites produced in brain systems derived from O$_2$. This radical, O$_2$, and OH are all products in the reaction shown in Scheme 2 when H$_2$O$_2$ is directly reduced by O$_2^{•−}$:

\[
2\text{H}_2\text{O}_2 + \text{O}_2^{•−} \rightarrow \text{H}_2\text{O} + \text{H}^+ + \text{O}_2 + \text{OH}^{-}
\]
\[ \text{O}_2^{•−} + \text{H}_2\text{O}_2 \leftrightarrow \text{O}_2 \text{ (or } \text{O}_2^{•}\text{)} + \text{OH}^{•} + \text{OH}^{−} \]  \textbf{Scheme (2)}

H\textsubscript{2}O\textsubscript{2} can cross cell membranes directly, whereas O\textsubscript{2}^{•−} crosses cell membranes through anion channels. Although H\textsubscript{2}O\textsubscript{2} cannot be classified as a radical because it does not contain unpaired electrons, it is still dangerous because it easily permeates cell membranes and can migrate from where it is first generated to other organic compartments. It can interact with the reduced forms of some metal ions (generally, Fe\textsuperscript{2+} and Cu\textsuperscript{+}) and decompose into the highly reactive OH\textsuperscript{•} and the OH\textsuperscript{−}, according to:

\[ \text{H}_2\text{O}_2 + \text{Fe}^{2+} \leftrightarrow \text{Fe}^{3+} + \text{OH}^{•} + \text{OH}^{−} \]  \textbf{Scheme (3)}

\[ \text{H}_2\text{O}_2 + \text{Cu}^{+} \leftrightarrow \text{Cu}^{2+} + \text{OH}^{•} + \text{OH}^{−} \]  \textbf{Scheme (4)}

The formation of OH\textsuperscript{•} requires reduced forms of metal ions, such as Fe\textsuperscript{2+} or Cu\textsuperscript{+}. The superoxide radical O\textsubscript{2}^{•−} can then give rise to Fe\textsuperscript{2+} or Cu\textsuperscript{+} by reducing Fe\textsuperscript{3+} or Cu\textsuperscript{2+} according to:

\[ \text{O}_2^{•−} + \text{Fe}^{3+} \leftrightarrow \text{O}_2 + \text{Fe}^{2+} \]  \textbf{Scheme (5)}

\[ \text{O}_2^{•−} + \text{Cu}^{2+} \leftrightarrow \text{O}_2 + \text{Cu}^{+} \]  \textbf{Scheme (6)}

The reaction in Scheme (2) is slow at physiological pH and would require steady-state concentrations of the reaction partners, which are much higher than those found in cerebral mitochondria to account for detectable amounts of the highly unstable radical.
As shown in Schemes 3-6, metal ions (M^{n+}) accelerate the reaction in Scheme (2) by catalyzing two intermediate reactions:

\[
O_2\cdot^- + M^{n+} \leftrightarrow O_2 + M^{(n-1)+} \quad \text{Scheme (7)}
\]

\[
M^{(n-1)+} + H_2O_2 \leftrightarrow M^{n+} + OH\cdot + OH^- \quad \text{Scheme (8)}
\]

Overall: \[O_2\cdot^- + H_2O_2 \leftrightarrow O_2 \text{ (or } ^1O_2) + OH\cdot + OH^- \quad \text{Scheme (9)}\]

In Scheme 9, \(O_2\cdot^-\) reduces redox-active transition metals (Fe^{3+}, Cu^{2+}), and generates oxygen or singlet oxygen. The reduced form of the metal subsequently reacts with \(H_2O_2\) to produce the oxidized form of the metal, the hydroxide ion, and the hydroxyl radical. In view of the evidence that in neural systems Scheme 2 proceeds very slowly, the hydroxyl radicals are possibly produced in the presence of a redox-active metal. An additional mechanism by which \(OH\cdot\) may be generated is supported by the observation that incubation of \(H_2O_2\) with \(Fe^{2+}\) and iodide ions (I\(^{-}\)) can generate a potent reactive molecule that is inhibited by scavengers of the hydroxyl radical (e.g., mannitol\(^{65}\) and ethanol\(^{66}\)).

Singlet oxygen\(^{63}\) is generated by the SOD in Scheme 2 or by the interaction with metal ions in Scheme 9 when one of the two unpaired electrons of \(O_2\) acquires sufficient energy to undergo spin inversion or both spin inversion and orbital transition. There are two distinct forms of singlet oxygen, \(\Delta\) and \(\Sigma\), which are dependent respectively on whether the excited electron forms an electron pair in the same orbital or remains unpaired in a different orbital. The \(\Delta\) form of singlet oxygen is more stable than the \(\Sigma\) form. Singlet


**Table 1.1.** Standard redox potentials for dioxygen species in water.

<table>
<thead>
<tr>
<th>Reaction</th>
<th>E° (V)</th>
</tr>
</thead>
<tbody>
<tr>
<td>O₂ + e⁻ → O₂•⁻</td>
<td>−0.33</td>
</tr>
<tr>
<td>O₂•⁻ + e⁻ + 2 H⁺ → H₂O₂</td>
<td>+ 0.89</td>
</tr>
<tr>
<td>H₂O₂ + e⁻ + H⁺ → H₂O + OH</td>
<td>+ 0.38</td>
</tr>
<tr>
<td>OH + e⁻ + H⁺ → H₂O</td>
<td>+ 2.31</td>
</tr>
<tr>
<td>O₂ + 2e⁻ + 2 H⁺ → H₂O₂</td>
<td>+ 0.281</td>
</tr>
<tr>
<td>H₂O₂ + 2e⁻ + 2 H⁺ → 2 H₂O</td>
<td>+ 1.39</td>
</tr>
<tr>
<td>O₂ + 4 H⁺ + 4 e⁻ → 2 H₂O</td>
<td>+ 0.815</td>
</tr>
</tbody>
</table>
Figure 1.1. Electronic states and standard reduction potentials for O₂. ⁶⁷
oxygen is highly electrophilic and can react with electron-rich biomolecules such as tryptophan, methionine, and molecules containing unsaturated double bonds.

The effect of physiological aging on brain SOD is controversial, though most reports describe some age-related decline, mainly of the Cu-Zn form\textsuperscript{68,69,70,71,72} probably related to the decline of SOD mRNA. However, other studies report no change of this form and an increase in MnSOD\textsuperscript{73,74} Thus, it would appear that higher levels of SOD are beneficial and lower levels of SOD detrimental. However, if the activity of SOD is increased without a concomitant enhancement of the activity of the enzymes which dispose of H\textsubscript{2}O\textsubscript{2} (i.e. glutathione peroxidase) and the concentration of reduced glutathione, then H\textsubscript{2}O\textsubscript{2} accumulates and reacts with O\textsuperscript{•−} and Fe\textsuperscript{3+} and/or reduced metal ions to form the very reactive OH\textsuperscript{•}. Thus the imbalance between SOD and H\textsubscript{2}O\textsubscript{2}-converting enzymes results in a toxic effect of SOD by OH\textsuperscript{•} generation, inducing DNA fragmentation, protein denaturation, and activation of the autocatalytic process of lipid peroxidation.

Down's syndrome (trisomy 21) has provided some interesting clues on the balance among SOD, ROS, and antioxidants. Human Cu,Zn-SOD is encoded by a gene located on chromosome 21 and Down patients have a 50\% increase in the activity of this enzyme secondary to gene dosage effect\textsuperscript{75} The increase in SOD, not accompanied by a concomitant adaptative rise in glutathione peroxidase,\textsuperscript{76} might induce oxidative damage to the CNS, including lipid peroxidation and this might explain some of the neurobiological abnormalities found in Down's syndrome, such as accelerated aging and AD-type neuropathology. This is supported by the results in an animal model of gene
dosage effect in transgenic mice carrying the human Cu,Zn-SOD gene.\textsuperscript{77,78} Cu,Zn-SOD
and its mRNA are preferentially expressed in the large pyramidal neurons of Ammon's
horn and granule cells of the dentate gyrus, which are susceptible to degenerative
processes in AD.\textsuperscript{79} Brain lipid peroxidation is also increased. In seeming contrast with
this finding, increased Cu-Zn SOD in transgenic mice makes the hippocampus more
resistant to the neurotoxicity induced by amyloid-\(\beta\).\textsuperscript{80}

The levels of Cu,Zn-SOD protein and mRNA in the vulnerable hippocampal
neurons of AD patients,\textsuperscript{81} their association with neurofibrillar degeneration (PHF),\textsuperscript{82} and
the observation that the cell distribution of Cu,Zn-SOD mRNA in the human
hippocampus is the same as amyloid mRNA\textsuperscript{83} suggest that high levels of enzymes for
ROS decomposition are needed to remove excess superoxide radicals which indicate that
ROS contribute to the degenerative processes leading to neuropathology in AD. Cu,Zn-
SOD activity is also upregulated in the temporal cortex and nucleus basalis Meynert\textsuperscript{84,85}
and in the fibroblasts of AD and Down patients.\textsuperscript{86} Cultured skin fibroblasts from both
familial and sporadic AD patients are more susceptible to ROS-induced damage then
from age-matched controls.\textsuperscript{87} Finally, high immunoreactivity for SOD and catalase in the
AD brain is associated with some neurofibrillar tangles and senile plaques.\textsuperscript{88} This
immunoreactivity is absent in tangle-free neurons of AD and all neurons of normal
control brains.
IV. METAL-BIDING TO Aβ AND STRUCTURE OF THE COMPLEX

The primary step in understanding metal-centered chemistry is the elucidation of the coordination of metal ions in its complexes. The geometry of metal complexes is often the determinant of reactivity. Once it was determined that transition metals could bind to Aβ from analysis of isolated plaques, more detailed investigation of the nature of the metal-Aβ interaction followed. Through the use of competitive binding in a pH gradient column chromatography, Cu$^{2+}$, Ni$^{2+}$, and Zn$^{2+}$ were determined to tightly bind to Aβ$_{1-42}$; shorter fragments (Aβ$_{1-16}$ and Aβ$_{1-28}$) could also be retained in the metal-chelate column but did not bind as tightly. Based on the pK$_a$ of 6.1 determined from the pH gradient elution, His residues were suggested as the ligands responsible for metal-binding. Further pH dependent binding studies using quantitative precipitation and turbidity revealed that Cu$^{2+}$ can induce aggregation of Aβ$_{1-40}$ and Aβ$_{1-42}$. Chemical modification of His residues in the peptide to N-carbethoxyhistidine extinguishes the metal-induced aggregation. These results indicated that His residues were important in metal binding, but did not provide insight into geometry nor the affinity of Aβ toward transition metals. Quantitative precipitation studies suggested that Cu$^{2+}$ binds with an attomolar dissociation constant to Aβ$_{1-42}$. This very low K$_d$, if correctly measured, would suggest that Aβ could actually remove metal from hemes, some of the tightest binding ligands in biological systems. This result however, erroneously explained the metal-bidning because the combination of multiple molecular events in terms of equilibrium can yield such a low metal-dissociation constant. Since quantitative
precipitation measures both the metal affinity to Aβ and also the affinity associated with peptide aggregation, the attomolar constant is thus a combination of different constants for the two processes that would not be quantified individually until chemical and spectroscopic methods were used to directly measure metal-peptide interactions in solution. Both methods yielded $K_d$ for metal-binding in the µM range for Aβ$_{1-16}$, Aβ$_{1-20}$, and Aβ$_{1-28}$.

In order to further elucidate the location of metal-binding in metal-Aβ complexes resonance Raman was employed to directly observe metal-peptide interactions. The Raman spectra demonstrate that three His residues in the N-terminal hydrophilic region provide primary metal binding sites and the geometry of the metal-Aβ complex is correlated with the metal binding mode. Zn$^{2+}$ binds to the N$\varepsilon$ atom of a His imidazole ring and the peptide was suggested to aggregate through intermolecular His(N$\varepsilon$)-Zn$^{2+}$-His(N$\varepsilon$) bridges. The N$\varepsilon$-metal ligation also occurs in Cu$^{2+}$-induced Aβ aggregation at mildly acidic pH. At neutral pH, however, Cu$^{2+}$ was shown to bind to N$\delta$, the other nitrogen of the His imidazole ring, and to a deprotonated amide nitrogen of the peptide main chain. The studies using electronic absorption spectroscopy and $^1$H NMR techniques have also shown the nature of the metal-Aβ interaction. The electronic spectrum of Cu$^{2+}$-Aβ$_{1-20}$ shows a typical tetragonally distorted octahedral environment that is consistent with many Cu$^{2+}$ complexes in solution. Co$^{2+}$ as a paramagnetic shift reagent in $^1$H NMR experiments revealed that three His residues coordinate all N$\varepsilon$ nitrogens from the imidazole rings of His6, His13, and His14. The results of the NMR...
study also revealed that neither Tyr10 nor the N-terminus bind to the metal as was previously suggested.\textsuperscript{95-97}

Through the use of CW-EPR techniques and line broadening observations in \textsuperscript{1}H NMR experiments, a model was suggested for the formation of Aβ dimers that resembled the structure of the active site of Cu,Zn-SOD.\textsuperscript{96} The suggested structure yields the metal:peptide stoichiometry of 1:1 and that two metal metal centers can be bridged through a third His residue, suggested to be His6.\textsuperscript{96} These results were further confirmed by using chemically modified (methylated) His residues and CW-EPR.\textsuperscript{98} The broadening of the EPR spectrum was proposed to be due to antiferromagnetically couple Cu\textsuperscript{2+} that could only arise from a bridging ligand interaction or a close proximity of two metal centers. However, the distinct features of Cu,Zn-SOD EPR are well known\textsuperscript{99} which do not resemble those of the CuAβ complexes. Indeed, the possibility of bridged metal-centers in CuAβ complexes is not preposterous which can be reasoned as an OH-bridged or even phosphate-bridged Cu centers (being that PBS is the preferred buffers in most of the experiments cited). The interpretation that the broadening of EPR signals in CuAβ samples is likely due to the aggregation of the metallopeptides in solution and not because of magnetic coupling, since the metal-induced aggregation of Aβ is a well known phenomenon.\textsuperscript{100-103}

Although the metal-binding properties of Aβ have been widely studied,\textsuperscript{25} questions about the correct stoichiometry were until recently not clarified. Recent studies using a number of different physical methods, ranging from \textsuperscript{1}H NMR to fluorescence and kinetics, have elucidated that each Aβ monomer of varying lengths can only bind one
Recent theoretical studies have shown that the seeds for dimerization occur between residues 16-22, causing β-sheet formation in smaller fragments that can range from dimmer to 16 peptides per fibril. It is thus possible for two individual metal centers to be close enough to each other due to dimerization and show binuclear metal-centered catalysis. Such structural motif may well explain the redox chemistry involved in etiology of AD.

V. GENERATION OF ROS BY METALLO-Aβ

The cytotoxic effects of Aβ, which accumulates in the brain in AD, have been studied extensively. Early studies suggested that the aggregation state of Aβ is related to its toxicity. It was shown that freshly solubilized Aβ exhibited little toxicity, whereas Aβ that had been aged for 7 days (forming aggregates in fibrillar states) was cytotoxic. However, the precise molecular mechanism by which Aβ mediates cell death has remained a matter of considerable dispute. There is general agreement that the production of ROS (see section II) and the influx of calcium ions into cells are both involved in toxicity, but it is still unclear how the generation of ROS might be related to Aβ. One possibility is that the aggregated peptide itself may be able to produce ROS directly in the forms of free radicals or H2O2.

It is generally accepted that Aβ needs to be in aggregated or partially aggregated states before it becomes toxic to cells. However, there is still no clear consensus on the precise nature of the toxic form of the peptide. Dimers, soluble oligomers, protofibrils
and annular protofibrils have all been implicated in amyloid toxicity. Furthermore, it is becoming more evident that not all amyloid fibrils are toxic to cultured cells. Recently a synthetic peptide corresponding to amino acid residues 1-15 of Aβ protein (Aβ 1-15) was shown to form amyloid fibrils in vitro that are completely non-toxic to cells, even at high concentrations. Some forms of synthetic Aβ are also reported to be non-toxic, as is the Aβ peptide mutant that contains a norleucine in the place of Met35. This leads to yet another area of research that is ongoing and still trying to elucidate which forms of amyloid are toxic. What underlying molecular structures and mechanisms can explain why some amyloids are toxic, whereas others are not.

Various other hypotheses have been put forward to explain the cytotoxic effects of Aβ in addition to the direct production of ROS from the peptide, including the formation of ion calcium channels in cell membranes by Aβ, interactions between Aβ and specific cell surface receptors, such as the RAGE (receptor for advanced glycation endproducts) or scavenger receptors, interactions between Aβ and intracellular target molecules such as ERAB (endoplasmic reticulum Aβ binding protein), and non-specific intercalation of aggregated forms of Aβ into membranes. These various interactions may not be related to the idea that the most important aspect of the toxicity of Aβ is due to its ability to generate ROS directly once bound to a redox-active metal. The toxic process could be the result of a combination of binding and/or attachment of Aβ aggregates to cell components, followed by the induction of oxidative damage.

According to the oxidative stress hypothesis, a spontaneous shower of Aβ-derived free radicals is produced upon incubation of Aβ in vitro, and it is this phenomenon which
rendered the cytotoxic properties of Aβ. Three main pathways can lead to the formation of free radicals from organic molecules: photolysis, thermal scission, and one-electron redox reactions. Mechanical stress would be a fourth possible route that may apply in some special cases. Of these, photolysis is not relevant to neurodegenerative and amyloid diseases. Thermolysis of most chemical bonds requires temperatures above 450 °C with the exception of peroxides and azo compounds. The temperature at which peroxides undergo unimolecular scission varies with the class of peroxide but can be between about 50 and 150 °C. Likewise, nitrogen can be eliminated from certain azo compounds (general structure R-N=N-R) over a similar temperature range. However, neither of these classes of bond is present in the parent Aβ peptide molecule. Perhaps the most important redox assisted bond scissions involve a metal ion that readily undergoes a one-electron transfer with one of the best known examples being the Fenton reaction:

\[ \text{M}^{n+} + \text{H}_2\text{O}_2 \rightarrow \text{M}^{(n+1)+} + \text{OH}^+ + \text{OH}^- \]

Metal assisted homolysis reactions can take place at much lower temperatures than the corresponding unassisted reaction. While such reactions in peptide molecules cannot be totally discounted at 37 °C, they do not provide an obvious route to the spontaneous generation of peptidyl free radicals from Aβ. However, it has been suggested on purely theoretical studies that peptidyl free radicals might be generated from Aβ by a mechanism involving mechanical stress, i.e physical contact among metallo-Aβ monomers in the plaque.
Electron paramagnetic resonance spectroscopy (EPR) is the preferred technique for the detection of free radicals and geometrical states around paramagnetic transition metal ions with unpaired electrons. EPR is highly sensitive and interpretation of the hyperfine structure associated with each spectrum allows the electronic nature of the radical to be established. However, the high reactivity and short life-time coupled with low steady-state concentrations of nearly all radicals prevent their direct detection in most cases. A simple experimental technique that is coupled with EPR is to employ spin-trapping, in which a reactive radical reacts with a nitrone or a nitroso compound to form a much more stable nitroxyl radical whose concentration rises well above the detection limit of the spectrometer. The selection of the appropriate spin-trap is important and depends upon the nature of the initial radical and on the objectives behind the experiment. Choice of incorrect spin-traps may yield data that are ambiguous and misinterpreted (discussed below).

The suggestion that Aβ itself might spontaneously generate free radicals was first made in 1994 when EPR spectra were observed during 6 hour incubation of Aβ 1-40 at 37 °C in phosphate buffered saline (PBS) in the presence of the spin-trap N-tert-butyl-α-phenylnitrone (PBN). The observation therein and further EPR spectra employing PBN as a spin-trap led to the “molecular shrapnel” hypothesis. The main feature of these EPR spectra was a 4-line pattern, and such a pattern is not characteristic of a true PBN adduct. In a similar experiment, Tomiyama et al. observed a spectrum consisting of the superimposition of 3-line, 4-line, and 6-line patterns during the incubation of Aβ 1-40 in the presence of PBN in deionized water. Again, the 3-line pattern is not characteristic of a
true PBN adduct. Under conditions of good spectral resolution, PBN adducts, without exception, consist of a 6-line pattern.\textsuperscript{135} Subsequently, Dikalov et al.\textsuperscript{136} reported that they were unable to detect any EPR spectra following the 6 h incubation of Aβ 1-40 and Aβ 25-35 with PBN. When these same peptides were incubated with a less pure sample of PBN they observed both 3-line and 4-line EPR spectra. These spectra were also observed in the absence of peptide, and were attributed to transition metal-catalyzed auto-oxidation of di-\textit{tert}-butylhydroxylamine and \textit{N}-\textit{tert}-butylhydroxylamine, present as impurities within their sample of PBN. It was proposed that these impurities were converted to their corresponding nitrooxyls, di-\textit{tert}-butylnitroxide and \textit{tert}-butylhydroaminoxyl. Further there is current evidence that Aβ and α-synuclein can generate ROS directly.\textsuperscript{118} The arguments presented herein could apply to proteins associated with some of the other protein conformational diseases mentioned above.

In some recent publications Bush and co-workers have reported that Aβ 1-40 forms H$_2$O$_2$ in the presence of both copper and iron and can reduce Fe$^{3+}$ to Fe$^{2+}$ as well as Cu$^{2+}$ to Cu$^{+}$.\textsuperscript{110,137-139} The possible formation of H$_2$O$_2$ during the incubation of Aβ in PBS could provide some interesting insights into the chemistry outlined. First, H$_2$O$_2$ is a strong oxidizing agent and, if present, would certainly promote the hydrolysis of PBN. Secondly, the presence of both H$_2$O$_2$ and Fe$^{2+}$ and/or Cu$^{+}$ would lead to the formation of hydroxyl radicals by Fenton chemistry. It has been found that low concentrations of metals are present even in chelex buffers, and that significant amounts of metals are also bound to the peptide itself.\textsuperscript{140} If formed, the hydroxyl radical would be trapped by PBN, but the resulting adduct is unstable and is not directly observed in aqueous solution.
There is good evidence that at pH 7 and above PBN rapidly transforms into the tert-butylhydroaminoxyl radical.\textsuperscript{141} This could also help to explain the origin of the characteristic 4-line spectrum of the tert-butylhydroaminoxyl radical obtained when Aβ is incubated in the presence of PBN. Consequently, it may be that the very weak EPR signals of tert-butylhydroaminoxyl observed by Butterfield and co-workers\textsuperscript{132,133,142-145} and by Monji \textit{et al}\textsuperscript{146,147} are resulted from the formation of H\textsubscript{2}O\textsubscript{2} in the presence of low concentrations of Fe and Cu rather than the spontaneous formation of peptide-derived radicals. A second scheme involving reactions of 2-methyl-2-nitrosopropane (MNP) and PBN explains the origins of the 3- and 6-line spectra in the presence of Aβ and ambient laboratory lighting. The possibility that H\textsubscript{2}O\textsubscript{2} might be formed in the immediate vicinity of the peptide is interesting since, as noted above, in the presence of the appropriate metal ions it would be readily converted to hydroxyl radicals \textit{via} Fenton chemistry. The hydroxyl radical is a strong oxidant that can attract hydrogen atoms from organic molecules with a rate constant close to the diffusion controlled limit.\textsuperscript{148} Because of its high reactivity the hydroxyl radical is very unselective and removes hydrogen atoms from primary, secondary and tertiary carbon atoms with almost equal facility. It also adds to C=C double bonds with a rate close to the diffusion controlled limit. Consequently, immediate oxidative damage would occur to any organic molecule in vicinity of the site of hydroxyl radical.

Bush and co-workers\textsuperscript{110,139} have reported results showing the production of H\textsubscript{2}O\textsubscript{2} in systems involving Aβ 1-40, Aβ 1-42 and low concentrations of Fe\textsuperscript{3+} and Cu\textsuperscript{2+} in PBS. Quantitative measurement of the H\textsubscript{2}O\textsubscript{2} levels was carried out using the standard
thiobarbituric acid reactive species (TBARS) assay and not spin-trapping. Their results are reasonably convincing, however, not in agreement with other studies where H$_2$O$_2$ levels appear to be lower than those reported by Bush, as double integration of DMPO hydroxyl radical adduct signals against calibration standards indicates the concentration of adducts is not higher than about 0.5 mM from 100 mM CuAβ 1-40.

A significant number of the aggregating polypeptides that have been implicated in neurodegenerative disease have been shown to bind to transition metal ions such as iron, zinc, copper or manganese. Another common emerging theme is that some of these polypeptides seem to function normally as antioxidants through structural or activity similarities to the active site of superoxide dismutase. This seems to apply to prion protein (PrP)$^{149}$ and the Aβ peptide.$^{150}$ When these proteins are in an altered pathological configuration, they could actually become pro-oxidants. If we take the example of Aβ, there is strong evidence supporting that binding of Cu$^{2+}$, Zn$^{2+}$, and possibly Fe$^{3+}$ to the peptide involves coordination with the three histidine residues at positions 6, 13 and 14, along with the tyrosine residue at position 10.$^{150}$ In the general case of biologically relevant redox-active transition metal ions, binding to peptide molecules would still leave available coordination sites, allowing oxygen to bind, possibly in the form of a peroxo bridge. Such bridges are common in a variety of metal complexes,$^{5,6,10,11,16}$ but in this case there is a further issue that should be considered. As the peptide molecules change conformation and begin to aggregate this could have the effect of twisting the bridging oxygens, resulting in a straining of the complex. However, in the dimeric and small polymeric forms of the soluble peptide the possibility of the formation of strong
electrophiles such as the Type-3 copper models (discussed below) cannot be discounted. The conversion of superoxide to H₂O₂ is a well established reaction. It is possible that the superoxide, and subsequently hydrogen peroxide, is formed as a byproduct of the aggregation and fibril assembly process. In other words, the amyloid-induced toxicity could actually be associated with the generation of ROS during the process of fibril extension and growth. In this case, a search for one or more types of special toxic aggregate may yield several variants of toxic intermediates.

It is assumed that Fenton chemistry can occur if peroxide is present in close proximity to Fe³⁺ resulting in the formation of hydroxyl radicals which will react with organic molecules in situ, including peptides. In this case a peptidyl radical may be detectable. Yanker and coworkers¹⁵¹ investigated the neurotoxicity of a range of peptides spanning the entire Aβ 1-40 sequence and concluded that 25-35 was the toxic domain. This report was supported by Pike et al¹⁰⁹ who observed the formation of stable aggregates and neurotoxicity in synthetic peptides containing residues 29-35. These results focused attention on the possibility that a particular amino acid residue within this sequence might play a key role. In particular the methionine 35 residue has attracted attention.¹⁴⁵ Substitution of this methionine by aspartate, serine and cysteine all resulted in peptides which were neither aggregated nor neurotoxic.¹⁵² Conversely, substitution by leucine, norleucine, lysine and tyrosine residues resulted in peptides which neither aggregated nor were neurotoxic. Significantly, substitution of the methionine residue by serine or cysteine led to peptides which were at least as neurotoxic as Aβ 25-35 itself suggesting that methionine is not unique in promoting these properties.¹⁵²
VI. TYPE-3 COPPER OXIDASE MODELS AND METALLO-ROS

Because the generation of activated forms of O₂ through metal-centered reactions has been implicated in the etiology of neurodegenerative disorders, it warrants the question: Is the ROS generated and then diffused or does it remain metal bound? Thus, even if the fields of Type-3 copper oxidase biomimetics and neurodegeneration by ROS seem unrelated, at least in terms of goals, they are certainly connected at the chemical level by the drive to understand copper-O₂ interactions.

The oxidation of organic substrates with molecular oxygen under mild conditions is of great interest for industrial and synthetic processes from an economical and environmental point of view. Although the reaction of organic substances with dioxygen is thermodynamically favorable, it is kinetically unfavorable due to the triplet ground state of O₂. In biological systems this problem is overcome by the use of copper- or iron-containing metalloproteins which serve as highly efficient oxidation catalysts. Some examples of Cu-oxidases/oxygenases are shown in Figure 1.

The catechol oxidases are Type-3 copper enzymes containing a binuclear copper center. Well-known representatives of these Type-3 copper proteins are hemocyanin, the O₂ carrier for arthropods and mollusks, and tyrosinase. Catechol oxidase belongs, like tyrosinase, to the polyphenol oxidases which catalyze the oxidation of phenolic compounds to quinone in the presence of O₂. Tyrosinase catalyzes the hydroxylation of tyrosine to DOPA and the oxidation of DOPA to dopaquinone with
Figure 1.2. Types of enzymes involved in O₂ metabolism.\textsuperscript{16}
electron transfer to O₂, while catechol oxidase only catalyzes the oxidation of catechols to quinones. This reaction is of significance in medicine and biology since catecholamine neurotransmitters like epinephrine, norepinephrine, and DOPA are part of the tyrosine metabolic pathway. Secondary reactions like melanin formation are downstream after oxidation of the catechol moiety of catecholamine substrates in the presence of polyphenol oxidases, which are also responsible for the brown color of injured plants.

The copper in isolated catechol oxidases was determined to be EPR-silent due to an antiferromagnetically coupled Cu²⁺–Cu²⁺ pair. The electronic absorption spectrum of oxy-catechol oxidase from Ipomoea batatas exhibits an intense absorption band at 343 nm and a weaker band at 580 nm, later found to be due to the peroxo complexes of hemocyanin and tyrosinase. These intense electronic transitions were assigned to a peroxo to Cu²⁺ charge transfer transitions with an O–O stretch vibrational band at 749 cm⁻¹ indicating a possible µ-η²:η² bridging mode of the peroxo group. X-ray absorption spectroscopy (XAS) investigations on the native met forms of catechol oxidases from Lycopus europaeus and Ipomeas batatas have revealed that the active site consists of a dicopper(II) center, in which the metal atoms are coordinated by four N:O donor ligands. Multiple scattering extended x-ray absorption fine structure (EXAFS) calculations have shown high significance for one or two coordinating histidine residues. The short metal–metal distance of 2.9 Å and the results of EPR investigations indicate that a µ-OH bridged dicopper(II) center resides in the active site in the met forms of the proteins. Model studies of synthetic analogues have furthered the understanding of the structural and chemical properties of these proteins. Current interest is focused
on the elucidation of the mechanism of copper complexes that show catecholase and phenolase activities with different structural and electronic features around the copper ions. In these studies, mono- or multinuclear complexes have been synthesized and the properties of the chelating ligands have been varied with respect to geometry, number, and nature of the donor atoms. There are several structural constraints for proper Type-3 oxidase activity, such as square-planar mononuclear Cu$^{2+}$ complexes exhibit only little catalytic activity while non-planar mononuclear Cu$^{2+}$ complexes show a high activity.\textsuperscript{172,173} It was also found that binuclear complexes can also catalyze oxidation reactions if the Cu···Cu distance is less than 5 Å. A steric match between substrate and complex is believed to be one of the determinants for reactivity where two metal centers have to be located in close proximity to facilitate binding of the two hydroxyl oxygen atoms of catechol prior to the electron transfer.\textsuperscript{172} This conclusion was supported by the observation that binuclear copper complexes are generally more reactive towards the oxidation of catechols than are the corresponding mononuclear species.\textsuperscript{174} It has been particularly challenging to synthesize structures where a substrate bridged compound can be resolved. Thus far only one case is known in which the crystal structure of a catalytically active binuclear Cu$^{2+}$ complex with a coordinated catecholato ligand has been solved.\textsuperscript{175} Two other examples of mononuclear square-planar copper complexes\textsuperscript{176} were also effective catalysts, demonstrating that geometrical effects are only one aspect of the complex activity. In mechanistic studies, the same authors pointed out that a narrow range of redox potentials for effective catalysis exists between ease of reduction by the substrate and subsequent reoxidation by O$_2$.\textsuperscript{177} Although some general structure-
reactivity patterns have been found, the oxidation chemistry of structurally well-characterized copper complexes is still not fully understood, especially regarding the parameters affecting the catecholase activity. For example, there appears to be no direct correlations between the rates of reaction and the redox potentials of these complexes. Exploration of the oxidation chemistry of well-characterized copper complexes together with a detailed understanding of the function of O₂ activating copper enzymes is expected to provide the basis for new catalytic oxidation systems for synthetic and industrial processes.

