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Two-electron Quenching of Dinuclear Ruthenium(II) Polypyridyl Complexes

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Two-electron Quenching of Dinuclear Ruthenium (II) Polypyridyl Complexes

A thesis submitted in partial fulfillment
of the requirements for the degree of
Master of Science in Chemistry

by

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ABSTRACT

A bridging ligand 5,5'-Bi-1,10-phenanthroline, diphen, was prepared using dichlorobis(triphenylphosphine)Ni(II), Ni(PPh₃)₂Cl₂ as catalyst with a yield of 40%. Yellow cubic crystals were able to obtain from the good purity product for single crystal analysis. The torsion angle between the planes of the subunit phenanthrolines is about 66 degrees.

A dinuclear ruthenium (II) polypyridyl complex, (phen)₂Ru(diphen)Ru(phen)₂⁴⁺, was synthesized by using polymeric ruthenium carbonyl compound as the entry point, diphen as the bridging ligand and 1,10-phenanthroline, phen, as the terminal legand. Brown needlelike crystals were precipitated from acetonitrile that were not suitable for single crystal diffraction.

The photochemistry of the dimer was investigated in regards to the oxidation and reduction of the ruthenium centers through a series of quenching reactions excited by visible light. The analogous monomeric complexes Ru(bpy)₃²⁺ and Ru(phen)₃²⁺ were used as comparisons. In the photoinduced oxidation with peroxydisulfate, S₂O₈²⁻, the dimer showed a higher Stern-Volmer quenching constant k_q than Ru(phen)₃²⁺. The dimer showed faster laser flash photolysis transients than Ru(bpy)₃²⁺. In the photoinduced reduction with ascorbate, no significant difference between the dimer and Ru(phen)₃²⁺.

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TABLE OF CONTENTS

CHAPTER 1 Introduction

1.1 Introduction	1
1.2 Conversion of Solar Energy to Alternative Fuels.....	6
1.3 Photophysical and Redox Properties of Ruthenium(II) Complexes.....	6
1.4 Applications of Ruthenium (II) Complexes and Laser Flash Photolysis.....	15
1.4.1 Laser Flash Photolysis.....	15
1.4.2 Reactions Between Ruthenium (II) Complexes and Proteins (Ru(II)-Fe(II)).....	18
1.4.3 Light-driven Oxidation of Water Catalyzed by Ruthenium (II) Complexes.....	25
1.4.4 Organic Synthesis Using Visible-light Triggered by Ru(bpy) ₃ ²⁺	28
1.5 Two and Four-electron Donors and Acceptors.....	30

CHAPTER 2 Experimental

2.1 Materials.....	32
2.2 Instrumentation and General Procedures.....	32
2.3 Photochemical Procedures.....	33
2.4 Syntheses.....	35
2.4.1 5,5'-Bi- 1,10-phenanthroline.....	35
2.4.2 Crystallization of 5,5'-Bi- 1,10-phenanthroline.....	35
2.4.3 [(phen) ₂ Ru(diphen)Ru(phen) ₂](PF ₆) ₄	36
2.4.4 Crystallization of [(phen) ₂ Ru(diphen)Ru(phen) ₂](PF ₆) ₄	37

CHAPTER 3 Results

3.1 Synthesis and Basic Characterization of 5,5'-Bi- 1,10-phenanthroline.....	38
3.2 Synthesis and Basic Characterization of [(phen) ₂ Ru(diphen)Ru(phen) ₂](PF ₆) ₄	48
3.3 Structure of 5,5'-Bi- 1,10-phenanthroline.....	59
3.4 Electrochemical Studies (CV).....	59
3.5 Oxidative Quenching of Ru(II) Dimer and Monomers by Peroxydisulfate.....	68
3.6 Reductive Quenching of Ru(II) Dimer and Monomers by Ascorbate.....	85
CHAPTER 4 Conclusions	
4.1 Synthesis of the Bridging Ligand.....	97
4.2 Crystal Structure of 5,5'-Bi- 1,10-phenanthroline.....	97
4.3 Synthesis of [(phen) ₂ Ru(diphen)Ru(phen) ₂](PF ₆) ₄ Complex.....	99
4.4 Quenching of [(phen) ₂ Ru(diphen)Ru(phen) ₂]Cl ₄ by Peroxydisulfate.....	99
4.5 Quenching of [(phen) ₂ Ru(diphen)Ru(phen) ₂]Cl ₄ by Ascorbate.....	102
REFERENCES.....	103

LIST OF TABLES AND FIGURES

Fig 1.1 Schematic of electron pathway in a DSC system.....	2
Fig.1.2 Structure of ruthenium based N-3 dye and thioicyanate-free dye.....	4
Fig.1.3 Absorption and emission spectrum for $\text{Ru}(\text{bpy})_3^{2+}$ in aqueous solution at room temperature along with the assignments for the various bands.....	8
Fig. 1.4 Excited-state decay pathway for $\text{Ru}(\text{bpy})_3^{2+}$	10
Fig. 1.5 Photoredox cycles of $\text{Ru}(\text{bpy})_3^{2+}$	13
Fig 1.6 Essential optical layout of a laser flash photolysis setup.....	16
Fig 1.7 Structure of a mitochondrion.....	19
Fig 1.8 Mitochondrial electron transport chain.....	19
Scheme 1.1 Photoredox reaction of ruthenium in the presence of external quenchers.....	23
Fig 1.9 Photocatalytic oxidation of water by $\text{Ru}(\text{bpy})_3^{2+}$	26
Fig 1.10 Examples for organic photosensitizers absorb ultraviolet light.....	28
Scheme 1.2 Traditional method for reductive dehalogenation.....	29
Scheme 1.3 Photoredox method for reductive dehalogenation.....	29
Scheme 1.4 Photoredox reduction of electron-deficient olefins.....	29
Scheme 3.1 Showing the homocoupling reaction used in the preparation of diphen.....	39
Figure 3.1 ^1H NMR spectrum of diphen in CD_3CN	42
Figure 3.2 ^1H NMR spectrum of 1,10-phenanthroline in CD_3CN	44
Figure 3.3 ^1H - ^1H COSY spectrum of diphen in CD_3CN	46
Figure 3.4 IR spectrum of neat $[\text{Ru}(\text{CO})_2\text{Cl}_2]_n$	50

Figure 3.5 IR spectrum of neat $[(\text{CO})_2\text{Cl}_2\text{Ru}(\text{diphen})\text{Ru}(\text{CO})_2\text{Cl}_2]$	50
Figure 3.6 IR spectrum of neat $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2](\text{PF}_6)_4$	52
Figure 3.7 IR spectrum of neat 1,10-phenanthroline.....	52
Figure 3.8 ^1H NMR spectrum of $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2](\text{PF}_6)_4$ in CD_3CN	55
Figure 3.9 UV-Vis spectrum of $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2](\text{PF}_6)_4$ in acetonitrile.....	57
Figure 3.10 UV-Vis spectrum of $\text{Ru}(\text{phen})_3\text{Cl}_2$ in 0.2 M $(\text{NH}_4)_2\text{SO}_4$	57
Table 3.1 Sample and crystal data for diphen.....	60
Table 3.2 Bond lengths (Å) for diphen.....	62
Table 3.3 Bond angles (°) for diphen.....	64
Figure 3.11 Cyclic Voltammogram of $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2](\text{PF}_6)_4$ in 0.1M TBAPF ₆ in acetonitrile versus Ag/AgCl.....	66
Figure 3.12 Cyclic Voltammogram of $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2](\text{PF}_6)_4$ under nitrogen in 0.1M TBAPF ₆ in acetonitrile.....	66
Table 3.4 Stern-Volmer quenching constants for the ruthenium complexes with $\text{S}_2\text{O}_8^{2-}$	70
Figure 3.13 Effect of $[\text{S}_2\text{O}_8^{2-}]$ on the emission intensity of the dimer in 0.2 M $(\text{NH}_4)_2\text{SO}_4$	71
Figure 3.14 Effect of $[\text{S}_2\text{O}_8^{2-}]$ on the emission intensity of $\text{Ru}(\text{phen})_3^{2+}$ in 0.2 M $(\text{NH}_4)_2\text{SO}_4$	73
Figure 3.15 Effect of $[\text{S}_2\text{O}_8^{2-}]$ on the emission intensity of $\text{Ru}(\text{bpy})_3^{2+}$ in 0.2 M $(\text{NH}_4)_2\text{SO}_4$	75
Figure 3.16 Plots of the relative emission intensities of $\text{Ru}(\text{bpy})_3^{2+}$, $\text{Ru}(\text{phen})_3^{2+}$ and the dimer versus the $[\text{S}_2\text{O}_8^{2-}]$ concentration. The solid lines and the associated equations were obtained using linear least squares and are the “best” fits.....	77

Figure 3.17 Comparison of transient absorbance change at 450 nm of a solution of dimer and Ru(phen)_3^{2+} in 0.2 M $(\text{NH}_4)_2\text{SO}_4$ following a short laser pulse.....	79
Figure 3.18 Comparison of transient absorbance change at 450 nm of a solution of dimer and $[\text{Ru(bpy)}_3]^{2+}$ in 0.2 M $(\text{NH}_4)_2\text{SO}_4$ following a short laser pulse.....	79
Figure 3.19 Changes in absorbance following steady state irradiation of Ru(phen)_3^{2+} in 0.2 M $(\text{NH}_4)_2\text{SO}_4$ containing 0.5 mM $\text{S}_2\text{O}_8^{2-}$	81
Figure 3.20 Changes in Absorbance following steady state irradiation of the dimer in 0.2 M $(\text{NH}_4)_2\text{SO}_4$ containing 0.5 mM $\text{S}_2\text{O}_8^{2-}$	81
Figure 3.21 UV-vis spectrum of the dimer oxidized by Cl_2	83
Figure 3.22 UV-vis spectrum of the $[\text{Ru(bpy)}_3]^{2+}$ oxidized by Cl_2	83
Table 3.5 Stern-Volmer quenching constants for the ruthenium complexes with ascorbate ...	86
Figure 3.23 Effect of [ascorbate] on the emission intensity of the dimer in argon purged 0.2 M $(\text{NH}_4)_2\text{SO}_4$	87
Figure 3.24 Effect of [ascorbate] on the emission intensity of $[\text{Ru(phen)}_3]^{2+}$ in argon purged 0.2 M $(\text{NH}_4)_2\text{SO}_4$	89
Figure 3.25 Effect of [ascorbate] on the emission intensity of $[\text{Ru(bpy)}_3]^{2+}$ in argon purged 0.2 M $(\text{NH}_4)_2\text{SO}_4$	91
Figure 3.26 Plots of the relative emission intensities of $[\text{Ru(bpy)}_3]^{2+}$, $[\text{Ru(phen)}_3]^{2+}$ and the dimer versus the [ascorbate] concentration. The solid lines and the associated equations were obtained using linear least squares and are the “best” fits.....	93
Figure 3.27 Representative absorption transient for ruthenium complexes with ascorbate.....	95
Table 4.1 Selected bond lengths (Å) and band angles (°) for diphen and phen.....	99

Figure 4.1 Left: The model of the diphen single-crystal structure. Anions are omitted for clarity.

Atom color code: carbon, grey; nitrogen, blue. Right: The structure for phen.....100

Chapter 1 Introduction

1.1 Introduction

In 1839 French physicist Edmond Becquerel showed that photoelectric conversion was possible by generating a current with the first photovoltaic cell¹. The cell consisted of layer of silver chloride placed in an acidic solution along with a platinum electrode. Modern photovoltaic devices are based primarily on semiconductor electrodes. These devices suffer from relatively high costs and low light-to-electric energy conversion efficiency. Many alternative devices are also being developed. Dye sensitized solar cells offer some advantages over solid state devices. They are potentially less expensive but suffer from poor stability among other things. In 1991, Michael Grätzel and Brian O'Regan reported a low-cost dye solar cell (DSC) based on a layer of the trimeric ruthenium complex, $[\text{Ru}(\text{bpy})_2(\text{CN})_2]_2\text{Ru}(\text{bpy}(\text{COO})_2)_2$ ². The conversion efficiency of the early cell was above 7 %.

Figure 1.1 is a schematic diagram of typical dye sensitized solar cell. In these cells the light is absorbed by the dye. The excited dye molecule transfers an electron to the TiO_2 semiconducting electrode. This forms the negative side of the cell much like an ordinary battery. The positive side involves the conversion of I_3^- to 3I^- by way of the returning flow of electrons. The structure of two of the most popular ruthenium based dyes is illustrated in Figure 1.2. The use of ruthenium(II) complexes containing polypyridyl ligands is a common theme and will be discussed in more detail shortly.

Fig 1.1 Schematic of electron pathway in a DSC system³

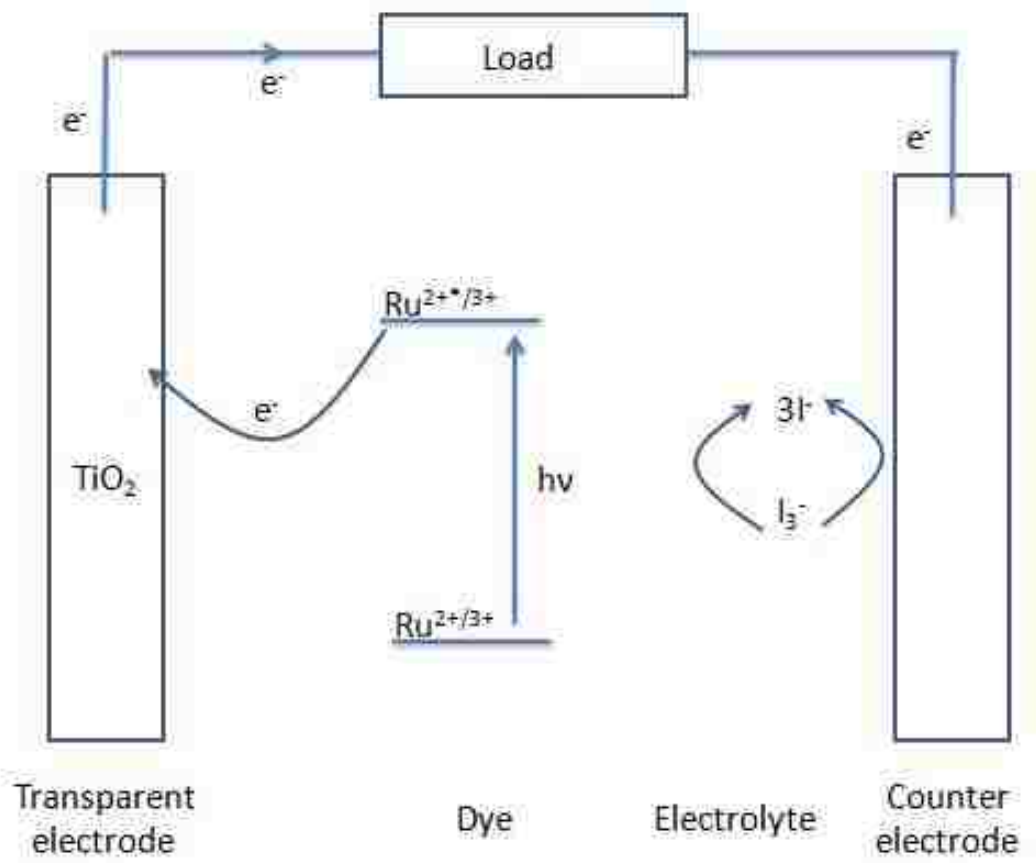
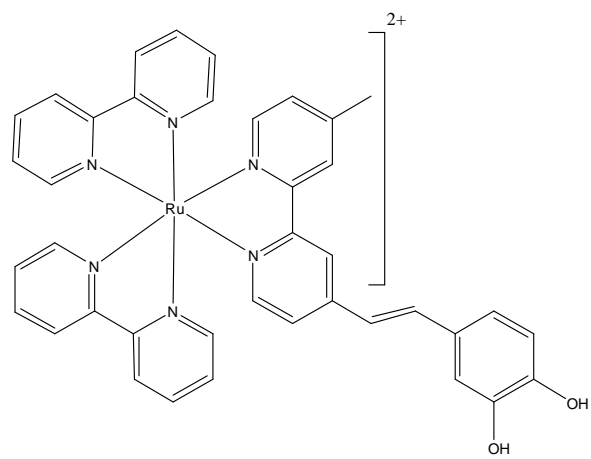
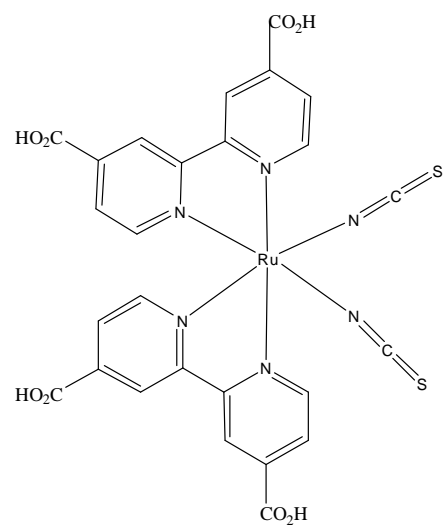


Fig.1.2 Structure of ruthenium based N-3 dye and thioicyanate-free dye



1.2 Conversion of Solar Energy to Alternative Fuels

Photovoltaic conversion of light into energy is one of many ways to capture solar energy.⁴ From a purely energetics standpoint it should be possible to use an appropriate dye to convert water into hydrogen and oxygen.⁵ The possibility is very appealing and many investigators have examined this problem. The ruthenium (II) complex containing three bipyridine ligands, $\text{Ru}(\text{bpy})_3^{2+}$, was among the first dyes extensively examined in this context.⁶ Unfortunately, despite a considerable effort the goal remains elusive.

One of the major stumbling blocks in the proposed application is the fact that the conversion of water to O_2 requires 4 electrons. The ruthenium complex is only capable of providing one electron for each photon absorbed. Unfortunately, the energetics of the process becomes unfavorable if the electrons can only be transfer one at a time because of the formation of high energy intermediates.⁷ The conversion of water to hydrogen likewise requires 2 electrons. Several researchers have tried using platinum based catalysts⁸ to collect electrons and convert water to hydrogen at the electrode surface.

1.3 Photophysical and Redox Properties of Ruthenium (II) Complexes

Ruthenium complexes have played a central role in a large variety of energy conversion schemes and photosensitive devices because of the unique set of characteristics.^{4,9} Ruthenium (II) forms a variety of polypyridyl coordination complexes. These complexes were first discovered almost 80 years ago.¹⁰ The simplest and most well-known complex of this family is (bipyridine)₃Ru(II) or $\text{Ru}(\text{bpy})_3^{2+}$, which has played a central role in the development of photochemical applications due to its remarkably chemical stability and its unique photo-physical properties¹¹. These properties have been applied to a variety of applications, such as artificial photosynthesis, photocatalytic production of hydrogen, dye-sensitized solar cells,

photon-induced switches, oxygen sensors, pH sensors, and molecular machines and devices. The Ru-ligand coordination is very strong. For example, $\text{Ru}(\text{bpy})_3^{2+}$ remains unreacted after refluxing in concentrated HBr. In addition, Ru(I), Ru(II) and Ru(III) complexes retain the bipyridyl coordination sphere.¹² This property is rare across the periodic table and critically important for sustained catalytic activity and electrochemically reversible redox processes (the $\text{Ru}^{1+/2+}$ and $\text{Ru}^{2+/3+}$ processes). These important electronic and photophysical properties are retained in hundreds of structurally modified derivatives of the parent complex. Such structural modifications enable researchers to tune both the properties for specific applications

The absorption and emission spectrum of $\text{Ru}(\text{bpy})_3^{2+}$ is shown in Fig.1.3. The bands at 185 nm and 285 nm are assigned as ligand-centred $\pi \rightarrow \pi^*$ transitions by comparison with the spectrum of the protonated bipyridine ligand¹³. Similarly, the two weak shoulders at 323 nm and 345 nm also with the twin peaks at 238 nm and 250 nm are assigned to the metal centred $d \rightarrow d$ transitions. The two intense bands at 240 nm and 450 nm which located in visible light range are spin-allowed MLCT $d \rightarrow \pi^*$ transitions. $\text{Ru}(\text{bpy})_3^{2+}$ shows a broad orange-yellow emission in solution at 293 K around 600 nm, and the rate of decay corresponds to an excited state lifetime approximately 600 nsec in aqueous solution at room temperature in the absence of oxygen.¹⁴

Fig.1.3 Absorption and emission spectrum for $\text{Ru}(\text{bpy})_3^{2+}$ in aqueous solution at room temperature along with the assignments for the various bands.

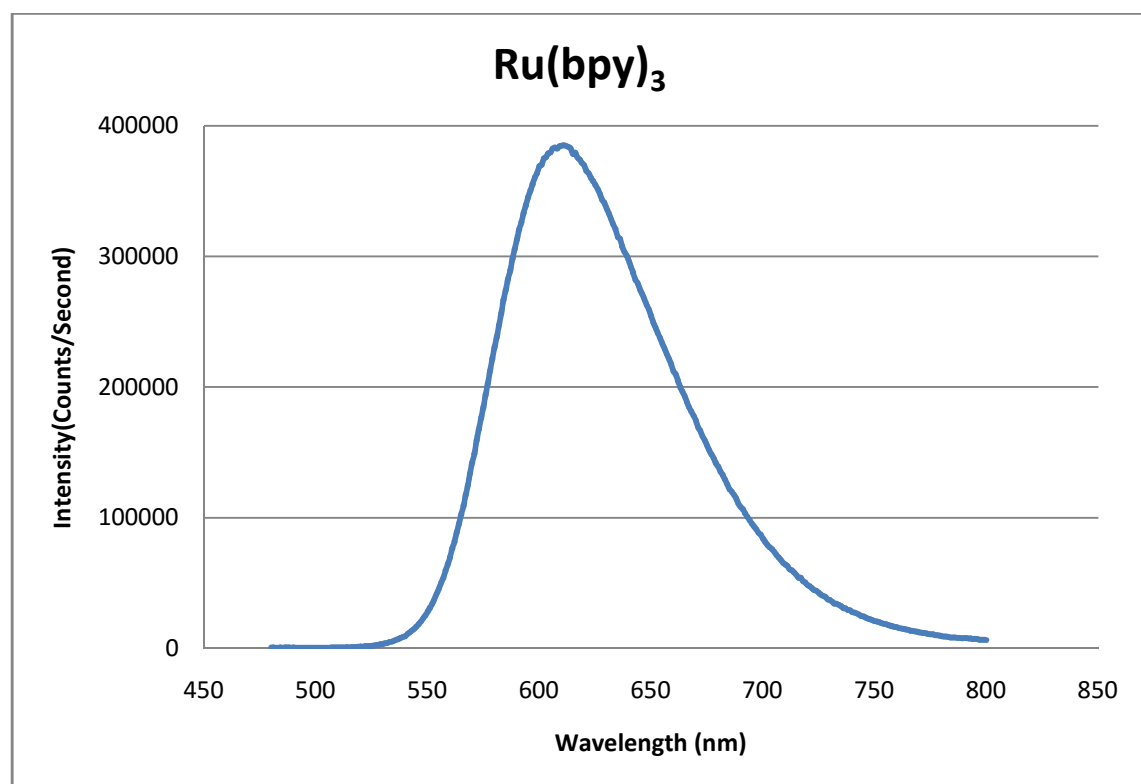
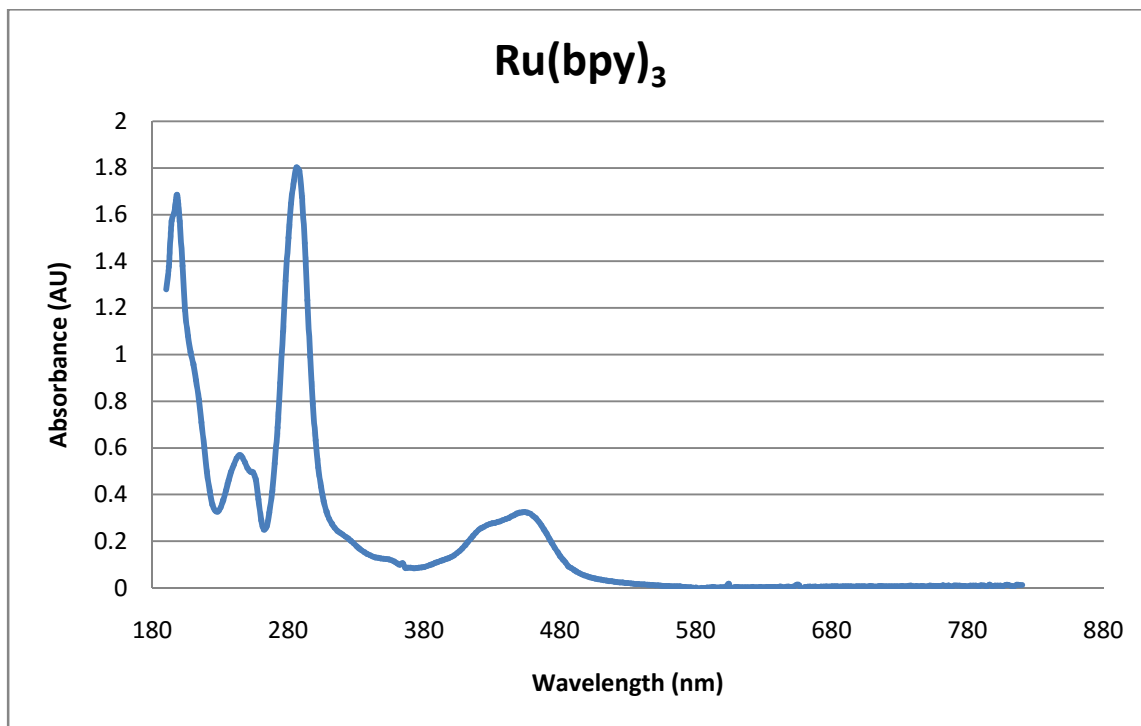
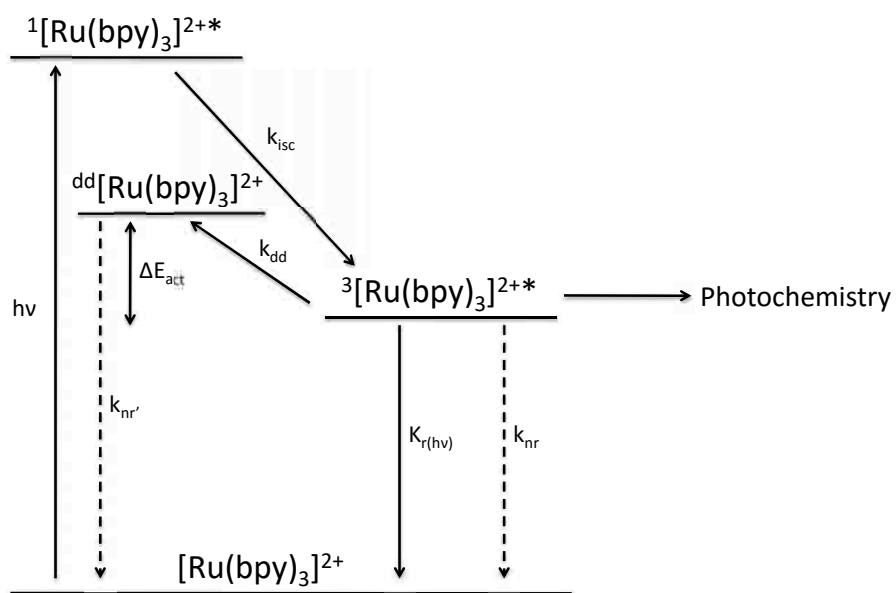


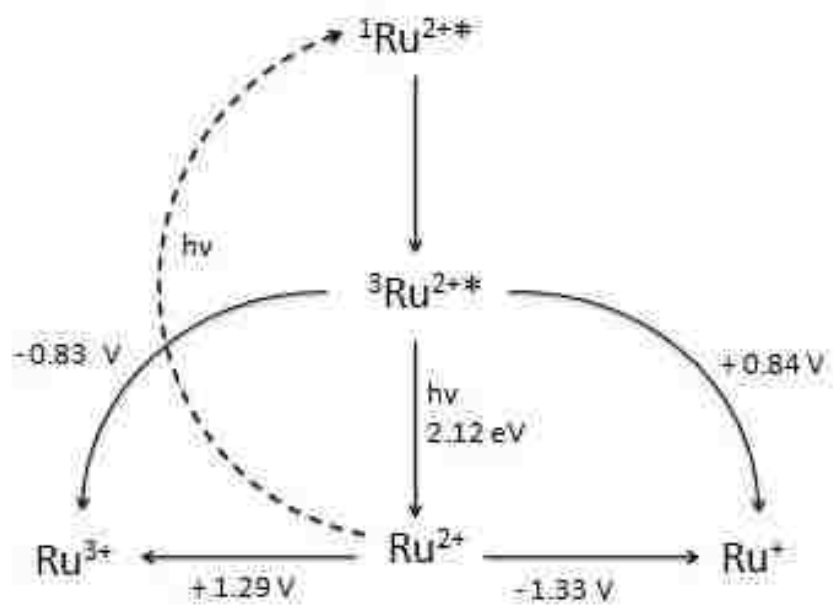
Fig. 1.4 Excited-state decay pathway for $\text{Ru}(\text{bpy})_3^{2+}$



The excited state decays involve several processes (Fig 1.4). By irradiating Ru complexes, for example $[\text{Ru}(\text{bpy})_3]^{2+}$, at the appropriate wavelength, a lowest photoexcited singlet metal-to-ligand charge transfer state ($^1\text{MLCT}$) is produced, $^1[\text{Ru}(\text{bpy})_3]^{2+*}$, which then undergoes intersystem crossing (k_{isc}) within 300 fs¹⁵, yielding a long-lived luminescent triplet excited state ($^3\text{MLCT}$), $^3[\text{Ru}(\text{bpy})_3]^{2+*}$ (600 ns to a few μs). The intersystem crossing yield in the few systems examined in detail is approximately 100%. This excited state exists long enough to transfer energy to another molecule by energy transfer or electron transfer. $^3[\text{Ru}(\text{bpy})_3]^{2+*}$ can also relax to the ground state through a non-radiative decay (k_{nr}) by emission of heat, or a radiative decay (k_{r}) by emission of light at about 600 nm, or thermal population of the nearby metal-centered d-d state, $^{\text{dd}}[\text{Ru}(\text{bpy})_3]^{2+}$. The d-d state rapidly decays back to the ground state and is the primary decay path for the excited state in the absence of other reagents.¹⁶

$\text{Ru}(\text{bpy})_3^{2+}$ in ground state tends to show no redox reactions. Once in excited state, however, $\text{Ru}(\text{bpy})_3^{2+*}$ can behave as both a strong oxidant or a strong reductant. One-electron transfer reactions result in the formation of ground state Ru(III) that is a strong oxidant (+1.29 V vs SCE in CH_3CN) or the formation of ground state Ru(I) a very strong reductant (-1.33 V vs SCE in CH_3CN). The standard reduction potentials for the ground and excited state of $\text{Ru}(\text{bpy})_3^{2+}$ are summarized in Figure 1.5.¹⁷

Fig. 1.5 Photoredox cycles of $\text{Ru}(\text{bpy})_3^{2+}$



1.4 Applications of Ruthenium (II) Complexes and Laser Flash Photolysis

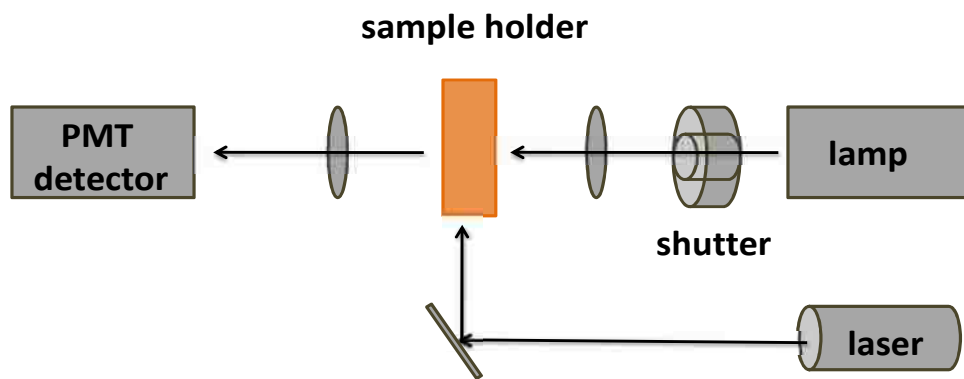
The redox properties of the excited states ruthenium polypyridyl complexes have led investigators to many applications. Of particular importance is the ability to monitor the rates of the electron transfer reactions.¹⁸ For several decades this ability has dominated the fundamental investigations of electron processes. Most notable was the verification of the basic tenants of Marcus Theory and development of basic understanding of electron transfer in metalloproteins and other systems in which the donor and acceptor are separated by significant distances.¹⁹

1.4.1 Laser Flash Photolysis

Laser flash photolysis has been used extensively to measure the rates of electron transfer reactions involving ruthenium polypyridyl complexes.²⁰ The technique is extremely well suited for these investigations. Laser flash photolysis allows for the measurement of reactions with almost no limit of the time resolution. Femtosecond measurements are currently common although most practical applications focus on reactions that occur in the microsecond or millisecond time range.²¹ The technique is based on the creation of reactants by using a short pulse of light. Typically, lasers have pulse widths in the nanosecond or picosecond range are able to produce reactants on this time scale. The pulse width is generally much shorter than the half-time of the chemical reaction. A simplified optical layout of the essential equipment is shown in Figure 1.7.

Most flash photolysis system use absorbance measurements to monitor the reaction. Transient absorbance changes are recorded on a fast digital oscilloscope (nsec sampling). Ruthenium complexes have excited states that last long enough to produce a good yield of product. Excitation and monitoring can be done with visible light which is best for most systems.

Fig 1.6 Essential optical layout of a laser flash photolysis setup



1.4.2 Reactions Between Ruthenium(II) Complexes and Proteins (Ru(II)-Fe(III))

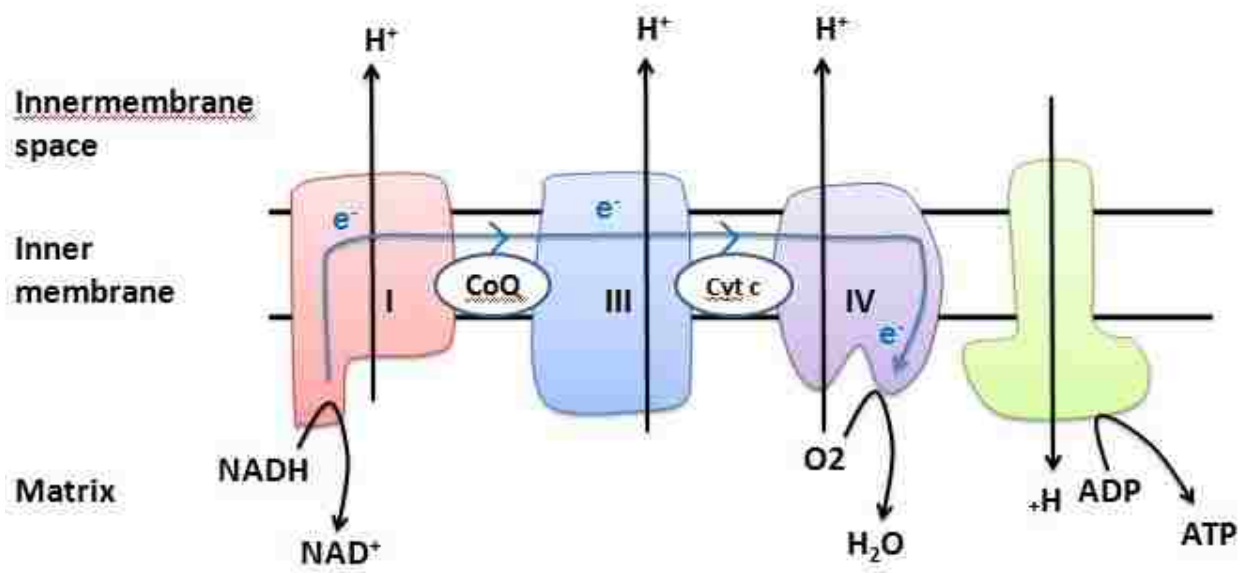
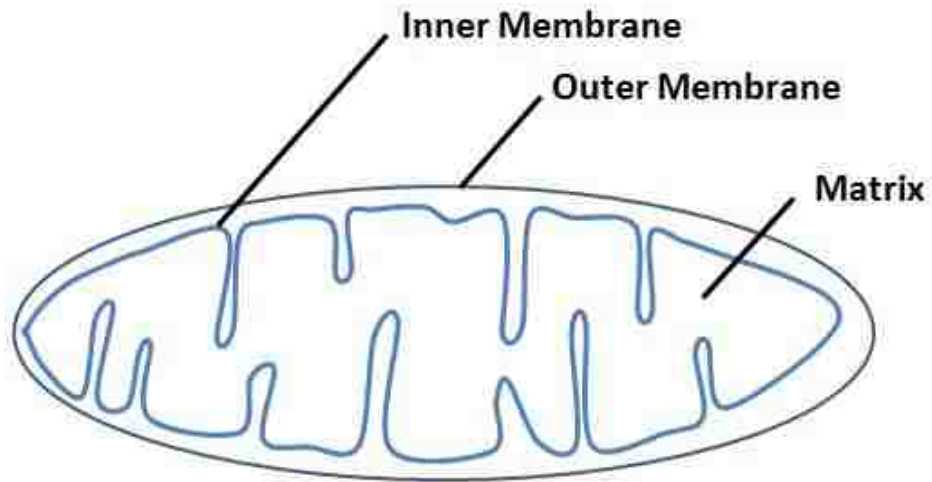
The process of aerobic cell respiration is a one of the most complicated and interested processes because it actually consists of many individual electron transfer reactions. The final step in respiration is called oxidative phosphorylation which takes place on the electron transport chain of the inner membrane of the mitochondria.²²

The electron transport chain is a series of proteins: complex I (or NADH dehydrogenase), complex II (succinate reductase), complex III (or cytochrome bc₁ complex)²³ and complex IV (or cytochrome c oxidase), as well as the final protein molecule known as ATP synthase.

Complex I is responsible for accepting the high energy electrons from NADH molecules produced in glycolysis and the citric acid cycle to ubiquinone. As the electrons move through complex I, four proton ions are pumped out the matrix into the inner membrane. Complex II converts succinate to fumarate and generates the FADH₂ molecules without pumping any proton ions. Ubiquinone is the electron carrier that shuttles these electrons from complex I and II onto complex III. Then complex III catalyzes the transfer of electrons from ubiquinone to cytochrome c.²⁴ The movement of electrons stimulates the movement of proton ions out of the matrix and into the inner membrane space of the mitochondria. Once the electrons end up on complex IV, this is where the diatomic oxygen is reduced by the electrons to form water. This series of reactions yields a proton gradient that is used by ATP synthase to generate ATP molecules.

Fig 1.7 Structure of a mitochondrion

Fig 1.8 Mitochondrial electron transport chain



During the process of this oxidation, cytochrome c is a relatively small water soluble molecule travels through the matrix fluid plays a crucial role as a one-electron shuttle between complex III and IV²⁵. Cytochrome c contains a single iron atom held by four nitrogens of a porphyrin ring, the nitrogen of histidine, and the sulfur atom of methionine. The iron atom is center of the redox activity and cycles between Fe(II) and Fe(III) form to accept one electron from Complex III and subsequently donate one electron to Complex IV.

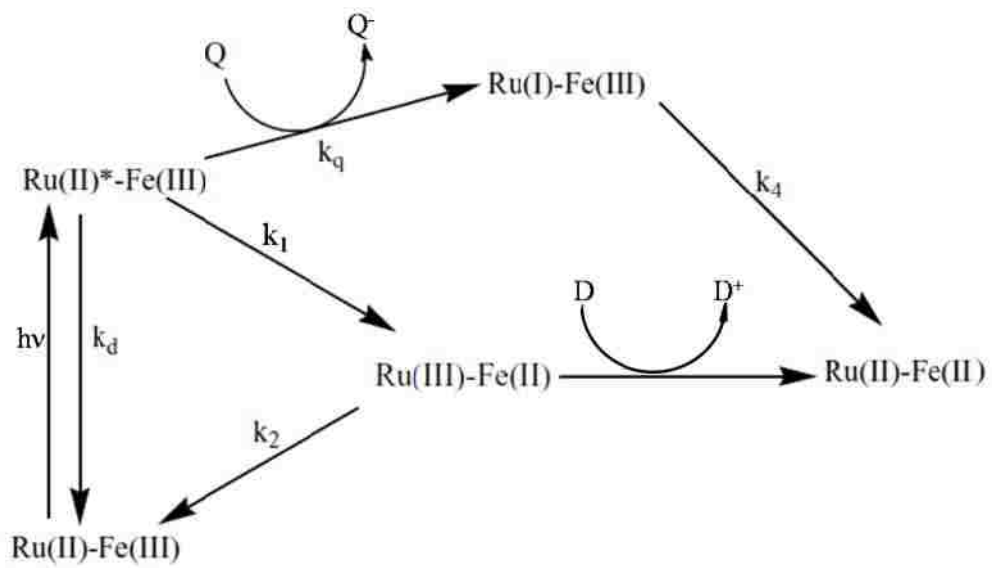
Measurement of the rate constants for electron transfer within the oxidative phosphorylation remained elusive until early 1900's because of the speed of the reactions. Rapid mixing techniques fell far short of the needed time resolution and flash-photolysis was not directly possible redox reactions could not be initiated by irradiation of the proteins alone. In early 1990, use of laser flash photolysis became possible through photoexcitation of ruthenium polypyridine complexes.²⁶

In the early research, ruthenium complexes were covalently bound to cytochrome c²⁷. Ru(II)-Fe(III) in Scheme 1.1 represents ruthenium complexes and the iron center of cytochrome c. Absorption of a short laser pulse leads to the formation of the Ru^{2+*} excited state, which rapidly reduces the iron center to produce Ru(III)-Fe(II). The use of an external sacrificial electron donor, D, prevents the thermal back reaction with the formation of Ru(II)-Fe(II). At this point the reduced cytochrome c can react with a metalloprotein of interest (cytochrome c oxidase, cytochrome peroxidase) after which the ruthenium components are reset to their original oxidation states.