Current understanding of the types of copper–dioxygen species or intermediates relevant to O₂-binding to or activation by copper proteins comes from the combination of biochemical/biophysical studies and coordination chemistry efforts, with the latter having played a significant role. Figure 2 on the next page provides a summary of structural types of nearly all now well established reactive intermediates of Cu-O₂ species. These are the starting point for describing the possible conformation of reactive intermediates based on copper chemistry. As seen by the large number of examples (Figure 2), many structural types exist which are perhaps far more than are known for heme or nonheme iron enzymes or complexes.

Synthetic bioinorganic copper(I)–O₂ chemistry has flourished in the last 25 years.¹¹,¹⁶,¹⁷⁸-¹⁸¹ These efforts to elucidate fundamental chemical aspects of Type-3 oxidases have been inspired by understanding of the diverse nature of the active-site chemistry of copper-protein O₂-carriers, monooxygenases, and oxidases.¹⁸²-¹⁸⁵ Enzymes with active sites having one, two, three, or four copper ions are known (Figure 1), and the
Figure 1.3. Structure of CuO$_2$ intermediates identified in biomimetic complexes.\textsuperscript{16}
variations in ligand environment and reactivity patterns are vast. The prototype for
dioxygen binding to copper, and subsequent the primary model for ligand design has
been the structure and reversible O₂-binding of arthropodal and molluscan blood oxygen
transporting protein, hemocyanin. The dicopper motif, also found in tyrosinase and
catechol oxidase, is of considerable interest with respect to the relationship of
dioxygen binding and activation. Formation of the copper-dioxygen adducts leads to
copper-mediated reduction of O₂ to superoxo (O₂⁻), peroxy (O₂⁻²), or O–O cleaved
products (copper-oxo, O⁻²), which constitute the active species responsible for substrate
oxidation. While there is ample literature for oxidative reactions by copper
complexes, it is only within the past few years and recent advances in low
temperature spectroscopy and reactivity that we have been able to assign a given organic
reaction to specific copper-O₂ derived species. In fact, a number of the CuO₂ species can
be placed into groups of reactivity types, for example as nucleophilic or electrophilic
reactive intermediates. Following is a brief summary of the known reactivity of
biomimetic complexes of Type-3 copper oxidases which is the basis for categorizations
of different CuO₂ intermediates responsible for reactivity.

The µ-η²:η² peroxo-dicopper(II) (Figure 1.2) complex was first elucidated with
x-ray crystallography in a synthetic system by K. Fujisawa and the late N. Kitajima, and subsequently confirmed for oxy-hemocyanin. This moiety was also detected in
tyrosinase and catechol oxidase with spectroscopic methods. It is likely, and now
generally accepted, that the µ-η²:η² peroxo-dicopper(II) core is closely involved in
aromatic hydroxylations, both in enzymes like tyrosinase and chemical systems. Notably,
through the research efforts of W.B. Tolman and co-workers there is an established high-valent bis-µ-oxo-dicopper(III) species,\textsuperscript{16,188} (Figure 1.2 O) that is thus far not found in biochemistry but only exists as an interesting example of how chemists can elucidate useful new chemical possibilities. A moiety such as this, or even containing more than two copper ions in a different overall redox state, is suggested to be responsible for the hydroxylation of methane in particulate methane monooxygenase (p-MMO), a membrane-bound copper-dependent enzyme found in methanogenic bacteria.\textsuperscript{192}

Mono and binuclear copper-hydroperoxo reactive intermediates are also known. Cu\textsuperscript{2+}-OOH and more recently Cu\textsuperscript{2+}-superoxide species, were suggested to be present in the copper-containing enzymes dopamine β-monooxygenase (DβM) and peptidylglycine α-hydroxylating monooxygenase (PHM).\textsuperscript{183,193,194} While DβM and PHM have two active-site copper ions, they are separated by a large distance (~11 Å) and the oxidative hydrogen-atom chemistry is thought to occur at the CuM site near the substrate binding pocket. The second metal center, dubbed CuH, has been suggested to assist the reaction by serving the role of an electron-transfer center to deliver the proper number of electrons to reduce O\textsubscript{2}. The chemistry of Cu-O\textsubscript{2} species is being pursued more heavily as of late and interesting new results from Klinman and co-workers will offer a new insight into both ligand design and mechanistic approaches to studying Type-3 oxidase systems.\textsuperscript{195} Binuclear peroxy-dicopper(II) complexes (Figure 1.2 TP and SP) were the first to be well characterized.\textsuperscript{16,190} In synthetic chemical systems, it is difficult to stabilize the principal reaction product of Cu\textsuperscript{+} and O\textsubscript{2} (Cu-O\textsubscript{2}), from coupling with another ligand-copper(I) complex to form a Cu\textsuperscript{2+}-(O\textsubscript{2}\textsuperscript{2−})-Cu\textsuperscript{2+} product.\textsuperscript{180} These compounds were originally
stabilized at low temperature (–80 °C) in organic solvents. Early examples are [Cu$_2^{2+}$ (XYLO$^-$)(O$_2$)]$^+$ which contained an asymmetrical end-on peroxo moiety, a µ-1,2-peroxo complex [{(TMPA)Cu$_{2+}$}$_2$(O$_2$)]$^{2+}$, (Figure 1.2 TP) and [Cu$_2^{2+}$ (N$_4$)(O$_2$)]$^{2+}$ (Figure 1.3). Several analogues of [{(TMPA)Cu$_{2+}$}$_2$(O$_2$)]$^{2+}$ and [Cu$_2^{2+}$ (N$_4$)(O$_2$)]$^{2+}$ have been characterized, showing variable stability and reactivity. These complexes were used to generate a reactivity profile based on the type of reactive intermediate. It was determined that [Cu$_2^{2+}$ (XYLO$^-$)]$^+$ reversibly binds O$_2$ and that the resulting peroxo complex is basic. This means that a catalyst that works through a nucleophilic substitution mechanism. It was also determined that the complex [Cu$_2^{2+}$ (XYLO$^-$)O$_2$]$^+$ does not oxygenate triphenylphosphine nor CO, but it releases O$_2$ and binds PPh$_3$ and CO. The peroxo complex however can be protonated to afford a µ-1,1-hydroperoxo complex, (Figure 1) or it can be acylated to form an analogous µ-1,1-acylperoxo complex. Further, [Cu$_2^{2+}$ (XYLO$^-$)(O$_2$)]$^+$ reacts as a nucleophile with CO$_2$, with the formation of a percarbonato species that thermally decomposes to the carbonato complex [Cu$_2^{2+}$ (XYLO$^-$)(CO$_3$)]$^+$. Phenols are not hydroxylated nor oxidatively coupled (forming a dimer product) by [Cu$_2^{2+}$ (XYLO$^-$)(O$_2$)]$^+$, but it was the first example of a model system in the area of Type-3 oxidase biomimetics to show reactivity consistent with general acid-general base chemistry. The same nucleophilic chemistry is observed in reactions of [{(TMPA)Cu$_{2+}$}$_2$(O$_2$)]$^{2+}$. With these preliminary results it became evident that the nature of the intermediate would determine reactivity.

By contrast, the µ-η$_2^-$:η$_2^+$ side-on peroxo complex [Cu$_2^{2+}$ (N$_4$)(O$_2$)]$^{2+}$ reacts differently with the same substrates tested for the end-on peroxo complexes yielding
O=PPh3 in a reaction with PPh3. Perhaps one of the simplest and most elegant example of a model compound for Type-3 oxidases was synthesized by Stack and co-workers; they showed that a side-on peroxo complex formed from a simple binucleating ligand, namely \([\{(L)Cu\}_2(O)_2\}]^{2+}\) (L=N,N-di-tert-butyl-N,N’-dimethylthelyenediamine Figure 1.4 B), also oxygenates PPh3. Further, \([Cu_2^{2+}(N_4)(O_2)]^{2+}\) does not protonate, acylate, nor react with CO2. The peroxo complex oxidizes phenols by H+ abstraction, leading to catalytic oxidative coupling chemistry under pre-equilibrium conditions (i.e. in the presence of excess phenol and dioxygen). Thus, a side-on peroxo complex is nonbasic or electrophilic in its reactivity toward these substrates. The electrophilic behavior of these model compounds has also been demonstrated in reactions of greater impact, as in hydroxylation of arenes (including phenols); this reactivity can have significant implications toward green chemistry, since phenols and chlorinated phenols are rather stable pollutants which can contaminate soil and water ways. Activation of the peroxo complex \([Cu_2^{2+}(XYLO^-)(O_2)]^+\) was also determined to be electrophilic with H+ or CO2, affording \([Cu_2^{2+}(XYLO^-)(O_2H)]^{2+}\) and the percarbonato species \([Cu_2^{2+}(XYLO^-)(CO_4)]^+\), respectively. These, in contrast to the original end-on peroxo complex \([Cu_2^{2+}(XYLO^-)(O_2)]^+\), readily convert PPh3 to O=PPh3. Thus, in biochemical systems, hydroperoxo-copper complexes are a possible reactive intermediate which may include percarbonato species, since CO2 as bicarbonate is certainly available in biological media. While carried out for a specific set of peroxo-dicopper(II) compounds, the trends observed in these early reactivity studies of reactive intermediate types in copper-dioxygen chemistry have generally been found to be consistent; however, there are some
exceptions.\textsuperscript{190,205,206} The end-on peroxo complexes with nucleophilic behavior have not yet shown oxidative chemistry comparable to the side-on complexes; however, Karlin and co-workers continue to pursue if this alternative structure can perform useful chemistry, such as in the Baeyer-Villiger oxidation (oxidation of ketones to carboxylic acids in the presence of peracids).

It is now known that binuclear copper enzymes such as tyrosinase and catechol oxidase adopt the side-on $\mu$-$\eta^2$-$\eta^2$-peroxo structure, as do model compounds like the Kitajima/Fujisawa complex, the Karlin complex [Cu$_2^{2+}$(N$_4$)(O$_2$)]$^{2+}$ and several other analogues.\textsuperscript{16} The great number of investigations in copper(I)–dioxygen reactivity have also revealed a novel binding motif in which the peroxo O–O bond is fully cleaved, after receiving 2 electrons from di-Cu$^{2+}$, resulting in a high-valent dicopper(III) bis-$\mu$-oxo (Cu$_2^{3+}$-O$_2$) species (Figure 1.2 O), first characterized by Tolman’s group,\textsuperscript{207} using tridentate substituted triazacyclononane as ligands, and also by Stack and co-workers,\textsuperscript{208,188} using ethylenediamine-based donors.\textsuperscript{16,188} The two isoelectronic species have distinct spectroscopic features and structural differences. The main structural difference is the shortened Cu–Cu distance of ~2.8 Å for the bis-$\mu$-oxo-dicopper(III) species when compared to the side-on peroxo-dicopper(II) Cu–Cu distance of ~3.5 Å. Theoretical calculations have predicted that there is a small difference in the free energies of the side-on peroxo Cu$_2^{2+}$-O$_2$ and the Cu$_2^{3+}$-O$_2$ species and a low barrier for interconversion,\textsuperscript{209-211} meaning that under experimental conditions (ambient temperature) the two forms are under equilibrium. This is experimentally observed in several systems
Figure 1.4. Early examples of Type-3 oxidase models by Karlin.\textsuperscript{196-198} and Tolman.\textsuperscript{207}
where the side-on peroxo-dicopper(II) and bis-µ-oxo-dicopper(III) species are found in a dynamic equilibrium, strongly influenced by the nature of the ligand such as denticity, N-alkyl vs. N-pyridyl donor, and steric effect, as well as reaction conditions, including concentration, counter ion effect, temperature, and solvent. These well-characterized Cu2O2-containing synthetic complexes are quite reactive, with oxidative capabilities ranging from alcohol/catechol oxidation, oxo transfer to phosphines and sulfides, to aromatic/aliphatic hydroxylations, and also oxidative N-dealkylations. The existence of two oxidative species coexisting in equilibrium complicates the determination of which form is the true reactive intermediate responsible for the reactivity. A brief summary and comparison of Karlin’s and Tolman’s efforts is discussed below, highlighting preferential reactivity patterns of side-on peroxo-dicopper(II) and bis-µ-oxo-dicopper(III) species.

Karlin’s group elucidated several examples of copper-promoted dioxygen activation. While stoichiometric in nature, this monooxygenase model system is a rare chemical system where an unactivated C–H bond substrate is rapidly hydroxylated under “very mild” conditions utilizing dioxygen (−80 °C and ≤1 atm O2). The dicopper(I) complex [Cu2+(XYL-H)]2+ reacts with O2 to form [Cu22+(XYL–)(OH)]2+ stoichiometrically. Isotope labeling experiments (using 18O) demonstrate the product phenol O-atom is derived from dioxygen, and the observed O2-consumption stoichiometry (Cu:O2 = 2:1) is consistent with monooxygenases. Detailed low temperature stopped-flow kinetic studies revealed the intermediate
[Cu$_2$]$^{2+}$(XYL-H)(O$_2$)]$^{2+}$. Its peroxo structure was confirmed by utilizing the electron withdrawing R=NO$_2$ derivative, which results in a decreased $k_{on}$ but even more diminished first order rate constant for the hydroxylation step, thus allowing for spectroscopic investigation. In resonance Raman studies, characteristics of [Cu$_2$]$^{2+}$ (NO$_2$-XYL-H)(O$_2$)]$^{2+}$ are consistent with a $\mu$-η$_2$-$\eta^-$-peroxo moiety, with the O–O vibrational stretch at 747 cm$^{-1}$. Analysis of spectroscopic and kinetic studies indicates that the side-on peroxo moiety is the reactive species responsible for oxygentation, supported by the observation that the O–O stretching disappears at the same rate as the phenolic C–O bond stretch at 1320 cm$^{-1}$ in [Cu$_2$]$^{2+}$(NO$_2$-XYLO–)(OH)]$^{2+}$. A number of results support the $m$-xylyl hydroxylation reaction where [Cu$_2$]$^{2+}$(R-XYL-H)(O$_2$)]$^{2+}$ acts as an electrophile, attacking the $\pi$-system of the arene substrate. Side-on peroxo complexes [Cu$_2$]$^{2+}$(R-XYL-H)(O$_2$)]$^{2+}$ are very similar to [Cu$_2$]$^{2+}$(N$_4$)(O$_2$)]$^{2+}$, shown to be electrophilic. Stopped-flow kinetic studies showed that rate constant for hydroxylation increases as the R substituent becomes more electron donating (Hammett plot $\rho = -2.1$), with deuterium substitution into the 2-H position of the substrate not affecting the first order rate constant for hydroxylation. The lack of a deuterium kinetic isotope effect is consistent with an electrophilic attack on the arene substrate $\pi$-system, precluding C–H bond cleavage in the rate-determining step. Methyl substitution into the xylyl 2-position where hydroxylation occurs for the parent compound [Cu$_2$]$^{2+}$(XYL-CH$_3$)]$^{2+}$ in a reaction with O$_2$, results in aromatic hydroxylation followed by a 1,2-migration of the methyl group. The process is similar to the “NIH shift”, observed previously in iron hydroxylases via isotope labeling, where a carbonium
on intermediate is obtained as a result of an electrophilic iron-oxo reactive intermediate, resulting in R migration.\textsuperscript{217} Computational studies indicate that the side-on peroxo moiety is indeed electrophilic; the xylyl hydroxylation reaction can be seen as derived from interactions of the empty $\pi^*/\sigma^*$ orbitals of peroxo O=O with the filled $\sigma$ orbitals in the arene substrate.\textsuperscript{216,218} The favorable proximity of the side-on peroxo moiety to the m-xylyl substrate, due to the ligand design and resulting intramolecular O$_2$ reaction with two copper(I) centers, is certainly important in promoting the aromatic hydroxylation reaction. In this sense, the system resembles reactions at enzyme active sites, where pre-organization of the substrate and reactive species occur.

Since the discovery and characterization of the model monooxygenase system $[\text{Cu}_2^{+}(\text{XYL-H})]^{2+}+\text{O}_2 \rightarrow [\text{Cu}_2^{2+}(\text{XYLO}^-)(\text{OH})]^{2+}$, a good number of analogous systems have been described.\textsuperscript{11,179} Tolman and co-workers,\textsuperscript{206} however, have shown that a bis-$\mu$-oxo-dicopper(III) complex is also capable of aromatic hydroxylation. This observation is opposite in trend to the general reactivity patterns established thus far, serving to demonstrate the general points that placing a reactive intermediate species in close proximity to a substrate by ligand design greatly enhances the probability of turnover, and that different types of intermediates will promote certain placements thus promoting different reactivities. Itoh and co-workers\textsuperscript{219} have demonstrated the importance of the side-on peroxo-dicopper(II) species as relevant to phenol $o$-hydroxylation both in synthetic systems and in the enzyme tyrosinase.\textsuperscript{220} Cresolase activity ($o$-hydroxylation of phenols, i.e cresol) was successfully mimicked with a Cu$^+$ complex possessing a tridentate ligand $N,N$-bis[2-(2-pyridyl)-
ethyl]-R,R-dideuteriobenzylamine (L^{Py2Bz}, Figure 1.4 A) that binds Cu$^+$ and reacts with O$_2$ to generate a low-temperature stable side-on peroxo-dicopper(II) species.$^{219,220}$ Addition of lithium salts of $p$-substituted phenols gives the corresponding $o$-hydroxylated catechols in high yields. Neither oxidation to the $o$-quinone nor C–C/C–O coupled dimerization was observed. With the use of $^{18}$O isotope labeling experiments, it was confirmed that the incorporated catechol oxygen atom is derived from O$_2$. Mechanistic studies revealed substrate saturation kinetics, consistent of formation of a phenolate/Cu$_2$-O$_2$ complex prior to the rate-determining oxygenation step. An observed increase in reaction rate with increasing electron donating ability of the X substituent ($p$-XC$_6$H$_4$O-Li) coupled with the lack of a deuterium kinetic isotope effect, suggests the side-on peroxo species reacts via an electrophilic aromatic substitution mechanism,$^{219}$ similar to that described for the Karlin system,$^{179}$ the dicopper-mediated $m$-xylyl aromatic hydroxylation. To compare and contrast, Itoh and co-workers$^{220}$ studied the analogous chemistry on tyrosinase. As it was also seen in the model reaction, oxygenation rates increased with increasing $p$-substituted phenol electron donating properties, clearly demonstrating that the enzyme acts as an electrophilic reactive intermediate. Hammett plots of the enzymatic reaction give a $\rho$ value of $–2.4$, similar to the model reaction ($\rho = –1.8$) and Karlin’s XYL hydroxylation ($\rho = –2.1$),$^{179}$ supporting the hypothesis that the phenolase activity of tyrosinase occurs by an electrophilic attack by a $\mu$-$\eta^2$:$\eta^2$-peroxo-dicopper(II) species.

Casella and co-workers$^{221,222}$ and Sayre and coworkers$^{223,224}$ have also contributed in the area of $o$-phenol hydroxylation chemistry and their papers should be noted.
Consistent with the conclusions described above, Casella’s group has also concluded that the side-on $\mu$-$\eta^2: \eta^2$-peroxo species is critical in this chemistry. Stack and co-workers showed that a side-on peroxo-dicopper(II) complex with a secondary amine ligand, N,N-di-tert-butylethylenediamine (Figure 1.4 B), also exhibits some phenolase and catecholase activity, producing mixtures of catechol and $o$-benzoquinone from 2,4-di-tert-butylphenolate as substrate. With addition of the neutral phenol, no oxidatively coupled dimer is produced, contrary to Karlin’s complex, $[\text{Cu}_2^{2+}(\text{N}_4)(\text{O}_2)]^{2+}$. Stack’s side-on peroxo complex is also able to perform other electrophilic types of reactions, converting PPh$_3$ to O=PPh$_3$, catechols to quinones, and benzyl alcohol and benzylamine to benzaldehyde and benzonitrile, respectively.

Bis-$\mu$-oxo-dicopper(III) species have been implicated in a number of different reactivities, including alcohol oxidation, phenol coupling, oxidative N-dealkylation, aliphatic hydroxylation, and oxygenation of phosphines or sulfides. Detailed mechanistic studies and theoretical calculations suggested that the bis-$\mu$-oxo core facilitates hydrogen-atom abstraction reactions. Tolman and co-workers studied intramolecular N-dealkylation reactions by bis-$\mu$-oxo-dicopper(III) complexes. The electrophilic nature of the bis-$\mu$-oxo core was elucidated by its lack of reactivity towards acids, while it can only act as an outer-sphere reactive intermediate in the presence of acid. The bis-$\mu$-oxo complex decomposes by oxidative N-deakylation, with one of the arms of the tri-substituted triazacyclononane ligand being cleaved, resulting in production of the primary amine and the aldehyde/ketone. Double crossover experiments using labeled $^{18}$O$_2$ confirm the carbonyl oxygen derives from the high-valent Cu$_2^{3+}$-O$_2$. 

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Figure 1.5. Type-3 model systems by Itoh\textsuperscript{219} (A) and Stack\textsuperscript{204} (B).
core and that the reaction is intramolecular. Primary kinetic isotope effects and Eyring activation parameters revealed that C–H bond cleavage is the rate-determining step. A Hammett study with derivatives of tribenzyl-substituted triazacyclononane ligands demonstrates that reaction rates increase with electron-donating substituents. A small observed negative $\rho$ value of $-0.4$ suggests the Cu$_2^{3+}$-O$_2$ core behaves as an electrophilic radical; similar $\rho$ values were reported for benzylic hydrogen-atom abstraction reactions using free radical reactive intermediates, as well as cytochrome P450 and porphyrin-derived high-valent metal-oxo species. Bis-µ-oxo-dicopper(III) complexes with pyridylalkylamine ligands have been used by Itoh et al. to model benzylic hydroxylation chemistry similar to dopamine-β-monooxygenase (DbM) activity. Their experiments using $^{18}$O$_2$ confirmed oxygen atom incorporation into the oxygenated alcohol, which in the experiment is trapped by coordinating to copper(II). Deuteration at benzylic positions revealed a large kinetic isotope effect of 35.4 at $-80 \, ^\circ$C. The activation parameters are consistent with intramolecular rate-determining hydrogen atom abstraction followed by rebound of a copper-bound OH group, analogous to cytochrome P450 monooxygenase chemistry. A Hammett plot based on experiments with varying substituents at the para position gives $\rho = -1.48$, consistent with benzylic hydrogen atom abstraction reactions. With the tridentate ligand L$^{Py2Bz}$, benzylic hydroxylation was also observed, but spectroscopic evidence, reveals the low-temperature dioxygen adduct formed is a side-on $\mu$-$\eta^2$-$\eta^2$-peroxo-dicopper(II) species. For this system, the deuterium kinetic isotope effects are much smaller than expected for a rate-determining hydrogen atom abstraction step, and the rate of hydroxylation is not affected by the $p$-substituent of
the ligand. Instead, the rate of reaction was found to be solvent dependent. From the combined observations, it was proposed that cleavage of the O–O bond to form the high-valent copper(III) bis-µ-oxo core was rate-limiting, which is then responsible for the hydrogen atom abstraction. Karlin and co-workers showed that copper(I)/O₂ reactivity with tridentate ligands give rise to mixtures of side-on peroxo-dicopper(II) and bis-µ-oxo-dicopper(III) complexes, in rapid equilibrium at –80 °C. These showed oxidative chemistry towards different substrates. With respect to dimethylaniline oxidative N-dealkylation, mechanistic studies using p-substituted and deuterated dimethylanilines suggest the reactive intermediate can operate via either a rate-limiting hydrogen-atom abstraction or electron-transfer pathway, depending on the R group in the complex [{Cu(R-Me-PY₂)}₂(O₂)]²⁺ and the ease of the substrate oxidation.

Itoh and co-workers have been successful in distinguishing the active intermediate for the radical coupling of neutral phenols. According to their spectroscopic features, the Cu₂O₂ adducts are the bis-µ-oxodicopper(III) and the side-on peroxo-dicopper(II) species (Figure 1.2), respectively. Both isomeric species react with neutral phenols to produce solely the C–C coupled dimers. Both Cu₂³⁺-(O)₂ and Cu₂²⁺-(O₂) species exhibit this same behavior. However, the rate constant for the reaction of the bis-µ-oxo species was two orders of magnitude greater than that for the peroxo complex. This could be due to two reasons: either the bis-µ-oxo is intrinsically a better reactive intermediate than the side-on peroxo because of a difference in redox potentials, or the (LPY₁)₂Cu₂³⁺-(O₂)₂ bis-µ-oxo complex is the true reactive intermediate, which would be in much higher concentration than a small amount of bis-µ-oxo isomer present in
equilibrium with the side-on complex \((L^{\text{Py2Bz}})_2\text{Cu}_2^{2+}\text{-(O}_2\)). The hypothesis that the bis-µ-oxo is the true reactive species is favored because if the side-on peroxo complex was able to couple phenols via the proton-coupled electron transfer mechanism, then the reaction with phenolates should also occur through an outer-sphere electron transfer mechanism to give the coupled phenol. However, this contradicts previous observations that the µ-\(\eta^2:\eta^2\)-peroxo-dicopper(II) complex reacts with phenolates to yield solely o-catechol products.

In the chemistry described for hydroperoxodicopper(II) (Figure 1.2 µ-1,1 hydroperoxo), enhanced electrophilic reactivity occurs due to protonation of less reactive and nucleophilic peroxo-dicopper(II) complex, at least with respect to triphenylphosphine oxygenation. While the latter reaction is not of great importance due to the ease of PPh\(_3\) oxygenation, protonation of peroxo-copper or other O\(_2\)-derived species may be important as a means of peroxo-copper activation\(^{233-235}\). In heme enzymes such as cytochrome P450 monooxygenase, protonation of a heme-peroxo leads to O–O heterolytic cleavage and generation of the high-valent ferryl species responsible for electrophilic oxidation reactions like C–H hydroxylation or epoxidation\(^{236}\). Activation to an alternate reactive intermediate appears to be present in heme oxygenase, where an O\(_2\)-derived heme-hydroperoxo species attacks the porphyrin meso-carbon substrate\(^{237,238}\). More unique examples of reactivity for binuclear hydroperoxo complexes which do not have the ArO\(^-\) bridging ligands have been reported\(^{239}\). Although structural descriptions are not as readily available, kinetic and/or spectroscopic probing shows such Cu\(^{2+}\text{-(–O}_2\text{H)-Cu}_2^{2+}\) intermediates are capable of arene hydroxylation. Thus, Karlin, Zuberbühler, and co-
workers\textsuperscript{239} report that reaction of a dicopper(II) complex with ligand XYL-H and hydrogen peroxide also leads to efficient arene hydroxylation. It is likely, according to reactivity, that a μ-η\textsuperscript{1}:η\textsuperscript{1}-OOH hydroperoxo-dicopper(II) species facilitates arene hydroxylation. Casella and co-workers\textsuperscript{240} reported a compound capable of performing double arene hydroxylation, again in a dicopper(II) plus hydrogen peroxide reaction; not only were kinetic parameters determined for the kinetics of substrate attack by an η\textsuperscript{1}:η\textsuperscript{1}-hydroperoxo-dicopper(II) intermediate, but the latter was also characterized by UV-Vis and EPR spectroscopies. Thus, dicopper(II)-hydroperoxo species, with or without an additional phenoxy bridging ligand, which could, be replaced by a water or hydroxo ligand in solution,\textsuperscript{240} can effect oxidation reactions, and further exploration of their chemistry should be considered.

The work involved in ligand design and low-temperature techniques leads to stable enough species to be characterized spectroscopically, but may prevent efficient reactivity, especially for hydrocarbon-based substrates. Thus, there is a general caveat in concluding that an observed organic substrate oxidative reaction actually occurs from a well-defined and fairly stable Cu\textsubscript{n}-O\textsubscript{2}(H) complex. The true reactive intermediate may be something other than what can be observed spectroscopically. It is possible that a transient mononuclear Cu\textsuperscript{3+}-oxide↔Cu\textsuperscript{2+}-O•\textsuperscript{−} species is the most likely candidate for the reaction of Type-3 oxidases both in the native proteins and their biomimetic models.\textsuperscript{188}
VII. CONCLUDING REMARKS

Considerable effort in trying to elucidate the nature of the cause of neurodegeneration in AD and the catalytic mechanism through which Type-3 copper oxidases catalyze their chemical transformations has yielded some common threads. First, it is known that oxygen radicals and reduced forms of oxygen collectively termed ROS are generated during normal metabolism. It is also known that the amyloid-β fibrils and plaques can generate those same ROS via a metal-centered mechanism. These activated forms of O₂ are no different than the metal-bound forms associated with Type-3 copper oxidases. It is expected that further analysis of metal complexes of amyloid-β can show that the chemistry proposed as the culprit in the etiology of AD is very much consistent with the reactivity of Type-3 copper oxidases and their biomimetic systems.
VI. LIST OF REFERENCES


49) K. Chaudrasekaran, J. Stoll, D.R. Brady, S.I. Rapoport, Localization of cytochrome oxidase (COX) activity and COX mRNA in the hippocampus and entorhinal cortex of


65) P.J. Benke, H. Levcovitz, J. Paupe, E. Tozman, Scavengers of free radical oxygen affect the generation of low molecular weight DNA in stimulated lymphocytes from


93) G.F.Z. da Silva, W.M. Tay, and L.-J. Ming, Catechol Oxidase-like Oxidation Chemistry of the 1–20 and 1–16 Fragments of Alzheimer's Disease-related β-


Irreversible Reduction of Dioxygen by Simple Peralkylated Diamine-Copper(I)

209) C.J. Cramer, B.A. Smith, W.B. Tolman, Ab Initio Characterization of the Isomerism
between the \( \mu-\eta^2:\eta^2 \)-Peroxo- and Bis(\( \mu \)-oxo)dicopper Cores. *J. Am. Chem. Soc.*

210) P. Spuhler, M.C. Holthausen, Mechanism of the aliphatic hydroxylation mediated
by a bis(\( \mu \)-oxo)dicopper(III) complex. *Angew. Chem. Int. Ed.* (2003), 42, 5961–
5965.

and Electronic Structural Studies of the Cu(III)$_2$ Bis-\( \mu \)-oxo Core and Its Relation to

212) K.D. Karlin, P.L. Dahlstrom, S.N. Cozzette, P.M. Scensny, J. Zubieta, Activation of
oxygen by a binuclear copper(I) compound. Hydroxylation of a new xylyl
binucleating ligand to produce a phenoxy-bridged binuclear copper(II) complex; x-
ray crystal structure of \( [\text{Cu}_2\{\text{OC}_6\text{H}_3[\text{CH}_2\text{N}(\text{CH}_2\text{CH}_2\text{py})_2]_2-2,6\}'(\text{OMe})]\) (py = 2-

213) K.D. Karlin, Y. Gultneh, J.C. Hayes, R.W. Cruse, J.W. McKown, J.P. Hutchinson,
J. Zubieta, Copper-mediated hydroxylation of an arene: model system for the
action of copper monooxygenases. Structures of a binuclear copper(I) complex and


231) J. Shearer, C.X. Zhang, L.Q. Hatcher, K.D. Karlin, Distinguishing Rate-Limiting Electron versus H-Atom Transfers in Cu$_2$(O$_2$)-Mediated Oxidative N-Dealkylations:


CHAPTER II. Cu^{2+}-AMYLOID-β COMPLEXES AS A REDOX-ACTIVE CATALYST TOWARD THE OXIDATION OF 1,2,3-TRIHYDROXY BENZENE‡

I. INTRODUCTION

Abnormal metal-ion homeostasis has been closely associated with several neurodegenerative diseases, including Parkinson’s, amyotrophic lateral sclerosis, Creutzfeldt-Jakob disease (i.e. mad cow disease), and Alzheimer’s disease (AD).1–3 Because high cytoplasmic concentrations of free metal ions are toxic and potentially lethal, intricate physiological pathways have evolved to transport and distribute metal ions to their targets which include enzymes and proteins.4 With aging, physiological processes responsible for accurate delivery of metal ions break down and “leakage” of free metal ions can cause toxic effects to cells.5,6 Divalent ions of redox-active transition metals have often been associated with oxidative stress and closely involved in the

chemistry of reactive oxygen species (ROS), including hydrogen peroxide and superoxide and hydroxyl radicals.\textsuperscript{7} Because increases in intracellular concentrations of metal ions is closely related to the effects of aging, oxidative stress, and AD, there is considerable interest in investigating the connection between malfunction of regulatory processes such as metal transport and the presence of ROS with the pathology of AD.