Subsequent research revealed that covalent bonding was not necessary for many of the questions under investigation. Electrostatic interactions with the ruthenium complexes with a

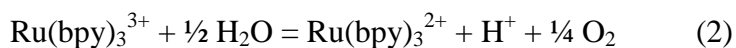
metalloprotein of interest, when correctly designed, provided good yields of proteins in reactive states. For examples, this approach worked very well in studies of the internal electron transfer reactions of multicentered metalloproteins such are Complex III and IV. In these applications the ruthenium complex essentially replaced cytochrome c as an electron donor or acceptor. Extensive design efforts ultimately showed that weakly coupled dimeric ruthenium complexes worked best. For example, the complex $[(bpy)_2Ru-qpy-Ru(bpy)_2]^{4+}$, where qpy is quatrapyridine was used extensively to examine the internal electron transfer reactions of cytochrome c oxidase.²⁸

Scheme 1.1 Photoredox reaction of ruthenium in the presence of external quenchers



1.4.3 Light-driven Oxidation of Water Catalyzed by Ruthenium (II) Complexes

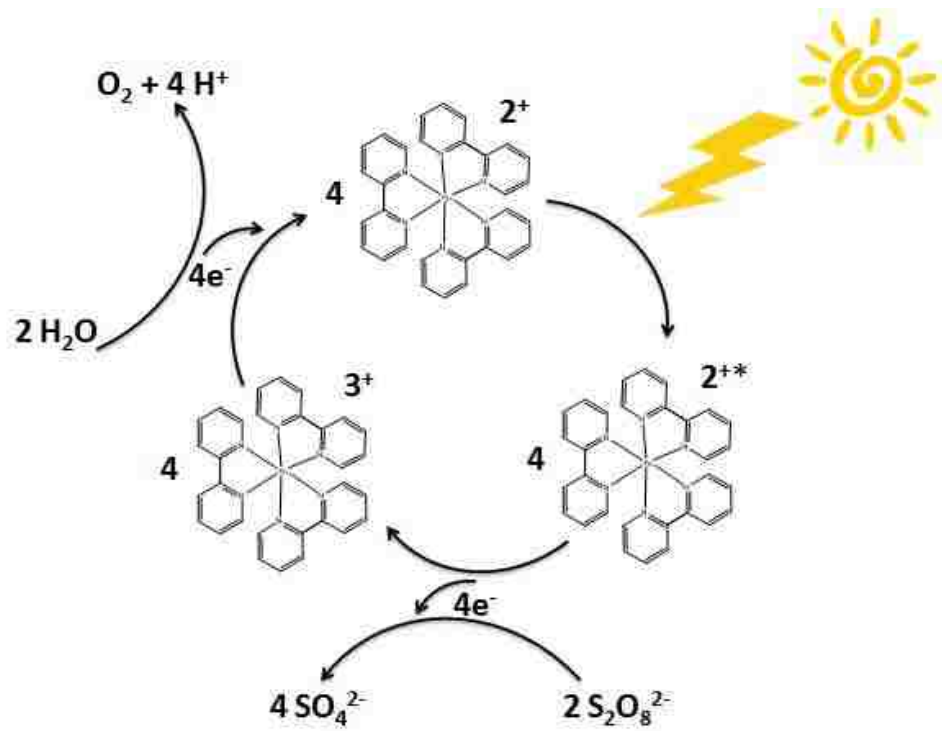
Oxidation of water to release O_2 is the terminal reaction of green plants, photosystem II (PSII) (eq 1).²⁹ Oxidation of water is extremely energy-demanding and occurs thermodynamically above pH ~ 5 with potentials, $E^0(O_2/H_2O_2) = 0.27$ V, $E^0(H_2O_2/H_2O) = 1.36$ V) (eq 2).³⁰ There are very few suitable molecules that can oxidize water by absorbing the energy from sunlight as it happens in the photosynthesis in green plants. However ruthenium (III) complexes have this ability, for example $Ru(bpy)_3^{3+}$.³¹



$Ru(bpy)_3^{2+}$ absorbs visible light to form the MLCT excited state $Ru(bpy)_3^{2+*}$. Then $Ru(bpy)_3^{2+*}$ is oxidized by an appropriate oxidative quencher such as $S_2O_8^{2-}$ to $Ru(bpy)_3^{3+}$. By accepting electrons from water, water is oxidized to O_2 and $Ru(bpy)_3^{3+}$ is reduced and returns back to the initial state $Ru(bpy)_3^{2+}$.

However, the evolution of O_2 from water is difficult in this four-electron redox process and $Ru(bpy)_3^{3+}$ decomposes rapidly. More research is need to develop more efficient and practical photosensitizer for water oxidation that could meet the following requirements: a high turnover number which reflects high stability; a high turnover frequency which reflects high activity and a low overpotential which means water oxidation can be driven by visible light³².

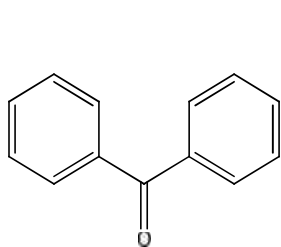
Fig 1.9 Photocatalytic oxidation of water by $\text{Ru}(\text{bpy})_3^{2+}$.



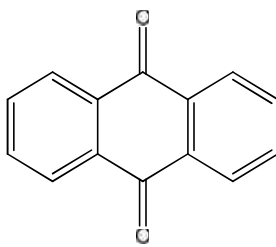
1.4.4 Organic Synthesis Using Visible-light Triggered by $\text{Ru}(\text{bpy})_3^{2+}$

Sunlight is inexpensive and endlessly renewable. Photosynthesis in green plants serves as natural ability to convert solar energy to chemical energy. This green chemistry has inspired the development of photoredox catalysts for organic synthesis. However, most organic catalysts absorb short wavelengths of light³³, for example ultraviolet light which is reactive for unsaturated bonds and some functional groups makes up a small amount (~3%) of sunlight. By utilizing the properties of ruthenium polypyridine complexes, strategies using combination of $[\text{Ru}(\text{bpy})_3]^{2+}$ and visible light for preparative organic chemistry have recently been developed.

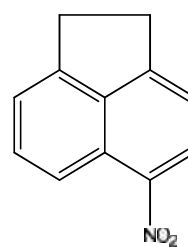
Fig 1.10 Examples for organic photosensitizers absorb ultraviolet light



Benzophenone (BP)

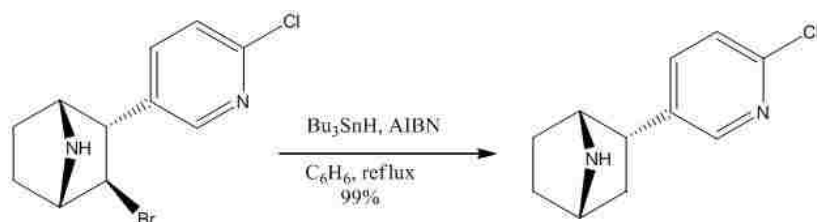


Anthraquinone (AQ)

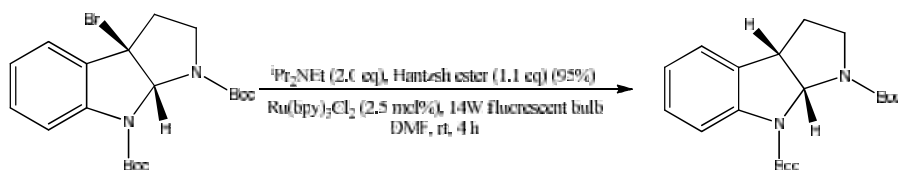


5-Nitroacenaphthen (NAN)

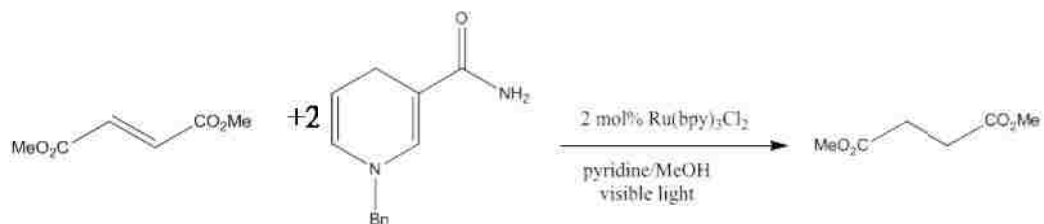
Scheme 1.2 Traditional method for reductive dehalogenation



Scheme 1.3 Photoredox method for reductive dehalogenation



Scheme 1.4 Photoredox reduction of electron-deficient olefins



Reductive dehalogenation is degradation of halogenated organic compounds by removal of a halogen substituent and simultaneous addition of a hydrogen atom. An example of commonly used traditional method is shown in Scheme 1.2 which involves trialkyltinhydride compound as the catalyst³⁴. However, trialkyltin is neurotoxic can cause impaired motor function³⁵. Subsequently a tin-free method using $[\text{Ru}(\text{bpy})_3]^{2+}$ as a visible-light photosensitizer irradiated by a 14 W fluorescent lamp under mild reaction conditions was developed (Scheme 1.3)³⁶. This simply reaction maintained good chemoselectivity and high yields of products.

$[\text{Ru}(\text{bpy})_3]^{1+}$ is a strong reductant can reduce some electron-deficient olefins. No reaction between several olefins and 1-benzyl-1, 4-dihydronicotinamide (BANH) is observed in the dark at room temperature. By adding catalytic quantities of $[\text{Ru}(\text{bpy})_3]^{2+}$ and under visible light, the excited state $\text{Ru}(\text{bpy})_3^{2+*}$ was formed and was reduced by accepting an electron from BANH to generate highly reducing $\text{Ru}(\text{bpy})_3^{1+}$. $\text{Ru}(\text{bpy})_3^{1+}$ reduced the carbon-carbon double bond to single bond (Scheme 1.4).³⁷

$\text{Ru}(\text{bpy})_3^{2+}$ has been investigated for more than four decades.³⁸ However, the utilizing of $\text{Ru}(\text{bpy})_3^{2+}$ to trigger organic synthesis started in 2008.³⁹ This is a new area in advanced organic synthesis and catalysis but application examples are still limited.

1.5 Two and Four-electron Donors and Acceptors

In some of the examples above it is clear that development of catalysts capable of delivering more than a single electron per photon would be beneficial. Direct 2-photon processes occur with low yields and only with an extremely high photon flux. Alternatively dimeric complexes can be envisioned in which the metal centers are very weakly coupled. In this case the metal centers would behave independently and it should be possible to excite both

centers with only modern levels of photon flux. In this case the probability of absorbing a second photon would be the same as absorbing the first photon.

Several dimeric ruthenium complexes have recently been developed that fit the criteria of weakly coupled metal centers. One has been used extensively in the investigation of metalloproteins and was mentioned above, [(bpy)₂Ru-qpy-Ru(bpy)₂]⁴⁺.²⁸ Another is closely related and easier to prepare, [(phen)₂Ru-diphen-Ru(phen)₂]⁴⁺. In both of these dimers the bridging ligands are sterically constrained out of planarity and presumably the lack of planarity leads to very weak coupling between the two ruthenium moieties. Extensive investigation of the photophysical properties supports the weak coupling. Perhaps the most compelling is the fact that excited state lifetimes of the dimers are longer than the monomers indicating that the ruthenium centers do not interact destructively.⁴⁰

Given the above observations it appears that the dimers are well suited to an investigation of possibility that both centers can be oxidized or reduced photochemically and thus be converted into two electron donors or acceptors. Following is a description of initial investigations of quenching reactions design to yield two electron redox products. In particular the investigation will focus on the reactions of the excited state with S₂O₈²⁻ and ascorbate. Reactions of Ru(bpy)₃²⁺ and Ru(phen)₃²⁺ with these reactants have been thoroughly investigated by previous workers. The quenching reaction with S₂O₈²⁻ proceeds with an irreversible quantum yield of two strongly suggesting that the dimers will yield two electron oxidized products. Likewise ascorbate, based on previous work should yield a two electron reduced product and possibly hydrogen.

Chapter 2 Experimental

2.1 Materials

2-Methoxyethanol was purchased from Acros Organics. Triphenylphosphine (PPh_3), N,N-dimethylformamide (DMF), ammonium hexafluorophosphate (NH_4PF_6), the zinc dust, and 5-chloro-1,10-phenanthroline (Cl-phen) were all purchased from Alfa Aesar. Nickelous chloride hexahydrate ($\text{NiCl}_2 \cdot 6\text{H}_2\text{O}$) and potassium cyanide (KCN) were produced by J.T. Baker.

Trimethylamine N-oxide (Me_3NO) was purchased from TCI. Ruthenium (III) chloride hydrate ($\text{RuCl}_3 \cdot \text{XH}_2\text{O}$) and 1,10-phenanthroline (phen) were purchased from Oakwood Chemical.

Hydrochloric acid (HCl), diethyl ether, acetonitrile, methanol, and acetonitrile- d_3 (CD_3CN) were produced by BDH. Formic acid (HCOOH) was produced by Mallinckrodt Chemicals. L-ascorbic acid (Vitamin C) sodium salt, ammonium persulfate ($(\text{NH}_4)_2\text{S}_2\text{O}_8$), ferrocene ($\text{Fe}(\text{C}_5\text{H}_5)_2$), and tetrabutylammonium hexafluorophosphate (NBu_4PF_6) were purchased from Sigma-Aldrich.

Ammonium sulfate ($(\text{NH}_4)_2\text{SO}_4$) was purchased from Macron Chemicals.

2.2 Instrumentation and General Procedures

Infrared (IR) spectra were recorded with a Shimadzu IRAffinity-1S FTIR Spectroscopy equipped with a Quest ATR accessory with a diamond crystal puck. Samples were placed directly the sample holder and the empty chamber was scanned as the background. Spectra were collected in the range of $600 - 4000 \text{ cm}^{-1}$ with 2 cm^{-1} resolution and 24 spectral scans averaged by the LabSolutionsIR operational software and then converted to Excel spreadsheets.

$^1\text{H-NMR}$ were recorded on a Bruker Avance 300 MHz Spectrometer. CD_3CN was used as solvent. NMR software TopSpin was used to analyze the spectra.

All ultraviolet-visible (UV/vis) absorption spectra were obtained using a Hewlett Packard 8452A Diode Array Spectrometer in conventional 1.0 cm quartz cells.

Fluorescence spectra were obtained with a Quanta Master™ 300 Plus from Photon Technology International. The excitation wavelength was 450 nm and the emission spectra were recorded over the range of 480-800 nm at room temperature.

CH Instruments Electrochemical workstation was used for cyclic voltammetry with a three electrode system. The counter electrode was a platinum wire and the reference electrode was a silver wire. The working electrode was a platinum disc with a 1 mm diameter. The supporting electrolyte was 0.1 M NBu₄PF₆ in acetonitrile. Ferrocene was used as reference. All measurements were carried out at a scan rate of 100 mV/s.

Full structural analysis of the diphen was performed by Collin McMillen, Managing Director of the Molecular Structure Center in Clemson University by single crystal X-ray diffraction using a Rigaku AFC8 diffractometer with a sealed tube used to generate graphite monochromated Mo K (= 0.71073 Å) radiation. The diffractometer was equipped with a Mercury CCD detector. The diphen crystal was mounted on a glass fiber using epoxy glue, and data were collected at 100 K by omega scans over 480 diffraction images. Data processing was performed using the Crystal Clear software package. The space groups were determined from the systematic absences, and the structures were solved by direct methods and refined by least-squares techniques using the SHELXTL software suite.

2.3 Photochemical Procedures

Kinetics of Quenching Back-Reactions. Solutions containing Ru(bpy)₃²⁺, Ru(phen)₃²⁺ or dimer were excited with a PhaserR flash-lamp pumped dye laser with LD490 dye which

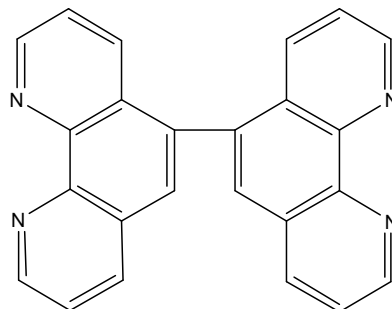
provided a 500 nsec pulse of 480 nm light. The probe beam was provided by a 100 watt tungsten lamp at right angle relative to the laser excitation. A shutter blocked the sample from irradiation by the probe prior to laser excitation. The probe beam was passed through a Kratos Schoeffel Instruments monochromator and the intensity monitored with a R446 photomultiplier. The signal was passed through a unity gain summing amplifier to allow subtraction of the steady state background signal. The transient absorbance changes were monitored by a LeCroy Model 6054 digital oscilloscope. The files were transferred to a PC for kinetic analysis.

In order to prevent possible changes in concentration due to the photoreaction, all the solutions were prepared and stored under darkroom conditions the same day as the experiment. However, no change of absorption was observed after storage of the solutions for a week with/without saturated nitrogen, which indicates the high stability of the ruthenium solutions. The ruthenium complexes concentrations employed in the quenching measurements varied from 22 to 60 μM , corresponding to an absorbance of 0.3-0.8 (ϵ 14000 $\text{M}^{-1} \text{cm}^{-1}$). The oxidative quenching measurements were performed with different concentrations of ammonium peroxodisulfate ($(\text{NH}_4)_2\text{S}_2\text{O}_8$) under air saturated at room temperature. The reductive quenching were performed with different concentrations of L-sodium ascorbic acid at pH =5 under nitrogen at room temperature.

2.4 Syntheses

2.4.1 5,5'-Bi-1,10-phenanthroline

NiCl₂·6H₂O (0.304 g, 2.34 mmol) and PPh₃ (2.44 g, 9.30 mmol) were dissolved in 20 ml dry DMF in a 100 ml RBF and stirred at 50 °C under nitrogen for 1h. Unpurified zinc dust (0.27 g, 4.41 mmol) was added to the resulting blue solution. The solution was stirred at 50 °C under nitrogen until it was dark red. Cl-phen (0.496 g, 2.30 mmol) in 10 ml of DMF with nitrogen protection was added to the dark red solution. The solution was stirred at 50°C under nitrogen overnight until it was dark green. The resulting mixture was evaporated through the rotary evaporator to yield a dark green solid. Then the dark green solid was boiled in 80 ml of water for 2h. The mixture was cooled to room temperature and filtered. The yellow solution was precipitated with 10 ml of a saturated aqueous NH₄PF₆ solution. The precipitate was filtered by a medium porosity fritted funnel and then were suspended in a solution of KCN (1.50 g, 23.0 mmol) in 40 ml of a methanol and H₂O (19:1, v:v) and refluxed for 4 h. The resulting brown solution was cooled completely and filtered. The yellow solid was washed with H₂O and ether, and then dried in a desiccators (Yield: 49%).

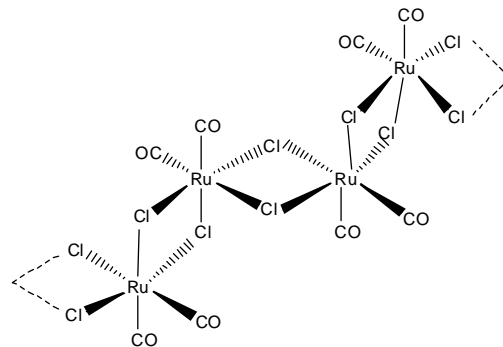


2.4.2 Crystallization of 5,5'-Bi-1,10-phenanthroline

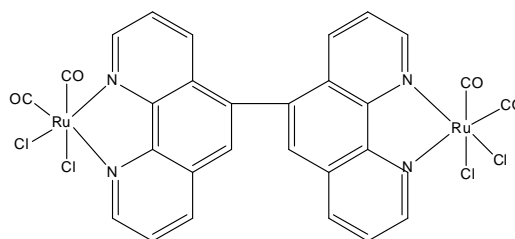
Diphen was dissolved in a minimum volume of hot CH₃CN in a RBF yielding a clear yellow solution. The solution was slowly cooled down to room temperature and allowed to stand undisturbed for 2 days. Yellow cubic crystals formed on the bottom.

2.4.3 [(phen)₂Ru(diphen)Ru(phen)]PF₆

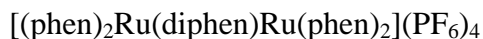
RuCl₃·XH₂O (2.02 g, 7.72 mmol) was dissolved in a 100 ml 1:1 (v:v) solution of concentrated HCl and HCOOH and refluxed for 24 h under nitrogen. The color of the solution changed from brown to yellow. The yellow solid was isolated through evaporation on a hotplate (Yield: 93%).



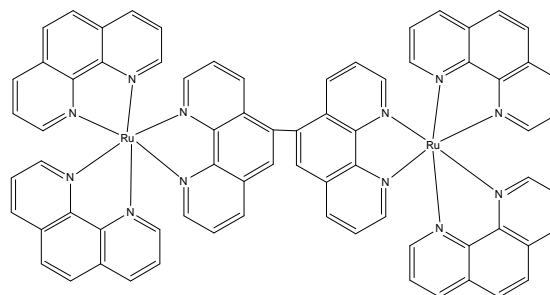
Diphen (0.462 g, 1.29 mmol) in 20 ml of hot 2-methoxyethanol was added to a solution of [Ru(CO)₂Cl₂]_n (1.47 g, 6.45 mmol) in 60 ml of hot 2-methoxyethanol in a 100 ml RBF and the resulting



deep red solution was boiled for 7 min and cooled. The solution was filtered by a medium porosity fritted funnel, and the orange solid was dried in a desiccators (Yield: 37%).



(CO)₂Cl₂Ru(diphen)Ru(CO)₂Cl₂ (0.200 g, 0.246 mmol), phen (0.426 g, 2.36 mmol), and Me₃NO (0.187 g, 2.49 mmol) were suspended in 70 mL of nitrogen purged 2-methoxyethanol and refluxed for 2 h under nitrogen (dark brown).



Me₃NO (0.05 g, 0.666 mmol) was added to the solution and continued refluxing for 1 h. The resulting solution was evaporated through the rotary. Then the oily solid was dissolved in 100 ml

of water. Then the solution was precipitated with 10 ml of a saturated aqueous NH_4PF_6 solution. The precipitate was filtered by a medium porosity fritted funnel and then dried in a desiccator.