The chemistry of redox-active metal complexes of $\beta$-amyloid peptide (A$\beta$) has been an area of intense focus in the study of AD. The aggregation of A$\beta$ within the neocortex is closely related to the pathology of AD and has been shown to be induced by metal binding.\textsuperscript{8,9} The A$\beta$ peptides are generated by the cleavage of the ubiquitous amyloid precursor protein (APP) by $\alpha$, $\beta$, and $\gamma$ secretases.\textsuperscript{10} A$\beta$ in the form of insoluble plaques contains up to mM amounts of Zn$^{2+}$, Cu$^{2+}$, and Fe$^{3+}$ in the neocortical region of the brain;\textsuperscript{7} however, the cause/effect connection of the metallo-A$\beta$ plaques with AD is still under debate.\textsuperscript{11} The metal coordination environment of the 1–40 and 1–42 peptides has been previously studied and their pH dependent aggregation reported.\textsuperscript{9,12} The results showed the metal binding seemed to be non-stoichiometric with approximately 3.5 metal ions per pair of aggregated peptides and a cooperative binding pattern as the amount of aggregates increases.\textsuperscript{7} Since A$\beta_{1-42}$ and A$\beta_{1-40}$ have been shown to bind Zn$^{2+}$ and Fe$^{3+}$ with nM and Cu$^{2+}$ with aM apparent dissociation constants by means of quantitative determination of the metal-complex precipitates,\textsuperscript{7} understanding of the metal-binding domain and its structure may shed light on the chemistry related to the neuropathology of AD.
Though the coagulation of the peptide plaques leaves little doubt that interaction with cytoplasmic molecules is unlikely, smaller fragments of the amyloid peptide are soluble and Aβ fibrils extend across membranes, exposing them to the cytoplasm. Recently, insulin degrading enzyme (IDE) has been shown to digest the longer Aβ peptides (40–42 amino acids) into smaller soluble fragments.\textsuperscript{13} Moreover, the cleavage of APP by α and β- secretases produces the Aβ\textsubscript{1-16} fragment of APP.\textsuperscript{14} These soluble fragments and intramembrane spanning fibrils still possess possible metal binding sites such as histidines, glutamate, aspartate, and tyrosine within the 1-20 fragment of Aβ in the sequence DAEFR\textsubscript{5} HDSGY\textsubscript{10} EVHN\textsubscript{15} KLVFF\textsubscript{20}. Redox chemistry of Aβ has been previously reported, wherein Met\textsubscript{35} was suggested as a “built-in” reducing agent required for the redox cycling hypothesis.\textsuperscript{15} The lack of sufficient data on the redox chemistry of and the oxidative stress caused by metallo-Aβ and the discrepancies in previous studies such as the presence or absence of free-radicals and the nature of the metal-Aβ interaction seem often to be the shortcoming in Aβ research. Understanding of the chemical processes associated with metallo-Aβ may provide insight into the upstream and/or downstream regulatory processes that lead to AD. Herein, we describe the oxidation chemistry of CuAβ in the presence and absence of H\textsubscript{2}O\textsubscript{2}, showing conclusive metal-centered pre-equilibrium kinetics toward the oxidation of a simple substrate and the oxidative cleavage of double-stranded plasmid DNA.
II. EXPERIMENTAL

The 1-20 and 1-16 fragments of Aβ were purchased from Aldrich (St. Louis, MO) or synthesized at the Peptide Center of the University of South Florida. The identities of the peptides have been confirmed with a Bruker MALDI-TOF mass spectrometer. The substrate 1,2,3-trihydroxybenzene (THB) was obtained from Sigma-Aldrich, 3-methyl-2-benzothiazolinone hydrazone hydrochloride monohydrate from Acros (Fairlawn, NJ), the plasmid pQE30Xa from Quiagen (Valencia, CA), EDTA, DMSO, mannitol, H2O2, Cu(NO3)2, ZnCl2, and Ni(SO4) from Fisher (Swanee, GA), and CoCl2 from Mallickrodt (Paris, KY). All plastic ware was demetallized with EDTA and extensively rinsed with 18.0-MΩ water to remove the chelator. The water used for the studies of DNA cleavage was autoclaved to remove ubiquitous nucleases.

DNA cleavage assay: The metal derivatives of Aβ were prepared by dissolving the peptide in 18.0-MΩ water and separated into aliquots followed by addition of corresponding metal ions at 1:1 stoichiometric ratio, which was further diluted into aliquots of working concentrations. The metal-complexes were freshly prepared in all experiments. A typical reaction toward DNA cleavage contained 150 ng of plasmid DNA, 4.0%, 2.0%, or 0.2% H2O2, and 5.0 µM of metallo-Aβ derivatives in 100 mM HEPES at pH 7.00 in a volume of 15.0 µL. A time-course experiment was performed and analyzed in a 1.0 % agarose gel stained with ethydium bromide, and then photographed on a transilluminator.

THB oxidation assay: Several concentrations of THB ranging from 0.10 to 5.0 mM were incubated with 7.5 µM of CuAβ and 1.60, 3.20, 16.20, 32.3 or 70.0 mM H2O2 and
buffered with 100 mM HEPES at pH 7.00 in a final volume of 1.0 ml. The formation of product was monitored at 420 nm ($\epsilon = 4,583 \text{ M}^{-1} \text{cm}^{-1}$) on a Varian CARY50 Bio-UV-Vis spectrophotometer for 5 minutes, and the rates determined by the change in absorbance over time. The background oxidation of THB was conducted in the same manner without CuAβ in the assay solution. Rates were fitted to appropriate rate laws and rate constants determined by the use of SigmaPlot 8.0.

The dependence of H$_2$O$_2$ on THB oxidation by CuAβ was determined by measuring the oxidation rate at several different concentrations of hydrogen peroxide with saturating concentration of THB (6.0 mM). The initial rates were determined and then fitted as a function of [H$_2$O$_2$] to an appropriate rate law to reveal the rate constant. Alternatively, the catechol oxidase assay was performed as previously reported with minor changes to fit current studies.¹⁶ Same molar concentrations of THB and 3-methyl-2-benzothiazolinone hydrazone (which serves as an ortho-quinone indicator) were mixed in 100 mM HEPES at pH 7.00 in the presence of 3.5 µM CuAβ. The red-adduct of the ortho-quinone product was monitored at 500 nm ($\epsilon = 32,500 \text{ M}^{-1} \text{cm}^{-1}$) and rates calculated. Auto-oxidation of THB was determined under the same conditions without Cu-Aβ.

*Metal Titration:* Apo-Aβ was dissolved in 100 mM HEPES at pH 7.00 to a final concentration of 1.0 mM. Cu$^{2+}$ binding was monitored by titrating the metal into the apo-Aβ solution and the electronic spectra collected after each addition of the metal. Cu$^{2+}$ binding was also determined through the activity of CuAβ complex toward THB. In this case, Cu$^{2+}$ was titrated into a fixed amount of the peptide in 100 mM HEPES at pH 7.00
in the presence of 10.0 mM of THB and 70.0 mM H₂O₂. The oxidation rates were
determined as a function of [Cu²⁺], then fitted to a simple equilibrium of metal:peptide =
1:1 or a cooperative binding pattern using the Hill equation.

NMR Spectroscopy: All the NMR spectra were acquired on a Bruker DPX250
spectrometer at ¹H resonance of 250 MHz. The metal-binding was monitored through the
changes in the NMR spectra. The peptides Aβ₁₋₁₆ and Aβ₁₋₂₀ and the paramagnetic shift
reagent Co²⁺ were prepared in d⁶-DMSO. The metal ion was gradually titrated into the
peptide and the paramagnetically shifted ¹H NMR signals detected. A typical spectrum
showing the paramagnetically shifted ¹H NMR signals consists of ~80,000 transients
from accumulation of several spectra of 10,000–20,000 transients with a recycle time of
~50 ms and a spectral window of ~250 ppm. Solvent exchangeable signals were
determined by adding a drop of D₂O into the sample which disappear after the addition.

Molecular mechanics: The primary sequence of Aβ₁₋₁₆ peptide was entered into
BioCAChE 6.0 (Fujitsu, Beaverton, Oregon) and the energy of its structure under
solvation using a simulated water droplet was minimized with the MM3 molecular
mechanics method. Histidine side chains were considered the ligands in the calculations
on the basis of the NMR data.

III. RESULTS AND DISCUSSION

Oxidative Double-Stranded DNA Cleavage: Though the bulk of the amyloid plaques
in AD brain is membrane bound, proteolytic processing of amyloid has been shown to
yield soluble fragments.¹² It has also been demonstrated that the neuropathology of AD
may directly affect DNA, eventually leading to apoptosis. As metal ions are involved in the formation of amyloid plaques, the oxidative activity of metallo-Aβ complexes against plasmid DNA was probed in vitro with gel electrophoresis. The oxidative activities of the Zn$^{2+}$, Ni$^{2+}$, and Cu$^{2+}$ complexes of Aβ$_{1-20}$ (ZnAβ$_{1-20}$, NiAβ$_{1-20}$, and CuAβ$_{1-20}$) toward the cleavage of plasmid DNA were determined by incubating several different concentrations of the complexes with plasmid DNA in the presence of 4.0 % H$_2$O$_2$ at room temperature for 30 minutes (Figure 2.1). Here, ZnAβ$_{1-20}$ serves as the control since Zn$^{2+}$ is oxidative inactive. The plasmid in the presence of metallo-Aβ$_{1-20}$ at lower concentrations shows a middle band that is not present in the reference (Lane R, Figure 2.1). Comparing the middle band with the DNA ladder gives an approximate size of 3.5 kbp, consistent with the size of linearized plasmid from the manufacturer. The activities of the derivatives follow the trend CuAβ$_{1-20}$ > NiAβ$_{1-20}$ > ZnAβ$_{1-20}$, demonstrating the involvement of metal in the oxidative cleavage of dsDNA.

One interesting result is shown in the H$_2$O$_2$ + DNA and the H$_2$O$_2$ + DNA + Aβ$_{1-20}$ control experiments. The 4.0 % H$_2$O$_2$ shows a significant damage toward plasmid dsDNA (Lanes 1, Figure 2.1), whereas Aβ$_{1-20}$ decreases the H$_2$O$_2$ damage of plasmid dsDNA and perhaps acts as a scavenger of ROS species (Lane 2, Figure 2.1). The role of Aβ as an antioxidant has been previously reported, wherein the presence of Met$_{35}$ was proposed to prevent lipid peroxidation while the M35L mutant showed reduced antioxidant activity. In a similar study, Aβ$_{1-40}$ was found to prevent the oxidation of the lipoproteins from cerebral spinal fluid and plasma. Moreover, Aβ$_{1-42}$ was shown to exhibit an
**Figure 2.1.** Concentration and metal-dependent assay of dsDNA cleavage. R is the reference plasmid DNA which shows supercoiled band (bottom) and nicked circular band (top); lane 1, DNA + 4% H$_2$O$_2$; lane 2, DNA + apo-A$\beta_{1-20}$ (200 µM) + 4% H$_2$O$_2$; lanes 3–6, ZnA$\beta_{1-20}$ (40, 80, 100, and 200 µM, respectively) + 4% H$_2$O$_2$; lanes 7–13, NiA$\beta_{1-20}$ (5, 10, 20, 40, 80, 100, and 200 µM, respectively) + 4% H$_2$O$_2$; CuA$\beta_{1-20}$ (at 5, 10, 20, 40, 80, 100, and 200 µM) + 4% H$_2$O$_2$ shows complete DNA cleavage (“blank” gel, not shown). All assays were incubated for 30 minutes. The middle band is the linearized plasmid detected at 3.5 kbp position, consistent with that obtained from the sequence of the plasmid pQE30Xa (Qiagen). Lanes 1’–6’ show the plasmid cleavage by Cu$^{2+}$ ions (5 µM) + 3.6% H$_2$O$_2$ at 10, 20, 40, 60, 90, 120 min. The standard DNA ladder starts with 1 kbp from the bottom with 1 kbp increment upward.
antioxidant activity more effective than ascorbate in cerebral spinal fluid. The antioxidant activity of Aβ was also demonstrated in the decrease of cytoplasmic amounts of 8-hydroxyguanosine, a major product of nucleic acid oxidation present in elevated amounts in the brains of AD patients. These observations implied that the production of Aβ could be related to prevention of oxidative stress. We have demonstrated here that even shorter fragments of Aβ without a Met can serve as a protective agent against oxidative damage of DNA, corroborating with some previous reports and supporting the hypothesis that apo-Aβ might be an effect of the oxidative stress in AD brains and might serve a specific purpose to scavenge ROS. This antioxidant activity is also observed in all concentrations of ZnAβ1-20 (Lanes 3–6, Figure 2.1), consistent with the lack of redox chemistry of Zn2+ and a protection role against dsDNA cleavage as in the case of apo-Aβ1-20.

Although Ni2+ is redox active and some of its complexes have been shown to exhibit oxidative damage toward DNA, NiAβ1-20 does not show such “chemical nuclease” activity, probably attributed to its low redox potential. Conversely, like apo-Aβ and ZnAβ1-20 discussed above, NiAβ1-20 shows a concentration-dependent protection against oxidative damage of dsDNA by H2O2, with better protection at higher concentrations while no significant protection at [NiAβ1-20] < 80 µM (Lanes 7–12, Figure 2.1). NiAβ has not been shown to be associated with AD pathology; however, it may serve as a structural and mechanistic probe in future studies of metallo-Aβ or similar systems.

The activity of 5.0 µM CuAβ1-20 (Lane 14, Figure 2.1) is exceedingly higher than that of ZnAβ1-20 at all concentrations tested in the presence of 4.0% H2O2 (Lanes 3–6, Figure
2.1) which effectively oxidizes the entire dsDNA plasmid sample into fragments that are too small to be resolved with the agarose gel electrophoresis (empty lanes not shown). Cu$^{2+}$ ion has been demonstrated in the literature to be active toward DNA cleavage in the presence of H$_2$O$_2$. The use of Cu$^{2+}$ (5.0 µM) in the presence of 3.6% H$_2$O$_2$ as control shows a much slower cleavage rate as the plasmid is not completely digested after 2 hours of incubation. In order to monitor the catalytic activity of CuAβ$_1$-20 toward plasmid dsDNA, the concentration of H$_2$O$_2$ was reduced to 2.0% and a time course experiment conducted (Figure 2.2). CuAβ$_1$-20 completely oxidizes plasmid DNA within 5.0 minutes in the presence of 2.0% H$_2$O$_2$, leaving only a faint streak in the gel (Lane 7, Figure 2.2). The ability of apo-Aβ$_1$-20 to act as a protector against oxidation of dsDNA in the presence of H$_2$O$_2$ is once again demonstrated here. Further reducing the concentration of H$_2$O$_2$ to 0.2% allows a clearer monitoring of plasmid cleavage patterns (Figure 2.3). Within 10 minutes of incubation, 5.0 µM of CuAβ$_1$-20 shows double-stranded DNA cleavage as evident in the appearance of a middle band approximately 3.5 kbp (Lane 6, Figure 2.3). Within 20-30 minutes, complete conversion of the supercoiled plasmid into linear and nick-circular conformations is observed, evident in the changes in the intensity of the different forms of the plasmid compared to the reference (Lanes 3 and 4, respectively, Figure 2.3). After 30 minutes, plasmid is cleaved into small pieces leaving a streak of oligonucleotide products (Lanes 1, 2, and 3). The different and quite opposite activities between apo-Aβ$_1$-20 and CuAβ$_1$-20 toward dsDNA damage may hint a physiological role of small fragments of apo-Aβ.
Figure 2.2. Time course reactivity assay of dsDNA cleavage. R is the reference plasmid DNA; lanes 1–7, DNA + 5.0 μM CuAβ₁-20 + 2% H₂O₂ (60, 50, 40, 30, 20, 10, 5 min, respectively); lanes 8-11, DNA + Aβ₁-20 (metal free) + 2% H₂O₂ (60, 40, 20, 10 min, respectively); lanes 12–15, DNA + 2% H₂O₂ (60, 40, 20, 10 min, respectively); and lanes 16–18, DNA + Aβ₁-20 (metal free, peroxide free). The standard DNA ladder starts with 1 kbp from the bottom with 1 kbp increment upward.
Figure 2.3. Time course reactivity assay toward the cleavage of 150 ng dsDNA. R is the reference DNA; lanes 1–6, DNA + 5.0 μM CuAβ_{1-20} + 0.2% H_{2}O_{2} (60, 40, 30, 20, 15, 10 min, respectively); lanes 7-10, DNA + 5.0 μM CuAβ_{1-20} (peroxide free 60, 40, 20, 10 min, respectively); and lanes 11-15, DNA + 0.2% H_{2}O_{2} (40, 30, 20, 15, 10 min, respectively). The standard DNA ladder starts with 1 kbp from the bottom with 1 kbp increment upward. The cleavage of plasmid by 5.0 μM Cu^{2+} ions and 3.6% H_{2}O_{2} is shown in Fig. 1, which exhibits a much lower activity.
To determine the role of the oxidizing agent in these reactions, the same concentration of CuA\(\beta\)\(_{1-20}\) was incubated with the plasmid in the absence of H\(_2\)O\(_2\) up to 60 minutes, which shows negligible cleavage (Lane 7, Figure 2.3). The low activity of CuA\(\beta\)\(_{1-20}\) without H\(_2\)O\(_2\) indicates a metal-centered activation of peroxide, such as the formation of a Cu\(_2\)-peroxo center found in many Cu complexes\(^{23,24}\) which subsequently results in oxidative damage to dsDNA. To distinguish the reaction pathways of oxidative DNA cleavage by H\(_2\)O\(_2\) in the presence and absence of CuA\(\beta\)\(_{1-20}\), a time-course experiment was established (Lanes 11-15, Figure 2.3). The reaction patterns of dsDNA cleavage in these two cases are clearly different. In the absence of A\(\beta\)\(_{1-20}\), dsDNA is cleaved into small fragments without formation of a linear intermediate as evident by the faint band at 2.0 kbp. The nature of the band is not clear at this stage and is not associated with A\(\beta\). The dsDNA cleavage by CuA\(\beta\)\(_{1-20}\) in the presence of H\(_2\)O\(_2\) is conformation-dependent, most active toward supercoiled dsDNA as evident by the accumulation of nicked-circular and linear forms with time in the reaction, likely due to the structural constraints of the supercoiled form. The accumulation of the linearized form (middle bands) is indicative of double-stranded DNA cleavage, rather than a random single-stranded cleavage, which is a key trigger that can result in cell apoptosis.\(^{25}\) The linearization of the plasmid via cleavage of double-stranded (ds) DNA is also characteristic of the cleavage pattern by DNA-recognizing agents such as Cu-bleomycin.\(^{26}\)

False regulation of metal homeostasis and ROS physiology is closely related to aging and oxidative stress,\(^{6}\) wherein apo-A\(\beta\)\(_{1-20}\) seems to serve as a scavenger of metal ions due
to its large affinity constant with metal ions and a protective agent against oxidative
damage of biological macromolecules by H₂O₂ based on the observations in this and
other studies. However, the presence of H₂O₂ can result in severe damage toward
dsDNA and presumably other redox-sensitive biomolecules as well by metallo-Aβ when
the metal ions are redox active as demonstrated herein.

**Kinetics and Mechanism of Oxidative Catalysis by CuAβ:** To gain further insight into
the mechanism for the oxidation activity of CuAβ₁-20 and its interaction with H₂O₂, the
catechol analogue 1,2,3-trihydroxylbenzene (THB) was utilized to provide detailed
kinetic information owing to its easily accessible oxidation state which also has been
utilized for investigation of oxidative activities of metal complexes. The oxidation rate
of THB by 7.5 µM CuAβ₁-20 was determined at different values of [THB] in the presence
of H₂O₂ at various concentrations (Figure 2.4), which reached saturation at high THB

\[
V_o = V_{background} + \frac{k_{cat}[CuAβ][THB]}{K'_{app} + [THB]}
\]  

(1)

concentrations. This saturation pattern suggests a possibility of pre-equilibrium kinetics.
The rate law for this reaction mechanism can be expressed as in Eq. (1), assuming that
the concentration of the intermediate THB-CuAβ₁-20 complex is much lower than that of
the unbound THB in which \( K'_{app} = (k_{-1} + k_{cat})/k_1 \) is the virtual dissociation constant and \( k_1 \)
and \( k_{-1} \) are the rate constants for the formation and dissociation, respectively, of the THB-
CuAβ complex. The data can be well fitted to Eq. (1), yielding \( k_{cat} = 0.00767 \text{ s}^{-1} \) and
\( K'_{app} = 1.67 \text{ mM} \), and a second-order rate constant \( k_{cat}/K'_{app} \) of \( 4.59 \text{ M}^{-1} \text{ s}^{-1} \) for the
reaction in the presence of 16.0 mM (0.0544%) $\text{H}_2\text{O}_2$. This represents a 724 fold increase in terms of the first-order rate constant when compared to the auto-oxidation of THB under the same reaction conditions in the absence of CuAβ1-20 (determined to be $k_0 = 1.06 \times 10^{-5}$ s$^{-1}$). A plot of $k_{\text{cat}}$ as a function of [H$_2$O$_2$] from Figure 2.4 shows that $k_{\text{cat}}$ reaches a plateau at high H$_2$O$_2$ concentrations (inset, Fig. 1.4). However, the $k_{\text{cat}}$ value does not reach zero at 0% H$_2$O$_2$ that is higher than $k_0$ of the auto-oxidation of the substrate. The oxidation reaction in the absence of H$_2$O$_2$ was further explored and discussed in a later section below. The plot seems slightly sigmoidal which indicates a possible presence of either a consecutive or a cooperative binding of H$_2$O$_2$ to the active center. Since catechol oxidation involves 2-electron transfer which matches with the two-electron reduction of H$_2$O$_2$ to yield two oxides, a consecutive mechanism is not fundamentally necessary for the reaction to take place. The data were fitted to the Hill equation to extract the Hill coefficient $\theta$ of 2.09 and $k_{\text{cat}}$ value of 0.00731 s$^{-1}$ at 0% H$_2$O$_2$ (close to the value of 0.0065 s$^{-1}$ directly measured in the absence of exogenous H$_2$O$_2$ discussed later), indicative of the presence of weak cooperativity and H$_2$O$_2$-independent oxidative catalysis.

Interestingly, the smaller fragment CuAβ1-16 showed more than 4-folds higher $k_{\text{cat}}$ of 0.0340 s$^{-1}$ for the reaction with the same concentration of H$_2$O$_2$. However, its catalytic efficiency is only twice higher than the larger fragment in terms of the second order rate constant $k_{\text{cat}}/K'_{\text{app}}$ (10.5 M$^{-1}$ s$^{-1}$), which suggests a participation of the last four C-terminal hydrophobic residues (LVFF) in the reaction pathway. The hydrophobic C-terminus may influence substrate binding and product release as reflected by the higher
Figure 2.4. Effect of peroxide concentrations on the rate of THB oxidation in the presence of 7.5 µM of CuAβ₁₋₂₀ in the presence of 1.6 (●), 3.2 (○), 16.2 (□), 32.3 (▼), 64.6 (♦), and 70.0 (△) mM H₂O₂ (HEPES buffer of 100 mM at pH 7.0 and 25.0 °C). The dotted line is CuAβ₁₋₁₆ in the presence of 16.2 mM H₂O₂. The inset shows the first order rate constant $k_{cat}$ as a function of hydrogen peroxide, wherein the solid trace is a fitting of the data to the Hill equation by taking into consideration of an activity at 0% H₂O₂.
\( K'_{\text{app}} \) and \( k_{\text{cat}} \) values for CuA\( \beta \)\(_{1-16} \) \( (K'_{\text{app}} = 3.23 \text{ mM}) \). This observation suggests that the C-terminus is close enough to the metal center to influence THB binding or a transition-state conformational change that affects both the binding of THB and the turnover of the reaction.

It has previously been documented that H\(_2\)O\(_2\) and other ROS generated by metallo-A\( \beta \) may play a role in the pathology of AD\(^{27,28,30} \). Since the local concentration of metallo-A\( \beta \) in an AD brain can reach sub-mM range\(^7 \), the above observation implies that a significant rate acceleration in redox reactions can be expected at a location where H\(_2\)O\(_2\) is produced. This rate enhancement in the brains of AD patients can be metabolically catastrophic. In the studies shown here, I have further specified the fate of H\(_2\)O\(_2\) in metallo-A\( \beta \)-associated redox reactions.

To further analyze the role of H\(_2\)O\(_2\) in the reaction pathway, a saturation profile was constructed with a fixed amount of the substrate THB at 6.0 mM. Under such conditions, the reaction reaches plateau at [H\(_2\)O\(_2\)] near 70.0 mM or 0.238% (Figure 2.5). The results can be well fitted to a pre-equilibrium kinetics (Eq. 2). This kinetics further corroborates a metal-centered mechanism as described above.

\[
V_O = V_{\text{background}} + \frac{k_{\text{cat}} [Cu\beta] [H_2O_2]}{K'_{\text{a,app}} + [H_2O_2]} \quad (2)
\]

The rate of acceleration against the background oxidation of THB at saturating [H\(_2\)O\(_2\)] is approximately 6,000 folds (background rate constant \( k_o = 1.14 \times 10^{-5} \text{ s}^{-1} \)) with \( k_{\text{cat}} = \)
Figure 2.5. H$_2$O$_2$ saturation profile of CuA$\beta_{1-16}$ (○) and CuA$\beta_{1-20}$ (●) activity at a fixed [THB] of 6.0 mM (HEPES buffer of 100 mM at pH 7.0 and 25.0 °C). The inset shows that DMSO (○) and mannitol (●) do not inhibit the oxidation of THB by CuAβ.
0.066 s\(^{-1}\) and \(k_{\text{cat}}/K'_{\text{app}} = 37.2 \text{ M}^{-1} \text{ s}^{-1}\). The hydrophobicity at the C-terminus does not affect the binding of H\(_2\)O\(_2\) as reflected by the similar apparent virtual dissociation constant \(K'_{\text{app}}\) between Cu\(\alpha\) and Cu\(\beta\) (17.3 and 16.6 mM, respectively). Qualitatively, since the \(k_{\text{cat}}\) value is small, the \(K'_{\text{app}}\) value is expected to be closer to the dissociation of the Cu\(\alpha\)-THB complex. The binding of each H\(_2\)O\(_2\) molecule in this case may not yield a complete turnover, suggesting that formation of the intermediate Cu-peroxo active center is not favorable under the experimental conditions which may well be the rate-determining step. This result corroborates with chemical model studies of catechol oxidase and tyrosinase in that a dinuclear Cu-peroxo intermediate is often short-lived and thermodynamically unstable.\(^{24,31}\)

Because the involvement of free radicals in the redox chemistry of metallo-\(\alpha\) had been implicated in previous reports,\(^7\) different amounts of DMSO and mannitol, two common scavengers for superoxide free radical and hydroxyl free radical,\(^{37,32}\) were added to the reaction solution separately with saturating concentrations of H\(_2\)O\(_2\) and THB. No noticeable effect on the reaction rates was observed under the experimental conditions (Figure 2.5, inset). However, this does not discount possible free radical generation since the free radicals may well be metal-centered and free-radical oxidation in solution may not be the predominant pathway in the oxidation of THB catalyzed by Cu\(\alpha\). We have established here a metal-centered oxidative catalysis by Cu\(\alpha\) and Cu\(\beta\) which can not only generate H\(_2\)O\(_2\) as noted previously\(^{30}\) but also activate H\(_2\)O\(_2\) for possible oxidation of biomolecules.\(^{33}\)
Since both H$_2$O$_2$ and THB can interact with the metal center and are considered “substrates” for CuA$\beta$, it is imperative to further narrow down the mechanism about how these two substrates interact individually with the metal center. For this purpose, the redox indicator 3-methyl-2-benzothiazolinone hydrazone (MBTH) was used to probe the oxidation product of THB in the absence of H$_2$O$_2$. MBTH is a common indicator used in catechol oxidase assay which forms a red adduct with the o-quinone products instantaneously. The rate for the oxidation of a catechol into its corresponding o-quinone can thus be easily monitored colorimetrically as the oxidation is the rate-limiting step. The rate for the oxidation of THB by CuA$\beta_{1-20}$ in the absence of H$_2$O$_2$ as a function of [THB] is not linear which can be well fitted to pre-equilibrium kinetics to give $k_{cat} = 0.0065$ s$^{-1}$ and $K^{app} = 2.0$ mM. The first order rate acceleration of THB oxidation here is 650 folds with respect to the auto-oxidation (i.e. $k_{cat}/k_o; k_o = 1.06 \times 10^{-5}$ s$^{-1}$). This oxidative reaction is much less significant in terms of rate acceleration than that in the presence of a saturating amount of H$_2$O$_2$ described above. Here, catechol is possibly oxidized by CuA$\beta$ in form of a dinuclear Cu$^{2+}$ center via 2-electron transfer to afford 2Cu$^+$ and o-quinone product. The reduced 2Cu$^+$ in turn can bind O$_2$ to form a dinuclear Cu$^{2+}$-peroxo center and follow the catalytic pathway as the CuA$\beta$/H$_2$O$_2$ system discussed above. Because a bi-substrate mechanism was implied from our results (i.e. both THB and H$_2$O$_2$ show saturation), further analysis of the data was performed. The Hanes analysis was used to minimize the error across the concentration range (Figure 2.6A). The virtual dissociation constant $K^{app}$ for both substrates cannot be resolved only from the primary nonlinear fitting without analyzing their combined effects.
important to determine the rates at varying amounts of H₂O₂ when holding THB constant and vice versa. The data in Figure 2.4 were fitted to a two-substrate random-binding mechanism according to Equation 3 (Figure 2.6A), wherein the binding of THB and H₂O₂ to CuAβ₁-20 was assumed to be random and in rapid-equilibrium with a subsequent ordered product release. Under these conditions, a simple conversion to a secondary plot

\[
\frac{[THB]}{V_o} = \left(1 + \frac{K'}{[H_2O_2]}\right) \frac{[THB]}{V_{max}} + \frac{K'_α}{V_{max}} \left(1 + \frac{K_i}{[H_2O_2]}\right)
\]

of the slope (slope = \( (1 + K'/[H_2O_2])/V_{max} \)) and the y-intercept (y intercept = \((K'_{app}/V_{max})*(1 + K_i/[H_2O_2])\)) in Figure 2.6B versus 1/[H₂O₂] yields \( K' \) and \( K'_α \), the true values for the virtual dissociation of H₂O₂ and THB, respectively, and the intrinsic affinity constant of H₂O₂ \( K_i \). Moreover, if any cooperativity is present in this bi-substrate reaction mechanism, it would be revealed by the ratio of \( K'_{app}/K' \). For a random equilibrium mechanism a ratio of \( K'_{app}/K' \) between 1 and 5 would suggest little cooperativity. In the oxidative catalysis by CuAβ₁-20, the \( K'_{app}/K' \) ratio is 2.85 for THB oxidation and \( K'_{app}/K'_α \) is 1.62 for H₂O₂ which indicates little cooperativity. It is important to note that based on the data alone it is difficult to distinguish between an ordered sequential-binding mechanism and the mechanism herein proposed. However, a random equilibrium phase is a sound assumption since both THB and H₂O₂ can interact with the metal centers separately.