2.4.4 Crystallization of $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})](\text{PF}_6)_4$

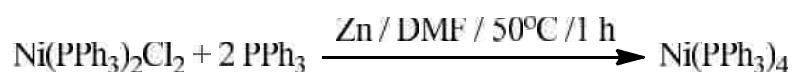
Dissolved dimer in a minimum volume of CH_3CN in a RBF until formed a clear brown solution. Set the RBF aside undisturbed to evaporate CH_3CN . Brown needle-shaped crystals were precipitated on the bottom.

Chapter 3 Results

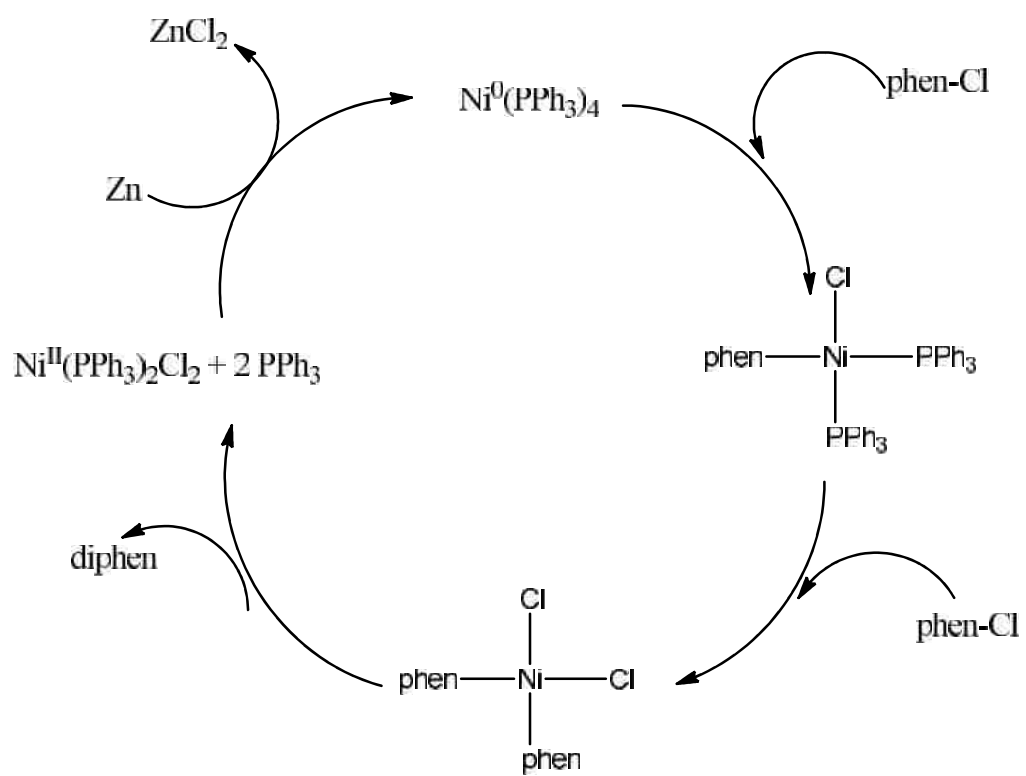
3.1 Synthesis and Basic Characterization of 5,5'-Bi-1,10-phenanthroline (diphen).

This investigation focused on photochemistry of $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2](\text{PF}_6)_4$ and a comparison to corresponding monomeric complexes $[\text{Ru}(\text{bpy})_3]^{2+}$ and $[\text{Ru}(\text{phen})_3]^{2+}$. The nature of the investigations required synthetic procedures that provided reasonable yields and most importantly were free from contamination of the dimeric complex with $[\text{Ru}(\text{phen})_3]^{2+}$ or $[\text{Ru}(\text{phen})_2(\text{diphen})]^{2+}$. $[\text{Ru}(\text{bpy})_3](\text{PF}_6)_2$ and $[\text{Ru}(\text{phen})_3](\text{PF}_6)_2$ have been thoroughly investigated and the synthesis of these complexes is well established.

The bridging ligand, diphen, was prepared using procedures previously reported⁴¹ and later optimized by Puckett⁴⁰. Briefly, diphen was prepared from 5-chloro-1,10-phenanthroline using a dichlorobis(triphenylphosphine)Ni(II), $\text{Ni}(\text{PPh}_3)_2\text{Cl}_2$, catalyst. Zinc metal is used to reduce $\text{Ni}(\text{PPh}_3)_2\text{Cl}_2$ in the presence of triphenylphosphine in oxygen-free DMF to obtain a red-brown slurry of $\text{Ni}(\text{PPh}_3)_4$ the catalyst for the reaction. The work-up procedures described in the previous chapter effectively eliminated triphenylphosphine from the product as indicated by NMR. Typical yields range from 30-50%.



Scheme 3.1 Showing the homocoupling reaction used in the preparation of diphen



Diphen is readily soluble in polar organic solvents such as acetonitrile and was amenable to analysis by proton NMR. The ^1H -NMR of diphen in deuterated acetonitrile is shown in Figure 3.1. Interpretation of the spectrum is complicated by the fact that there are basically two chemically different phenanthroline moieties that are very similar. Comparison of the spectrum obtained with diphen to that of 1,10-phenanthroline shown in Figure 3.2 supports the presence of two spectrally dissimilar rings. The protons at the 4 and 7 positions are spectrally the most dissimilar. The close proximity of the rings of the adjacent phenanthroline shift the resonance of the 4 proton to 7.9 ppm whereas the resonance for the 7 proton remains at 8.5 ppm similar to that observed with the unsubstituted phenanthroline. The remaining assignments were based on the comparison to the spectrum of 1,10-phenanthroline. The resonances centered at 9.15 ppm correspond to the protons at the 9 and 2 positions. Cross coupling in the COSEY spectrum of diphen shown in Figure 3.2 indicates that the resonance at 9.2 ppm corresponds to the proton at the 9 position (and the 9' position). The COSEY spectrum indicates that the resonance for the proton at the 3 position occurs at 7.5 ppm while the resonance for the proton at the 8 position appears at 7.8 ppm, again similar to the unsubstituted phenanthroline.

Figure 3.1 ^1H NMR spectrum of diphen in CD_3CN

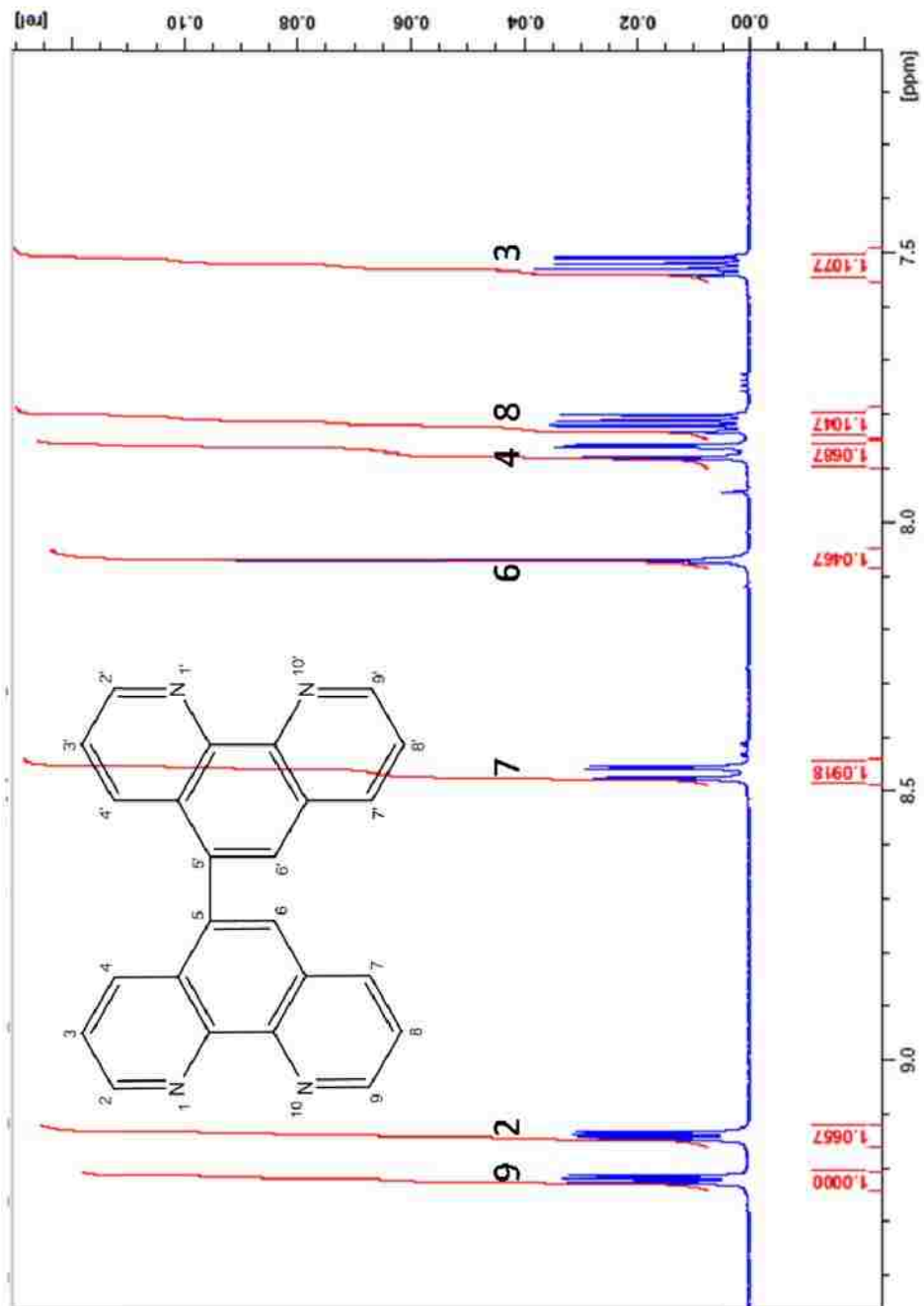


Figure 3.2 ^1H NMR spectrum of 1,10-phenanthroline in CD_3CN

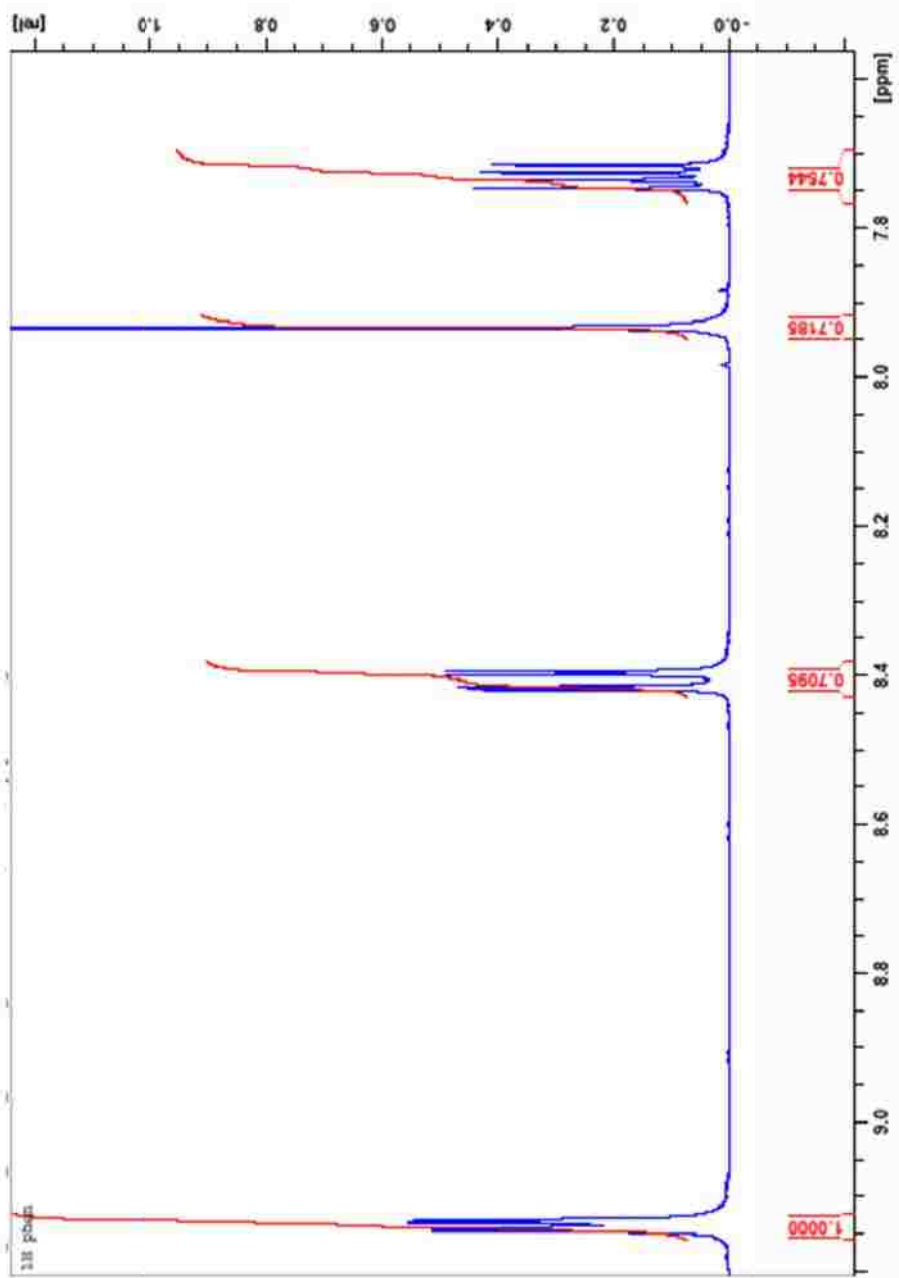
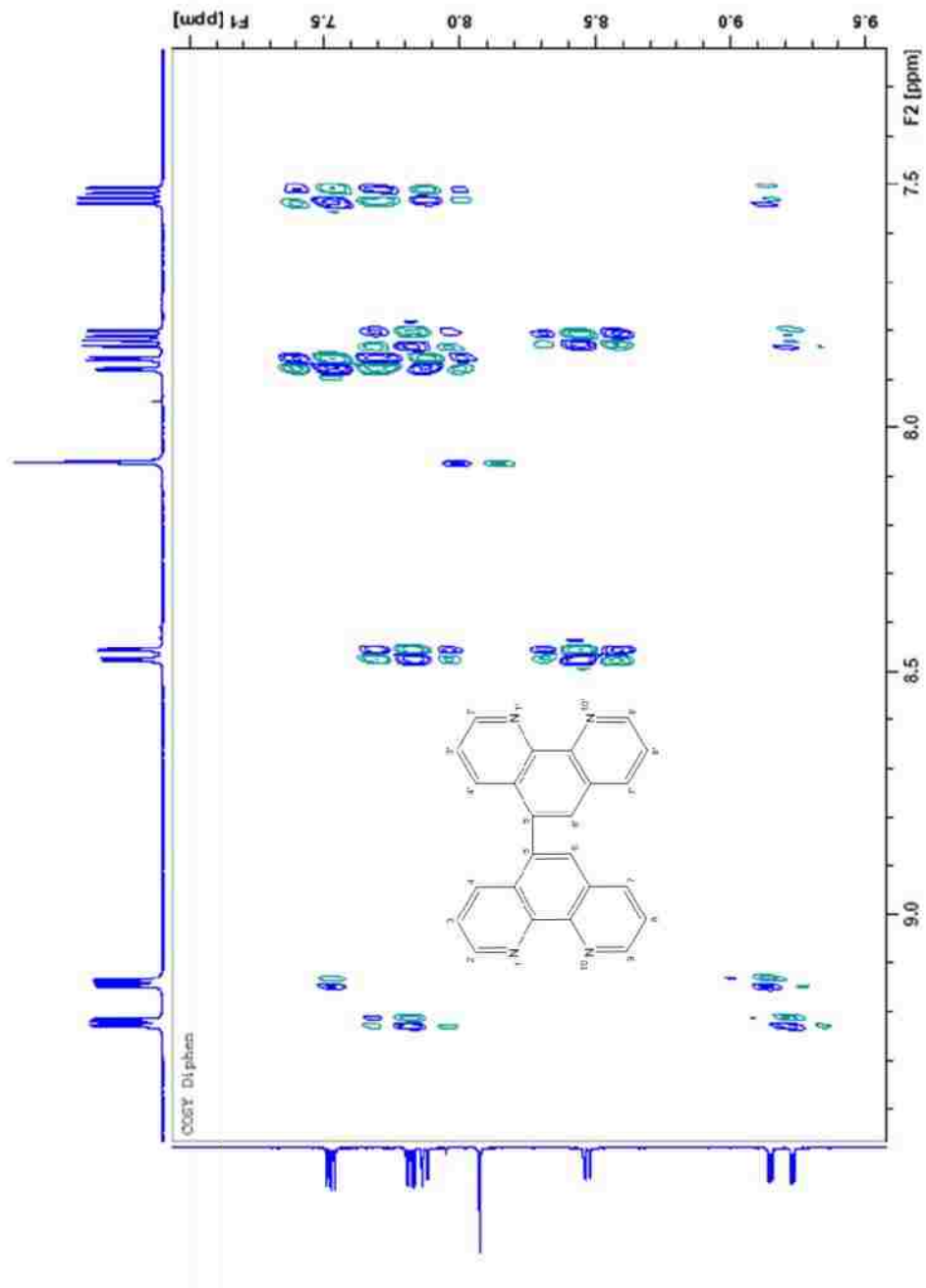
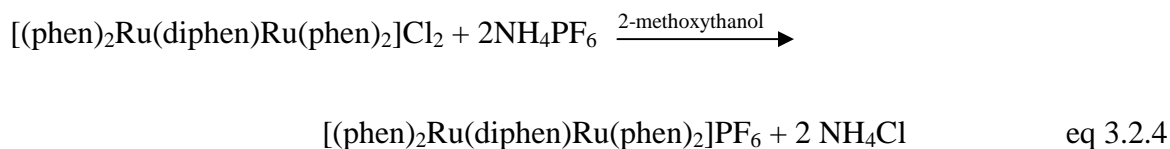
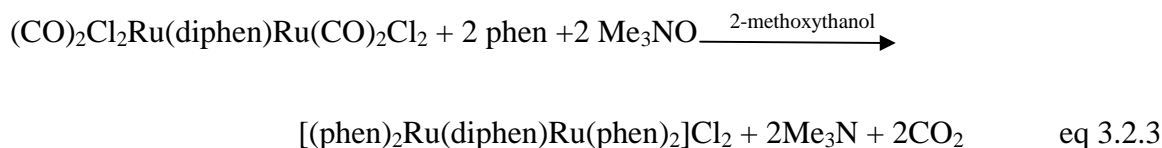
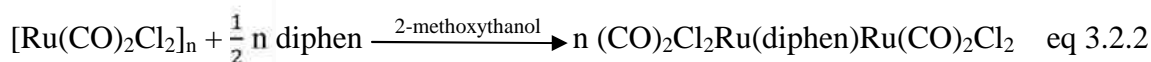
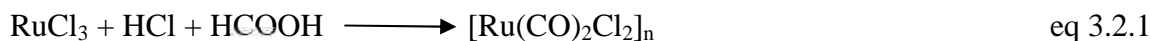


Figure 3.3 ^1H - ^1H COSY spectrum of diphen in CD_3CN



3.2 Synthesis and Basic Characterization of $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2](\text{PF}_6)_4$

$[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2](\text{PF}_6)_4$ was synthesized using a method similar to that used to prepare heteroleptic ruthenium polypyridyl complexes and later elaborated by Puckett⁴⁰ for the preparation of dimeric complexes. The following reactions summarize the chemistry



The reaction described by eq 3.2.1 was originally developed by Aguirre⁴² and has been used extensively as an entry point into the preparation of ruthenium polypyridyl complexes⁴³. The diphen and phenanthroline were added stepwise in separate reactions. The final product was recovered by precipitation as indicated by eq. 3.2.4.

$[\text{Ru}(\text{CO})_2\text{Cl}_2]_n$ is a common starting material used to prepare ruthenium polypyridine complexes. The synthesis is simple and proceeds in high yield of 90%. Although the structure of this complex is not well characterized, it is reproducible and it has characteristic carbonyl vibrational bands that are easily monitored by IR spectroscopy. The IR spectrum of $[\text{Ru}(\text{CO})_2\text{Cl}_2]_n$ shows three absorption bands at 2011 cm^{-1} , 2063 cm^{-1} and 2139 cm^{-1} typical of metal complexes containing multiple carbonyl groups.

Reaction of $[\text{Ru}(\text{CO})_2\text{Cl}_2]_n$ with diphen in 2-methoxyethanol results in the precipitation of a product in a few minutes at elevated temperatures consistent with the formulation $[(\text{CO})_2\text{Cl}_2\text{Ru}(\text{diphen})\text{Ru}(\text{CO})_2\text{Cl}_2]$. The IR spectrum of the product reveals two absorption bands at 1978 cm^{-1} and 2054 cm^{-1} . The number of bands and corresponding energies are typical of the asymmetric and symmetric stretching bands found in monomeric metal complex containing two carbonyl groups in the cis geometry.

$[(\text{CO})_2\text{Cl}_2\text{Ru}(\text{diphen})\text{Ru}(\text{CO})_2\text{Cl}_2]$ was used without further purification. Addition of trimethylamine-N-oxide was used to convert the carbonyl groups to carbon dioxide which allowed the incorporation of four equivalents of 1,10-phenanthroline to produce $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2](\text{PF}_6)_4$. No absorption bands typical of metal carbonyls in the region around 2000 cm^{-1} were evident in the IR spectrum indicating the replacement of all of CO groups during the reaction.

Figure 3.4 Attenuated total reflectance IR spectrum of neat $[\text{Ru}(\text{CO})_2\text{Cl}_2]_n$

Figure 3.5 Attenuated total reflectance IR spectrum of neat
 $[(\text{CO})_2\text{Cl}_2\text{Ru}(\text{diphen})\text{Ru}(\text{CO})_2\text{Cl}_2]$

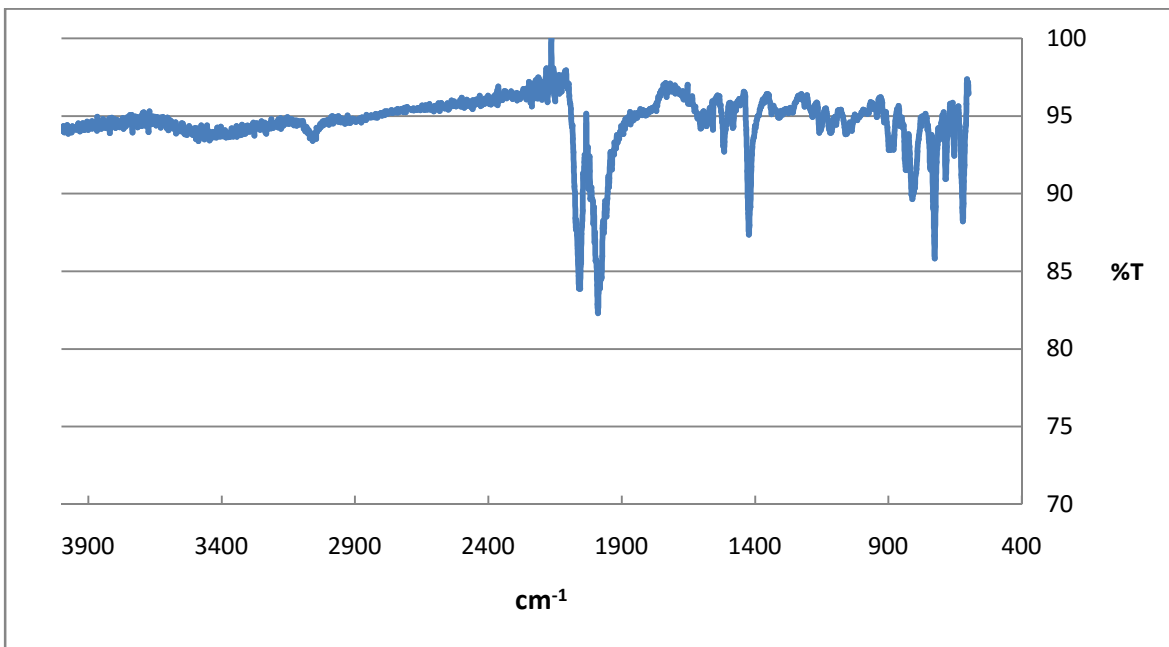
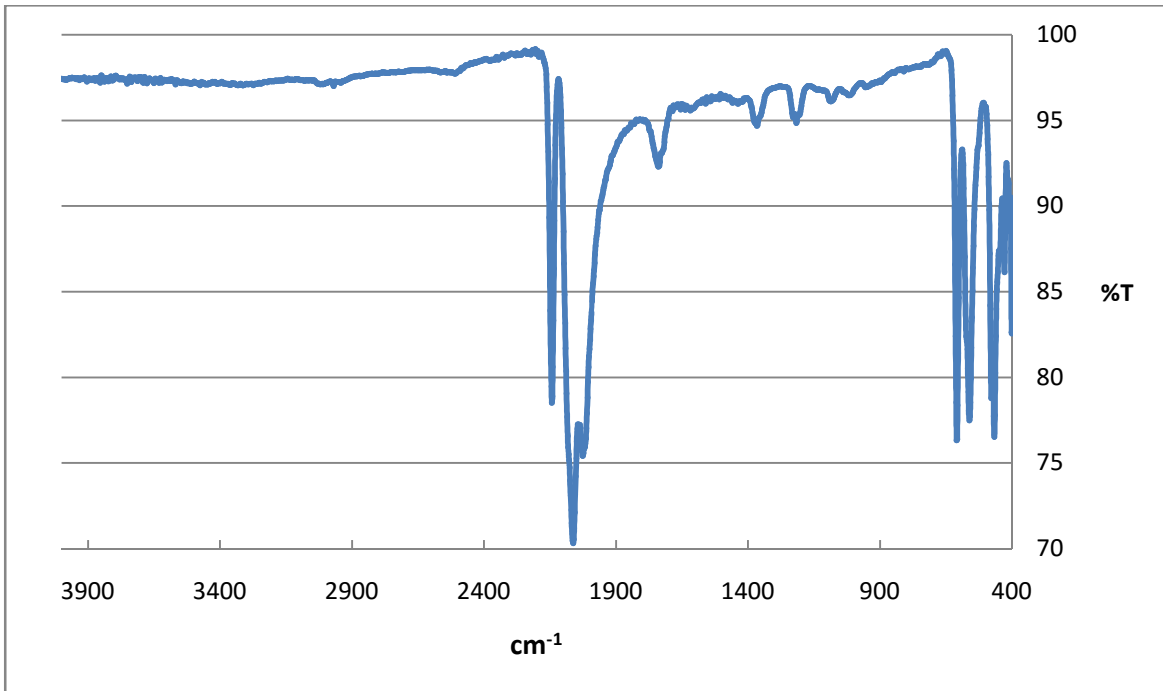
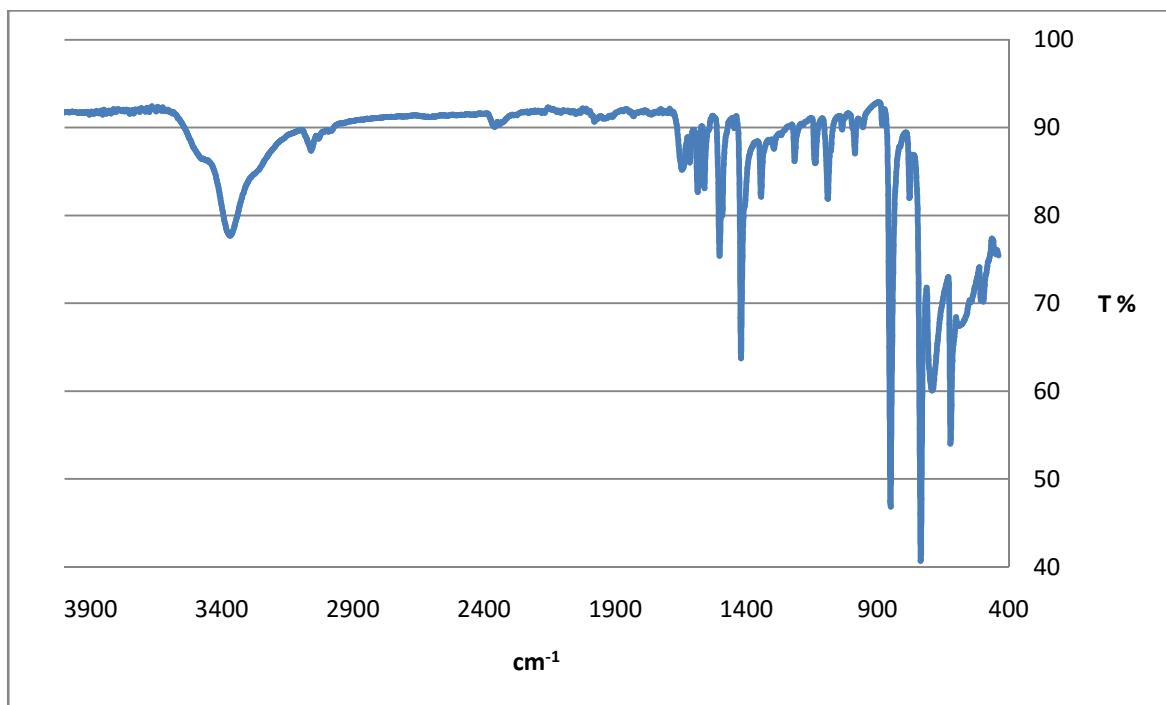
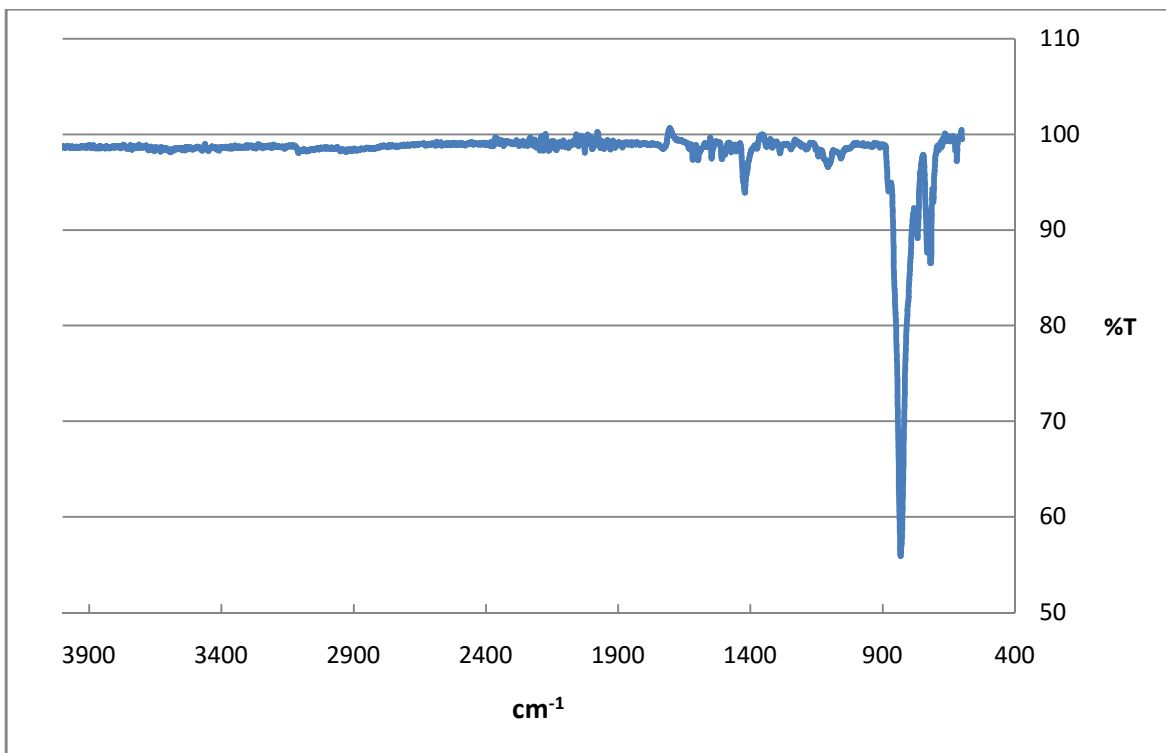


Figure 3.6 Attenuated total reflectance IR spectrum of neat
[(phen)₂Ru(diphen)Ru(phen)₂](PF₆)₄

Figure 3.7 Attenuated total reflectance IR spectrum of neat 1,10-phenanthroline



The $^1\text{H-NMR}$ of dimer in deuterated acetonitrile is shown in Figure 3.8. Interpretation of the spectrum is challenging in light of the fact that there are many similar but not identical heterocyclic rings. The spectrum shown in Figure 3.8 lacks the resolution to reveal the dissimilarities. However, the general features are consistent with the target complex. Comparison of the spectrum obtained with dimer to that of diphen clearly illustrates the structure is highly symmetric. The resonances centered at 9.1 ppm correspond to the protons at the 9 and 2 positions. The protons at the 4 and 7 positions are spectrally closer than it shows in diphen. The protons at 5 and 6 positions are identical. The resonances for the 3 and 8 protons occur at 7.8 ppm, again similar to the unsubstituted phenanthroline. The peaks indicated with X are impurities. Integration of the peaks is consistent with the target structure.

The UV-Visible absorption spectrum of the product and the corresponding monomer $[\text{Ru}(\text{phen})_3]^{2+}$ are shown in Figures 3.9 and 3.10 respectively. Comparison of the two spectra indicates that spectroscopically the two complexes are nearly identical showing a λ_{max} value of 448nm.

Figure 3.8 ^1H NMR spectrum of $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2](\text{PF}_6)_4$ in CD_3CN

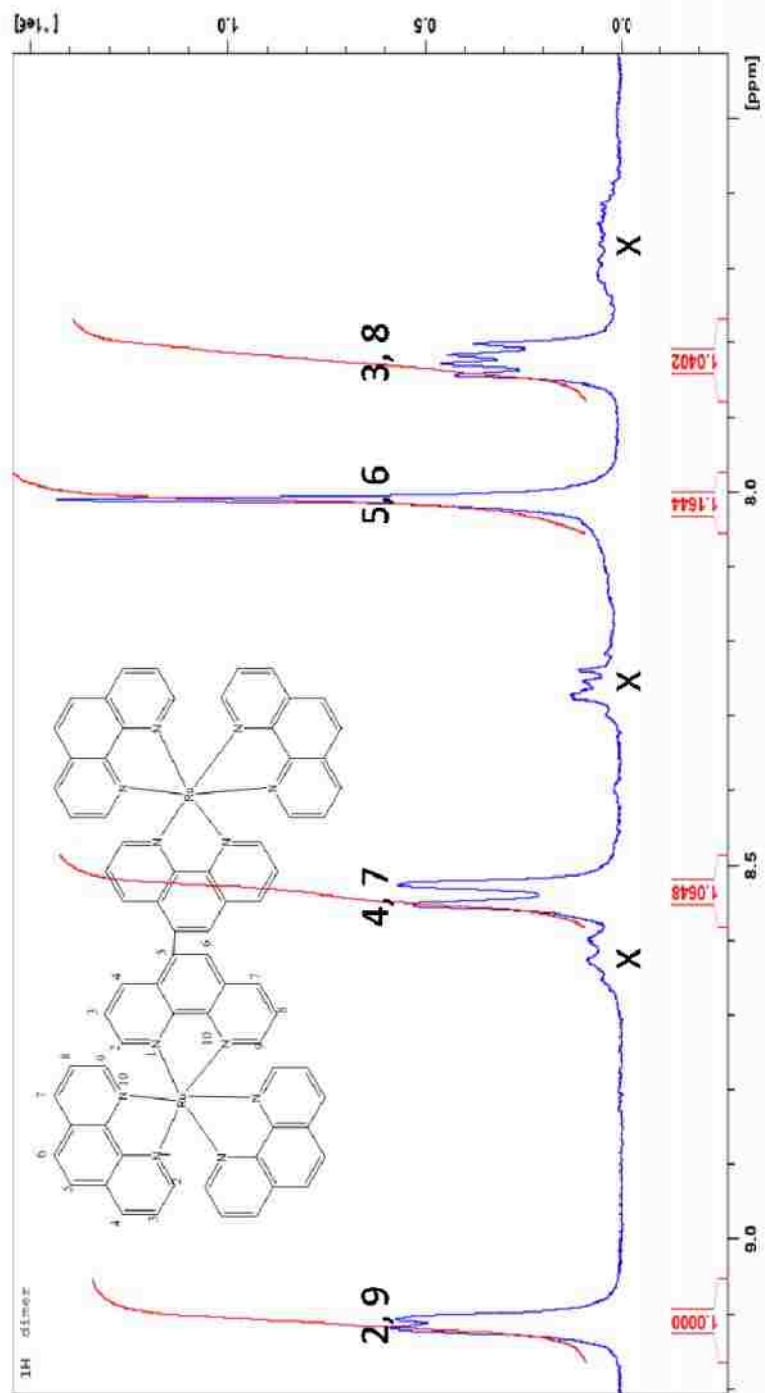
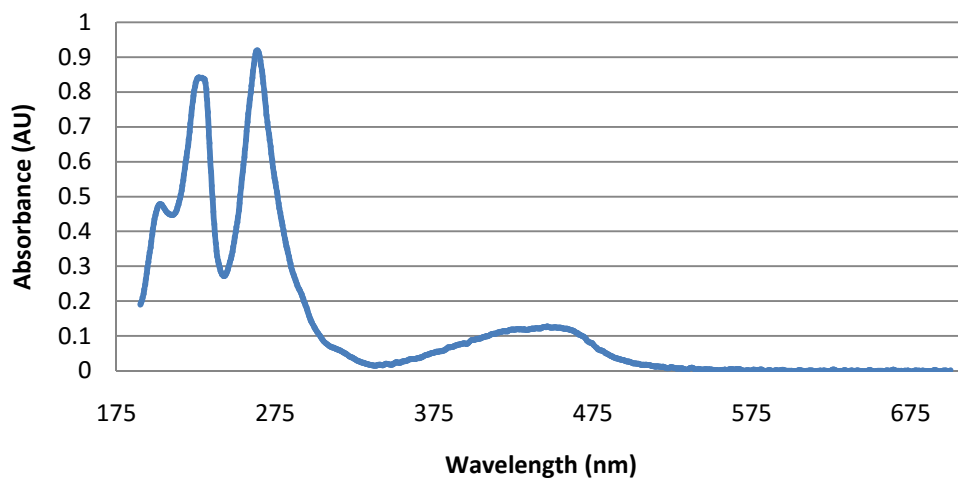


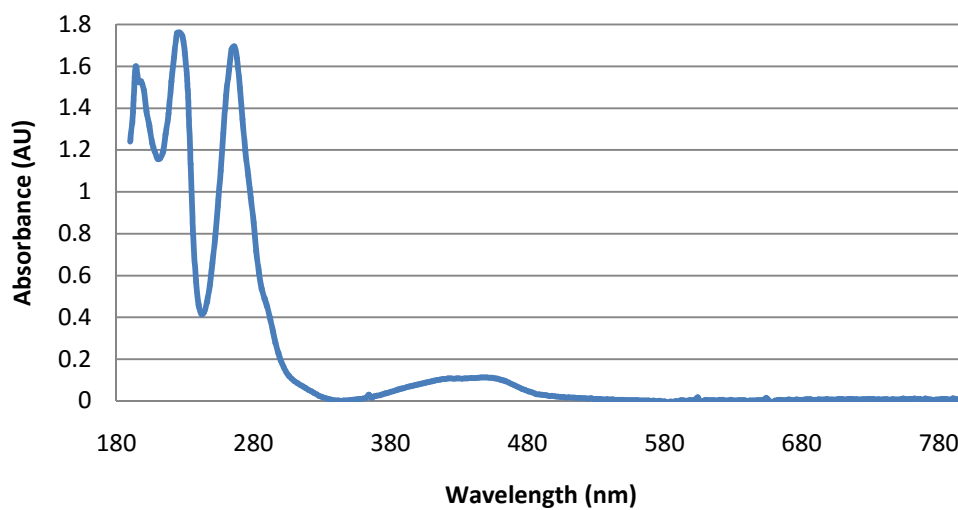
Figure 3.9 UV-Vis spectrum of $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2](\text{PF}_6)_4$ in acetonitrile

Figure 3.10 UV-Vis spectrum of $\text{Ru}(\text{phen})_3\text{Cl}_2$ in 0.2 M $(\text{NH}_4)_2\text{SO}_4$

$[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2](\text{PF}_6)_4$



$\text{Ru}(\text{phen})_3$



3.3 Structure of 5,5'-Bi-1,10-Phenanthroline

Small golden cubic crystals of diphen were obtained by slow evaporation of an acetonitrile solution. Full structure analysis of diphen crystals was performed by single crystal X-ray diffraction. Plots of the structure of diphen single molecule are shown in Figure 4.1. Selected bond lengths and angles are reported in Table 3.1-3.3.

3.4 Electrochemical studies (CV)

$$E_{1/2} = \frac{1}{2} (1.015 \text{ V} + 0.973 \text{ V}) = 0.994 \text{ V}$$

The redox potential for $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2](\text{PF}_6)_4$ is 0.994 V vs Ag/AgCl, which is similar with $\text{Ru}(\text{phen})_3^{2+}$ with a redox potential of 0.858V vs Ag/AgCl.

$[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2](\text{PF}_6)_4$ exhibits three consecutive cathodic couples (ligand-based reductions) at -0.343, -0.807, -1.095 V vs Ag/AgCl.

Table 3.1 Sample and crystal data for diphen.

Chemical formula	C ₂₆ H ₁₇ N ₅	
Formula weight	399.45 g/mol	
Temperature	100(2) K	
Wavelength	1.54178 Å	
Crystal size	0.151 x 0.188 x 0.202 mm	
Crystal system	monoclinic	
Space group	C 1 2/c 1	
Unit cell dimensions	a = 16.7253(7) Å	= 90°
	b = 11.2482(5) Å	= 120.6180(10)°
	c = 11.5844(5) Å	= 90°
Volume	1875.52(14) Å ³	
Z	4	
Density (calculated)	1.415 g/cm ³	
Absorption coefficient	0.685 mm ⁻¹	
F(000)	832	

Table 3.2 Bond lengths (\AA) for diphen.