In our proposed reaction mechanism, only when both THB and H₂O₂ substrates bind to the metal-center can productive turnover be observed with second-order rate constants of 8.66 and 15.6 M⁻¹s⁻¹ for the oxidation of THB by CuAβ₁-20 and CuAβ₁-16, respectively, in
the presence of H$_2$O$_2$ (whereas molecular O$_2$ serves as the second substrate in the absence of H$_2$O$_2$). This pathway differs from the previously proposed mechanism in the redox cycling of metallo-Aβ wherein the presence of the thioether group of Met$_{35}$ was accounted for the reduction of the metal center. The results presented here indicate substrate-mediated reduction of the metal center (since Met is absent in the studies) as well as oxidation of the substrate by metal-activated H$_2$O$_2$. However, our data do not discount the possibility of the involvement of Met in the reductive pathway in Aβ$_{1-40/42}$. Regardless, the redox chemistry of CuAβ presented here shows an important mechanism for possible destructive actions in Alzheimer’s disease.

Taken together, the metal-centered redox cycle of CuAβ action in this study seems to match the mechanism of the dinuclear Cu-containing catechol oxidase, wherein the oxidation of the substrate takes place both in the presence and absence of H$_2$O$_2$. Since the oxidation of catechols is a two-electron transfer process, the involvement of a dinuclear Cu center is thus a preferred pathway as in the case of the enzyme. In the presence of THB and H$_2$O$_2$, the dinuclear μ-$\eta^2,\eta^2$-peroxo-Cu$_2^{2+}$-THB transition state is eventually formed by assembling two metal centers together via the bridging peroxo as in the case of many mononuclear Cu$^{2+}$ complexes (Figure 2.7, steps A–C), which is followed by 2-electron transfer from the bound peroxide to the bound catechol (likely through the metal center) to yield Cu$_2^{2+}$-μ-OH and o-quinone to complete a catalytic cycle (step D). Here, the complexes oxy-Cu$_2^+$, Cu$_2^{2+}$-μ-(η$^2,\eta^2$-peroxo), Cu$_2^{2+}$-μ-(η$^1,\eta^1$-).
Figure 2.6. Hanes analysis of the kinetic data from Figure 4. The plots in A yield the apparent virtual dissociation constant for THB. The re-plot of the slope (●) and y-intercept (○) from A reveals dissociation constants for THB ($K'_a = 0.41$ mM) and H$_2$O$_2$ ($K_i = 17.3$ mM) in a bi-substrate reaction.
peroxo), and Cu$^{3+}$-($\mu$-oxo)$_2$ (B) are isoelectronic$^{24}$ which are not distinguishable in our study. In the absence of H$_2$O$_2$, the oxidation of the bound THB is achieved by 2-electron transfer to the dinuclear Cu$^{2+}$ center to yield Cu$^+$ (steps F–G) which is followed by O$_2$ and THB binding to regenerate the THB-Cu$^{2+}$-η$_2$-peroxo transition state (steps H, B, and C). The binary and the ternary complexes then follow the same pathway as the case in the presence of H$_2$O$_2$ for another turnover. H$_2$O$_2$ is also generated according to this mechanism under reducing conditions (steps E and I), which has been previously observed$^{30}$ and can serve as a competing reaction pathway toward the oxidation of catechols (steps C, D).

The mechanism of oxidative “chemical nucleases” has been thoroughly studied and reviewed.$^{36}$ According to the studies of some simple chemical nucleases such as Cu-1,10-phenanthroline, a reduced state of the metal center (by a reducing agent) is required for catalysis in the presence of O$_2$. In our experiments, however, the absence of a reducing agent to convert Cu$^{2+}$ to Cu$^+$ and the use of H$_2$O$_2$ as the oxidation agent suggest a different oxidative pathway. Moreover, the free radical scavenger$^{37}$ dimethyl sulfoxide (DMSO) did not inhibit the reaction, suggesting the absence of free radicals to induce the oxidative damage. On the basis of the results shown here, we propose a η$_2$-peroxo-bridged dinuclear Cu$^{2+}$ active center for CuAβ$_{1-20}$ as observed in a number of chemical model systems$^{23}$ and in catechol oxidase, tyrosinase, and hemocyanin.$^{38,39}$ The nature of this transient η$_2$-peroxo species and its attack on the substrates, although thoroughly
Figure 2.7. Proposed mechanism for the oxidation of THB by CuAβ in the presence (steps A–D) and absence of H2O2 (steps F–H and B–D). The production of H2O2 under reduction conditions (steps E) is also consistent with this mechanism.
studied, still has key mechanistic questions to be answered, such as the true structure of reaction intermediates, the role of substrate in the reaction mechanism, rate determining steps in catalysis, and specificity of metal cofactor for the function of enzymes like catechol oxidase, tyrosinase, peptidylglycine monooxogengase, methane monooxygenase, fatty acid desaturase, and ribonucleotide reductase.40–42

It has been proposed that substrate accessibility in the active site after O2 binding plays a key role in the action of these proteins.40,43 Consequently, a reversible O2 binding has been demonstrated in hemocyanin because of the lack of substrate accessibility, wherein bulky substrates such as aromatic systems and the ribose moiety of DNA may not easily gain access to the O2-binding active center of the proteins. However, studies of catechol oxidase and tyrosinase have shown the production of hydroxylated phenols and ortho-quinones, reflecting that substrates bind directly to the dinuclear Cu2-η2-O2 active center which enables a direct attack on the substrates by the peroxo unit.44,24

Metal Binding and Structure: Detailed information about the metal-binding ligands and geometry of the metal site is needed to gain further insight into the metal-centered redox chemistry and to elucidate any structure-function correlation important for the action of metallo-Aβ. Since activity is an excellent probe for monitoring reaction mechanism, it is thus chosen as a probe for the determination of the metal-binding stoichiometry of metallo-Aβ. Upon introduction of Cu2+ to Aβ, oxidative activity can be measured as described above. It is evident from the data that metal binding reaches saturation at slightly above 1:1 ligand-to-metal ratio (Figure 2.8). Despite a previous
Figure 2.8. Cu²⁺ titration toward Aβ₁-20 (▲) and Aβ₁-16 (●) monitored with their oxidation activities (100 mM HEPES buffer at pH 7.0 and 25.0 °C). The dotted curves show the best fittings to a noncooperative M+L ↔ ML equilibrium. Solid curves are the nonlinear fittings to the Hill Equation which yield Hill coefficients of 1.94 and 3.27 for Cu²⁺ binding to Aβ₁-16 and Aβ₁-20, respectively.
EPR study which indicate the binding of two Cu$^{2+}$ ions to Aβ, our result indicates the active species is a 1:1 CuAβ complex. Both a noncooperative binding equilibrium (a quadratic pattern) and a cooperative equilibrium (a sigmoidal pattern) were used to fit the data. It is evident from the fitting that the shorter CuAβ$_{1-16}$ fits equally well to both binding patterns with a metal-to-ligand stoichiometry of 1 to 1, whereas CuAβ$_{1-20}$ seems to fit better the cooperative binding pattern. The binding of Cu$^{2+}$ to Aβ$_{1-16}$ gives a Hill coefficient $\theta$ of 1.94, while the binding to Aβ$_{1-20}$ shows a higher cooperativity with $\theta$ of 3.27. This result is consistent with previous reports of cooperative metal binding to the entire Aβ determined on the basis of quantitative precipitation. The results presented here further indicate the presence of cooperativity in the oxidative activity as well as metal-binding. The higher C-terminal hydrophobicity of Aβ$_{1-20}$ may influence intermolecular interactions, resulting in a more apparent cooperativity. The data were also analyzed to determine whether or not there were possible inactive dimer conformations of this metallopeptide by plotting activity as function of the square root of metal ion concentration as previously described. However, the data do not reflect the existence of such equilibrium in this reaction pathway, adding supporting evidence to the dinuclear metal-centered redox mechanism herein proposed. Dissociation constants ($K_d$) for metal binding to Aβ can be extrapolated from both fits with values of 3.96 and 4.30 μM for CuAβ$_{1-16}$ and CuAβ$_{1-20}$, respectively. Since activity serves as the probe here, the values obtained above are thus the intrinsic dissociation constants attributable to the active CuAβ complexes and are not affected by coagulation equilibrium for Aβ. The intrinsic dissociation constant for metal binding in CuAβ$_{1-40}$ is likely to be in the same
range of ~4 µM for CuAβ₁-20 and CuAβ₁-16 owing to their probably similar metal-binding configuration. Indeed, dissociation constants in the range of ~0.1 µM for CuAβ₁-28 and ~2 µM for CuAβ₁-40 and CuAβ₁-42 were determined with ligand-competition⁴⁵ and direct fluorescence measurement.⁴⁸ An apparent dissociation constant \( K_{d_{app}} \) of 0.50 pM for Cu\(^{2+}\) binding to Aβ₁-40 was determined based on the formation of CuAβ₁-40 coagulates,⁴⁶ which can be dissected into the intrinsic metal dissociation constant \( K_{Cu} \) of ~4 µM and the dissociation constant of CuAβ₁-40 coagulates \( K_{co} \) ~0.13 µM (i.e., \( K_{d_{app}} = K_{Cu} \times K_{co} \)). The much smaller dissociation constant of 6.3 aM for Cu\(^{2+}\) binding to Aβ₁-42 would thus afford an apparent dissociation constant of CuAβ₁-40 coagulates in the range of 1.6 pM. A recent report indicates that trace amounts of metal ions can significantly affect Aβ coagulation, it is thus suspected that the dissociation constants may be underestimated based on the coagulation.⁴⁹ I report in this dissertation a direct and reliable means for the determination of metal binding to soluble Aβ fragments which is not complicated by the formation of the coagulation as previously observed⁴⁶ that can be influenced by other factors, such as trace amount of metal ions.⁴⁹

To further investigate the metal-coordination environment, the electronic spectrum of CuAβ₁-20 was obtained (Figure 2.9). The spectrum reveals a typical type-2 copper center with d-d transitions showing \( \lambda_{\max} \) at 610 nm (107 M\(^{-1}\) cm\(^{-1}\)), clearly distinguishable from the near IR absorption at 820 nm for aqueous Cu\(^{2+}\) solutions. This absorption is consistent with that of the “CuH₂L” species of acetyl-Aβ₁-16 with three coordinated His side chains in a potentiometric study⁵⁰ (617 nm and 117 M\(^{-1}\) cm\(^{-1}\)) and another study⁴⁵ (610 nm and ~50 M\(^{-1}\) cm\(^{-1}\) which seems to be too low an absorptivity⁵¹). The result
Figure 2.9. Electronic spectra of Aβ1-20 with 0.5 equivalents (solid trace) and 1.0 equivalent Cu$^{2+}$ (dashed trace) referenced against apo Aβ1-20 (100 mM HEPES buffer at pH 7.0). The dotted trace is the difference spectra of 1.4 and 1.0 equivalents of Cu$^{2+}$ in Aβ1-20, showing no further increase in the d-d transition after the binding of one equivalent of Cu$^{2+}$.
agrees well with a tetragonally distorted octahedral environment\textsuperscript{51,52} for the Cu\textsuperscript{2+} in CuAβ\textsubscript{1-20}. Upon addition of more than one equivalent of metal, the spectrum does not change. This is consistent with the results when activity is used as the probe to monitor metal binding (Fig. 2.8), wherein one equivalent Cu\textsuperscript{2+} is determined to bind to one peptide. It is also worth noting that there are no intense transitions in the near-UV range that can be possibly assigned to Tyr-to-Cu\textsuperscript{2+} charge-transfer transitions as observed in the Cu\textsuperscript{2+}-substituted proteases astacin and serralysin.\textsuperscript{53}

The metal coordination chemistry was also investigated by the use of Co\textsuperscript{2+} as an NMR probe. Co\textsuperscript{2+} has been well demonstrated to be an excellent probe for the investigation of metal-binding sites in a number of metalloproteins, including Zn and Cu proteins.\textsuperscript{54,55} Although Aβ\textsubscript{1-20} has four additional hydrophobic amino acids on the C-terminus, the conformations of the two peptides in d\textsuperscript{6}-DMSO are similar as they show nearly identical \textsuperscript{1}H NMR spectra (Figure 2.10). The signals due to L\textsubscript{16}VFF\textsubscript{20} side chains in Aβ\textsubscript{1-20} are clearly observed when compared with the spectrum of Aβ\textsubscript{1-16}, wherein LV are seen at ~0.6 ppm and FF ~7.2 ppm. This similarity reflects their similar configuration. There are two solvent exchangeable signals in the 14–16 ppm range (imidazole N-H signature chemical shifts\textsuperscript{56}) with a 1:2 ratio in intensity, corresponding to the three His side chains (insets, Figure 2.10 A and B).

Upon Co\textsuperscript{2+} titration, the intensities of these solvent exchangeable His-imidazole signals gradually decreased which was accompanied by the appearance of three far-downfield paramagnetically shifted signals in the region of 40–80 ppm as shown here for Aβ\textsubscript{1-20} (Figure 2.10C). These far-shifted signals are also solvent exchangeable and
correspond to the chemical shift of the solvent exchangeable signal of a paramagnetic Co\textsuperscript{2+}-bound imidazole group of a histidine residue (which cannot be due to dipolar shift of unbound His residues since the octahedral Co\textsuperscript{2+} center is expected not to possess magnetic anisotropy) as observed in many Co\textsuperscript{2+}-substituted metalloproteins.\textsuperscript{54,55} These three solvent exchangeable NH signals further confirms the involvement of all three histidine residues in Aβ\textsubscript{1-20} for metal binding, consistent with previous Raman spectroscopic studies,\textsuperscript{9} and is indicative of the absence of a bridging histidyl imidazole (in contrast to previously suggested\textsuperscript{9,12}) which would result in the loss of an imidazole NH signal.

Tyr\textsubscript{10} was suggested to be a possible ligand for Cu\textsuperscript{2+} and Fe\textsuperscript{3+} binding,\textsuperscript{9,12,57} but was suggested not to be a ligand in other studies.\textsuperscript{50} The \textsuperscript{1}H NMR signals of Tyr\textsubscript{10} (the two asterisked doublets centered at \~6.7 ppm in Figure 2.10 do not show any noticeable change upon the addition of the paramagnetic Co\textsuperscript{2+} ion. The binding of a Tyr-phenol group to Co\textsuperscript{2+} is expected to exhibit paramagnetically shifted \textsuperscript{1}H NMR signals of the bound phenol group outside the diamagnetic region as previously observed.\textsuperscript{58} This result indicates that this Tyr is not a metal-binding ligand, consistent with the lack of charge-transfer transitions for a possible Tyr-Cu\textsuperscript{2+} binding as described above. Our results also do not support the binding of the N-terminal amino group to the metal as previously suggested.\textsuperscript{45} This binding mode for paramagnetic Co\textsuperscript{2+} would show far upfield-shifted NH\textsubscript{2} signal(s), a downfield-shifted C\textsubscript{α}H proton, and slightly upfield-shifted C\textsubscript{β}H\textsubscript{2} protons owing to spin polarization, which were not observed.\textsuperscript{55}
Figure 2.10. $^1$H NMR spectra of Aβ$_{1-20}$ (A) and Aβ$_{1-16}$ (B) in DMSO. The insets show the imidazole N-H solvent exchangeable signals of His which disappear upon addition of a drop of D$_2$O (remain intact with same amount of H$_2$O). These signals disappear upon addition of paramagnetic Co$^{2+}$, with concomitant appearance of three solvent exchangeable hyperfine-shifted signals in the far-downfield region as shown here for Co-Aβ$_{1-20}$ (C). The asterisked signals in A and B are due to Tyr ring protons which remain the same upon Co$^{2+}$ binding.
Molecular mechanical calculations have been applied to determine the structure of Aβ1-16 and its metal-binding domain. The energies for different metal-binding modes have been calculated by the use of the MM3 force field and a simulated water droplet to solvate the peptide. Binding of Cu$^{2+}$ to His$_{13}$ and His$_{14}$ yields the lowest energy of $-385$ kcal/mol as compared to all other possible binding modes in the peptide. The binding to all three His side chains yields a distorted octahedral geometry (with 3 open coordination sites presumably occupied by water molecules) and a slightly higher energy at $-363$ kcal/mol (Figure 2.11). The energy difference between these two metal-binding modes may be low enough to be easily overcome at room temperature. Extensive H-bonding are observed in this calculated structure, particularly Glu3-Arg5-Asp7 H-bonding interactions may stabilize the structure to a great extend (dotted lines in Fig. 2.11). The energies for Cu$^{2+}$ binding to His$_{6/13}$ and His$_{6/14}$ are much higher at $-125$ and $-210$ kcal/mol, respectively, and are not likely to be the metal binding modes for Aβ. A histidine-bridged dimer form of the peptide previously proposed$^{12}$ was also calculated which gave an unacceptably high overall energy of 52,500 kcal/mol. The binding of Tyr$_{10}$ along with the histidine residues is also highly unfavorable which puts undue stress on the phenol ring causing it to pucker and the peptide backbone to distort, with a high overall energy of 570 kcal/mol. The recently suggested N-terminal binding mode (along with the binding of the 3 histidines) has also been calculated to give an unfavorable overall energy of 147 kcal/mol. Since Cu$^+$ can easily adopt a trigonal coordination sphere,$^{52}$ a calculation with a fixed trigonal coordination was performed which yielded an energy of $-318$ kcal/mol. The low energy difference between octahedral and trigonal
Figure 2.11- Proposed metal coordination and solution structure of CuAβ_{1-16} in a relaxed-eye stereo view based on NMR study of Co^{2+} binding and molecular mechanics calculations. According to the molecular mechanical calculations both the two-histidine (H13/H14) and three-histidine binding patterns are stable, whereas NMR study suggests the latter binding pattern. The dotted lines are H-bonds, which may prevent further bending of the peptide to allow the binding of the N-terminus to the metal as recently reported.\textsuperscript{45}
geometries rationalizes the redox cycle of the Cu center in the catalysis of catechol oxidation. The binding of the 3 His side chains to the metal renders one side of the metal center to have an open coordination sphere which can possibly bind H₂O₂ or O₂ to form the dinuclear bridging $\eta^2$-peroxo center described above.

IV. CONCLUDING REMARKS

The results presented here have added further insight and support to the structure and chemistry of metallo-Aβ which may assist better understanding of the neuropathology of Alzheimer’s disease. A complete redox cycle for the action of CuAβ has been proposed from the kinetic studies which is consistent with the mechanism proposed for the dinuclear Cu catechol oxidase. The results in this report, however, do not resolve the cause/effect debate about the role of Aβ in AD, but add more information to the chemistry of metallo-Aβ. As a cause for AD, I have shown and quantified redox chemistry of CuAβ in this dissertation which can serve as a catalyst both in the absence and presence of H₂O₂ to cause severe oxidative damages in the brains of AD patients. As an effect of AD, Aβ can be reasoned to be present as a regulator toward metal ion homeostasis due to its considerable metal affinities and its protective property toward oxidative DNA damage in the absence of Cu²⁺. In the latter case, abnormal homeostasis of redox-active metal ions can leach the metal ions to yield metallo-Aβ that can undergo redox destruction of biomolecules. I have presented data herein to revise the redox chemistry of the methionine-centered hypothesis by showing a metal-centered catalysis
as a significant contribution to the oxidative damage in the pathology of the neurodegenerative AD. The fate of H$_2$O$_2$ generated by CuAβ in the presence of a reducing agent previously observed or an electron-donating substrate shown here has also been established and quantified with exogenous addition of this oxidant. Further studies currently under way that focus on the structure-activity relationship of metallo-Aβ are expected to shed light on the roles of metal ions and Aβ in AD and with hope to provide useful information for treatment and prevention of Alzheimer’s disease.
IV. LIST OF REFERENCES


21) C.J. Burrows, V. Lepentsiotis, J. Domagala, I. Grgic, R. van Eldik, J.G. Muller, Mechanistic information on the redox cycling of nickel(II/III) complexes in the
presence of sulfur oxides and oxygen. Correlation with DNA damage experiments. 

*Inorg. Chem.* (1999), 38, 3500-3505


25) M. Stanulla, J. Wang, D.S. Chervinsky, S. Thandla, P.D. Aplan, DNA cleavage within the MLL breakpoint cluster region is a specific event which occurs as part of higher-order chromatin fragmentation during the initial stages of apoptosis. *Mol. Cell. Biol.* (1997), 17, 4070–4079.


33) See Chapter IV.

34) V. Leskovac, Comprehensive Enzyme Kinetics, Kluwer/Plenum, Boston, MA (2002).


40) N.J. Blackburn, F.C. Rhames, M. Ralle, S. Jaron, Major changes in copper coordination accompany reduction of peptidylglycine monooxygenase: implications

41) L. Que Jr., W.B. Tolman, Bis(μ-oxo)dimetal "diamond" cores in copper and iron complexes relevant to biocatalysis. *Angew. Chem.* (2002), 41, 1114-1137.


CHAPTER III. CATECHOL OXIDASE AND PHENOL MONOOXYGNEASE

ACTIVITIES OF CuAβ₁₋₂₀

I. INTRODUCTION

Over the past few years an enormous effort has been directed toward the investigation of the metal-dependent mechanisms that lead to the neuropathology of Alzheimer’s disease (AD). The self-assembled metallo-β-amyloid (Aβ) peptide fibrils are the hallmark of this disease and have been attributed to Fe³⁺ and Cu²⁺-centered generation of H₂O₂ under reducing conditions which has been postulated to be of significant importance in connection with neuropathy in AD. However, an area of oversight has been the detailed chemical processes associated with the neuropathology of AD, besides the general acclaimed “ROS” (reactive oxygen species, including H₂O₂) assault. Hence,

‡ This work has been published: G.F.Z. da Silva, L.-J. Ming, Alzheimer’s Disease Related Copper(II)-β-Amyloid Peptide Exhibits Phenol Monooxygenase and Catechol Oxidase Activities. Angew. Chem. Int. Ed. (2005), 44, 5501-5504.
a better understanding of metal-centered redox chemistry and the mechanism for the generation of ROS and their fate can provide insight into potential strategies for prevention and treatment of AD.

Several examples of redox chemistry in biological systems are known to be associated with di- or multi-nuclear “Type-3” Cu oxidases, which might be related to the redox activity of Cu\textsuperscript{II}-A\textbeta.\textsuperscript{1-4} A number of chemical model systems targeting Type-3 copper centers have successfully been demonstrated to contain highly active isoelectronic copper-dioxygen species (i.e. Cu\textsuperscript{II}\textsubscript{2}-µ-η\textsuperscript{1}·η\textsuperscript{1}-peroxo, Cu\textsuperscript{II}\textsubscript{2}-µ-η\textsuperscript{2}·η\textsuperscript{2}-peroxo, and Cu\textsuperscript{III}\textsubscript{2}-bis-µ-oxo) responsible for copper-dependent oxidation and hydroxylation reactions.\textsuperscript{6-9} Despite the extensive modeling studies, peptide mimics of these enzymes have apparently been excluded from the studies. Cu\textsuperscript{II}-A\textbeta seems to fill the gap since it is a natural-occurring Cu-peptide complex demonstrated to exhibit oxygen-associated redox chemistry, although details about its oxygen binding and activation mechanisms is lacking. I present herein results which bring together two distinct fields of research, AD and Type-3 copper centers. The results elucidate that the Cu\textsuperscript{II} complex (CuA\textbeta\textsubscript{1-20}) of the icosapeptidyl metal-binding domain of A\textbeta (DAEFR\textsuperscript{5} HDSGY\textsuperscript{10} EVHHN\textsuperscript{15} KLVFF\textsuperscript{20}) exhibits metal-centered redox chemistry consistent with the mechanisms of the Type-3 copper enzymes phenol monooxygenase (e.g., tyrosinase) and catechol oxidase.
II. EXPERIMENTAL

Phenol Monooxygenase assay: Phenol hydroxylation and oxidation and catechol oxidation were performed as previously reported\textsuperscript{10} with minor changes to fit current studies. Same molar concentrations of phenol or catechol and 3-methyl-2-benzothiazolinone hydrazone (MBTH, which serves as an ortho-quinone indicator) were mixed in 100 mM HEPES at pH 7.00 in the presence of 5.0 µM (phenol hydroxylation/oxidation) or 0.50 µM CuAβ\textsubscript{1-20} (catechol oxidation). The red-adduct of the ortho-quinone product was monitored at 500 nm (ε = 32,500 M\textsuperscript{-1}cm\textsuperscript{-1}; cf. Figure 3.1) Auto-oxidation of phenol and catechol were determined under the same conditions without CuAβ\textsubscript{1-20}. The 1-20 fragment of Aβ was synthesized at the Peptide Center of the University of South Florida. The identity of the peptide has been confirmed with a Bruker MALDI-TOF mass spectrometer.

The rates were determined on a Varian CARY50 spectrophotometer equipped with a temperature controller. The rate law for pre-equilibrium kinetics (Eq. 1), which Cu\textsuperscript{II}-Aβ follows, is shown in Eq. 2, where \( k_{cat} \) and \( K'_{app} \) are the first-order rate constant and virtual dissociation constant, respectively, \( S \) is the substrate and \( v_{background} \) is the auto-oxidation rate.

\[
\text{Cu}^{\text{II}}\text{-Aβ} + S \xrightleftharpoons[k_{-1}]{k_1} \text{Aβ-Cu}^{\text{II}}\text{-S} \xrightarrow[\text{cat}]{k_{cat}} \text{Cu}^{\text{II}}\text{-Aβ} + \text{Prod.} \tag{1}
\]

\[
v = v_{background} + \frac{k_{cat}[CuAβ][S]}{K'_{app} + [S]} \tag{2}
\]
Figure 3.1. The production of \( o \)-quinone from phenol in the absence of \( \text{H}_2\text{O}_2 \) is monitored by the increase in the absorption due to its adduct with MBTH.
The Hill equation can serve as a general model for cooperativity as reflected by the Hill coefficient $\theta$ which deviates from one in the presence of cooperativity (Eq. 3), wherein $K_x$ is the intrinsic dissociation constant.

$$ \frac{V_v}{V_{\text{max}}} = \frac{[\text{CuA}\beta]^\theta}{K_x + [\text{CuA}\beta]^\theta} $$

(3)

In case of a bi-substrate catalysis, such as phenol hydroxylation/oxidation and catechol oxidation in the presence of $\text{H}_2\text{O}_2$, both substrates can interact with the metal center independently. The data are fitted to the Hanes plot (Eq. 4) to yield true values of substrate dissociation constants $K'$.\textsuperscript{19}

$$ \frac{[S]}{V_0} = \frac{1 + \frac{K'}{[\text{H}_2\text{O}_2]}}{V_{\text{max}}} \left( \frac{[S]}{V_{\text{max}}} \right) + \frac{K'}{V_{\text{max}}} \left( 1 + \frac{K_i}{[\text{H}_2\text{O}_2]} \right) $$

(4)

III. RESULTS AND DISCUSSION

*Catechol Oxidase activity:* The metal-centered redox chemistry of CuA$\beta_{1-20}$ was probed using catechol and the much more inert phenol as substrates.\textsuperscript{10} The oxidation of catechol under aerobic conditions reaches a plateau at low-mM concentrations (Fig. 3.2). This saturation profile fits nicely to pre-equilibrium kinetics (Eqs. 1 and 2), affording $k_{cat} = 0.154 \text{ s}^{-1}$, $K'_{app} = 0.35 \text{ mM}$ and a significant second-order rate constant $k_{cat}/K'_{app} = 440 \text{ M}^{-1} \text{ s}^{-1}$ (● Fig. 3.2). Since the formation of quinone from catechol is a two-electron oxidative process, the reaction is expected to follow the two-electron dinuclear reaction pathway for catechol oxidase,\textsuperscript{11} wherein the binding of catechol to the active-site di-Cu$^{II}$
Figure 3.2. Saturation profile for the oxidation of phenol (solid square), deuterated phenol (square), and catechol (solid circle) in the absence of H$_2$O$_2$. 
center results in the reduction of the center to yield di-Cu$^1$ with concomitant production of o-quinone. The reduced di-Cu$^1$ center can bind dioxygen to afford the active peroxo-bridged di-Cu$^{II}$ center, which can further oxidize a subsequently bound substrate. H$_2$O$_2$ can also be generated in this reaction pathway from the peroxo-bridged di-Cu$^{II}$ center in the presence of a reducing agent, such as the substrate itself. This H$_2$O$_2$ production pathway under reduction conditions is consistent with previous observations in AD studies.$^3$ This catechol oxidase-like mechanism has also been observed in kinetic studies in several chemical model systems$^{12}$ and in polyphenol oxidation by CuA$\beta$$_{1-20}$.$^{13}$ It is noteworthy that recent density functional theory (DFT) results reported mixed valence Cu$^{II}$-Cu$^I$ transition states,$^{14}$ corroborating with the reduction pathway for the Cu center.

$Zn^{2+}$ dilution: I show in Chapter II that the Cu$^{II}$:A$\beta$$_{1-20}$ stoichiometry to be in 1:1 ratio for oxidative activity with three N$_{\epsilon}$-coordinated imidazole histidine rings as metal-binding ligands.$^{13}$ Since activity is an excellent probe for determining stoichiometry, gradual replacement of Cu$^{II}$ in CuA$\beta$$_{1-20}$ with redox-inactive Zn$^{II}$ can serve as a practical method for addressing the nature of the metallo-active center by virtually “silencing” the active sites through the dilution with Zn$^{II}$. A linear correlation between the activity and the extent of of Cu$^{II}$ in the Zn$^{II}$ dilution should be observed for simple 1:1 metal binding if there is no cooperativity and/or interactions between the active site Cu$^{II}$ from different A$\beta$ strands. Conversely, a sigmoidal activity profile was observed as a function of mole fraction of Cu$^{II}$ in A$\beta$$_{1-20}$ toward the oxidation of the catechol derivative 3,5-di-tert-butyl catechol (DTC, $k_{cat} = 0.411$ s$^{-1}$ and $K'_{app} = 0.781$ mM) which can be nicely fitted to the Hill equation (Eq. 3) with a Hill coefficient $\theta$ of 2.40 and $r^2 = 0.99$ (Figure 3.3A, solid
trace). The data clearly cannot be fitted well to a quadratic equation for 1:1 non-cooperative binding mode ($r^2 = 0.91$; dashed trace, Figure 3.3A). These results imply a possible presence of a cooperative dinuclear active Cu$^{II}$ center during the catalytic oxidation of catechol by CuAβ$_{1-20}$, consistent with the catalytic cycle of catechol oxidase.$^{11}$

Effect of H$_2$O$_2$: The presence of the ROS H$_2$O$_2$ (25 mM) significantly enhances the turnover and catalytic efficiency of CuAβ$_{1-20}$ toward catechol oxidation (Figure 3.4), yielding $k_{cat} = 0.531$ s$^{-1}$ and $K'_{app} = 0.342$ mM and a significant second-order rate constant $k_{cat}/K' = 1.51 \times 10^3$ M$^{-1}$s$^{-1}$ from the Hanes plot (to give $K'$; Eq. 4, Figure 3.5A)$^{15}$ for a random bi-substrate reaction, wherein the bindings of the two substrates H$_2$O$_2$ and catechol are independent of each other. It is worth noting that the second-order rate constant for catechol oxidation is only about 20 times lower than that of the catechol oxidase from gypsywort.$^{16}$ This pathway is consistent with the “peroxide shunt” in the action of catechol oxidase in the presence of H$_2$O$_2$, wherein a Cu$^{II}$-$\mu$-$\eta^2$-$\eta^2$-peroxo intermediate is proposed to be the active species (cf. Fig. 2.7 A-D).$^6$ The oxidation of catechol to form $o$-quinone in the absence and presence of H$_2$O$_2$ exhibits remarkable $3.25 \times 10^5$ and $1.12 \times 10^6$ folds, respectively, rate acceleration in terms of the first-order rate constant $k_{cat}$ compared to that of aerobic auto-oxidation of catechol without CuAβ$_{1-20}$ determined to be $k_o = 4.74 \times 10^{-7}$ s$^{-1}$. 
Figure 3.3. Oxidative activity of CuAβ_{1-20} toward the oxidation of DTC (A) and phenol (B) in the absence of H_{2}O_{2} as a function of Cu^{II} mole fraction with a constant Cu^{II} + Zn^{II} at pH 7.0 and 25 °C. The solid traces are fittings to the Hill equation and the dotted traces are fittings to a quadratic binding pattern with metal:ligand = 1:1.
Figure 3.4. The effect of H$_2$O$_2$ on the first order rate constant $k_{cat}$ toward the oxidation of phenol (solid square) and catechol (solid circle).
Figure 3.5. Hanes plot analysis of kinetic data from Figure 3.4.
**Phenol hydroxylation:** Owing to their inertness, metal-centered hydroxylation of phenol and derivatives, particularly polychlorophenols, poses some challenging tasks in chemical synthesis and environmental detoxification and remediation\(^{17}\) and may provide further insight into the action of those metalloenzymes for arene monooxygenation.\(^{8,18}\) In addition to catechol oxidation described above, phenol was observed to be hydroxylated and oxidized by CuA\(^{\beta1-20}\) in the presence of saturating amount of H\(_2\)O\(_2\) (> 50.0 mM), wherein the formation of o-quinone exhibits rate constants of \(k_{\text{cat}} = 0.213\) s\(^{-1}\) and \(K'_{\text{app}} = 1.31\) mM, and \(k_{\text{cat}}/K' = 457\) M\(^{-1}\)s\(^{-1}\) from the Hanes plot (Figs. 3.1 and 3.2). This result represents a remarkable rate acceleration of \(4.6 \times 10^6\) folds for the hydroxylation/oxidation of phenol to form o-quinone in terms of \(k_{\text{cat}}\) compared to that of aerobic auto-oxidation of phenol (measured to be \(k_o = 4.6 \times 10^{-8}\) s\(^{-1}\)). This reaction is expected to take place following a similar dinuclear mechanism as in the action of the di-Cu enzyme tyrosinase toward hydroxylation and oxidation of tyrosine, wherein the active center is believed to contain dinuclear \(\mu-\eta^2:\eta^2\)-peroxo-Cu\(II\)\(_2\) on the basis of spectroscopic studies.\(^{8,14}\)

Cooperativity for the oxidation of both catechol and phenol in terms of \(k_{\text{cat}}\) is observed upon titration of H\(_2\)O\(_2\) as reflected in the sigmoidal activity profile with respect to H\(_2\)O\(_2\) (Figure 3.4). The data from the oxidation of catechol and phenol by H\(_2\)O\(_2\) fit nicely to the Hill equation (Eq. 3), yielding Hill coefficients \(\theta = 2.23\) and 1.78, respectively. Moreover, fitting of the rates to a random bi-substrate reaction mechanism yields corrected \(K'\) values and a cooperativity index based on the ratio \(K'_{\text{app}}/K'\) (Fig. 3.5 B), wherein a ratio of 1.70 for H\(_2\)O\(_2\) in both phenol and catechol oxidations and 2.07 and
2.10 for phenol and catechol, respectively, also suggest a small cooperativity and their independent binding to the active center.

Of particular interest is the observation of scarcely reported\textsuperscript{19} Cu\textsuperscript{II}-centered hydroxylation and oxidation of phenol \textit{aerobically without H}_2\textit{O}_2. The production of \textit{o}-quinone from phenol catalyzed by CuAβ\textsubscript{1-20} follows pre-equilibrium kinetics, yielding \( k_{\text{cat}} = 3.90 \times 10^{-3} \text{ s}^{-1} \) and \( K'_{\text{app}} = 1.23 \text{ mM} \) (Figure 3.2) which represents a first-order rate acceleration of \( 8.67 \times 10^{4} \) folds with respect to aerobic auto-oxidation of phenol. The \( k_{\text{cat}} \) value is lower than that for catechol oxidation, indicating that the hydroxylation step here must be the rate-limiting step. Otherwise, these two reactions would have similar \( k_{\text{cat}} \) values attributed to the oxidation of a bound catechol upon hydroxylation of a bound phenol. Moreover, the use of deuterated phenol as substrate shows significant kinetic isotope effect (KIE), wherein \( k_{\text{cat}} \) values of \( 1.12 \times 10^{-3} \) and \( 0.0442 \text{ s}^{-1} \) are obtained in the absence (\( \square \) Fig. 3.2) and presence of 100 mM H\textsubscript{2}O\textsubscript{2} which represent KIE of 3.46 and 4.77, respectively. The results indicate that hydroxylation and breakage of the \textit{o}-C–H bond of phenol is the rate-limiting step, which is followed by a faster step to form \textit{o}-quinone. The different KIE values for phenol hydroxylation in the presence and absence of H\textsubscript{2}O\textsubscript{2} suggest that the rate-determining step in these two cases may be different and/or possibly involve additional pathways. The \( K'_{\text{app}} \) values are not significantly different between phenol and deuterated phenol (1.31 and 1.23 mM for the latter in the presence and absence of H\textsubscript{2}O\textsubscript{2}, respectively), suggesting that \( k_{\text{cat}} \) does not significantly contribute to the magnitude of \( K'_{\text{app}} \) and that they may have a similar binding mode. The mechanistic reasoning (Figure 3.6) for the conversion of phenol to \textit{o}-quinone may be
**Figure 3.6.** Proposed mechanism for aerobic hydroxylation and oxidation of phenol in the absence of H$_2$O$_2$, wherein the binding of phenol and reduction of the metal center is a key step (A). The binding of dioxygen and formation of superoxide (B) is proposed to be assisted by a dinuclear center.
attributed to that the CuII/I redox equilibrium can be achieved upon phenol binding, followed by electron transfer to afford CuI and phenol radical which was suggested to be stabilized through resonance structure with the free radical situated at the ortho and para positions\textsuperscript{19a} (step A). This intermediate is then attacked by dioxygen followed by electron transfer to possibly form a CuII-superoxide center which may be further stabilized by a dinuclear center (step B). The free radical was proposed previously to be attached directly by triplet dioxygen,\textsuperscript{19a} which however is not symmetrically favorable. Coupling between the bound phenol radical and superoxide radical at an ortho position can then be expected to be a favorable step (step C), which is followed by transfer of electrons and an oxygen atom to afford the final quinone product (step D). An involvement of a dinuclear center for the catalysis is possible as discussed below. Since the hydroxylation and oxidation of phenol is a multi-electron transfer process, the involvement of two metal centers is suspected. An activity profile is obtained for phenol oxidation similar to the case of the “ZnII dilution” experiment for catechol oxidation. The data can be fitted equally well to the Hill equation (Eq. 3) to afford θ value of 1.80 ($r^2 = 0.98$) and a quadratic equation for single metal-binding ($r^2 = 0.98$), consisting with either a mononuclear oxidation\textsuperscript{19} or a cooperative mechanism involving a dinuclear center,\textsuperscript{14} or a combination of both pathways.

\textit{DCC optical titration:} To monitor substrate binding, a slow substrate 4,5-dichlorocatechol (DCC which is approximately 200 times slower in terms of $k_{\text{cat}}$ than catechol) was titrated to 0.2 mM CuA\textsubscript{β1–20} in the presence or absence of 100 mM H\textsubscript{2}O\textsubscript{2} and the electronic spectrum collected at 25 °C and pH 7.0 (Figure 3.4). Similar spectra
Figure 3.7. Optical titration of DCC to 0.2 mM CuAβ in the presence of 100.0 mM H$_2$O$_2$ in 100 mM HEPES at pH 7.0. Analogous spectra were obtained in the absence of H$_2$O$_2$. The inset shows the change in absorbance at 437 nm as a function of equivalents of DCC added, consisting with the formation of a 1:1 adduct.
were obtained, indicating that H₂O₂ was not involved in DCC binding to Cu²⁺Aβ under the experimental conditions. The absorption at 437 nm increases upon addition of DCC and reaches saturation at >1.2 equivalents of its addition, which can be nicely fitted to single-substrate binding mode to yield a dissociation constant of 0.24 mM. The result provides a direct evidence for catechol binding to the metal center, consistent with the observation in a chemical model²⁰ and the mechanism proposed above.

**IV. CONCLUDING REMARKS**

In conclusion, we have established the catalytic activities of CuAβ₁₋₂₀ toward the relatively inactive (according to their $k_o$ values) catechol and phenol in the presence and absence of H₂O₂ in aqueous solutions near physiological conditions. The reaction patterns are consistent with the mechanisms carried out by Type-3 copper centers as observed in catechol oxidase and tyrosinase and their dinuclear model systems.⁸,¹⁸ These results are unique thus far in metal-centered redox chemistry related to Alzheimer’s disease, and expected to offer further insight into the neuropathology of this disease since it has been suspected to connect with, in addition to many other factors, the oxidation of mono- and diphenol-containing neurotransmitters such as dopamine, epinephrine, norepinephrine, and serotonin.²¹,²² Furthermore, the connection of this highly reactive Cu-oxygen chemistry with Alzheimer’s disease can better define the role of metallo-Aβ in the neuropathology of this disease and possibly lead to different treatment strategies toward this disease.
V. LIST OF REFERENCES


CHAPTER IV. METALLO-ROS IN ALZHEIMER’S DISEASE: METAL-CENTERED OXIDATION OF NEUROTRANSMITTERS BY Cu$^{2+}$-β-AMYLOID PROVIDES AN ALTERNATIVE PERSPECTIVE FOR THE NEUROPATHOLOGY OF ALZHEIMER’S DISEASE‡

I. INTRODUCTION

The generation of reactive oxygen species (ROS), including superoxide, peroxide, and free radicals, is associated with normal redox metabolic pathways as side-tracks which can be regulated through the action of superoxide dismutase, catalase, and some reducing agents under homeostasis.$^{1,2}$ However, long-term effects of such oxidative chemical imbalance in normal and disease states can be expected. ROS are often considered the culprits responsible for the devastating effects of oxidative stress,$^3$ concerning cancer, aging, heart diseases, arthritis, diabetes, and the etiology of neurodegenerative disorders such as Parkinson’s disease and Alzheimer’s disease (AD)$^4$. AD affects primarily the elderly which causes considerable distress of the patients and emotional sufferings of their families and close friends. One mechanism

proposed for the neurodegeneration in AD focuses on amyloid-β peptides (Aβ) and their metal complexes through formation of plaques and fibrils and generation of the ROS H2O2 and free-radicals.5–9

Aggregation of Aβ of 40 or 42 amino acids (DAEFR HDSGY10 EVHHQ KLVFF20 AEDVG SNKGA30 IIGLM VGGV340 IA) in the brain is the hallmark in AD neuropathology induced by metal binding10–12 and is usually found as metal-containing plaques and insoluble fibrils. Similar pathological effects are also found in transgenic mouse models with human Aβ.13,14 Moreover, soluble fragments of Aβ can be generated in vivo by insulin degrading enzyme as well as α and β-secretases.15,16 Nevertheless, the cause or effect connection of the metallo-Aβ plaques with AD is still under debate.17–19 Despite immense endeavor in Aβ research, the potential risk of metal-centered oxidative catalyses by metallo-Aβ has been overlooked.20 The Cu II complexes of metal-binding domains of Aβ (CuAβ) have recently been demonstrated to exhibit metal-centered oxidative catalysis, consistent with Type-3 copper oxidases.21,22 To verify the bio-relevance of this metal-centered catalysis, we have determined the oxidation of several catecholamine and indoleamine neurotransmitters catalyzed by CuAβ1–40 and two metal-binding N-terminal fragments CuAβ1–16 and CuAβ1–20 under various biomimetic conditions. The studies described herein are expected to provide chemical basis for better understanding the etiology of AD.

II. EXPERIMENTAL

The peptides Aβ1–16(20) were synthesized at the Peptide Center of the University of South Florida, and confirmed with a Bruker MALDI-TOF mass spectrometer. Aβ1–40 was purchased from Biopeptide Co., LLC (San Diego, CA). The Cu II complex
of Aβ<sub>1-40</sub> was prepared according to literature procedures<sup>23</sup> and concentration determined with a standard BCA assay. The CuAβ stock solutions were quickly aliquoted to prevent concentration deviations caused by aggregation.

**Kinetic assays:** Kinetic measurements were performed and analyzed as previously described<sup>21,22</sup> by the use of MBTH as an o-quinone indicator (ε<sub>product</sub> = 32,500 M<sup>-1</sup> cm<sup>-1</sup> for phenol and catechol<sup>24</sup> and 28,900, 27,200, and 27,500 M<sup>-1</sup> cm<sup>-1</sup> for Dopa, dopamine, and epinephrine/norepinephrine, respectively,<sup>25</sup> and 35,200 M<sup>-1</sup> cm<sup>-1</sup> for serotonin and 5-hydroxytryptophan). The data are analyzed with the bi-substrate Hanes plot for experiments in the presence of H<sub>2</sub>O<sub>2</sub> to yield the apparent K<sub>S</sub> and intrinsic K<sub>Si</sub> dissociation constants for the neurotransmitters and H<sub>2</sub>O<sub>2</sub>.<sup>37</sup>

**III. RESULTS AND DISCUSSION**

Dopamine has been directly linked to the neurodegenerative Parkinson’s disease.<sup>26</sup> Moreover, symptoms of Parkinson’s disease have also been recognized in AD patients.<sup>27</sup> Thus, disturbance of dopamine metabolism may be closely associated with AD. Dopamine is effectively oxidized aerobically to dopaquinone by CuAβ<sub>1-40</sub> at pH 7.0, reaching saturation at high dopamine concentrations (Fig. 4.1; Table 1) which is consistent with pre-equilibrium-like kinetics. There is an apparent cooperativity in the catalysis that is not usually expected in simple monomeric systems, which may be due to the tendency of the peptide to coagulate and/or the formation of a dinuclear center during catalysis. Fitting the results to the Hill equation gives kinetic constants k<sub>cat</sub> = 7.48 × 10<sup>-4</sup> s<sup>-1</sup> and k<sub>cat</sub>/K<sup>-1</sup> = 2.77 M<sup>-1</sup> s<sup>-1</sup> and a Hill coefficient θ = 1.48. This reaction shows 85-fold rate enhancement relative to
Figure 4.1. Oxidation of catechol (○) and dopamine (●) by 1.47 µM CuAβ1-40 in 100 mM HEPES at pH 7.0 and 25°C. Dashed traces are fittings to simple pre-equilibrium kinetics while the solid traces are the fitting to the Hill equation. The inset shows the effect of H₂O₂ on $k_{\text{cat}}$ which also show the presence of cooperativity.
<table>
<thead>
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<th>CuAβ</th>
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<th>$K^*$ (mM)</th>
<th>$k_{cat}/K^*$ (M$^{-1}$s$^{-1}$)</th>
<th>$k_{rel}$</th>
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</table>

*Table 2.* Kinetic parameters for dopamine oxidation by Aβ$^\alpha$ in 100 mM HEPES buffer at pH 7.0 and 25 °C. $^b$ in the presence of 20 mM H$_2$O$_2$
auto-oxidation of dopamine. The shorter CuAβ₁–20 and CuAβ₁–16 do not exhibit the apparent cooperativity and exhibit 16 and 37-fold, respectively, higher activity than CuAβ₁–40 toward dopamine oxidation in terms of $k_{\text{cat}}$. Catechol oxidation shows the same catalytic trend (Fig. 4.1; Table 1), with CuAβ₁–40 showing cooperativity ($\theta = 2.7$) and CuAβ₁–16 exhibiting the highest activity. The results herein indicate that the soluble CuAβ fragments may be more severe in causing oxidative stress in the brain of AD patients than the coagulation-prone CuAβ₁–40. Recent studies also suggested pathological significance of soluble forms of Aβ.²⁸

Collectively termed catecholamines, dopamine, epinephrine, norepinephrine, and Dopa are catechol-containing neurotransmitters (whereas serotonin and its precursor 5-hydroxytryptophan are indole-containing) which are involved in cognitive, behavioral, physical, physiological, and psychological functions.²⁹ Oxidation of these molecules may cause severe alteration in bioactivity, eventually leading to neuronal death.³⁰ Metabolic malfunctions of neurotransmitters are known phenomena in the physiology of AD³¹,³² and have also been suggested to be related to the neuropathology of this disease.³¹–³⁴ However, a chemical mechanism for the neurotransmitter malfunction in AD is still unknown. These catecholamine neurotransmitters are found to be effectively oxidized aerobically to their respective $\alpha$-quinone products by CuAβ₁–20 (Fig. 4.2, Table 1). Herein, CuAβ₁–20 is shown to significantly accelerate aerobic oxidation rate of these neurotransmitters by 333–2,420 times in terms of $k_{\text{cat}}$ relative to auto-oxidation rate constant $k_o$. 
Figure 4.2. a) Catecholamine oxidation by 2.5 µM CuAβ1-20 in the absence of H₂O₂ toward dopamine (●), (+/−)-epinephrine (▲) and (−)norepinephrine (■), L-DOPA (○) and D-DOPA (□). The solid lines are the best fit to an enzyme-like pre-equilibrium kinetics). b) The dependence of the first-order rate constant on H₂O₂ toward the oxidation of dopamine (●), (+/−)-epinephrine (○), and (−) norepinephrine (▼).
Compared to that of catechol, the auto-oxidation rates of these neurotransmitters are faster by nearly 10 times in terms of $k_o$. However, their oxidation rates by CuAβ are ~130–980-fold slower than that of catechol, reflecting their higher resistance to oxidation by CuAβ by ~$10^3$–$10^4$-fold relative to catechol oxidation. The relatively higher stability against oxidative damage entitles these molecules better suited for the purpose of neurotransmission. Dopa shows a lower reactivity than other catecholamines, likely due to the inductive effect of the carboxyl group on the side chain. Chirality of these neurotransmitters does not appear to play a role in this oxidation chemistry (Table 2).

The ROS H$_2$O$_2$ has been commonly suggested to be a culprit causing the oxidative stress in AD.$^{4,6}$ However, this ROS alone at 50.0 mM does not significantly affect the oxidations of neurotransmitters (footnote b, Table 2). Conversely, oxidations of these neurotransmitters by CuAβ in the presence of >15 mM (>0.051%) H$_2$O$_2$ under the same conditions exhibit significant rate enhancement ($k_{rel}$, Table 2; Fig. 4.3), e.g., 312-fold relative to auto-oxidation rate constant $k_o$ for dopamine oxidation by CuAβ$_{1-40}$ with $k_{cat} = 0.0056$ s$^{-1}$ and $k_{cat}/K^* = 21$ M$^{-1}$ s$^{-1}$. The oxidation of dopamine is dependent on H$_2$O$_2$ and reaches a plateau at H$_2$O$_2 > 15$ mM with $\theta = 3.57$ (Fig. 1, inset), which indicates that the ROS H$_2$O$_2$ can bind to CuAβ to afford a "metallo-ROS" in a cooperative manner. Oxidation of catechol without the side chain by Cu-Aβ$_{1-40}$ exhibits a similar kinetic pattern (Fig. 4.1; Table 2), confirming the cooperative nature of this oxidative catalysis.

The results conclude that "metallo-ROS" is much more reactive than free ROS alone as far as H$_2$O$_2$ is concerned. Once again, the shorter CuAβ$_{1-16(20)}$ fragments exhibit higher activity than CuAβ$_{1-40}$ toward dopamine oxidation in the presence of
Figure 4.3. H$_2$O$_2$ effect on oxidation of dopamine (a), (+/-)epinephrine (b), and (-)norepinephrine (c) catalyzed by 3.15 µM CuAβ$_{1-20}$ in the presence of H$_2$O$_2$ (from bottom, 0, 1.0, 2.0, 4.0, 8.0, 16.0, and 20.0 mM). (d–f) Hanes plot analysis of catecholamine oxidation in the presence of H$_2$O$_2$ (from plots a–c). The lines are the best fits to a bi-substrate mechanism (g–i) Secondary plots of the Hanes plots from d–f which reveal apparent and intrinsic dissociation constants $K'$ values, by plotting of the slope (●) and y-intercept (ο) from plots d–f as a function of 1/H$_2$O$_2$. 
H$_2$O$_2$ (Table 1). This observation once again suggests pathological significance of soluble Aβ fragments in AD. Herein, CuAβ$_{1-20}$ is able to significantly accelerate oxidation rates of catechol and catecholamine neurotransmitters by 1,350 to $2.24 \times 10^5$ times in terms of $k_{\text{cat}}$ relative to $k_o$ (Table 2). It is worth noting that $k_{\text{cat}}/K^*$ of 1,690 M$^{-1}$ s$^{-1}$ for catechol oxidation by CuAβ$_{1-16}$ approaches enzyme-like catalytic efficiency which is 5.3% of the activity of the catechol oxidase (32,000 M$^{-1}$ s$^{-1}$) from gypsywort (*Lycopus europaeus*).$^{36}$

Since both H$_2$O$_2$ and the neurotransmitters can bind to the metal active-center, the data are analyzed with a two-substrate random-binding mechanism according to the Hanes equation (Fig. 4.3)$^{37}$ to afford the apparent and intrinsic dissociation constants $K_S$ and $K_{i(S)}$ (0.34 and 0.23 for catechol and 0.52 and 0.22 for dopamine, respectively). The ratios of $K_S/K_{i(S)}$ are greater than one, indicating that the neurotransmitters and H$_2$O$_2$ are affecting the binding of each other in this bi-substrate reaction mechanism.$^{37,38}$

**Influence of micelles:** Aβ$_{1-40}$ and fragments are found in various cellular environments, including soluble forms in the cytosol and insoluble forms as membrane-bound plaques. Herein, the detergent sodium dodecyl sulfate (SDS) is used to approximate the amphiphilic nature of cell membrane. Structures of short Aβ fragments in the presence of SDS differ considerably from that in the absence of this micelle-forming surfactant.$^{39}$ The activities of CuAβs in the presence of micelles may offer insight into the nature of the structure and activity of Aβ. The rate constant $k_{\text{cat}}$ of CuAβ$_{1-20}$ for the oxidation of dopamine is largely affected by the soluble form of SDS (4.5 times) and noticeably influenced (80%) by micelles with the critical micelle concentration (CMC) of SDS.
<table>
<thead>
<tr>
<th>Substrate</th>
<th>$k_{\text{cat}}$ (10^{-3} \text{s}^{-1})</th>
<th>$K'$ (mM)</th>
<th>$k_{\text{cat}}/K'$ (M^{-1}\text{s}^{-1})</th>
<th>$k_{\text{rel}}^b$</th>
<th>$k_{\text{cat}}$ (10^{-3} \text{s}^{-1})_{\text{sat. H}_2\text{O}_2}$</th>
<th>$k_{\text{cat}}/K'$ (M^{-1}\text{s}^{-1})_{\text{sat. H}_2\text{O}_2}$</th>
<th>$k_{\text{rel}}^d$_{\text{sat. H}_2\text{O}_2}$</th>
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<tbody>
<tr>
<td>Catechol (by Aβ1–16)</td>
<td>154</td>
<td>0.35</td>
<td>440</td>
<td>3.25 × 10^5</td>
<td>531</td>
<td>1.510</td>
<td>2.24 × 10^5</td>
</tr>
<tr>
<td>(by Aβ1–40)</td>
<td>280</td>
<td>0.31</td>
<td>900</td>
<td>5.92 × 10^5</td>
<td>1,150</td>
<td>1,690</td>
<td>7.64 × 10^5</td>
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<tr>
<td>Dopamine</td>
<td>0.87</td>
<td>0.19</td>
<td>4.6</td>
<td>1,850</td>
<td>11.2</td>
<td>28</td>
<td>1.96 × 10^4</td>
</tr>
<tr>
<td>(by Aβ1–16)</td>
<td>11.6</td>
<td>0.90</td>
<td>13</td>
<td>1,320</td>
<td>99</td>
<td>190</td>
<td>5,530</td>
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<tr>
<td>(by Aβ1–40)</td>
<td>28</td>
<td>0.31</td>
<td>90</td>
<td>3,180</td>
<td>230</td>
<td>340</td>
<td>1.27 × 10^4</td>
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<tr>
<td>(±) Epinephrine</td>
<td>0.75</td>
<td>0.27</td>
<td>2.8</td>
<td>85</td>
<td>5.61</td>
<td>11</td>
<td>312</td>
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<tr>
<td>(–) Epinephrine</td>
<td>3.1</td>
<td>0.60</td>
<td>5.2</td>
<td>2,420</td>
<td>24</td>
<td>34</td>
<td>9,420</td>
</tr>
<tr>
<td>(–) Norepine</td>
<td>2.9</td>
<td>0.61</td>
<td>4.8</td>
<td>2,270</td>
<td>22</td>
<td>54</td>
<td>8,630</td>
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<tr>
<td>(+) Norepine</td>
<td>2.7</td>
<td>0.52</td>
<td>5.2</td>
<td>1,330</td>
<td>17</td>
<td>32</td>
<td>6,940</td>
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<tr>
<td>(–) Dopa</td>
<td>2.8</td>
<td>0.57</td>
<td>4.9</td>
<td>1,380</td>
<td>18</td>
<td>39</td>
<td>7,350</td>
</tr>
<tr>
<td>(+) Dopa</td>
<td>1.1</td>
<td>0.34</td>
<td>3.2</td>
<td>333</td>
<td>9.11</td>
<td>27</td>
<td>1,360</td>
</tr>
<tr>
<td>Phenol (by Aβ1–16)</td>
<td>1.2</td>
<td>0.32</td>
<td>3.8</td>
<td>364</td>
<td>9.03</td>
<td>28</td>
<td>1,350</td>
</tr>
<tr>
<td>(by Aβ1–40)</td>
<td>3.9</td>
<td>1.23</td>
<td>3.2</td>
<td>8.67 × 10^4</td>
<td>213</td>
<td>170</td>
<td>4.16 × 10^6</td>
</tr>
<tr>
<td>(by Aβ1–16)</td>
<td>6.4</td>
<td>0.59</td>
<td>11</td>
<td>1.41 × 10^5</td>
<td>340</td>
<td>58</td>
<td>6.64 × 10^6</td>
</tr>
<tr>
<td>(by Aβ1–40)</td>
<td>6.4</td>
<td>0.45</td>
<td>14</td>
<td>8.25 × 10^3</td>
<td>43.9</td>
<td>97.5</td>
<td>5.64 × 10^4</td>
</tr>
<tr>
<td>5-hydroxy-Trp serotonin</td>
<td>6.7</td>
<td>1.47</td>
<td>4.5</td>
<td>7.75 × 10^3</td>
<td>30.4</td>
<td>67.6</td>
<td>3.28 × 10^4</td>
</tr>
<tr>
<td>(by Aβ1–16)</td>
<td>26</td>
<td>0.63</td>
<td>41</td>
<td>3.03 × 10^4</td>
<td>250</td>
<td>380</td>
<td>2.69 × 10^5</td>
</tr>
<tr>
<td>(by Aβ1–40)</td>
<td>0.28</td>
<td>0.33</td>
<td>0.84</td>
<td>324</td>
<td>7.8</td>
<td>24.4</td>
<td>8.41 × 10^3</td>
</tr>
</tbody>
</table>

**Table 3.** Kinetic parameters for the oxidation of catecholamines to o-quinone by Aβ1–20, except those indicated. Intrinsic dissociation constant for the CuAβ-S complex. b The background self-oxidation rate constant $k_o$ is calculated based on the MBTH detection of o-quinone formation in the absence of CuAβ at 25 °C in 100.0 mM HEPES pH 7.0 under aerobic conditions.
around 8 mM (Fig. 4.4);¹xi whereas that of CuAβ1–40 is slightly influenced by soluble SDS (2-fold) and greatly affected by micellar SDS (8-fold).

Under our experimental conditions, SDS micelles do not influence the self-oxidation rates of neurotransmitters. The shortest CuAβ1–16 is only slightly affected (85% enhancement) in the presence of saturating amount of SDS and shows only 14% enhancement in correspondence with CMC. The results indicate that the plaque-forming CuAβ1–40 is expected to exhibit more significant oxidation chemistry when it is “solubilized” and incorporated into a hydrophobic environment; whereas the soluble CuAβ1–16 and CuAβ1–20 are already more powerful oxidation agents than CuAβ1–40 in aqueous environments (Table 1). The results seem to also corroborate the “opposing activities” proposal for neuroprotection via proteolysis and aggregation,¹xii wherein the highly active small Aβ1–16(20) fragments are supposed to be eliminated by the former process whereas the activity of Aβ1–40 is much decreased by the latter process by forming aggregates. The local concentrations of metallo-Aβ plaques can reach mM range,⁴ which can cause significant oxidative damages of proximal areas on the brain. Consequently, the devastating metallo-ROS chemistry due to CuAβ1–40 and fragments in the brain of AD patients can have a widespread effect in different cellular environments, particularly in hydrophobic membranes that significantly enhance the metal-centered oxidative activities as demonstrated herein.

Influence by reducing agents: The NAD(P)+/NAD(P)H ratios vary according to changes in metabolism and is species/tissue-dependent,⁴xlii thus are expected to affect the redox property of CuAβ in vivo. Changes in the homeostatic levels in terms of NAD(P)+/NAD(P)H may reflect the neurochemical status under oxidative stress.⁴xliv Metabolic changes have been
Figure 4.4. Effect of SDS on the oxidative activity of CuAβ1-16 (●), CuAβ1-20 (○), and CuAβ1-40 (▼) in 100.0 mM HEPES pH 7.0 and 25 °C. The data show influence of the activity by both the monomeric and the micellar forms of SDS with CMC of 8 mM.
noted to be associated with several age-associated diseases, including neurodegenerative diseases. It has been previously reported that these ratios are managed according to spatial and temporal constraints in the brain under oxidative stress. The oxidation activity of CuAβ1-20 is lowered by NAD(P)H. As the ratios of NAD(P)⁺/NAD(P)H decrease, the activity in terms of $k_{cat}$ toward dopamine oxidation decreases by 2.4 and 1.9 times, respectively (Fig. 4.5). Based on the proposed mechanism for catechol oxidation by Cu-Aβ1–20, the inhibitory effect of NAD(P)H is due to shifting in equilibrium toward H₂O₂ generation under reducing conditions. The more pronounced inhibition caused by NADPH as compared to NADH might be attributed to the phosphate group in NADPH. Indeed, phosphate has been observed to be a competitive inhibitor toward the oxidation of dopamine, showing $K_i = 4.7$ mM (Fig. 4.6). The ratio of free NAD⁺/NADH has been under debate which nevertheless has recently been suggested to be around 600, consistent with the value based on potentiometric measurement. The relatively smaller availability of free form of NADH suggests that this biological reducing agent may not significantly influence the metal-centered oxidative catalysis of CuAβ under physiological conditions.