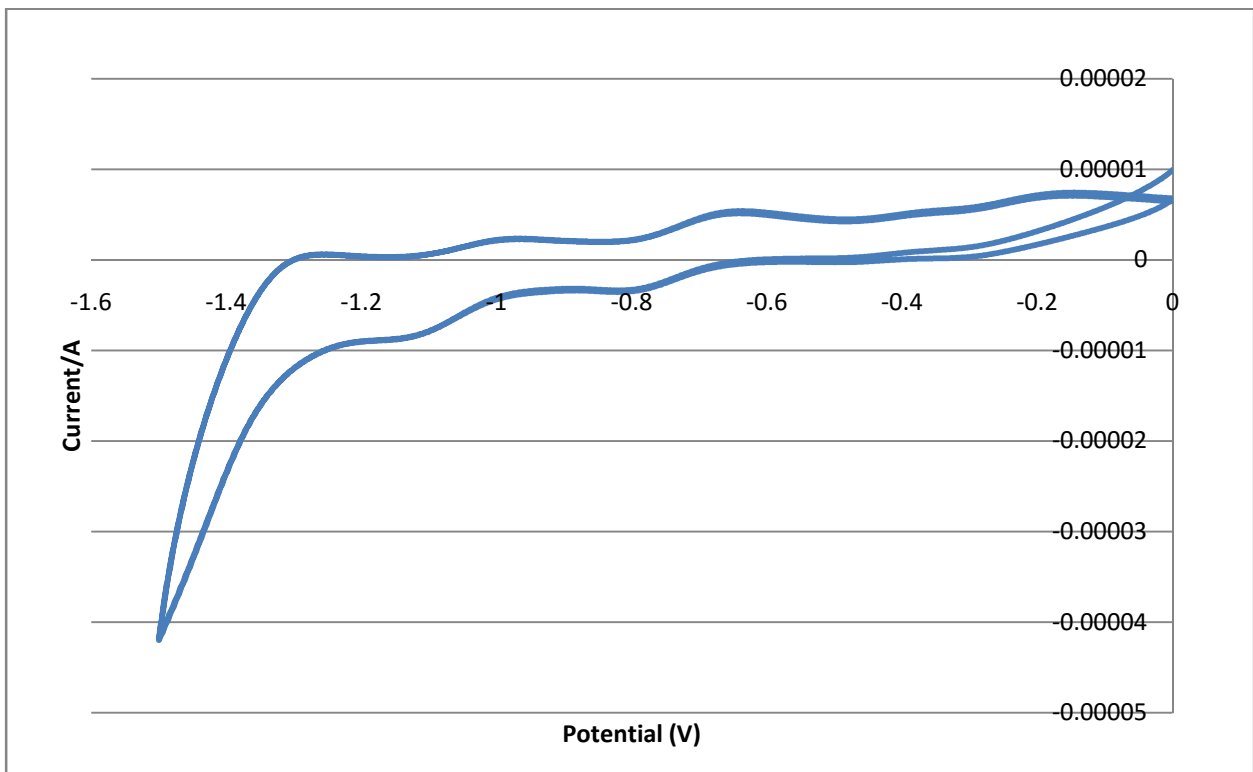
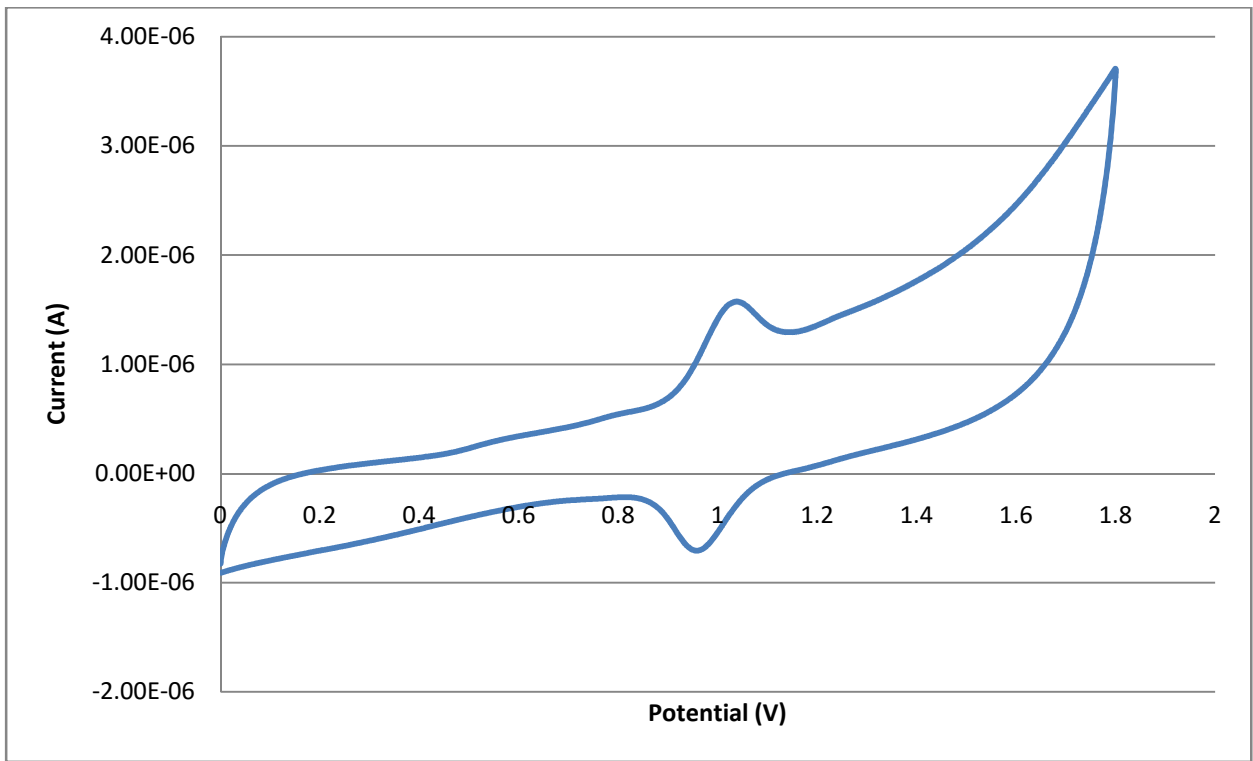
N1-C1	1.325(2)	N1-C5	1.361(2)
N2-C10	1.321(2)	N2-C6	1.357(2)
C12-C11	1.357(2)	C12-C7	1.451(2)
C12-C12	1.494(3)	C6-C7	1.412(2)
C6-C5	1.453(2)	C4-C5	1.409(3)
C4-C3	1.412(2)	C4-C11	1.435(2)
C3-C2	1.369(3)	C3-H3	0.95
C11-H11	0.95	C7-C8	1.412(3)
C8-C9	1.371(3)	C8-H8	0.95
C1-C2	1.403(3)	C1-H1	0.95
C9-C10	1.405(3)	C9-H9	0.95
C10-H10	0.95	C2-H2	0.95
C13A-C13A	1.67037(4)	C13A-H13A	0.98
C13A-H13B	0.98	C13A-H13C	0.98
N14-C13A	1.11703(4)	N13-C14	1.11719(3)
C14-C15	1.15875(5)	C15-H15A	0.98
C15-H15B	0.98	C15-H15C	0.98

Table 3.3 Bond angles ($^{\circ}$) for diphen

C1-N1-C5	117.25(16)	C10-N2-C6	117.99(16)
C11-C12-C7	119.79(16)	C11-C12-C12	120.56(13)
C7-C12-C12	119.65(13)	N2-C6-C7	122.81(16)
N2-C6-C5	117.99(16)	C7-C6-C5	119.20(16)
C5-C4-C3	117.89(17)	C5-C4-C11	120.01(16)
C3-C4-C11	122.04(16)	C2-C3-C4	118.68(17)
C2-C3-H3	120.7	C4-C3-H3	120.7
C12-C11-C4	121.51(16)	C12-C11-H11	119.2
C4-C11-H11	119.2	C6-C7-C8	117.06(16)
C6-C7-C12	120.17(16)	C8-C7-C12	122.77(17)
C9-C8-C7	120.02(18)	C9-C8-H8	120.0
C7-C8-H8	120.0	N1-C5-C4	122.99(16)
N1-C5-C6	117.83(16)	C4-C5-C6	119.16(16)
N1-C1-C2	123.78(17)	N1-C1-H1	118.1
C2-C1-H1	118.1	C8-C9-C10	118.23(17)
C8-C9-H9	120.9	C10-C9-H9	120.9
N2-C10-C9	123.86(17)	N2-C10-H10	118.1
C9-C10-H10	118.1	C3-C2-C1	119.39(17)
C3-C2-H2	120.3	C1-C2-H2	120.3
C13A-C13A-H13A	109.5	C13A-C13A-H13B	109.5
H13A-C13A-H13B	109.5	C13A-C13A-H13C	109.5
H13A-C13A-H13C	109.5	H13B-C13A-H13C	109.5
N13-C14-C15	168.7410(10)	C14-C15-H15A	109.5
C14-C15-H15B	109.5	H15A-C15-H15B	109.5
C14-C15-H15C	109.5	H15A-C15-H15C	109.5
H15B-C15-H15C	109.5		

Figure 3.11 Cyclic Voltammogram of $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2](\text{PF}_6)_4$ in 0.1M TBAPF₆ in acetonitrile versus Ag/AgCl

Figure 3.12 Cyclic Voltammogram of $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2](\text{PF}_6)_4$ under nitrogen in 0.1M TBAPF₆ in acetonitrile.



3.5 Oxidative Quenching of Ru(II) Dimer and Monomers by Peroxydisulfate

The quenching reactions of $\text{Ru}(\text{bpy})_3^{2+}$, $\text{Ru}(\text{phen})_3^{2+}$ and the dimer in aqueous solutions by $\text{S}_2\text{O}_8^{2-}$ were investigated using traditional Stern-Volmer techniques steady-state irradiation and laser flash photolysis. The monomeric complexes have been previously investigated in detail and were used for comparison. The focus of the laser experiments was twofold; monitor the rate of the back-reaction and measure, by comparison, the quantum yield for the quenching reaction. The steady-state irradiation experiments were also done to provide information about the quantum yield for product formation.

In the fluorescence measurements, the ruthenium complexes concentrations are enough to provide an absorbance of around 0.8, and the concentrations of $\text{S}_2\text{O}_8^{2-}$ were varied from 0 to $3.0 \times 10^{-4} \text{ mol L}^{-1}$. When excited with a wavelength of 450 nm, the ruthenium complexes have a strong fluorescence emission peaked at 605 nm. The addition of $\text{S}_2\text{O}_8^{2-}$ caused a gradual decrease in the fluorescence intensity of ruthenium complexes and the emission had a red-shift for the dimer and $[\text{Ru}(\text{phen})_3]^{2+}$. This suggested that the microenvironment around the dimer and $[\text{Ru}(\text{phen})_3]^{2+}$ was changed after the addition of $\text{S}_2\text{O}_8^{2-}$.

Stern-Volmer plots for $\text{S}_2\text{O}_8^{2-}$ quenching of ruthenium complexes fluorescence are shown in Figure 3.17. The possible quenching reactions were analyzed by the Stern-Volmer equation:

$$\frac{I_0}{I} = 1 + k_q \tau_0 [Q] = 1 + K_{sv} [Q]$$

Where I_0 and I are the steady-state fluorescence intensities in the absence and presence of $\text{S}_2\text{O}_8^{2-}$ quencher, respectively, k_q the bimolecular quenching rate constant, τ_0 the life time of fluorescence in absence of $\text{S}_2\text{O}_8^{2-}$, K_{sv} the Stern-volmer quenching constant, and $[Q]$ the

concentration of $S_2O_8^{2-}$. Hence the above equation could be applied to determine k_q by linear regression of a plot of I_0/I against. The result shows that the quenching constant (Figure 3.17 and Table 3.1), k_q , for the dimer is higher than $[Ru(phen)_3]^{2+}$.

A representative absorption transient showing the formation of Ru(III) from ruthenium(II) complexes are shown in Figure 3.17 and 3.18. Absorbance changes $A = 0.0946$ for dimer, which is about two times as $A = 0.0474$ for $Ru(phen)_3^{2+}$. Absorbance changes A are 0.1410 and 0.0988 for dimer and $Ru(bpy)_3^{2+}$, respectively. The slow down curves showed in the monomers $Ru(phen)_3^{2+}$ and $Ru(bpy)_3^{2+}$ show that the formation of Ru(III) is slower than form dimer.

A 300 watt, 120 volt bulb “sunlamp” with no filtering was also used for photochemical reactions. The photochemical reaction of the dimer and $S_2O_8^{2-}$ was monitored via UV-vis spectroscopy. The spectra can be seen in Figure 3.20. Figure 3.19 for $Ru(phen)_3^{2+}$ is used as a comparison. The peak at 448 nm, which is correlates to the Ru(II) concentration, decreases over time. This is the expected result of a quench reaction where $Ru(II)^*$ is oxidized by $S_2O_8^{2-}$ to Ru(III). Absorbance changes A at 30 s are 0.5837 and 0.05145 for dimer and $Ru(phen)_3^{2+}$, respectively. With the time consuming, the absorbance changes of $Ru(phen)_3^{2+}$ become much smaller, where as the dimer almost maintain the same, which means the dimer excited state is more stable.

Oxidation reaction of the dimer was performed with fresh produced chlorine, and the product was detected with UV-vis spectroscopy showed in Figure 3.21. Figure 3.22 for $Ru(bpy)_3^{2+}$ is used as a comparison. A new peak shows at 622 nm only for the dimer, which means different product is being formed from $Ru(bpy)_3^{2+}$.

A representative absorption transient showing the rapid formation of Ru(I) and the subsequent slow back reaction that returns the system to the initial state is shown in Figure 3.27. Absorbance changes ΔA are 0.0231, 0.0289 and 0.0199 for $\text{Ru}(\text{bpy})_3^{2+}$, $\text{Ru}(\text{phen})_3^{2+}$ and dimer, respectively.

Table 3.4 Stern-Volmer quenching constants for the ruthenium complexes with $\text{S}_2\text{O}_8^{2-}$

pH	Ruthenium complexes	τ_0 (ns)	K_{sv} ($\times 10^3 \text{ M}^{-1}$)	k_q ($\times 10^9 \text{ M}^{-1} \text{ s}^{-1}$)
4	$[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2]^{2+}$	1470	9.719	6.612
	$[\text{Ru}(\text{phen})_3]^{2+}$	1000	2.236	2.236
	$[\text{Ru}(\text{bpy})_3]^{2+}$	590	7.050	11.95

Figure 3.13 Effect of $[\text{S}_2\text{O}_8^{2-}]$ on the emission intensity of the dimer in 0.2 M $(\text{NH}_4)_2\text{SO}_4$

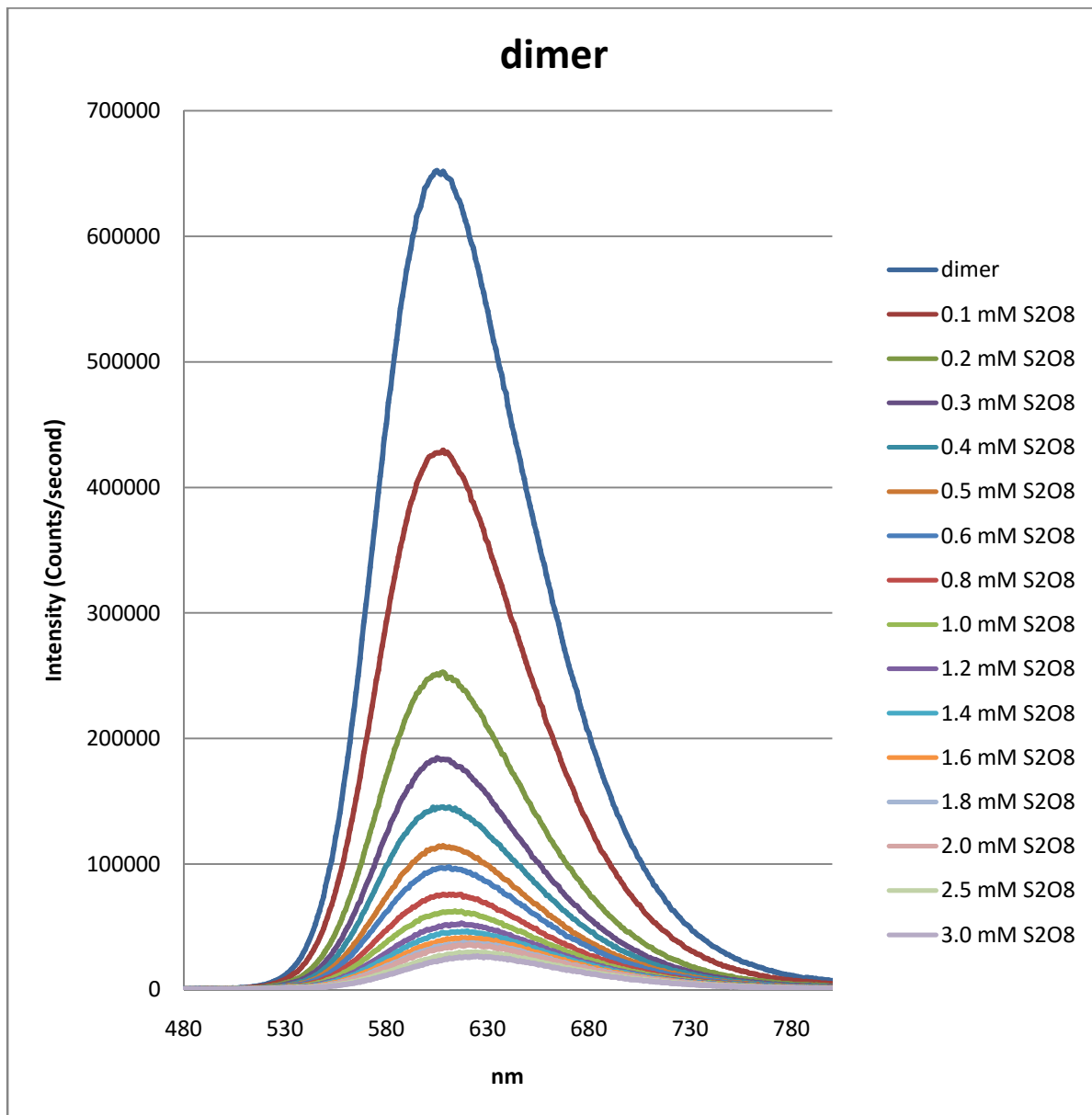


Figure 3.14 Effect of $[\text{S}_2\text{O}_8^{2-}]$ on the emission intensity of $\text{Ru}(\text{phen})_3^{2+}$ in 0.2 M $(\text{NH}_4)_2\text{SO}_4$

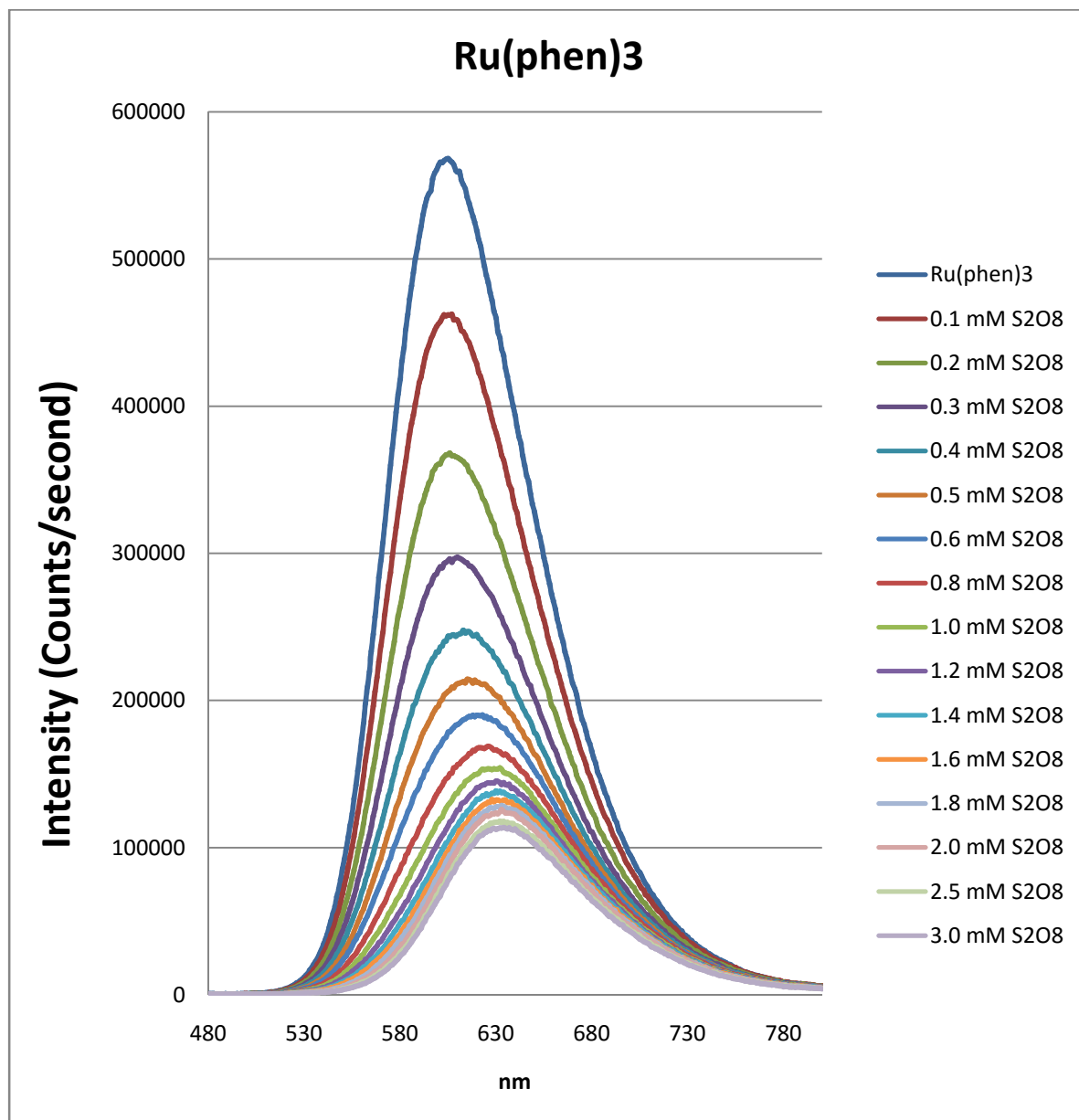


Figure 3.15 Effect of $[\text{S}_2\text{O}_8^{2-}]$ on the emission intensity of $\text{Ru}(\text{bpy})_3^{2+}$ in 0.2 M $(\text{NH}_4)_2\text{SO}_4$.