To reveal the mechanism of the full-length Aβ1–40, the slow substrate 4,5-dichlorocatechol (DCC) was titrated into a solution of CuAβ1–40. The observation of a charge transfer transition at 438 nm (Fig. 4.7) is indicative of DCC binding to the CuII center (with an affinity constant of $6.4 \times 10^5$ M⁻¹, dotted trace; Fig. 4.6) analogous to its binding to the soluble CuAβ1–20. Cooperativity is apparent in DCC binding (solid
**Figure 4.5.** Influence of NAD(P)$^+$ (○) and NAD$^+$ (●) on the oxidative activity of CuAβ1–20 toward dopamine oxidation with a fixed total concentration of NAD(P)$^+$ + NAD(P)H = 10.0 mM.
Figure 4.6. Phosphate inhibition toward catechol oxidation by CuAβ1-20 in 100 mM HEPES at pH 7.0. Phosphate concentrations are 0, 10, 20, and 40 mM (from bottom).
trace; Fig. 4.7), probably due to the coagulation nature of Aβ1–40. The stoichiometry of DCC:CuAβ1–40 = 1:2 verifies the dinuclear nature of the catalysis. The activation profile during CuII titration to Aβ1–40 is sigmoidal with a Hill coefficient θ = 2.68 (Fig. 4.8) which further support a dinuclear catalysis. Cooperativity is not usually expected in mononuclear catalysis, wherein a linear correlation of the activity with CuII is expected until one equivalent is reached.

The oxidation of catecholamine neurotransmitters by CuAβ1–20 and the stoichiometry for DCC binding are consistent with the action of dinuclear catechol oxidase. The mechanism of this enzyme thus serves as a working model for the metal-centered oxidation of catecholamines by CuAβ. Under aerobic conditions, the catechol moiety binds to a di-CuII center (Fig. 4.9) and is oxidized via 2-electron transfer to afford di-CuI and o-quinone product (B). Di-CuI then binds O2 to form the metallo-ROS μ-peroxo-Cu2 center (C and D) after 2-electron transfer from di-Cu2+ to O2, which may bind (E) and oxidize (F) another substrate. In the presence of an electron donor such as NAD(P)H (H), the oxidation of neurotransmitters is inhibited. Herein, H2O2 is formed to certain extents which can produce other types of ROS and may exacerbate the oxidative destruction by going through the peroxide shunt (G) upon forming the highly reactive metallo-ROS μ-peroxo-di-CuII intermediate. The metal-bound H2O2 as a metallo-ROS in equilibrium of three possible iso-electronic speciesxlvii (D) has been demonstrated herein to be a more potent oxidative agent than H2O2 alone to cause neurodegeneration.

The full-length plaque-forming CuAβ1–40 also shows significant metal-centered oxidative activity toward the relatively more inert phenol (Fig. 4.10; Table 2),
Figure 4.7. a) Optical titration of DCC into 0.05 mM CuAβ1–40. Inset shows change of intensity at 438 nm upon DCC binding to a di-CuII center. Both were in 100.0 mM HEPES at pH 7.0.
**Figure 4.8.** Activity titration of Cu$^{II}$ into 2.5 micro-M Aβ$_{1-40}$ under saturating conditions of catechol.
Figure 4.9. Mechanism for the oxidation of catecholamine neurotransmitters and the cause of neurodegeneration by CuAβ. The metal-bound H₂O₂ as a metallo-ROS is shown in D.
showing $k_{\text{cat}} = 0.439$ and $1.43 \text{ s}^{-1}$ (with a small cooperativity $\theta = 1.6$) and $k_{\text{cat}}/K' = 0.351$ and $1.14 \text{ M}^{-1} \text{ s}^{-1}$, respectively, in the absence and presence of 50.0 mM H$_2$O$_2$.

The activity reaches saturation at high H$_2$O$_2$ (Fig. 4.10), reflecting binding of H$_2$O$_2$ to the metal to afford metallo-ROS. Phenol hydroxylation and oxidation exhibits the same trend of reactivity with the shortest CuA$\beta_{1-16}$ showing the highest activity (Table 2). Herein, oxidation of phenol is dramatically enhanced by $6.64 \times 10^6$ and $1.41 \times 10^5$ times by CuA$\beta_{1-16}$ with and without 50.0 mM H$_2$O$_2$, respectively.

Serotonin can be hydroxylated and oxidized by CuA$\beta_{1-16,20,40}$ into its quinone form with significant rate enhancements ranging from $324$ to $3.03 \times 10^4$ and $8.41 \times 10^3$ to $2.69 \times 10^5$-fold, respectively, relative to $k_o$ in the absence and presence of a saturating amount of H$_2$O$_2$ (Fig. 4.11; Table 2). As in the case of catecholamine oxidation, the oxidation of serotonin by CuA$\beta_{1-40}$ is slower than that by CuA$\beta_{1-16,20}$. Moreover, the precursor of serotonin 5-hydroxy-Trp can also be effectively hydroxylated and oxidized by CuA$\beta_{1-20}$ (Table 2). The apparent and intrinsic dissociation constants $K_S$ and $K_{i(S)}$ for phenol are determined to be 1.23 and 0.54 mM, respectively, from the Hanes plots, indicating H$_2$O$_2$ binding to the metal center decreases the binding of serotonin. The reactions may follow the tyrosinase mechanism$^\text{xlviii}$ for the hydroxylation and oxidation of phenol, serotonin, and 5-hydroxy-Trp. However, Cu$^{II}$-A$\beta$ is still highly active without H$_2$O$_2$ in hydroxylation reaction, whereas the di-Cu$^{II}$ met-form of tyrosinase is not. The large oxidation enhancements of the two indoleamines by CuA$\beta$ suggest that their oxidation possibly taking place in the brain of AD patients may alter serotonin-mediated physiological functions, including sleep disorder, mood change, and anxiety often associated with AD patients.$^\text{xlix}$
Figure 4.10. (a) Aerobic oxidation of serotonin by 1.47 μM CuAβ1–40 in 100.0 mM HEPES at pH 7.0 and 25 °C. Dashed traces are fittings to a pre-equilibrium kinetics while the solid traces are fittings to the Hill equation, showing the presence of cooperativity. (b) Oxidation of serotonin by CuAβ1–40 in the presence of H2O2 as in (a).
Figure 4.11. (a) Saturation profile of phenol oxidation by 1.47 µM CuAb1-40 in 100.0 mM HEPES at pH 7.0 and 25°C. Dashed line is the fitting to a pre-equilibrium kinetics while the solid line is the fitting to the Hill equation ($k_{\text{cat}} = 4.39 \times 10^{-4}$, $K_m = 1.25$ mM, $k_{\text{cat}}/K_m = 0.35$ M$^{-1}$s$^{-1}$, $\theta = 1.60$). (b) The effect of H$_2$O$_2$ on the first order rate constant $k_{\text{cat}}$. Cooperativity is not very apparent in this case, showing $\theta = 1.33$. 
Cell culture experiments reveal that catecholamines can exacerbate the oxidative stress caused by Aβ. However, the metal-centered oxidative catalysis has been overlooked. Oxidation of catecholamines is known to generate neurotoxic quinone products that are involved in protein modification (e.g., covalent modification of dopamine transporter) and polymerizing tau protein into fibrils. Age-related deficits of both dopamine and norepinephrine have been implicated in the vulnerability of noradrenergic neurons in the hippocampus, which suffers significant damage in AD. Moreover, loss of noradrenergic neurons is linked to degradation of the locus ceruleus, which is rich in dopaminergic neurons that shows severe lesion in AD. Hence, a possible mechanism for reduction in neurotransmitter-regulated alertness response, delay-period activity, sleep cycle, mood stabilization, short-term memory, cognition, attention and problem solving capability, satisfaction feeling, and coordination of physical movement experienced by AD patients may be due to excessive oxidation of neurotransmitters, hinting at a possible neuropathological role of metallo-ROS associated with CuAβ.

IV. CONCLUDING REMARKS

In conclusion, the full-length plaque-forming CuAβ1-40 found in the brain of AD patients has been demonstrated to exhibit significant activities toward the oxidation of neurotransmitters with or without H2O2 which can be further enhanced by interacting with membranes. The results suggest that imbalance of neurotransmitter metabolism
can be created near $\text{A}\beta_{1-40}$ plaques in AD. Moreover, the smaller fragments CuA$\beta_{1-16,20}$ show higher activities than CuA$\beta_{1-40}$ toward the oxidation of neurotransmitters. This observation suggests that small fragments of A$\beta$, due to their soluble nature, can significantly disturb neurotransmission in a more systematic manner in the brain of AD patients and thus may play an important role in neuropathology of this devastating disease.
IV. LIST OF REFERENCES


35) The off-the-shelf H$_2$O$_2$ solution in the U.S. is 3%.


CHAPTER V. METHIONINE-35 IS NOT A REDUCING AGENT FOR THE METAL-CENTERED OXIDATION CHEMISTRY OF Cu$^{2+}$-β-AMYLOID—KINETIC AND EPR STUDIES

I. INTRODUCTION

Oxidative stress\(^1\) has been a key topic of research concerning cancer, aging, heart diseases, arthritis, diabetes, and neurodegenerative disorders such as Parkinson’s and Alzheimer’s diseases\(^2\) (AD). Mechanisms proposed for the neurodegeneration in AD brains generally focus on the amyloid-β peptide (Aβ),\(^2\) a proteolytic product of 40–42 amino acids of the ubiquitously distributed amyloid precursor protein (APP), and its interaction with redox-active metal ions. The recent “Aβ cascade hypothesis” suggested that Aβ aggregates trigger a complex pathological cascade which leads to neurodegeneration in AD,\(^3\) including generation of H$_2$O$_2$,\(^4,5\) free-radical induced oxidation,\(^6-8\) and the involvement of Met35 as a reducing agent\(^9\) in the redox chemistry of metallo-Aβ. A central focus of the neuropathology of AD thus has been the effects of
redox-active transition metal ions and reactive oxygen species (ROS), such as superoxide, hydroxyl free radical, and H$_2$O$_2$. Although physiological processes responsible for dealing with ROS can be up-regulated to tackle variations in oxidative stress, long-term negative effects of such oxidative chemical imbalance such as that taking place in the brains of AD patients can be expected. Some AD treatment strategies have targeted the metal center in Aβ to prevent peptide aggregation and ROS generation. However, comparatively little effort has been focused on the metal-centered chemistry associated with the bound metal ions, such as the detailed coordination chemistry and the reactivity of the metal–bound ROS in CuAβ.

Transgenic mouse models with human Aβ show similar effects as AD patients, including Aβ aggregation and loss of memory. Rodent Aβ has been shown to exhibit redox activity in vitro that was attributed to ROS generation via Met35, even though the metal-binding domain was mutated in rodent Aβ (i.e., His13→Arg). Since Aβ activity and aggregation in AD brains is sequence-specific and metal-dependent, it is a priority to establish the targets of redox activity that can contribute to the physiological and cognitive effects of AD. I have shown in Chapters II-IV that the Cu$^{2+}$ complexes of Aβ and its soluble fragments (Aβ$_{1-16}$ and Aβ$_{1-20}$) showed considerable activities toward the oxidation of phenol, polyphenol, catechol, and neurotransmitters to form o-quinones which challenges the redox role of Met35 that is not present in the fragments. The capability of Aβ to bind copper$^{21}$ with subsequent H$_2$O$_2$ generation under reduction conditions$^5$ and catechol oxidation$^{18-20}$ hint at the possible formation of a reactive μ-η$^2$:η$^2$-peroxo-di-Cu$^{2+}$ species (or its isoelectronic counterparts, oxy-di-Cu$^+$, μ-η$^1$:η$^1$-...
peroxo-di-Cu$^{2+}$, and bis-µ-oxo-di-Cu$^{3+}$)\textsuperscript{22} which is the active species in action of the type-3 dinuclear copper oxidases\textsuperscript{23} such as catechol oxidase.

In this chapter I present kinetic and spectroscopic investigations of the oxidation chemistry of CuAβ\textsubscript{1-20} and the influence of methionine and reducing agents on the oxidation chemistry. The results support metal-centered oxidative stress and shed light on the mechanistic role of Met35 and reducing agents in the redox chemistry of CuAβ.

II. EXPERIMENTAL

The 1-20 fragment of Aβ was synthesized by the use of the Fmoc chemistry at the Peptide Center of the University of South Florida, and the identity of the peptide confirmed with a Bruker MALDI-TOF mass spectrometer. Dopamine, ascorbic acid, and glutathione (GSH) were obtained from Sigma-Aldrich (St. Louis, MO), 3-methyl-2-benzothiazolinone hydrozone hydrochloride monohydrate (MBTH) from Acros (Fairlawn, NJ), and H\textsubscript{2}O\textsubscript{2}, EDTA, L-methionine, and Cu(NO\textsubscript{3})\textsubscript{2} from Fisher Scientific (Swanee, GA). All plastic ware and glassware were demetallized with EDTA and extensively rinsed with 18.0-MΩ deionized water.

The catechol oxidase assay toward dopamine was performed as previously reported\textsuperscript{20,24}. Same equivalent of a substrate dopamine and the o-quinone indicator MBTH were mixed in 100 mM HEPES at pH 7.00 in a final volume of 1.0 mL. The MBTH-adduct of o-quinone was monitored at 505 nm ($\varepsilon = 27,200$ M$^{-1}$ cm$^{-1}$) and 25 °C for 5–10 minutes with a Varian CARY50 Bio-UV-Vis spectrophotometer equipped with
a CARY PCB-150 Water Peltier temperature regulation system, and the rates determined by the change in absorbance over time. Rates were fitted to appropriate rate laws and rate constants determined with non-linear regression (SigmaPlot 8.0), such as the enzymatic Michaelis-Menten-like kinetics. The effects by H₂O₂, ascorbate, GSH, and L-Met were determined similarly in the presence of different amounts of the corresponding reagent and the inhibition/dissociation constants determined accordingly. Binding of L-Met to CuAβ₁₋₂₀ was performed by direct titration of Met into CuAβ₁₋₂₀ in 100 mM HEPES at pH 7.0 and monitored with the CARY50 spectrophotometer.

Electron paramagnetic resonance (EPR) experiments were performed on a Bruker Elexsys E580 cw/pulsed X-band spectrometer at the University of Florida with professor Alexander Angerhofer. For a typical cw EPR spectrum, the field was set wide enough to reveal a possible low-field transition for magnetically coupled systems with a microwave frequency of 9.4 GHz, field modulation typically around 2 G, and time constant of 40-80 ms at ~5–6 K. The g and A tensors were obtained with numerical fittings using the “easyspin” toolbox for Matlab. Electron relaxation times were measured with the standard two-pulse Hahn’s spin-echo method or the inversion-recovery methods, wherein the signal intensity as a function of time is fitted to simple or double exponential decay to obtain the relaxation time. ESEEM (electron spin echo envelope modulation) spectra were recorded with the usual π/2-τ- π/2-T- π/2 pulse sequence with a π/2 pulse of 20 ns in order to study ¹⁴N and ²H nuclei in the ligand sphere. A typical ESEEM trace consisted of 1024 points taken at time intervals of 8 ns. The transient was first base-line corrected by subtracting the ordinary T₁ exponential decay function and any remaining constant
baseline offset, then zero-appended to 2048 points. A Hamming window function was then applied to the time-domain spin-echo envelope, followed by Fourier transformation to afford the frequency-domain spectrum.

An ESEEM spectrum can reveal those nuclei having super-hyperfine coupling with the Cu$^{2+}$ center, including coordinated His side chains and water, which can thus serve as a very useful tool for the investigation of the Cu$^{2+}$ center in CuAβ. The theoretical background$^{27}$ of ESEEM is summarized herein. In the case of coordinated His side chains, the super-hyperfine coupling arises from the electron-nuclear interactions and the nuclear quadrupole interactions (NQI) of the remote non-coordinated nitrogen ($^{14}$N, I = 1) on the imidazole ring of coordinated His side chains. At X-band, three zero-field nuclear quadrupole resonance lines $\nu_\pm$ and $\nu_o$ are observed which can be determined from the NQI lines in the ESEEM spectrum (Eq. 1 and 2), wherein $e^2qQ$ is the quadrupole coupling constant and $\eta$ the asymmetry parameter ($\eta = 0$ for a complete axial symmetry and 1 for a pure rhombic symmetry).

$$\nu_\pm = \frac{1}{4}(e^2qQ)(3 \pm \eta) \quad (1)$$
$$\nu_o = \frac{1}{2}(e^2qQ)\eta \quad (2)$$

The $e^2qQ$ and $\eta$ values can thus be obtained from the NQI lines in the ESEEM spectra since $(\nu_+ - \nu_o)/3 = \frac{1}{4}(e^2qQ)$. In the case of coordinated water, the super-hyperfine coupling arises from the electron-nuclear interactions and the NQI of the coordinated water molecules upon deuteration. The deuterium Zeeman interaction at X-band is much
larger than the isotropic component of the electron-nuclear coupling. Thus, the
deuterium peak in the ESEEM spectrum is found at the deuterium Zeeman frequency and
split slightly by the electron-nuclear coupling.

III. RESULTS

Dopamine is oxidized by CuAβ1–20 in the absence of H2O2 with rate constant of $k_{\text{cat}} = 0.0104$ s$^{-1}$ and $K_m = 0.89$ mM (trace •, Fig. 5.1), consistent with the observations
discussed in Chapter IV.$^{20}$ Addition of L-Met to the reaction solution significantly
increases the activity (Fig. 5.2). The reaction in the presence of saturating amount of Met
seems to induce slight cooperativity which can be fitted to the Hill equation with a Hill’s
coefficient $\theta = 2.3$ (Solid trace Figure 5.1). The activity reaches a plateau at high [Met]
(Fig. 5.1), indicating direct Met binding to the active center of CuAβ. Fitting of the $k_{\text{cat}}$
values as a function of [Met] to a simple equilibrium of $[\text{CuAβ} + \text{Met} \rightleftharpoons \text{Met-CuAβ}]$
gives an affinity constant of 1,900 M ($K_d = 0.53$ mM). Met follows a non-essential
activation pattern toward dopamine oxidation by CuAβ1–20 as shown in the Lineweaver-
Burk plot (Fig. 5.1), i.e., both CuAβ-substrate and Met-CuAβ-substrate complexes are
active. From the data, the dissociation constant for the activation $K_A$ can be obtained to
be 0.087 mM, and the dissociation constant for the ternary [CuAβ1–20-Met-dopamine]
complex $K_{TS}$ is 0.125 mM. In the presence of H2O2, the activity is further
Figure 5.1. Saturation kinetics curves of dopamine oxidation by CuAβ fitted with Michaelis-Menten kinetics (dotted traces) and with Hill equation (solid traces).
Figure 5.2. Effect of L-Met on the first order rate constant toward dopamine oxidation.
Figure 5.3. Lineweaver-Burk plot of showing the non-essential activation pattern for L-Met toward dopamine oxidation by CuAβ
enhanced. Herein, both Met and H2O2 can activate dopamine oxidation which are not exclusive of each other and show a combined effect, i.e., the $k_{\text{cat}}$ value of 0.180 s$^{-1}$ in the presence of saturating amount of Met and H2O2 (Fig. 5.4) is a combination of those in the saturating amount of Met (0.088 s$^{-1}$ for Met activation) and H2O2 (0.099 s$^{-1}$ for H2O2 activation), indicating their independent activation mechanisms. As opposed to Met, the reducing agents ascorbic acid and GSH act as inhibitors toward dopamine oxidation by CuAβ (Fig. 5.5), with the former being a competitive inhibitor ($K_i = 66 \mu$M) while GSH a non-competitive inhibitor ($K_i = 53 \mu$M).

The electronic spectrum of CuAβ exhibits an intense charge-transfer transition at 316 nm upon addition of Met (Fig. 5.6), indicative of direct Met binding to the Cu$^{2+}$ center. Such charge-transfer transition is not observed in CuAβ$_{1-40}$, suggesting that Met35 does not interact with the Cu$^{2+}$ center. The d-d transition of CuAβ$_{1-20}$ at 600 nm is not significantly affected in the presence of saturating amounts of Met (Fig. 5.6; inset). The change in the intensity of the charge transfer transition as a function of [Met] can be fitted to the 1:1 binding pattern of [CuAβ + Met $\rightleftharpoons$ Met-CuAβ] to yield a dissociation constant $K_d = 0.25$ mM. This value is comparable to that obtained from the activity profile discussed above, indicating that direct Met binding to the Cu$^{2+}$ center affords the enhancement of CuAβ activity.

The coordination of CuAβ and its binding with Met have been investigated with EPR spectroscopy. The EPR spectrum of CuAβ$_{1-20}$ (Figure 5.7) can be attributed to a typical tetragonally distorted Cu$^{2+}$ center, i.e., an elongation along the z axis due to the Jahn-
Figure 5.4. Rate of dopamine oxidation by CuAβ_{1-20} in the presence of saturating amounts of L-Met and H_{2}O_{2}. 
Figure 5.5. (A) Glutathione (10.0 µM, 20.0 µM, and 40.0 µM from bottom) inhibition and (B) ascorbic acid inhibition (0.0, 0.12, 0.55, and 0.95 mM from bottom) toward dopamine oxidation by Cu-Aβ_{1-20} in 100.0 mM HEPES pH 7.0.
Figure 5.6. Optical titration of L-methionine to 0.2 mM CuAβ. The insets show the low energy d-d transition and the saturation curve fitted to 1:1 ligand:metal stoichiometry.
Teller effect, which can be slightly better fitted to a rhombic than an axial magnetic $g_z = 2.266$, $g_x = 2.057$, and $g_y = 2.080$ and $A_z = 547$, $A_x = 30.6$, and $A_y = 51.1$ MHz (Fig. 5.7) or $g_{//} = 2.268$, $g_{\perp} = 2.064$, $A_{//} = 547$, and $A_{\perp} = 51.1$ MHz. These values are consistent with those reported for several Cu$^{2+}$ complexes of Aβ and fragments, i.e., $g_{//} = 2.265–2.269$, $g_{\perp} = 2.059–2.062$, and $A_{//} = 465–577$ MHz (from 166-206 G). Upon addition of Met, the EPR spectrum changes only slightly to $g_z = 2.257$, $g_x = 2.055$, and $g_y = 2.072$ and $A_z = 564.9$, $A_x = 35.8$, and $A_y = 46.0$ MHz (Fig. 4A) or $g_{//} = 2.257$, $g_{\perp} = 2.059$, $A_{//} = 565$, and $A_{\perp} = 35.8$ MHz, as opposed to the dramatic change in the electronic spectrum.

The Cu$^{2+}$ center in CuA$\beta_{1-20}$ and its interaction with Met are further investigated with pulsed EPR. The X-band ESEEM spectrum of CuA$\beta_{1-20}$ (Fig. 5.8) reveal three $^{14}$N NQI lines at $\nu_0 = 0.33$, $\nu_- = 1.11$ (shoulder), and $\nu_+ = 1.45$ MHz (Eqs. 1 and 2), the double-quantum transitions at ~4 Mz, and the combination lines at 2.33, 2.95, and 3.45 MHz. From which, $e^2qQ$ is obtained to be 1.71 MHz, a value typical of a coordinated His, and $\eta$ calculated to be 0.39 (Eqs. 1 and 2). The $\nu_-$ line (a shoulder) can be better determined once the values of $e^2qQ$ and $\eta$ are determined from $\nu_0$ and $\nu_+$. At least one coordinated water is also revealed which attributes to the deuterium modulation at 2.29 MHz when the spectrum was acquired from a sample in D$_2$O buffer excited at 3391 G (dashed trace, Fig. 5.8), consistent with $^2$H resonance of 2.21 MHz at this field. This small discrepancy may be attributed to the presence of a small super-hyperfine coupling. An ESEEM spectrum for Met-bound CuAβ in D$_2$O buffer was acquired under the same conditions (dotted trace, Fig. 5.8) to investigate the status of the coordinated water upon
Figure 5.7. X-band EPR spectra of CuAβ₁₋₂₀ in the presence and absence of L-Met and simulated spectra with rhombic g tensors (dashed traces).
Figure 5.8. ESEEM spectra of 1.0 mM CuAβ₁⁻²⁰ in 100.0 mM HEPES buffer in H₂O (solid trace) and in D₂O (dashed trace) at pH(D) 7.0 and after addition of 8.0 equivalents Met (dash-dot trace).
Met binding. The $^{14}$N NQI lines change only slightly upon Met binding, found at $\nu_0 = 0.15$, $\nu_\nu = 1.22$ (shoulder), and $\nu_+ = 1.37$ MHz which afford $\epsilon^2 q Q = 1.73$ MHz and a small $\eta = 0.17$. Upon Met binding, the deuterium line is observed at 2.19 MHz when excited at 3351 G, right at the $^2$H resonance frequency at this field which reflects a negligible super-hyperfine coupling.

IV. DISCUSSION

The oxidation of catechol and phenol and their derivatives by CuAβ in the presence and absence of H$_2$O$_2$ was demonstrated to be consistent with the mechanism of type-3 copper oxidases such as di-Cu catechol oxidase (Fig. 5.9).$^{19}$ Therein, the catechol-containing substrate like dopamine binds to a di-Cu$^{2+}$ active center (step $i$) under aerobic conditions and is oxidized via 2-electron transfer to afford a di-Cu$^+$ active center (C) and o-quinone product (step $ii$). The reduced di-Cu$^+$ can bind O$_2$ (step $iii$) to form a $\mu$-peroxo-Cu$^{2+}$ center (D) as demonstrated in both enzyme and chemical model systems, which may bind a substrate and followed by oxidation of the substrate (steps $iv$ and $v$). The presence of a reducing agent such as ascorbic acid (or Met$^{35}$ as previously suggested$^9$) can thus facilitate the aerobic pathway to yield the di-Cu$^+$ active center (C) ready for O$_2$ binding. In the meantime, a reducing agent can also result in H$_2$O$_2$ production through the reduction of the $\mu$-peroxo-Cu$^{2+}$ center (step $vi$). An alternative “short-cut” route, also known as the “peroxide shunt” pathway in heme-containing peroxidase,$^{29}$ for the oxidation to take place is to form the $\mu$-peroxo-Cu$^{2+}$ intermediate
**Figure 5.9.** Proposed mechanism for the catechol oxidase-like activity of CuAβ toward the oxidation of dopamine. X indicates an endogenous ligand or a bound Met.
by direct binding of H$_2$O$_2$ to the di-Cu$^{2+}$ center (step vii), followed by substrate binding (step iv) to form the $\mu$-peroxo-Cu$^{2+}$$_2$-substrate ternary complex (E) which then undergoes oxidation. The ternary complex E can also be formed upon H$_2$O$_2$ binding to the substrate-bound intermediate (B; step viii), representing a random bi-substrate mechanism.

Met$_{35}$ in A$\beta$ has been suggested to be a reducing agent$^9$ responsible for the initiation of the redox cycling of the Cu$^{2+}$ center in CuA$\beta$ and leads to H$_2$O$_2$ production (i.e., step vi).$^5$ The oxidation of the thioether moiety of Met to its sulfoxide form in A$\beta$ has been implicated in aggregation, lipid peroxidation, and redox chemistry in association with the metal center.$^9$ The activity profile of [Met] (Fig. 5.1) and the optical spectrum (Fig. 5.6) indicate direct Met binding to CuA$\beta$, rather than outer-sphere interaction, which potentially can reduce the Cu$^{2+}$ center as proposed previously. Herein, the amount of di-Cu$^{+}$ (C) increases which in turn forms a larger amount of the $\mu$-peroxo-Cu$^{2+}$$_2$ center upon O$_2$ binding (D; step iii), but fewer amount of di-Cu$^{2+}$ (A) would be present for substrate binding as it is reduced by Met. However, once CuA$\beta$ (A) is reduced, it cannot follow the peroxide shunt pathway anymore. The observation that H$_2$O$_2$ can still significantly enhance the activity in the presence of a saturating amount of Met (Fig. 5.2) suggests the Cu$^{2+}$ center is not significantly reduced by Met, which is demonstrated by the detection of the S = 1/2 EPR features (Fig. 5.7). Moreover, the difference in electrode potentials between Cu$^{2+}$-A$\beta$ and Met is 0.71 V,$^{30}$ which gives a dramatic 68.5 kJ Gibbs free energy that is equivalent to a negligibly small equilibrium constant of 9.8 x 10$^{-15}$ for one-electron reduction of Cu$^{2+}$-A$\beta$ by Met at 298 K.
The non-exclusive nature of Met and H₂O₂ binding and their additive activations indicate that there are two pathways for the oxidation of dopamine in the presence of Met and H₂O₂, presumably the pathways \textit{i}–\textit{v} and \textit{vii}–\textit{iv}–\textit{v} with a bound Met (the \textit{“X”} ligand in Fig. 5.9), wherein the enhancement of the activity with Met alone is supposed to be due to a non-redox mechanism that fine-tunes the reduction potential of the Cu²⁺ to favor the oxidative catalysis. The sigmoidal activity profiles for dopamine oxidation in the presence of Met (Fig. 5.1) and from the concerted action of both H₂O₂ and Met (Fig. 5.4) reflect possible presence of cooperativity, such as the formation of the dinuclear active center \textbf{D}. Despite the lack of a Met and any redox-active amino acid, the fragments CuA₁–16 and CuA₁–20 exhibit significant metal-centered oxidative activity\textsuperscript{18–20} which indicates the redox role of Met\textsubscript{35} might have been overstretched.

The oxidation state of the Cu center in CuA₁–20 upon Met binding can be concluded from the electronic and EPR spectra, wherein the remaining d-d transition at 600-nm (Fig. 5.6) and EPR features (Fig. 5.7) indicate that Cu²⁺ is not reduced by Met as opposed to previous suggestions for the reductive role of Met\textsubscript{35} in CuA₁–40.\textsuperscript{9} Moreover, the charge transfer transition is consistent with a thio-to-Cu²⁺ charge-transfer transition observed in Cu²⁺-methionine complexes.\textsuperscript{31} The well resolved EPR spectral features at g~2 (Fig. 5.7) upon Met binding confirm the d⁹ electron configuration of the Cu²⁺ center with S = 1/2. The lack of \( \Delta M_S = \pm 2 \) transition (i.e., between the \( M_S = -1 \) and +1 levels in an antiferromagnetically coupled di-Cu²⁺ center) in the EPR spectrum at low field indicates magnetic coupling between the Cu²⁺ centers may not exist. Thus, the previously proposed His-bridged dinuclear Cu,Zn-superoxide dismutase center for
CuAβ_{1−28}^{32} is not present in the case herein. The small change in the EPR spectrum upon Met binding indicates the charge transfer transition must be due to the binding of the thioether of Met through the non-magnetic d_{z^2} orbital at an axial position of CuAβ_{1−20}. Otherwise, more significant changes in the g and A tensors would be observed as reported for the binding of thio-groups to Type-2 Cu proteins.\textsuperscript{33}

The ESEEM spectrum of CuAβ_{1−20} (Fig. 5.8) confirms the binding of Cu^{2+} to Aβ through His side chains via the magnetic d_{x^2−y^2} orbital in equatorial positions which gives rise to the quadrupole coupling with the remote non-coordinated $^{14}$N on the coordinated His imidazole ring. There are at least two coordinated His side as reflected by the combination lines since a single coordinated His does not give rise to these lines.\textsuperscript{33g,34} The small $\eta$ values indicate that the Cu$^{2+}$ center is only slight rhombic, which has already been observed in the CW EPR spectra (Fig. 5.7). The detection of a deuterium line at 2.29 MHz suggests the presence of coordinated water (as D$_2$O), presumably in an equatorial position via the magnetic d_{x^2−y^2} orbital.\textsuperscript{27} This signal is not much affected upon Met binding, once again suggesting Met binding to an axial position. It is interesting to note that the double-frequency peak at 4.6 MHz vanishes upon Met binding. This indicates that at least one of the weakly coupled deuterium atoms in the vicinity of the Cu$^{2+}$ is replaced by the Met ligand.

Since the binding of Met to CuAβ does not reduce the Cu$^{2+}$ center, the enhancement in activity must be attributed to a change in the reduction potential of the Cu$^{2+}$ center. The axially coordinated Met ligand in blue copper proteins has been suggested to play a role in controlling the reduction potential of the protein since mutation of this Met results
in a significant change in the potential. Based on the results herein, I propose the binding of methionine to Cu$^{2+}$ in CuA$\beta$ may modulate the Cu$^{2+}$/Cu$^{+}$ potential to favor the redox catalysis, yet is not necessarily involved directly in the redox chemistry as a reducing agent as previously proposed. The oxidation of Met35 previously observed may follow the metal-centered mechanism as a catechol or phenol substrate via the reactive di-Cu$^{2+}$-peroxo intermediate to result in oxygenation reaction, ie., the coordinated Met “X” serves as a substrate via step iv and v in the mechanism without the catechol substrate (Fig. 5.9).

The catalytic pathway of CuA$\beta$ is altered under reduction conditions (Fig. 5.9), wherein the reaction is locked into a H$_2$O$_2$-producing cycle (iii and vi). The competitive inhibition of ascorbate toward the oxidation of dopamine by CuA$\beta$ (Fig. 5.5A) may be because of possible chelation and reduction of the Cu$^{2+}$ center by ascorbate. GSH may bind to and reduce the Cu$^{2+}$ center as a monodentate ligand which does not prevent substrate from binding to the Cu$^{2+}$ center to form the inactive inhibitor-CuA$\beta$-substrate complex, thus exhibiting non-competitive pattern (Fig. 5.5B). The inhibitory effect of the reducing agents toward metal-centered catalysis is consistent with the proposed mechanism wherein H$_2$O$_2$ is generated (step vi), also reported previously. Thus, the ability of CuA$\beta$ to oxidize biologically relevant molecules such as dopamine is highly dependent on the redox state of its environment. Although reducing agents can potentially inhibit the oxidation of neurotransmitters by CuA$\beta$, the production of H$_2$O$_2$ may still exacerbate the situation of oxidative stress.
Metabolic malfunctions of catecholamines neurotransmitters have been suggested to be related to the neuropathology of AD. Of these neurotransmitters, dopamine has been directly linked to the neurodegenerative Parkinson’s disease and has been proposed to be associated with adult neurogenesis in the subventricular zone. I discuss in chapter IV that CuAβ can catalyze oxidation of catecholamine neurotransmitters such as dopamine to generate neurotoxic quinone products. Dopamine quinone can result in polymerization of tau protein into fibrils and covalent modification of dopamine transporter which directly affects dopamine uptake. A possible mechanism for reduction in the delay-period activity and short-term memory, lack of attention, and change in mood and motivation experienced by AD patients may probably be partially due to long-term oxidation of dopamine by CuAβ. Hence, the acceleration of dopamine oxidation via metal-centered mechanism that can be modulated by small molecules such as reducing agents, H₂O₂, and Met may well hint at the significant role of CuAβ in oxidative stress in the brain of AD patients.

V. CONCLUDING REMARKS

Taken together, the results herein present more structural information about the metal center and further support for metal-oxygen/peroxo-centered redox chemistry of CuAβ and provide additional mechanism for the oxidative stress in the brain of AD patients. The role of Met35 has been redefined and the dual mechanistic character of reducing agents (i.e., inhibition and H₂O₂ generation) in the redox cycle of CuAβ is
clearly defined. The overall picture of AD neuropathology is likely to be composed of the pieces of information uncovered thus far, including generation of ROS, metal-dependent aggregation of Aβ, and the largely overlooked metal-centered degradation of biomolecules. Treatment and prevention strategies hence must address all of these pathways, including inhibitions toward H₂O₂ production and oxidative damage of neurotransmitters and other biomolecules by the di-copper-peroxo active center.
VI. LIST OF REFERENCES


Structural implications derived from the analysis of electron paramagnetic resonance
spectra of natural and artificial copper proteins. *Arch. Biochem. Biophys.* (1974), 165,
691–708. (g) F. Jian, J. Peisach, L.-J. Ming, L. Que Jr., V.J. Chen, Electron Spin
Echo Envelope Modulation Studies of the Cu(II)-Substituted Derivative of
Isopenicillin N Synthase, a Structural and Spectroscopic Model. *Biochemistry* (1991),
30, 11437–11445.

34) S.S. Eto, J. Dubach, G.R. Eaton, G. Thurman, D.R. Ambruso, Electron Spin Echo
Envelope Modulation Evidence for Carbonate Binding to Fe(III) and Cu(II)

35) (a) H. Li, S.P. Webb, J. Ivanic, J.H. Jensen, Determinants of the relative reduction
potentials of type-1 copper sites in proteins. *J. Am Chem. Soc.* (2004), 126, 8010-
8019. (b) J.F. Hall, L.D. Kanbi, R.W. Strange, S.S. Hasnain, Role of the axial ligand
in type 1 Cu centers studied by point mutations of Met148 in rusticyanin.
*Biochemistry* (1999), 38, 12675-12680.

36) (a) W. Fu, H. Luo, S. Parthasarathy, M.P. Mattson, Catecholamines potentiate
amyloid β-peptide neurotoxicity: involvement of oxidative stress, mitochondrial
dysfunction, and perturbed calcium homeostasis. *Neurobiol. Disease* (1998), 5, 229-
243. (b) J.I. Friedman, D.N. Adler, K.L. Davis, The role of norepinephrine in the
pathophysiology of cognitive disorders: potential applications to the treatment of


CHAPTER VI. THE ASTACIN FAMILY OF ENDOPEPTIDASES AND EMBRYOGENESIS

I. INTRODUCTION

The astacin family of zinc-dependent endopeptidases is a class of enzymes ubiquitously distributed across all phyla and part of the superfamily of metzincins.\(^1\) Approximately thirty members of the astacin family have been characterized at the protein level, including meprins, bone morphogenetic protein-1 (BMP-1), and tolloid while several others have been identified through gene sequencing, including a large number in *Caenorhabditis elegans*.\(^2\) The signature active site sequence for this family of enzymes is \textbf{HEXXHGXXHEXXRxDR} (Figure 6.1), where one Zn\(^{2+}\) atom coordinates to three histidines, a tyrosine, and a water molecule (Figure 6.2).\(^3\) Most members share common expression features such as the pre- and pro-enzyme sequences immediately located NH\(_2\)-
Figure 6.1. Active site motif common to all astacins.$^2$
Figure 6.2. Relaxed eye stereo view of the active-site structure of astacin with transition state analogue (PDB ID 1QJI).
terminal to the protease domain. Several members contain one or two copies of epidermal growth factor EGF-like domains, and complement-like domains (Clr, Cls) near the COOH-terminus. The shuffling of different domains in relation to the catalytic (protease or astacin) domain creates a variety of proteins with several different structures and functions.

Embryo development is an area which remains unresolved in a number of more complex animals, including human embryogenesis and other representatives of subphylum vertebrata due to the inherent complexities of deuterostome embryogenesis. The challenge of resolving proteolytic signaling, that is likely responsible for modification of the extra cellular matrix (ECM), important in embryogenesis events, has partially been due to the lack of convenient model systems; a particular challenge is the isolation of ECM components in pure form in mice and human models. Sea-urchins hence are a good model system for the study of morphogenesis due to the facility of isolation of pure and large amounts of protein components of the ECM. Another advantage of the sea-urchin model is that its genome has recently been sequenced which will facilitate the connection of upstream genetic events with its downstream message and proteolytic mechanisms. The presence of several astacin family enzymes in the development of the sea-urchin embryo (i.e. suBMP, SpAN, envelysin, and BP10) makes it a target to study similar proteolytic processes in higher organisms, especially those containing enzymatic processes analogous to tolloid and BMP-1, two enzymes proposed to be similar to BP10 in structure. BP10 is an interesting enzymatic model system because it contains similar
Figure 6.3. Phylogeny of the astacin family catalytic domains. 6
domains as other complex astacins such as tolloid and BMP-1, but is unique in domain arrangement which will be discussed further.\textsuperscript{5}

Originally characterized by the use of immunoblotting techniques and sequence analysis of the gene and m-RNA transcript,\textsuperscript{5,6} BP10 has remained an unstudied member of this class of enzymes. The focus of research has shifted to other members of the astacin class of enzymes (astacin itself being a crayfish digestive enzyme and hence only a novel prototype in catalytic mechanism) such as BMP-1 and tolloid due to their potential role in unraveling developmental information about human embryos (BMP-1) and the facility of handling a well established model system (tolloid from \textit{Drosophila}). However, from an evolution standpoint, echinoderms (Echinodermata, the deuterostomes) are more closely related to humans than a fruitfly (Arthropoda, the ecdysozoans).\textsuperscript{7} BP10 hence is a good candidate as a functional model system of BMP-1 and tolloid. The transcription of the BP10 gene is transiently activated around the 16- to 32-cell stage and the accumulation of BP10 m-RNA is limited to a short period at the blastula stage. Temporarily, the highest BP10 activity is detected approximately 1.5 hours after expression of the sea urchin hatching enzyme (envelysin) reaches a maximum.\textsuperscript{5} The BP10 transcripts are spatially expressed and only detected in a limited area of the blastula in the animal half of the embryo. The protein is first detected in early blastula stages. Its level reaches a peak in late cleavage, and declines abruptly before ingress of primary mesenchyme cells and remains constant in late development.\textsuperscript{5} The likely role of zymogen activator has been assigned to BP10, since the presence of an EGF domain is a highly conserved motif in proteolytic cascades or activation of precursors.\textsuperscript{8} However, it was
accurately described\textsuperscript{5} that blocking BP10 activity prior to hatching with the use of an antibody resulted in deformed embryos, whereas after hatching the embryos developed normally when treated with the same antibody, though basal levels of the enzyme can be detected after hatching.

The morphogenetic studies conducted by Runnström and Horstadius determined that developmental processes occur across a gradient in where the ectodermal structures are controlled by the animal axis and the mesodermal (skeleton) is controlled by the vegetal pole. This established model has been challenged by recent studies.\textsuperscript{9,10} Hence it becomes clear that sole analysis of gene structure and transcriptional levels is not enough to provide irrefutable evidence about animal development. A multi-disciplinary approach is needed where the skills of a developmental biologist and the detailed information about structure/function of enzymes that a biochemist can provide are needed in order to fully understand such questions.

The BP10 protease is constructed of identical domains as BMP-1 and tolloid, and is described as a factor involved in cell differentiation in mid to late blastula. Human BMP-1 has been characterized as a splice variant of the tolloid gene which is required for correct dorsal-ventral patterning of the \textit{Drosophila} embryo. In preliminary studies, BP10 has not been shown to affect axis formation; hence the structure/function paradox does not hold true for BP10, BMP-1, and tolloid. How do similar global mechanisms of development differ at the molecular level? What are the mechanisms of evolution
Figure 6.4. Domain organization of proteins from the astacin/CUB/EGF subfamily.
involved in animal development across the deuterostomes? And finally, what are the regulatory networks during embryogenesis in humans, sea-urchins, and *Drosophila*?

BMP-1 has been assigned the role of a procollagen C-protease and its minimal domain structures necessary for activity and secretion have been identified. However BMP-1 is also able to cleave non-collagenous substrates such as dentin and laminin, hinting at a wide distribution of biological regulation factors that may be attributed to this enzyme. BP10 may serve a similar purpose in the sea urchin embryo. The developmental regulation of BP10 expression shows that it is an important and necessary event in embryogenesis. Recent studies have also shown that another astacin family enzyme found in sea urchins, SpAN, is able to regulate BMP signaling. BP10 may be part of the signaling events which involve the enzymes suBMP, SpAN and BP10 itself, or perhaps play the role as a less discriminate activator of other important molecular events.

Since BP10 is homologous to tolloid and BMP-1 (Figure 6.4), it is possible that the sea urchin enzyme is part of a regulatory set of interactions with polypeptides of the TGF-β family. The involvement of BP10 in a process similar to that of BMP-1 is unlikely and most likely resembles tolloid in function. However, the high similarity in sequence and structural motifs makes BP10 an important comparison model in a lower species. The mammalian BMP-1-related proteases are all capable of activating the TGF-β protein growth differentiation factor (myostatin), by freeing it from a noncovalent latent complex with its cleaved prodomain. Similarly, BMP-1 and the splice variant mTLL-1 but not tolloid or TLL-2 is able to free the TGF-β morphogens BMP-2 and BMP-4 from latent complexes with the extracellular antagonist chordin. Thus, BMP-1/tolloid-like
proteases may orchestrate formation of the ECM with signaling by various TGF-β-like proteins in morphogenetic events.

BP10 is also a unique system that may contribute important information about the relationship of structural features such as EGF-Ca^{2+} binding domain and its CUB domains and the catalytic domain of these proteases (Figure 6.2). It is unique to BP10 that the EGF-Ca^{2+} domain is located between the catalytic domain and the proposed regulatory sequences of CUB1 and CUB2. Often, these EGF-Ca^{2+} domains are located between CUB sequences. CUB domains have been implicated both in activity and regulation in BMP-1.\textsuperscript{11} The kinetic properties of BP10 should then compare and contrast nicely with those of other BMP-1 and tolloid-like proteins. Ca^{2+} may have a synergistic effect on the reaction catalyzed by BP10 due to its close proximity to the catalytic domain.

From a protein evolution viewpoint, BP10 contains modules proposed to be involved in the domain shuffling (i.e evolution through exon shuffling). Characterization of BP10 with simultaneous analysis of its gene structure will add further insight into this mechanism. Moreover, once other sea urchin enzymes involved in embryogenesis, such as SPAN and suBMP are overexpressed and characterized, and their gene structures fully characterized, a better model of module domain shuffling protein evolution can be presented, since BP10, SPAN and suBMP are part of the same family of enzymes, acting in the same organism, and involved in some functional role during embryogenesis.

In the case of astacin, BMP-1, tolloid, and BP10, the presence of a metal-coordinated tyrosine is a rather unusual mechanistic motif due to the reduction of Lewis
acidity induced by the coordination of a negatively charged phenolate to the metal center. This is not the case in astacin where the coordination of the Tyr-phenolate under physiological conditions does not affect its catalysis under neutral conditions. A metalloctriad mechanism has been proposed based on kinetic, optical, and EPR studies on the Cu\(^{2+}\) derivatives of astacin and serralysin. Further analysis of the physical characteristics of astacin-type enzymes using spectroscopically-active metal substitutes in concert with classical enzyme kinetics (i.e. pH profiles, inhibition studies, etc.) analysis of splice variants of recombinant BP10 (whole enzyme and astacin domain truncation) will add further insight into the catalytic mechanism of these enzymes. Concomitant with physical studies on the native system, active site mutants will allow for comparison both in kinetic parameters and the physical properties of BP10. Previous site directed mutagenesis studies have been conducted; however, the choice of mutations have shed no actual mechanistic insight toward the catalytic mechanism of astacins, other than establishing that the conserved glutamate and tyrosine residues are catalytically important. The status of the coordinated tyrosine has been proposed to be an inhibitory process in the metal centered hydrolysis of peptide bonds. In this proposed metalloctriad mechanism (Figure 6.5), the active-site Zn\(^{2+}\) coordinated by three His, tyrosine, and a water molecule can be activated via detachment of the phenolate with a concomitant
Figure 6.5. Proposed metallotriaid mechanism in astacin and serralysin.²²
general base deprotonation via a glutamate residue. The metal bound OH⁻ is able to
attack and hydrolyze the scissile bond after Zn²⁺ creating electrostatic strain in the
peptide bond by interaction with the carbonyl of the scissile bond. The confirmation of
this mechanism can shed valuable insight into numerous enzymatic processes across all
members of the astacin family, from the numerous astacin-like enzymes in *C.elegans* to
the astacin/EGF/CUB members in higher organisms.
IV. LIST OF REFERENCES


CHAPTER VII. OVEREXPRESSION AND CHARACTERIZATION OF
BLASTULA PROTEASE 10 (BP10) FROM PARACENTROTUS LIVIDUS

I. INTRODUCTION

Blastula Protease 10 (BP10) is a metalloenzyme assigned to the astacin family of Zn-dependent endopeptidases involved in sea urchin embryogenesis. It contains conserved structural motifs consistent with astacin, tolloid, and bone morphogenetic protein 1 (BMP-1). Astacin, a gut enzyme, and serralysin, a bacterial enzyme, have been proposed to carry out hydrolysis via a “metallotriad” mechanism that involves a metal-coordinated tyrosine. It has not been determined if the more structurally complex members of this family involved in eukaryotic development share this mechanism. The recombinant BP10 has been overexpressed in E.coli, its metalloenzyme nature confirmed, and its catalytic properties characterized through kinetic studies. BP10 shows significant

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1 This work has been published: G.F.Z. da Silva, R.L. Reuille, L.-J. Ming, B.T.Livingston J. Biol. Chem. 2006 281, 10737-10744.
hydrolysis toward gelatin both in its native Zn-containing form and copper derivative.
The copper derivative of BP10 shows a remarkable 960 % rate-acceleration toward the hydrolysis of the synthetic substrate N-benzoyl-arginine-p-nitroanilide when compared to the Zn form. The enzyme also shows calcium-dependent activation. These are the first thorough mechanistic studies reported on BP10 as a representative of the more structurally complex members of astacin-type enzymes in deuterostomes which can add supporting data to corroborate the metallotriad mechanism proposed for astacin.

The astacin family of zinc-dependent endopeptidases is a class of enzymes ubiquitously distributed across all phyla and part of the superfamily of metzincins.\textsuperscript{1} Approximately thirty members of the astacin family have been characterized at the protein level;\textsuperscript{2} including meprins, bone morphogenetic protein-1 (BMP-1), and tolloid, while several others have been identified through gene sequencing, including a large number in \textit{Caenorhabditis elegans}.\textsuperscript{3} The signature of the primary sequence active site motif for this family of enzymes is $\text{HEXXHGFXHEXXRXDR}$, where one $\text{Zn}^{2+}$ ion coordinates with three histidines, a tyrosine, and a water molecule.\textsuperscript{4, 5} Most members of this family share common domain structures such as the pre- and pro-enzyme sequences immediately located N-terminal to the protease domain. Several members contain one or two copies of epidermal growth factor EGF-like domains, and complement-like domains (Clr, Cls) near the C-terminus.\textsuperscript{2} The shuffling of different domains in relation to the catalytic protease domain creates a variety of proteins with different structures and functions.
Originally isolated and characterized as a developmentally regulated gene in sea urchin embryos, BP10 protein has remained uncharacterized. It shares sequence similarity with other members of the astacin family of enzymes. The simplest member of the family, astacin from crayfish digestive fluid is a digestive enzyme and hence a novel prototype in catalytic mechanism important in development such as BMP-1 in vertebrates and tolloid in *Drosophila*. Whereas BP10 is a good candidate as a functional model system of BMP-1 and tolloid. The BP10 protease is constructed of identical domains as BMP-1 and tolloid, but has different arrangement of these domains. The transcription of the BP10 gene is transiently activated around the 16- to 32-cell stage and the accumulation of BP10 mRNA is limited to a short period at the blastula stage. Temporarily, the highest BP10 activity is detected approximately 1.5 hours after expression of the sea urchin hatching enzyme (envelysin) reaches a maximum. The BP10 transcripts are detected in a limited area of the blastula. The protein is first detected in early blastula stages, its level peaks in late cleavage, declines abruptly before ingestion of primary mesenchyme cells, and remains invariable in late development. A likely role of zymogen activator has been suggested for BP10, since the presence of an EGF domain is a highly conserved motif in proteolytic cascades or activation of precursors. Blocking BP10 activity prior to hatching with the use of an antibody resulted in abnormal embryos.

BP10 has a unique arrangement of structural features such as EGF-Ca$^{2+}$ binding domain, two adjacent CUB domains, and a catalytic domain that is highly homologous to astacin. In particular, the EGF-Ca$^{2+}$ domain is located between the
catalytic domain and the proposed regulatory CUB sequences. More often, these EGF-Ca\(^{2+}\) domains are located between CUB sequences. CUB domains have been implicated in activity and regulation in BMP-1.\(^9\) Astacin family enzymes and serine proteases have been implicated in remodeling the pericellular space in sea urchin embryos, which is composed of the extracellular matrix and transmembrane proteins.\(^10,11\) Moreover, several studies have reported gelatinase and collagenase activities from enzymes located in the sea urchin egg and embryo which were characterized as metalloenzymes due to inactivation with EDTA and 1, 10-phenanthroline.\(^12-15\)

Beside the interesting distribution of astacin-like enzymes across phyla and the numerous functional roles of these enzymes, mechanistic questions about these highly conserved hydrolases domains still remain to be answered. Within the metzincin superfamily of enzymes, minor differences of active site function have been observed, which are likely to account for different substrate specificities.\(^16\) In the case of astacin as well as BMP-1, tolloid and BP10, a metal-coordinated tyrosine is a rather unusual metal-binding motif due to the reduction of Lewis acidity induced by the coordination of the negatively charged phenolate to the metal center.\(^17,18\) However, the coordination of the Tyr-phenolate does not seem to affect astacin catalysis under neutral conditions. The status of the coordinated tyrosine has been proposed to be an inhibitory process in the metal centered hydrolysis of peptide bonds.\(^5\) A metallotriad mechanism has been proposed for astacin and serralysin based on kinetic and spectroscopic studies of the native enzymes and their Cu\(^{2+}\) derivatives.\(^5,19\) In the proposed metallotriad mechanism, the active-site Zn\(^{2+}\) coordinated by three His, a tyrosine, and a water molecule can be
activated via detachment of the Tyr-phenolate with a concomitant deprotonation of the
coordinated water assisted by a glutamate residue. The metal-bound OH\(^-\) is able to
hydrolyze the scissile bond with Zn\(^{2+}\) creating electrostatic strain in the peptide bond by
interaction with the carbonyl group of the scissile bond. I discuss in this chapter the
overexpression and thorough mechanistic study of recombinant BP10, a model system
for astacin-type developmentally regulated metalloenzymes. Further analysis of BP10
will add insight into the catalytic mechanism of members of the astacin family of
enzymes, and the degree to which the mechanism is conserved among the enzymes
found in deuterostomes.

II. EXPERIMENTAL

The expression vector pQE30Xa, Ni-NTA agarose, mouse anti-His primary
antibody were from Qiagen (Valencia, CA), XL1-Blue chemically competent \textit{E. coli}
from Invitrogen (Carlsbad, CA), Rosetta Blue chemically competent \textit{E. coli} and Factor
Xa removal kit were from Novagen (San Diego, CA), all primers were from Integrated
DNA Technologies (Coralville, IA), all modifying and restriction enzymes were from
Promega (Madison, WI), Eppendorf Perfect plasmid preparation kit was from Eppendorf
(Westbury, NY), BM purple phosphatase substrate was from Roche (Indianapolis, IN),
EDTA, ZnCl\(_2\), Cu(NO\(_3\))\(_2\), Ca(NO\(_3\))\(_2\), glycerol, ninhydrin, guanidine hydrochloride,
bovine serum albumin, sodium dodecyl sulfate, Triton X-100, Tween 20, imidazole,
NaH\(_2\)PO\(_4\), Na\(_2\)HPO\(_4\), Tris-HCl, acrylamide, bis-acrylamide, TEMED, ammonium
persulfate, NaN₃, dimethyl sulfoxide (Me₂SO), sodium citrate, acetic acid, guanidine hydrochloride, and propanol were from Fisher (Swanee, GA). Type A porcine gelatin 300 bloom, N-benzoyl-arginine-\(p\)-nitroanilide (BAPNA), urea, isopropyl-\(\beta\) thiogalactopyronoside (IPTG), phenylmethyl sulfonyl fluoride (PMSF), benzamidine, urea, lysozyme, bicinechinonic acid, arginine-hydroxamate, and HEPES, CAPS, TAPS, and MES buffers were from Sigma-Aldrich (St. Louis, MO), 1,10-phenanthroline was from Acros (Fairlawn, NJ). All reagents were of enzyme or molecular biology grade when available, all others were reagent grade. All glassware and plasticware were extensively rinsed with EDTA to remove metal contaminants and thoroughly washed with 18 MΩ water to remove the chelator. All buffers contained Chelex resin to remove metal contaminants. All spectrophotometric measurements were performed on a Varian Cary 50 Bio-Spectrophotometer equipped with a PCB-150 water Peltier thermostable cell holder.

Overexpression, purification, and refolding of recombinant BP10: The cDNA coding for *Paracentrotus lividus* BP10 cloned into the pBluescript plasmid (pBP10) was a generous contribution from Christian Gache and Thierry Lepage (Unité de Biologie Cellulaire, Center National de la Recherche Scientifique et Université de Paris VI, Station Marine, 06230 Villefranche-sur-Mer, France). PCR primers coding for both 5’ and 3’ regions were designed according to the proposed full length BP10 to subclone the cDNA into the pQE30Xa overexpression vector. The 5’ primer: 5’-PO₄-AAAATACCTTTCCCTTTGATGT-3’ codes for the first 9 codons in the proposed nucleotide sequence in BP10 and is 5’ phosphorylated for blunt-end cloning.
using the *Stul* restriction site in pQE30Xa; 3’ primer was designed for cloning into the *Xma*I restriction site 5’-**AATTCCGGGTTA**GTTCA\text{GACGAGGATCTC GGGT-3’} (bold = extra base pairs for melting temperature optimization, underlined = *Xma*I restriction site, bold underlined = stop codon). The PCR product coding for BP10 was digested with *Xma*I and cloned into the pQE30Xa vector. The BP10 construct was transformed into Rosetta Blue competent cells. The colonies overexpressing recombinant BP10 were screened using a colony lift protocol according to Qiagen without modifications, where the production of BP10 was monitored using a mouse anti-His tag primary antibody. The active colonies were picked, propagated in liquid media to OD\text{600} = 0.4. IPTG was added to a final concentration of 1.0 mM and the culture grown at 30 °C and 300 rpm for 4.5 hours. The bacteria containing recombinant BP10 were pelleted at 4,000×g at 4 °C and resuspended in cell wall lysis buffer (50.0 mM NaH\text{2PO4}, 100.0 mM NaCl, 10.0 mM imidazole, 2.0 mM benzamidine, 2.0 mM PMSF, pH 8.0) containing 1.7 mg/mL lysozyme and incubated on ice lightly shaking for 60 minutes, then sonicated 6×10 seconds with 10 second intervals. The cultures were pelleted at 10,000×g at 4 °C for 20 minutes and the inclusion bodies were resuspended in an urea buffer (8.0 M urea, 10.0 mM Tris, 100.0 mM NaH\text{2PO4}, 1.0 % Triton X-100, 2.0 mM benzamidine, 2.0 mM PMSF, pH 8.0) and incubated at 37 °C at 200 rpm for 60 minutes. The solubilized inclusion bodies were pelleted at 10,000×g at 4 °C for 20 minutes, and 1.0 mL of Ni-NTA agarose was added to the supernatant. The Ni-NTA slurry was gently shaken at room temperature for 45 minutes then added to a gravity fed column and the recombinant BP10 eluted using a pH gradient, with buffers containing 6.0 M urea, 10.0 mM Tris,
100.0 mM NaH₂PO₄, at pH’s 6.3, 5.9, and 4.5. The recombinant BP10 was completely eluted at pH 4.5 and was immediately titrated to pH 7.4 using 0.5 M NaH₂PO₄. Overexpression and purification were monitored on a time-dependent basis using 12.5% SDS-PAGE according to Laemmli and Western blot techniques using the anti-His-tag antibody.

The concentration of recombinant BP10 was determined using standard BCA assay with a BSA standard curve. The recombinant protein was diluted 40 times by volume using 50.0 mM Tris with 50 mM NaCl then dialyzed extensively with several changes against phosphate buffered saline (PBS) containing 50 μM ZnCl₂ for 48 hours at 4 °C. Recombinant BP10 was concentrated either under 18 psi N₂ using an YM3 Amicon membrane or an Amicon Centricon YM3. Final BP10 concentrations were checked using BCA. The His-tag fusion was removed using a Factor Xa His-tag removal kit from Novagen according to instruction.

**Circular Dichroism (CD) studies:** CD spectra of urea-denatured, and folded Zn-BP10 and Cu-BP10 were collected in PBS using a 0.1 cm cell with a resolution of 0.5 nm. All absorbance readings were converted to molar ellipticity and the percent helicity and sheet content calculated. The α-helical content was calculated according to published methods.²⁰

**Preparation of the copper derivative of BP10:** During the refolding of urea-denatured BP10, 1.0 mM 1,10-phenanthroline was added to the 50.0 mM Tris 50.0 mM NaCl pH 7.5 buffer, then extensively dialyzed against PBS containing 0.5 M guanidine. The guanidine-containing buffer was exchanged through dialysis with PBS buffer, and
then with PBS containing 50 μM Cu(NO₃)₂. Protein concentrations were determined with BCA assay.

Gelatin Zymogram: Gelatin was incorporated into a polyacrylamide gel matrix according to standard protocols with modifications to fit current studies as listed below.²¹ A volume of 1.25 mL of 1.4 M Tris at pH 8.8, 0.50 mL of 5.0 mg/mL gelatin solution in water, 25.0 μL 10 % (w/v) APS, 200 μL 10% (w/v) SDS, 25.0 μL TEMED, 2.0 mL water, and 1.25 mL 30:1 acrylamide:bis-acrylamide were mixed and allowed to polymerize in a mini-gel minus a stacking gel. Several concentrations of BP10 were mixed with non reducing gel-loading dye and incubated for 15 minutes at room temperature (standard Laemmli protocol minus mercaptoethanol or dithiothreitol²²). These BP10 samples were loaded into each lane of the gel and run at 200 V and 4 °C until the dye front reached the bottom of the plate. The running buffer did not contain SDS. The gel was washed two times in 0.25 % Triton X-100 for 15 minutes with gentle shaking, then incubated for 10 hours in 50.0 mM Tris pH 7.50, 1.0 μM ZnCl₂ 0.5% Triton X-100, 0.02 % NaN₃, and 2.0 mM CaCl₂. After ten hours the gels were stained with 0.1% Coomassie brilliant blue in 40% propanol for 1.0 hour. The gel was destained in a 7% acetic acid solution to reveal the digestion, and then photographed on a light table with a digital camera. The enzyme with gelatinase activity is shown as unstained bands.

Gelatinase Assay: The gelatinase activity of recombinant BP10 was monitored using a detection method for α- amino groups with ninhydrin as an indicator of peptide hydrolysis according to a standard protocol with modifications to fit current studies as
described below. A 5.0 mg/mL gelatin solution was prepared in H2O and heated to 55 °C for 15 minutes until completely dissolved. A ninhydrin detection solution was prepared by mixing 9.0 mL glycerol, 3.0 mL of 0.5 M sodium citrate at pH 5.50, and 3.75 mL of 1.0% (w/v) ninhydrin solution in 0.5 M sodium citrate buffer. Gelatin and 1.0 µM BP10 were mixed in PBS and incubated at room temperature. A 50.0 µL sample was taken from the reaction at several time points and mixed with 950 µL of ninhydrin detection solution then boiled for 12.0 minutes. The absorbance at 570 nm was determined using a sample containing undigested stock gelatin (the same gelatin used for the experiment, incubated under the same conditions minus BP10) mixed with the same detection assay as the blank. The first order rate constant $k_{obs}$ was determined from an exponential curve fit. The molar absorptivity of Ruhemann’s purple ($\varepsilon_{570} = 22,000$ M$^{-1}$cm$^{-1}$) also allowed for monitoring the substrate dependent hydrolysis of gelatin. Several dilutions of a 10.0 mg/mL stock solution of gelatin were used according to the aforementioned protocol and rates were fitted as a function of substrate concentration according to the Michaelis-Menten equation, yielding $k_{cat}$ and $K_m$ parameters.