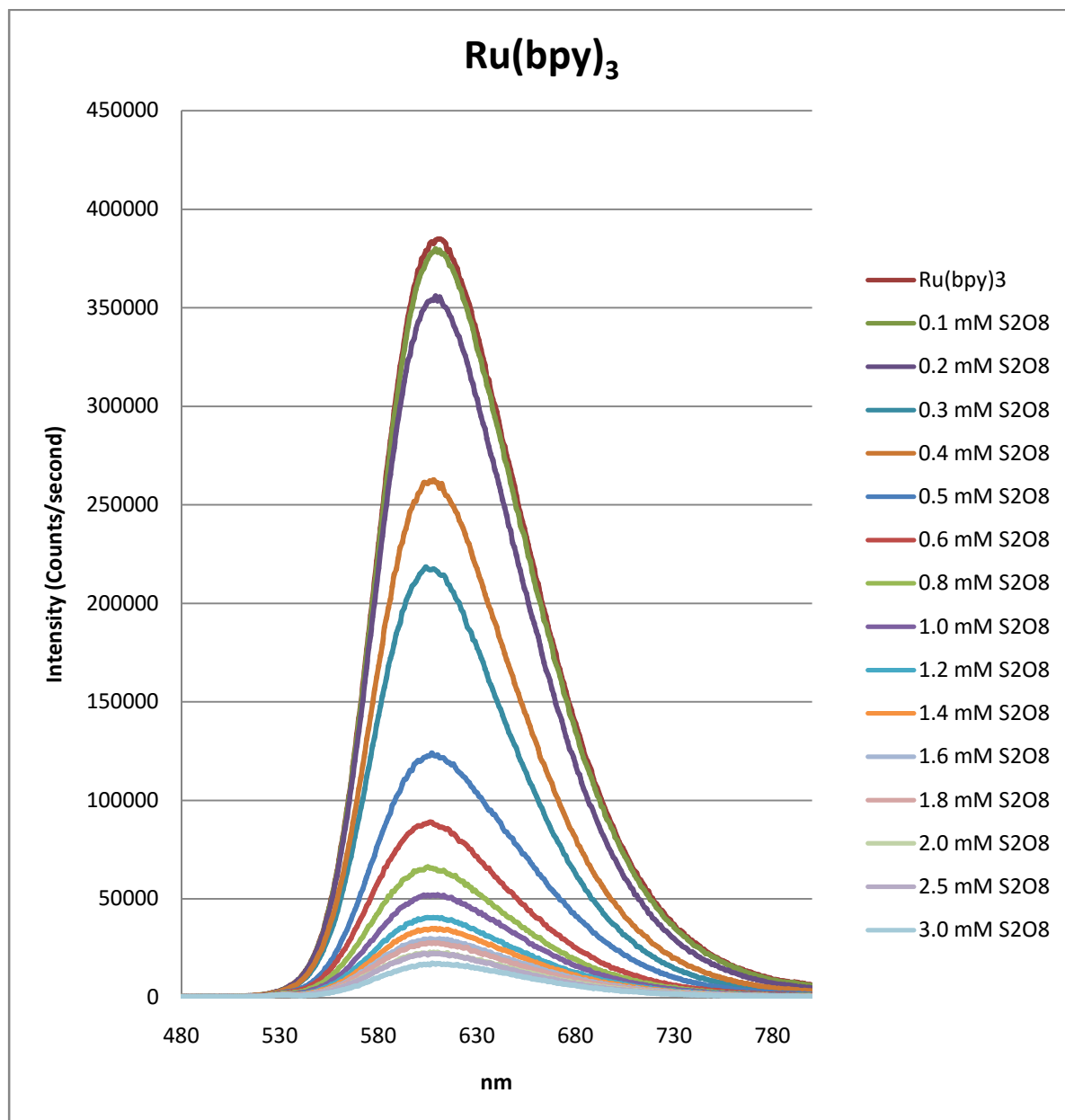


Figure 3.16 Plots of the relative emission intensities of $\text{Ru}(\text{bpy})_3^{2+}$, $\text{Ru}(\text{phen})_3^{2+}$ and the dimer versus the $[\text{S}_2\text{O}_8^{2-}]$ concentration. The solid lines and the associated equations were obtained using linear least squares and are the “best” fits.

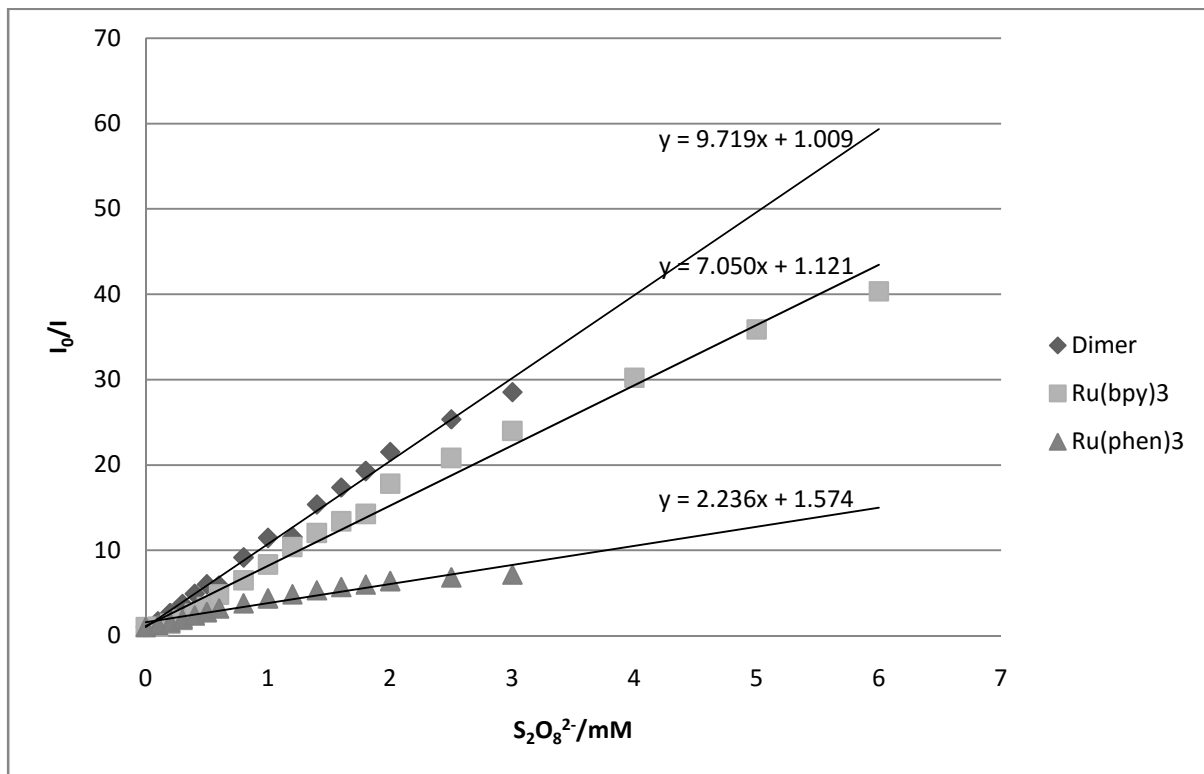


Figure 3.17 Comparison of transient absorbance change at 450 nm of a solution of dimer and $\text{Ru}(\text{phen})_3^{2+}$ in 0.2 M $(\text{NH}_4)_2\text{SO}_4$ following a short laser pulse.

Figure 3.18 Comparison of transient absorbance change at 450 nm of a solution of dimer and $\text{Ru}(\text{bpy})_3^{2+}$ in 0.2 M $(\text{NH}_4)_2\text{SO}_4$ following a short laser pulse.

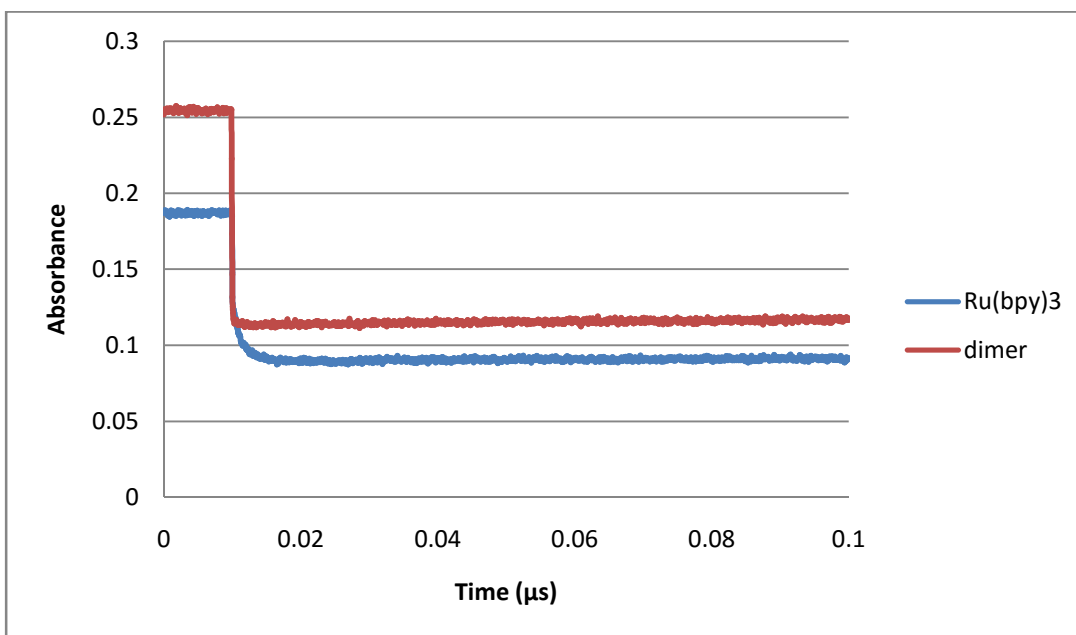
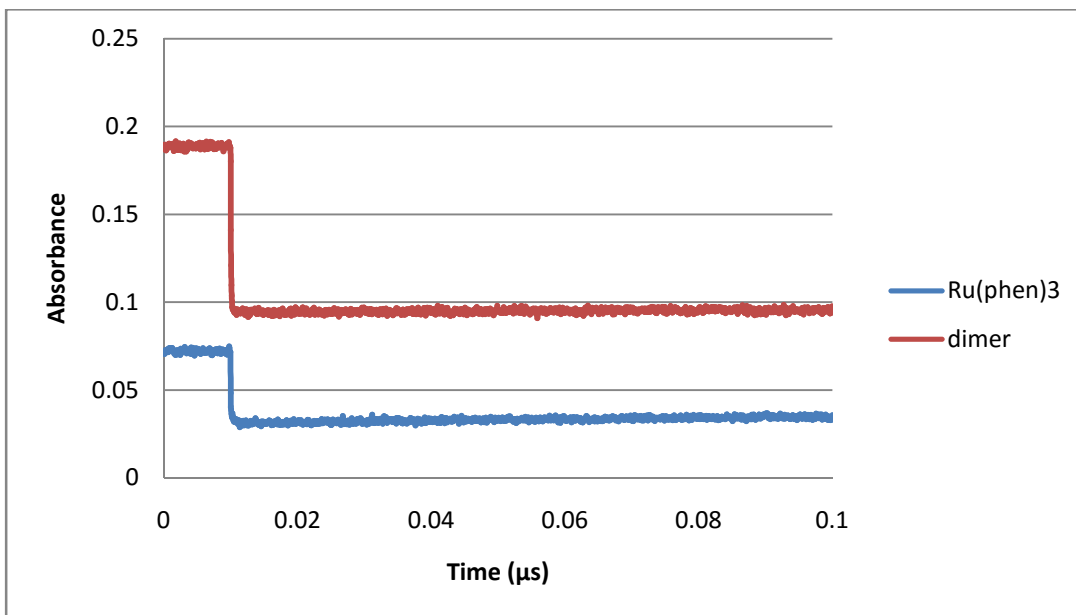


Figure 3.19 Changes in absorbance following steady state irradiation of $\text{Ru}(\text{phen})_3^{2+}$ in 0.2 M $(\text{NH}_4)_2\text{SO}_4$ containing 0.5 mM $\text{S}_2\text{O}_8^{2-}$.

Figure 3.20 Changes in Absorbance following steady state irradiation of the dimer in 0.2 M $(\text{NH}_4)_2\text{SO}_4$ containing 0.5 mM $\text{S}_2\text{O}_8^{2-}$.

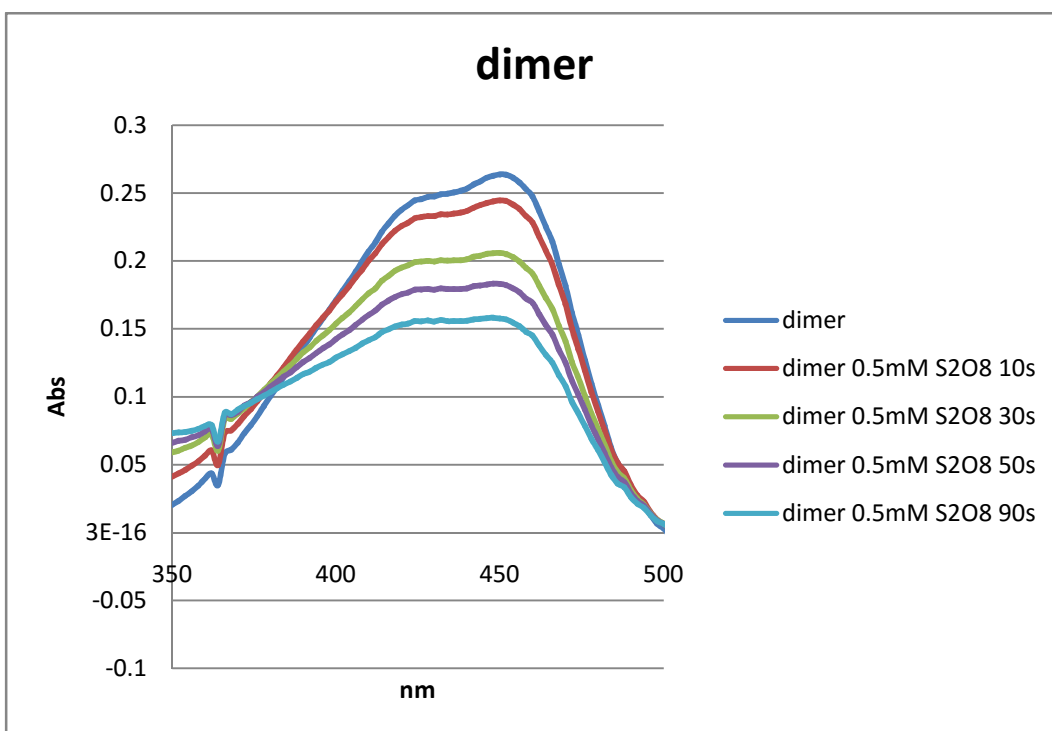
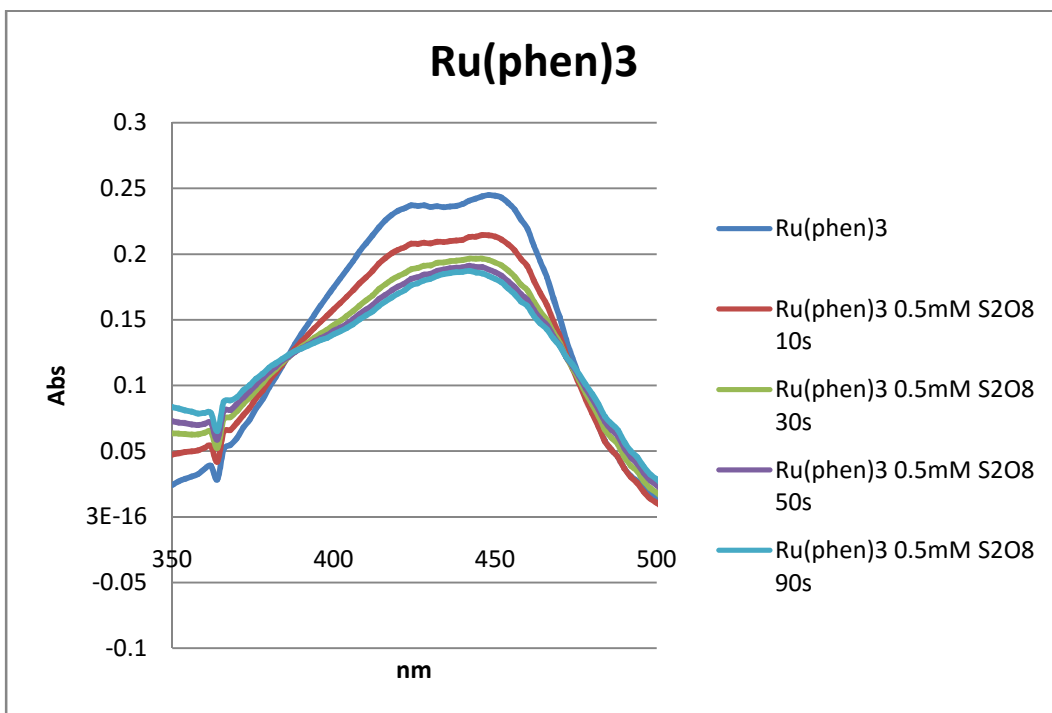
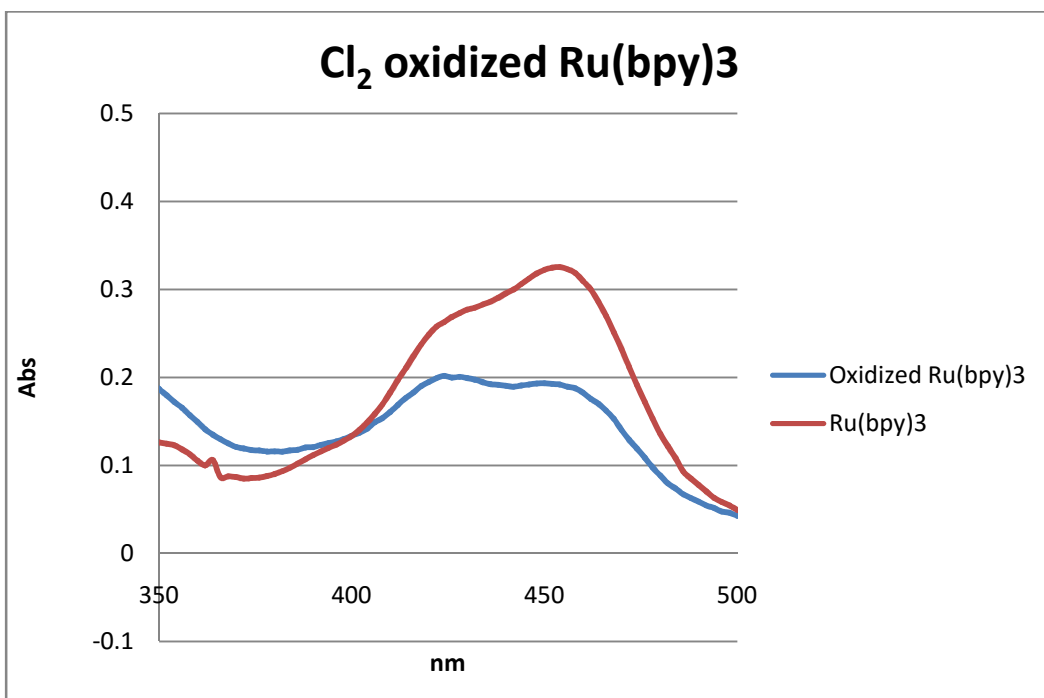
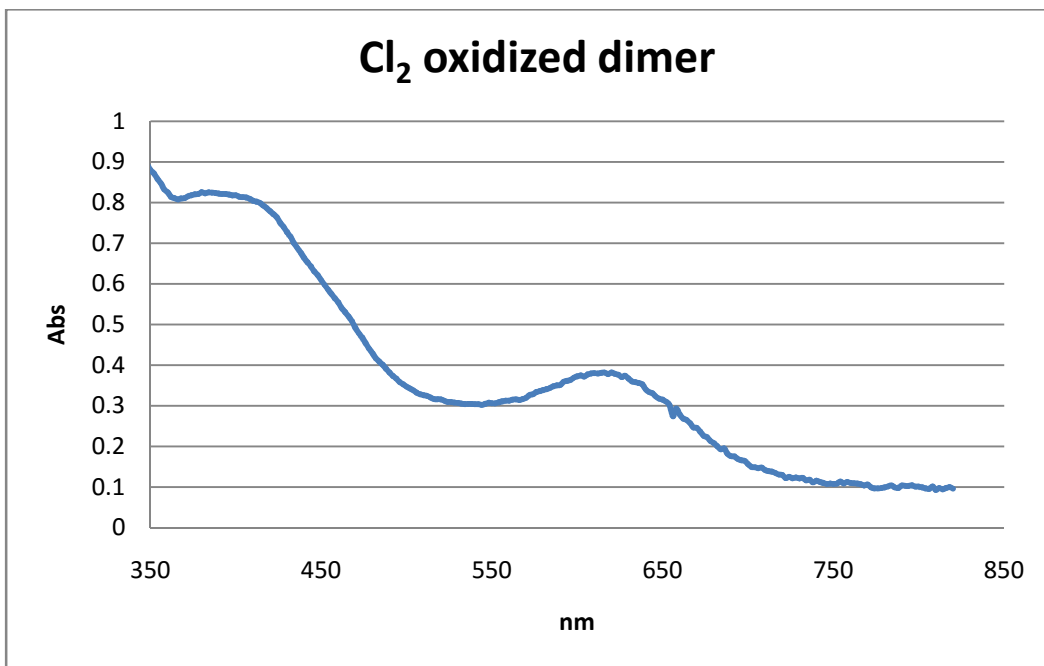


Figure 3.21 UV-vis spectrum of the dimer oxidized by Cl₂

Figure 3.22 UV-vis spectrum of the Ru(bpy)₃²⁺ oxidized by Cl₂



3.6 Reductive Quenching of Ru(II) Dimer and Monomers by Ascorbate

The quenching reactions of $\text{Ru}(\text{bpy})_3^{2+}$, $\text{Ru}(\text{phen})_3^{2+}$ and the dimer in aqueous solutions by ascorbate were investigated using traditional Stern-Volmer techniques and laser flash photolysis. The monomeric complexes have been previously investigated in detail and were used for comparison.