*Hydrolysis of BAPNA by Zn-BP10 and Cu-BP10*: BAPNA stock solutions were made in Me2SO and then diluted with 50.0 mM HEPES 50.0 mM NaCl pH 7.50. Less than 2% Me2SO by volume was present in each assay and found not to interfere with kinetic measurements. Several concentration of BAPNA were incubated with 2.17 µM BP10 and rates determined colorimetrically from the release of the p-nitroaniline product ($\varepsilon_{405} = 10,150$ M$^{-1}$cm$^{-1}$). Kinetic parameters were determined by non-linear fitting to the Michaelis-Menten equation.
Calcium activation assays: Gelatin was extensively dialyzed against an EDTA solution and then extensively dialyzed to remove the chelator. Calcium was carefully titrated under substrate saturation conditions to determine its effect on the hydrolysis of gelatin and BAPNA. Once saturating concentrations of calcium were determined, new kinetic parameters were obtained using sufficient (1.0 mM) calcium in all buffers.

Inhibition studies: the effect of two inhibitors, 1,10-phenanthroline (OP) and arginine-hydroxamate (Arg-NHOH) toward the hydrolysis of BAPNA were determined by running Michaelis-Menten kinetics under several concentrations of each inhibitor. Inhibition constants were determined according to the inhibition patterns for OP and Arg-NHOH respectively.

pH profiles: The pH profiles for Zn- and Cu-BP10 catalyses were constructed by monitoring gelatin and BAPNA hydrolysis under several different pH’s using 50.0 mM buffers containing 50.0 mM NaCl and 1.0 mM CaCl$_2$. The following buffers were used: acetate (pH 5.0), MES (pH 5.5-6.5), HEPES (pH 7.0-8.0), TAPS (pH 8.5-9.0), CAPS (pH 9.5-11.0). The pH-dependent kinetic parameters were determined by non-linear fitting and pK$_a$ values were obtained from fitting the data to a two-ionization process.

Electronic spectrum of Cu-BP10: The electronic spectrum of a 20.0 µM Cu-BP10 was monitored from 350 to 800 nm and the tyrosine to copper charge transfer transition was observed at 454 nm. The quenching of this ligand to metal charge transfer transition (LMCT) was monitored spectrophotometrically upon addition of Arg-NHOH.

Homology modeling and substrate docking: The primary sequence of BP10 was overlaid over the crystal structure of serralysin (PDB ID 1SAT) using BioCAChé 6.1.10
(Fujitsu, Beaverton, OR). The structure was energy-minimized using MM3 and molecular dynamics calculations in a simulated water box. The substrate BAPNA was docked into the active site using the standard procedure (PF5) in BioCAChe.

III. RESULTS AND DISCUSSION

Overexpression and refolding of recombinant BP10: The recombinant enzyme was efficiently overexpressed, though it was insoluble and contained within inclusion bodies (Figure 7.1, Lane 3). Urea solubilization proved to be an efficient method for extracting the enzyme from inclusion bodies, coupled with a fusion His-tag at the N-terminus to BP10 that allows for efficient purification using Ni-NTA agarose. The overexpression and purification yields an average of 0.7 mg/mL of total recombinant BP10 after a pH gradient elution from the Ni-NTA agarose column (Figure 7.1). The protein overexpression was monitored on a time course using SDS-PAGE and Western blotting (Figure 7.2B). The detection of the His-tag using an anti-His antibody upon induction with IPTG proved a sensitive and consistent method for monitoring the overexpression of the recombinant protein.

Upon removal of the chaotrophic reagent urea through extensive dialysis, the protein was refolded and activity could be monitored after removal of the His-tag. It was determined empirically that a 40 fold dilution was important since at higher concentrations the protein coagulates and precipitates out of solution. Recombinant BP10 showed no signs of degradation after refolding (Figure 7.2A). Upon removal of the
Figure 7.1. SDS-PAGE gel (12.5 %) during purification of recombinant BP10. Lane 1, MW marker; lane 2 total soluble protein after 4 hour induction after lysozyme and sonication; lane 3, total protein solubilized with 8.0 M urea from inclusion bodies; lane 4, flow-thru unbound proteins from Ni-NTA column; lanes 5-6, buffer C wash; lanes 7-9, buffer D at pH’s 6.3, 5.9, and 4.5 gradient elution, respectively. Lane 9 shows a homogenous band at 66 kDa, assigned to recombinant BP10.
Figure 7.2 (A) SDS-PAGE (12.5 %) showing intact BP10 after refolding. Lane 1, MW marker; lane 2, is total urea solubilized protein; lane 3, recombinant BP10 after refolding. (B) Western blot showing the time course of overexpression. Lane 1, uninduced sample; lane 2, 1.0 hr; lane 3, 2.0 hr; lane 3, 2.0 hr; lane 4, 3.0 hr; lane 5, 4hr. (C) Gelatin zymogram showing concentration-dependent substrate hydrolysis by BP10; (Lanes 1-5) 0.25µM, 0.50 µM, 1.0 µM, 1.50 µM, 1.75 µM BP10.
His-tag using a Factor Xa removal kit from Novagen, the protein undergoes autohydrolysis and remained stable only for a few days at 4 °C. However, the protein fused with the His-tag is stable indefinitely at 4 °C. Hence, the His-tag was used as an efficient method for long-term storage of recombinant BP10 and removed only prior to running experiments.

**CD Spectra of ZnBP10 and CuBP10:** The CD spectrum of urea-denatured BP10 (Figure 3, dotted trace) does not show the random-coil expected in the presence of a high concentration of chaotropic reagents. The CD spectrum of urea-denatured BP10 shows an overall β-sheet-shape with blue shifts observed in the minimum and a red shift in the maximum of the spectrum. The characteristic minimum at 200 nm for random-coil peptides is not present, with only bathochromic shifts resulting in the denaturation of BP10. This resistance to complete denaturation to a random-coil conformation may account for the efficient refolding of BP10 in the absence of reducing agents, due to partially formed secondary structures. The helical content of BP10 is 5.8 % which is consistent with the large content of sheet-like structures (i.e. β-barrel) in CUB domain-containing proteins.25 According to sequence homology, only the helices present in the astacin-domain should be present accounting for the low helical content of BP10. The CD spectra of Zn-BP10 and Cu-BP10 are virtually identical, suggesting no major conformational change in the overall structure of the protein due to metal substitution.

**Kinetics of gelatin hydrolysis:** Recombinant BP10 was not able to hydrolyze casein, a commonly used substrate for endopeptidases. Porcine gelatin however proved to be a good substrate to monitor proteolytic activity of BP10 with gelatin zymograms (Figure 7.2C) and a ninhydrin detection protocol (Figures 7.4 and 7.5), establishing the proposed role of BP10 as a protease. The formation of a colored ninhydrin-α-amino acid
conjugate with a known molar absorptivity was a convenient method to monitor gelatin hydrolysis. The saturation kinetics can be fit to the Michaelis-Menten equation, to yield $k_{\text{cat}} = 0.013 \text{ s}^{-1}$, $K_m = 51.3 \text{ µM}$, and the second order rate constant $k_{\text{cat}}/K_m = 253.4 \text{ M}^{-1}\text{s}^{-1}$.

Interestingly, BP10 shows a Ca$^{2+}$-dependent activation (Figure 7.5, inset) yielding $k_{\text{cat}} = 0.77 \text{ s}^{-1}$, $K_m = 46.5 \text{ µM}$, and $k_{\text{cat}}/K_m = 16,740 \text{ M}^{-1}\text{s}^{-1}$ at saturating [Ca$^{2+}$]. Other gelatinases found in the sea urchin embryo have shown a Ca$^{2+}$-dependent activation.$^{26,27}$ However, whether or not the Ca$^{2+}$ bound to the EGF domain is a cofactor in protein-protein interaction or signaling is not known for BP10 and cannot be determined in these studies. Further analysis of the rate constants can shed insight into the role of Ca$^{2+}$ in catalysis. The $K_m$ values are similar in the presence and absence of Ca$^{2+}$, while the $k_{\text{cat}}$ value for the catalysis in the presence of Ca$^{2+}$ is much larger than in the absence of Ca$^{2+}$. Since $K_m$ is defined as $(k_{-1}+k_{\text{cat}})/k_1$, with $k_1$ and $k_{-1}$ the rate constants for substrate binding and dissociating from the enzyme-substrate (ES) complex, a significant increase in $k_{\text{cat}}$ with a constant $K_m$ value in the presence of Ca$^{2+}$ reflects a smaller dissociation $k_{-1}/k_1$ of the ES complex. This observation reveals that Ca$^{2+}$ affects BP10 catalysis by enhancing substrate binding to the enzyme and by lowering the activation energy (i.e., a larger $k_{\text{cat}}$ value). The activation of BP10 by Ca$^{2+}$ is the first report of the effect of Ca$^{2+}$ on the activity of astacin family metalloproteases, although Ca$^{2+}$ dependent gelatinases have been identified$^{26}$ in sea urchins; Mg$^{2+}$ showed no effect in activation of BP10. BP10 contains an EGF-Ca$^{2+}$ domain that could mediate the effect of calcium on catalysis.
Elucidation of the role of the EGF domain and Ca\(^{2+}\) on BP10 activity may shed interesting insight once similarly functioning and structurally conserved enzymes have been characterized across all phyla.

The kinetic parameters for Zn-BP10 toward gelatin hydrolysis were determined

\[
k = \frac{k_{\text{lim}}}{\left(1 + \frac{H^+/K_{a_1}}{1 + \frac{K_{a_2}}{H^+}}\right)} \quad \text{Equation (1)}
\]

between pH’s 4.5-11.0. Plots of \(k_{\text{cat}}\) and \(k_{\text{cat}}/K_m\) against pH exhibit bell shaped curves (Figure 7.6 A,D) indicating the presence of two ionizable groups in the catalytic mechanism of ZnBP10 and can be fitted to Equation (1). The two ionization constants from the \(k_{\text{cat}}\) vs. pH profile yield pK\(_{a_1}\) = 5.94 and pK\(_{a_2}\) = 10.2 which can be assigned to the deprotonation of a Zn-bound water and to the coordinated tyrosine respectively upon substrate binding which is consistent with previous reports for serralysin.\(^5\) The role of the coordinated Tyr is further addressed below.

**Kinetics of BAPNA hydrolysis:** The synthetic substrate BAPNA is a good substrate for BP10. This activity was previously observed for serralysin and is the only synthetic tripeptide of a series of di- and tripeptide mimics, including glycine-, alanine-, valine-, leucine-, glutamate-, lysine-, arginine-, trialanine-, and succinyl-trialanine-p-nitroanilide that was hydrolyzed by BP10. The kinetics of BAPNA hydrolysis (Figure 7.7) fit Michaelis-Menten kinetics to yield \(k_{\text{cat}} = 0.079 \text{ s}^{-1}, K_m = 0.66 \text{ mM}\) and \(k_{\text{cat}}/K_m = 120 \text{ M}^{-1}\text{s}^{-1}\) in the presence of Ca\(^{2+}\), and \(k_{\text{cat}} = 1.83 \times 10^{-3} \text{ s}^{-1}, K_m = 1.55 \text{ mM}\) and \(k_{\text{cat}}/K_m = 1.18 \text{ M}^{-1}\text{s}^{-1}\) in the absence of Ca\(^{2+}\) (Figure 7.7 B). Like in gelatin hydrolysis by BP10,
Figure 7.3. CD spectra of Zn-BP10 in PBS with 8.0 M urea pH 7.4 (dotted trace), and renatured Zn- (solid trace) and Cu-BP10 in PBS pH 7.4 (dashed trace).
Figure 7.4. First order kinetics of gelatin hydrolysis by Zn$^{2+}$ (●) and Cu$^{2+}$ (○) derivatives of BP10 in the presence of 1.0 mM Ca$^{2+}$ at pH 7.5. The solid traces are the best fit to a pseudo-first-order rate law, which affords the rate constant $k_{obs}$ for each derivative.
Figure 7.5. Gelatin hydrolysis by 1.0 µM ZnBP10 in 50.0 mM HEPES pH 7.5, 50 mM NaCl, and 1.0 mM Ca(NO₃)₂. The solid line is the best fit to the Michaelis-Menten equation. The inset shows a Ca²⁺-dependent activation upon titration of Ca²⁺ to 60 µM ZnBP10. The solid line in the inset is the best fit to a loose-binding equilibrium.
Ca\(^{2+}\) activates BP10 by lowering the activation energy and enhancing substrate binding as reflected in the rate constants and discussed above. The higher $k_{\text{cat}}$ and lower $K_m$ for gelatin hydrolysis than BAPNA also indicate a higher affinity of gelatin than BAPNA binding to BP10, reflecting the endopeptidase nature of BP10.

The pH profile for BAPNA hydrolysis by Zn-BP10 (Figure 7.6 B, E) compares well with that of gelatin, showing a bell shaped curve that can be fitted to Equation (1). The ionization constants also fall within experimental range, yielding $pK_{a1} = 5.39$ and $pK_{a2} = 9.18$ for the first order rate constant $k_{\text{cat}}$, and $pK_{a1} = 5.83$ and $pK_{a2} = 8.98$ for $k_{\text{cat}}/K_m$.

The inhibition of Zn-BP10 by the metal chelator 1,10-OP toward the hydrolysis of BAPNA at pH 7.5 shows a noncompetitive pattern which is consistent with metal removal from metalloenzymes and gives $K_i = 7.85$ µM (Figure 7.8). The mixed-type inhibition by Arg-NHOH (Fig. 7.8 B) is a good indication of a combination of specific interaction along with metal chelation afforded by the hydroxamate moiety. The inhibition pattern for Arg-NHOH can be fitted to Equations (2) and (3) to yield $K_{ic} = 0.20$ mM and $K_{iu} = 0.90$ mM, representing the specific inhibition constant for the dissociation of the enzyme-inhibitor complex (EI) and the catalytic inhibition constant for the dissociation of the inhibitor from the enzyme-substrate-inhibitor complex (ESI), respectively.\(^{28}\)

\[
\frac{V_{\text{app}}}{K_m} = \frac{V_{\text{max}}}{K_m} \frac{K_m}{1 + [I]/K_i} \quad \text{Equation (2)}
\]

\[
V_{\text{app}} = V_{\text{max}} \frac{1 + [I]/K_i}{1 + [I]/K_{iu}} \quad \text{Equation (3)}
\]
Mechanistic studies of the copper derivative of BP10 (Cu-BP10): The spectroscopically inert Zn\(^{2+}\) ion found in astacins and several other metallohydrolases offers a poor probe for the metal coordination environment in the active site. Thus spectroscopically active metal derivatives of metalloenzymes can offer detailed insight into the catalytic mechanisms and structure within the active site.\(^{29}\) The formation of the Cu-BP10 derivative is evident by the intense Tyr-to-Cu\(^{2+}\) LMCT at 454 nm, analogous to that in Cu\(^{2+}\)-astacin and Cu\(^{2+}\)-serralysin. The activity of the Cu\(^{2+}\)-substituted BP10 (Cu-BP10) is considerably higher than Zn-BP10 in terms of \(k_{\text{cat}}\) (0.76 s\(^{-1}\)) and \(k_{\text{cat}}/K_m\) (5430 M\(^{-1}\) s\(^{-1}\)) toward the hydrolysis of BAPNA, reflecting a \(~960\ \%\) increase in activity in terms \(k_{\text{cat}}\) and \(~485\ \%\) in terms of \(k_{\text{cat}}/K_m\). This is a rather unusual characteristic of metal derivatives of Zn enzymes,\(^{29}\) since most Cu\(^{2+}\) derivatives of Zn\(^{2+}\)-enzymes are inactive. Increased activity of a Cu\(^{2+}\) derivative has been observed in serralysin, another astacin family member able to hydrolyze gelatin and BAPNA.\(^{5}\) Moreover, the overall proteolytic activity of Cu-BP10 toward gelatin hydrolysis is \(~20\%\) of that of the Zn derivative, which is also greater than many metal-substituted metallo-hydrolases previously reported.\(^{29}\)

Metal-centered hydrolysis relies on the Lewis acidity of metal ions, which can lower the pK\(_a\) of metal-bound water molecules by greater than 10\(^7\) fold, generating a metal-hydroxide at neutral pH that can perform nucleophillic attack on the scissile peptide bond. The versatility of metal-centered hydrolysis has been widely demonstrated in synthetic Cu\(^{2+}\) model systems which show proficient peptidase activities and phosphodiesterase activities.\(^{30-32}\) Conversely, Cu\(^{2+}\) derivatives of metallohydrolases are generally inactive or exhibit considerably lower activities\(^{29,33}\) despite the comparatively
high Lewis acidity of Cu$^{2+}$. Few examples of metal-substituted metallohydrolases in literature show considerable activation, with serralysin$^5$ and astacin$^{34}$ being the most significant representatives of metal-substituted metallohydrolases that are activated by Cu$^{2+}$. The poor activation of metallohydrolases observed in Cu$^{2+}$ derivatives may be attributed to Jahn-Teller distortion which can reduce the nucleophilicity of the metal bound water if positioned in the axial coordination of the metal center. The ability to use Cu$^{2+}$ as a viable probe for mechanistic studies is characteristic for the astacin family of metalloenzymes. It is noteworthy that the metal ligands found in the astacin family are a unique example of metal-phenolate coordination in metallohydrolases wherein the coordinated Tyr plays a “switch-off” role in catalysis$^{5,19}$ and may be involved in the unique Cu$^{2+}$-activation observed in astacin family enzymes thus far characterized. The analysis of BP10 is consistent with this mechanism of hydrolysis discussed below.

From analysis of kinetic parameters $k_{\text{cat}}$ and $K_m$, there is an obvious requirement of the metal center for catalysis. $K_m$ value for the hydrolysis of BAPNA by Cu-BP10 in the presence of Ca$^{2+}$ ($K_m = 1.32$ mM) is not significantly different (200 %) from the Zn form (0.66 mM) when compared to the 960 % increase in $k_{\text{cat}}$ of Cu-BP10 compared to the native Zn-BP10. Once again, as previously discussed, a small change in $K_m$ concomitant with a large increase in $k_{\text{cat}}$ suggests a lowering of the activation energy and an increase in the affinity for the substrate in the ES complex. The hyperactive Cu-BP10 suggests that the metal center must be involved in catalysis, most importantly in the turnover of the ES complexes to the product with a high $k_{\text{cat}}$ value.
**Figure 7.6.** pH dependence of $k_{\text{cat}}$ and $k_{\text{cat}}/K_{m}$ for the hydrolysis of gelatin by Zn-BP10 (A, D), hydrolysis of BAPNA by Zn-BP10 (B, E). The solid traces are the best fit to the equation

$$k = \frac{k_{\text{lim}}}{\left[1 + [H^+] / K_{a1}\right] \left[1 + K_{a2} / [H^+]\right]}$$

to afford the two pK$_a$ values reported in the text.
Figure 7.7. (A) Hydrolysis of L-BAPNA by Zn-BP10 in 50.0 mM HEPES, pH 7.5 in the presence of 50mM NaCl, and 1.0 mM Ca(NO₃)₂. (B) Same as in (A) in the absence of Ca²⁺.
Because of the low concentrations available for BP10 (above \( \sim 70 \) \( \mu \)M the protein precipitates), electronic spectra of the \( \text{Cu}^{2+} \) derivative was used to study the metal coordination environment in the active site. Upon substitution of \( \text{Cu}^{2+} \) during the refolding protocols, a 20 \( \mu \)M sample can show significant ligand to metal charge transfer transition (LMCT) in the visible range. Unfortunately, the low energy d-d transition bands for tetragonally distorted octahedral \( \text{Cu}^{2+} \) have very low molar absorptivity values (in the order of 100 \( \text{M}^{-1}\text{cm}^{-1} \)) and are too noisy to distinguish in the spectrum (Figure 7.9). The large LMCT (\( \varepsilon = 1220 \text{M}^{-1}\text{cm}^{-1} \)) observed at 454 nm are due to the tyrosinate-to-\( \text{Cu}^{2+} \) charge transfer transition as determined previously using Cu-astacin.\(^{34}\) This observation serves as support of EPR studies of Cu-astacin where \( g_{\|} > g_{\perp} \) spectral features suggests that the metal center is tetragonaly distorted with a weak axial ligand.\(^{19}\) This is further proof of a similar astacin-like active site structure for BP10.

The inhibition of \( \text{Zn}^{2+} \) and \( \text{Cu}^{2+} \) derivatives of BP10 by Arg-NHOH at pH 7.50 (Figure 7.8, 7.10) displays a mixed type pattern as observed in serralysin,\(^{19}\) wherein the inhibitor is able to bind both the enzyme and the ES complex. The mixed type inhibition yields two different inhibition constants for the dissociation of the inhibitor form the EI and EIS complexes, yielding \( K_{ic} = 1.58 \) \( \mu \)M and \( K_{iu} = 3.93 \) \( \mu \)M. The significantly different inhibition constants for Zn-BP10 and Cu-BP10 are good evidence that the inhibitor binds directly to the metal center in the active site. Mixed-type inhibition is often a good indicator of an alternative site for inhibitor and substrate binding in the ES complex to afford an ES-I and ES-S ternary complexes.\(^{28}\)
Figure 7.8. Inhibition of Zn-BP10 toward L-BAPNA hydrolysis in 50.0 mM HEPES pH 7.5, 50.0 mM NaCl, 1.0 mM Ca(NO₃)₂ by 1,10 phenanthroline (A) and by Arg-NHOH (B). Inhibitor concentrations are as follows from bottom to top: (A) 0, 0.5, 1.0, and 2.0 mM; (B) 0, 0.25, 0.5, and 1.0 mM.
The influence on the LMCT centered at 454 nm can serve as an indicator for inhibitor binding directly to the metal center (Figure 7.9). The quenching of the LMCT band upon inhibitor binding indicates that the metal-coordinated Tyr is detached upon inhibitor binding, a phenomenon observed in the studies of Cu-astacin and Cu-serralysin. The gradual decrease of the LMCT intensity upon inhibitor binding can be described according to Scheme (1), assuming that the binding of one equivalent of inhibitor per active site metal results in concomitant detachment of the coordinated Tyr.

\[
\text{BP10(bound Tyr) + Arg-NHOH} \rightleftharpoons \text{Arg-NHOH-BP10 + detached Tyr}
\]

When fitting the quenching of the Tyr-to-Cu\(^{2+}\) charge transfer with respect to inhibitor concentration according to Scheme (1) without including [H\(^+\)] gives an apparent association constant of \(2.9 \times 10^3\) M\(^{-1}\) for Arg-NHOH binding to Cu-BP10 at pH 8.5 (Figure 7.9, inset). The specific inhibition constant \(K_{ic}\) for the inhibition of Cu-BP10 by Arg-NHOH at pH 7.5 (Figure 7.10) can be converted into an apparent association constant of \(6.33 \times 10^5\) M\(^{-1}\), greater than that at pH 8.5, indicating that the protonation of the metal-coordinated Tyr at lower pH assists the binding of Arg-NHOH.

The kinetic parameters \(K_m, k_{cat}\), and \(k_{cat}/K_m\) for the hydrolysis of BAPNA by Cu-BP10 were determined between pH’s 5.0-9.5 and exhibit bell shaped curves that can be fitted to Equation (1) to give \(pK_{a1} = 5.48\) and \(pK_{a2} = 7.98\) for the pH dependence of \(k_{cat}\) and \(pK_{a1} = 5.83\) and \(pK_{a2} = 7.99\) for the pH dependence of \(k_{cat}/K_m\) (Figure 7.11). This describes a two ionization mechanism for the catalytic turnover of the substrate by BP10. The similar crystal structures of different metal derivatives of astacin suggest that the...
Figure 7.9. Optical titration of Arg-NHOH to Cu-BP10. The inset shows the decrease in the change of the molar absorptivity ($\Delta \varepsilon$) as a function [Arg-NHOH].
**Figure 7.10.** Inhibition of Cu-BP10 by Arg-NHOH. Inhibitor concentrations are as follows from bottom to top: 0, 1.5, and 3.0 µM.
Figure 7.11. pH profile of BAPNA hydrolysis by Cu-BP10.
coordination sphere of Zn\(^{2+}\) and Cu\(^{2+}\) derivatives of BP10 would be similar. This would possibly attribute the different ionization constants pK\(_{a1}\) and pK\(_{a2}\) for each metal derivative to the Lewis acidity of each metal, but not due to different metal environments within the active site of BP10. The low pK\(_{a2}\) approaching pK\(_{a1}\) for Cu-BP10 causes a considerable decrease in the catalytic efficiency, from the intrinsic value of 851 M\(^{-1}\)s\(^{-1}\) to the maximum fitted value of 580 M\(^{-1}\)s\(^{-1}\) which means that only 64% of Cu-BP10 is active at pH 7.0. Conversely the intrinsic and fitted values for \(k_{cat}/K_m\) of Zn-BP10 differ only slightly with the intrinsic value of 119.7 M\(^{-1}\)s\(^{-1}\) and the fitted value of 125.1 M\(^{-1}\)s\(^{-1}\). This is consistent with the coordination sphere of the two metal derivatives of BP10 being similar, and suggest that the differences in ionization constants is due to the effects on the Lewis acidity of each metal.

Ionizable groups must be coordinated or in very close proximity of the metal in the active site of an enzyme to be influenced by the metal ion, as reflected by a change of pK\(_a\) values. In astacin the crystal structure suggests that Tyr and Glu as well as a water molecule are bound to the metal center. This “metallotriad” framework of M–OH…–OOC is similar to the “catalytic triad” of serine proteases Ser–OH…His …–OOC, where the water nucleophile is sandwiched by a Lewis acid (the active site metal) and a Lewis base (the carboxylate in Glu) to serve in a general-acid/general-base catalytic mechanism. This metal-centered triad has also been confirmed in other metallobyldrolases including serralysin,\(^5\) thermolysin,\(^35\) matrilysin,\(^36\) and carboxypeptidase A.\(^37\) We utilized the Cu\(^{2+}\) derivative of BP10 to determine if a similar mechanism is utilized by this enzyme.
The 454 nm LMCT band does not show full intensity at neutral and lower pH values (~150 M$^{-1}$cm$^{-1}$ at pH 6.0 versus 1255 M$^{-1}$cm$^{-1}$ at pH 8.5), indicating that they are pH dependent. The Tyr(phenolate)-to-Cu$^{2+}$ CT in Cu-BP10 changes with pH in a sigmoidal manner (Figure 7.12). Thus, the change can be described by a single ionization process$^5$ for the ionization of the coordinated Tyr$_{249}$ and simultaneous binding to the active site Cu$^{2+}$ to give $\varepsilon_u = 109$ and $\varepsilon_b = 1182$ M$^{-1}$cm$^{-1}$ and a pK$_a$ value of 6.87 for the deprotonation of Tyr$_{249}$, where $\varepsilon_u$ and $\varepsilon_b$ are the molar absorptivities due to the background and the Cu-bound Tyr$_{249}$. However, the data do not fit well to a single-ionization model (Fig. 7.10, dotted trace) and the pK$_a$ value of 6.86 is not a close match to the $k_{cat}/K_m$ pK$_{a2}$ value of 7.99.

Since proposed models of astacin-type enzymes show that the Tyr side chain is H-bonded to the metal-coordinated water, the deprotonation of the phenolate moiety and subsequent binding to the active-site metal should be affected by the ionization of the coordinated water and reflected in the CT intensities. The data is much better fitted to a two-ionization process described previously$^5$ with a fixed pK$_{a1} = 5.83$ (from the $k_{cat}/K_m$ pH profile of Cu-BP10, (Figure 7.11B) to afford pK$_{a2} = 7.42 \pm 0.15$, $\varepsilon_u = 26 \pm 15$, $\varepsilon_w = 591 \pm 33$, and $\varepsilon_b = 1425$ M$^{-1}$cm$^{-1}$ (Figure 7.10 solid trace). Although the data can be reasonably fitted to a two-ionization process, the similar $\lambda_{max}$ throughout the titration suggests that it is likely to have only one species that affords the LMCT instead of two.

Taken together our data is most consistently described as a single species ionization involved in the LMCT, assigned to a Tyr$_{249}$ to Cu$^{2+}$ charge transfer. Herein, the increase in the LMCT due to Tyr binding corresponds to the decrease in activity, reflecting the
Figure 7.12. The change in intensity of the LMCT transition of Cu-BP10 at 454 nm as a function of pH. The data is much better fitted to a two ionization processes (solid trace) than to a single ionization process (dashed trace). The dashed bell-shaped curve is the best fit for the $k_{cat}/K_m$ vs. pH profile of CuBP10 from Figure 7.11B.
inhibiting role of Tyr249 and its role as a “catalytic switch”.

Homology Modeling: The catalytic domain of the astacin family of metallocydrases is highly conserved across all sequenced enzymes. Because BP10 and serralysin are both able to cleave BAPNA effectively, homology modeling by the use of serralysin as the template for the catalytic domain and substrate binding was performed (Figure 7.11). Molecular mechanics calculations (MM3) and molecular dynamics were used to arrive at the final structure which shows Tyr249 within H-bonding distance of the metal-bound water and to the guanidine group of arginine in BAPNA. A predominant hydrophobic interaction is also observed between Trp165 and the benzoyl group of the substrate. The p-nitroanilide moiety of BAPNA is exposed to solvent which likely facilitates product release after the cleavage of the substrate.

IV. CONCLUDING REMARKS

BP10 is the first member of the tolloid-like enzymes to be characterized with extensive kinetics and spectroscopic methods. The studies show that BP10 is a metallocydrase with a hydrolytic mechanism consistent with other astacin-like proteases. The influence of Ca$^{2+}$ toward the catalysis of gelatin and BAPNA by BP10 offers the important insight that Ca$^{2+}$-signaling can serve an important function in regulation of the proteolytic events in embryogenesis. The studies of the Cu-derivative support the metallocydrad mechanism previously proposed for astacin and serralysin, with the involvement of the metal bound Tyr residue in catalysis. Through homology
Figure 7.13. Stereo view of the BP10 active site based on the crystal structure of serralysin (PDB ID 1SAT) as the homology template with the use of molecular mechanics (MM3) and molecular dynamics calculations. The substrate BAPNA (ball-and-stick structure) is docked into the active site, with its carbonyl of the scissile bond pointing toward the active site metal (larger sphere), the benzoyl moiety of BAPNA interacting with Trp$_{165}$, and the guanidinium group of the Arg moiety interacting with Tyr$_{249}$.
modeling we observed the conserved astacin catalytic motif and that the active-site Tyr may not only play a role in substrate binding via detachment from the metal during a “resting” state, but also may assist in stabilization of the ES complex by directly interacting with the guanidine group of BAPNA via a charge interaction or H-bonding.

The understanding of the detailed mechanism of peptide hydrolysis by BP10 and revealing of the substrate specificity *in vivo* in future studies are important first steps in unraveling the proteolytic events during sea urchin embryogenesis.
IV. LIST OF REFERENCES


ABOUT THE AUTHOR

Giordano F.Z. da Silva was born in 1977 in the city of Rio de Janeiro, Brazil. Giordano graduated Magna Cum Laude from St. Petersburg High School and attended St. Petersburg Junior College, graduating Magna Cum Laude in 1998 with an Associate of Arts degree with emphasis in the natural sciences. In 1999 he enrolled at the University of South Florida and graduated in 2002 with a Bachelor of Arts. Giordano joined the graduate program in the Chemistry Department at USF in 2002 working in Dr. Li-June Ming’s MetalloBiomolecule Interest Group (MBIG) and Dr. Brian T. Livingston in the Department of Biology. In the five years in the graduate program Giordano has presented his work at the local, state, national, and international levels; taught a bioinorganic workshop at the National Chiayi University in Taiwan; published peer-reviewed top-tier journal articles, and received several awards, including the Theodore Ashford and George Bursa departmental awards and the USF Successful Latino Award.