In the fluorescence measurements with ascorbate, the ruthenium complexes concentrations are enough to provide an absorbance of around 0.3, and the concentrations of ascorbate were varied from 0 to 0.1 mol L⁻¹. When excited with a wavelength of 450 nm, the ruthenium complexes have a strong fluorescence emission peaked at 605 nm. The addition of ascorbate caused a gradual decrease in the fluorescence intensity of ruthenium complexes and the emission had no shift.

Stern-Volmer plots for ascorbate quenching of ruthenium complexes fluorescence are shown in Figure 3.26. The result shows that the quenching constant (Figure 3.26 and Table 3.5), k_q , for the dimer is comparable to the k_q value for $\text{Ru}(\text{phen})_3^{2+}$. This indicates that the monomeric complex and the dimeric complex behave similarly.

Table 3.5 Stern-Volmer quenching constants for the ruthenium complexes with ascorbate

pH	Ruthenium complexes	τ_0 (ns)	K_{sv} ($\times 10^3 M^{-1}$)	k_q ($\times 10^8 M^{-1}s^{-1}$)
5	$[(phen)_2Ru(diphen)Ru(phen)_2]^{2+}$	1470	0.492	3.347
	$[Ru(phen)_3]^{2+}$	1000	0.324	3.324
	$[Ru(bpy)_3]^{2+}$	590	0.021	0.356

Figure 3.23 Effect of [ascorbate] on the emission intensity of the dimer in argon purged 0.2 M (NH₄)₂SO₄

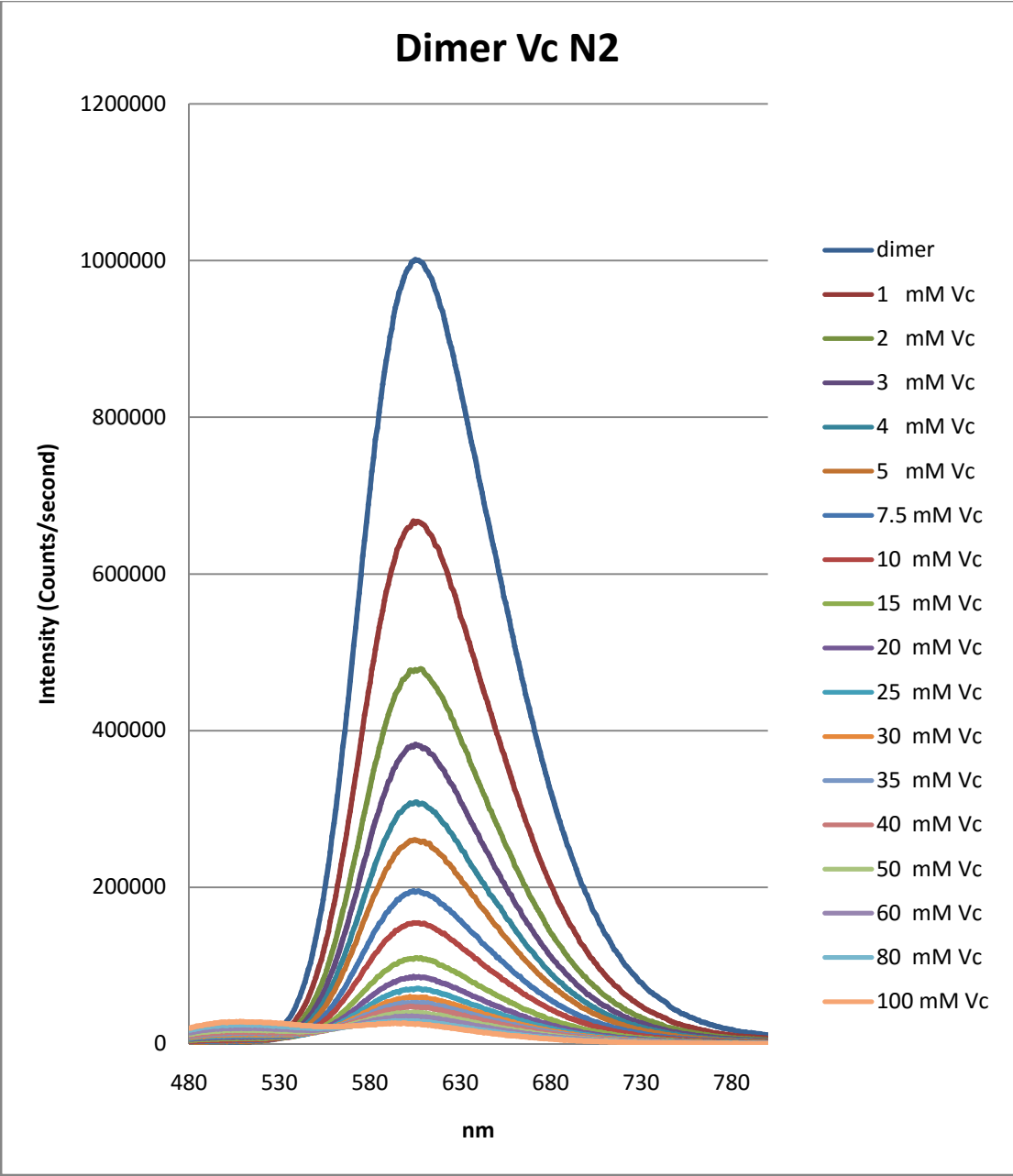


Figure 3.24 Effect of ascorbate on the emission intensity of $\text{Ru}(\text{phen})_3^{2+}$ in argon purged 0.2 M $(\text{NH}_4)_2\text{SO}_4$.

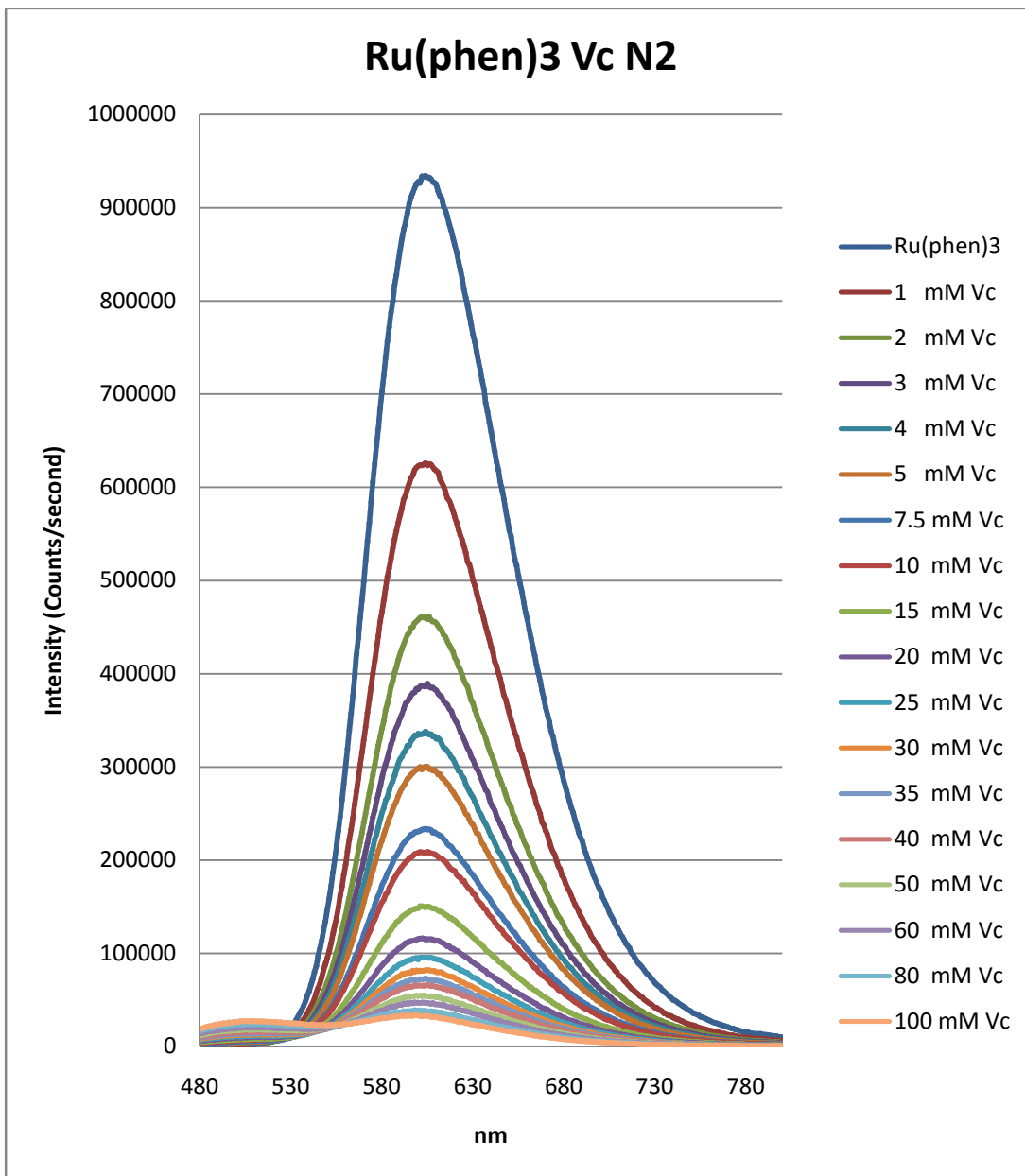


Figure 3.25 Effect of ascorbate on the emission intensity of $\text{Ru}(\text{bpy})_3^{2+}$ in argon purged 0.2 M $(\text{NH}_4)_2\text{SO}_4$.

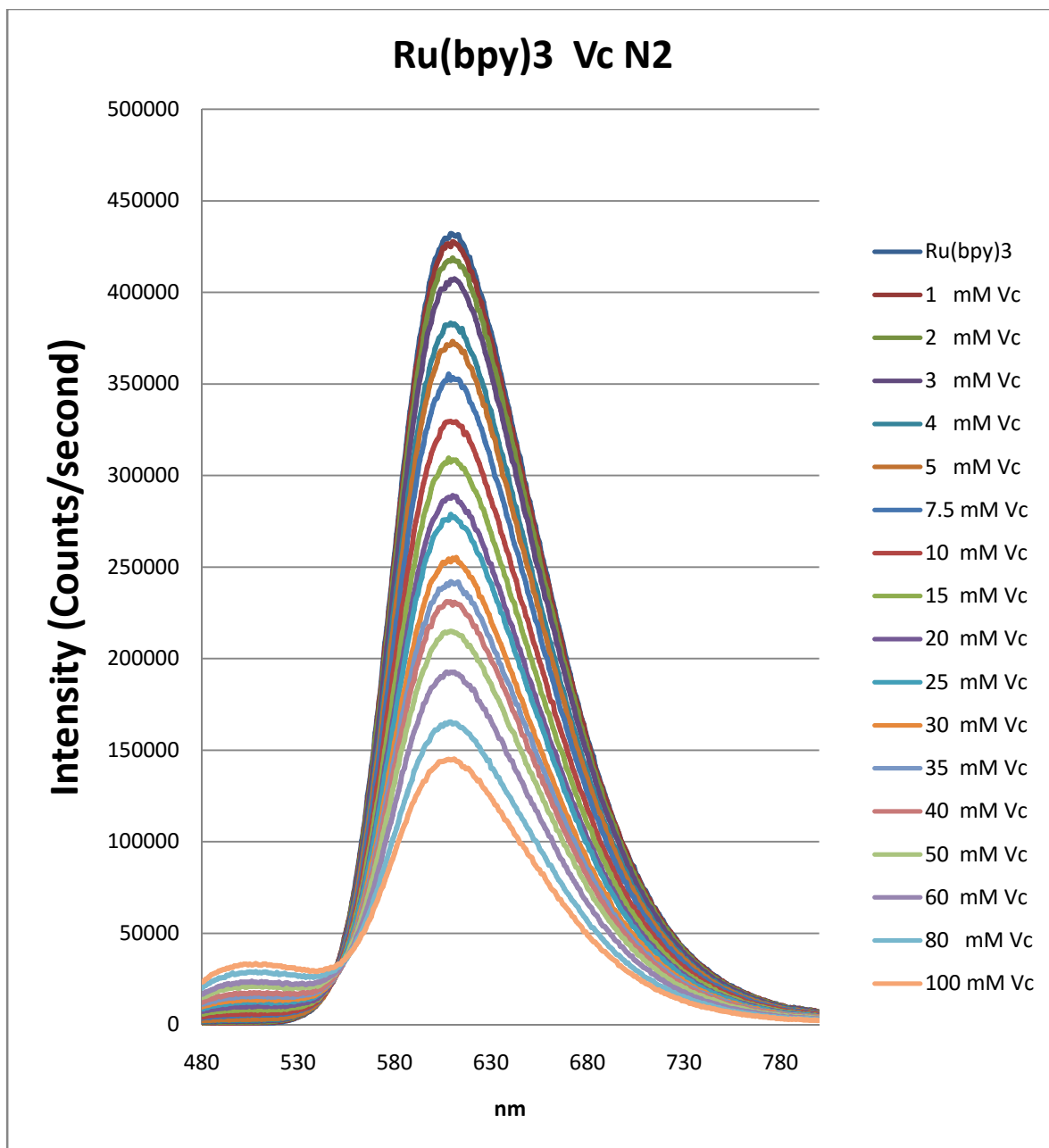


Figure 3.26 Plots of the relative emission intensities of $\text{Ru}(\text{bpy})_3^{2+}$, $\text{Ru}(\text{phen})_3^{2+}$ and the dimer versus the ascorbate concentration. The solid lines and the associated equations were obtained using linear least squares and are the “best” fits.

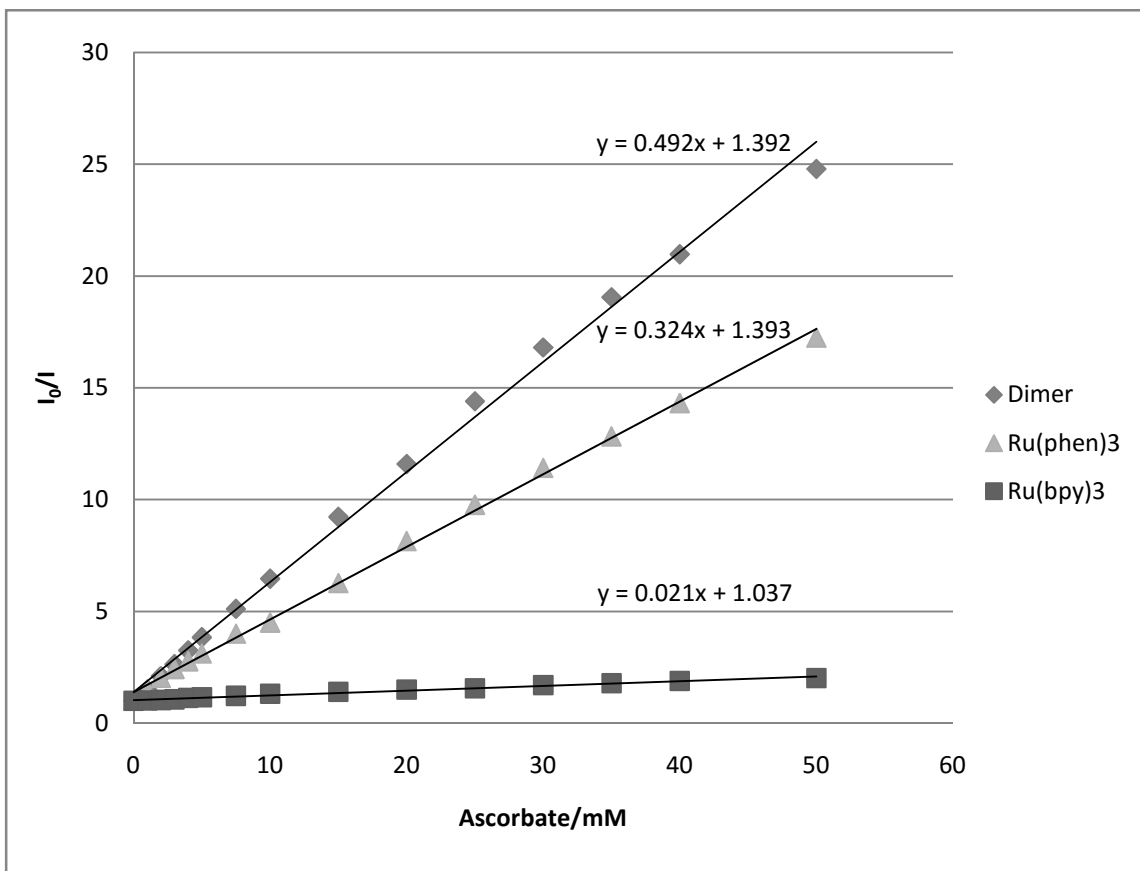
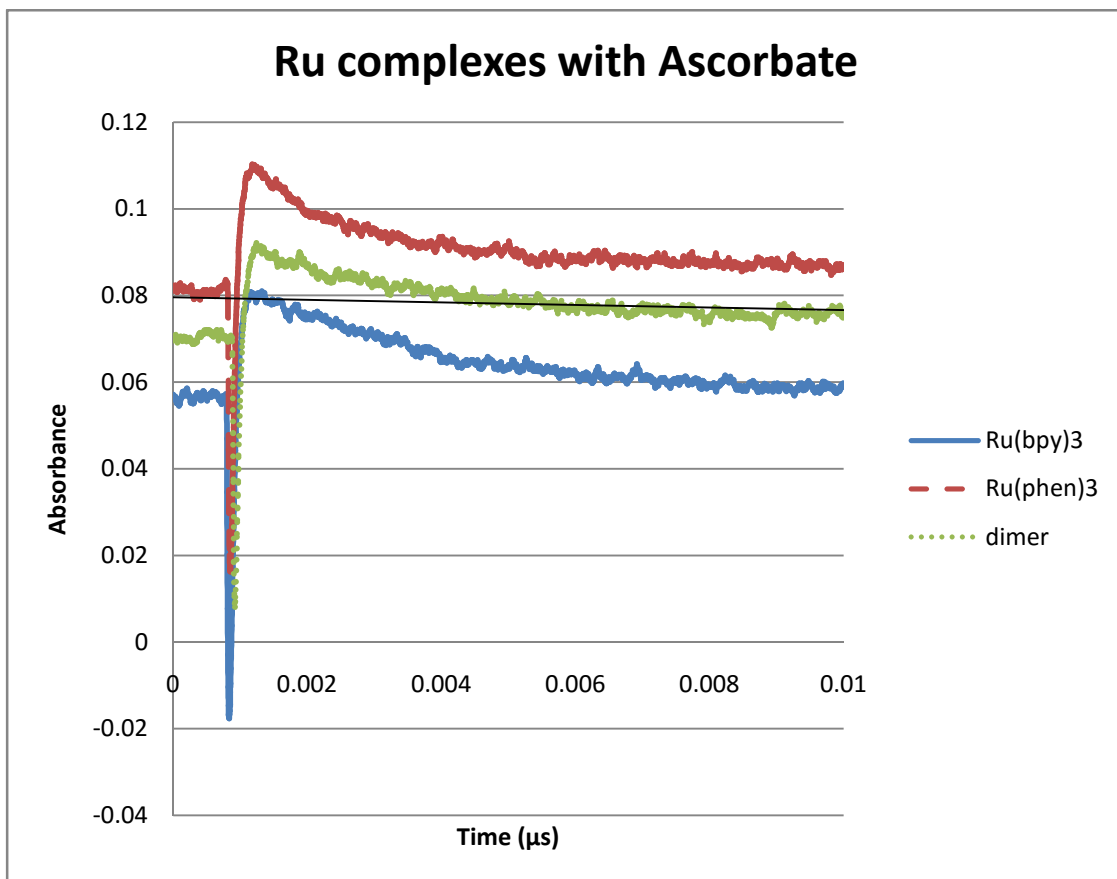


Figure 3.27 Representative absorption transient for ruthenium complexes with ascorbate



Chapter 4 Conclusions

4.1 Synthesis of the Bridging Ligand

The bridging ligand was prepared using a method recently optimized by Puckett. Additional optimization pointed out the critical nature of the oxygen sensitivity of the reaction and led to a reliable procedure with an average yield of approximately 40%. In addition work-up conditions were found that provided a product that was completely free from triphenylphosphine which had been a problem in past work. With a good source of product available several attempts were made to obtain crystals suitable for single crystal diffraction. Fortunately, the efforts proved successful and good quality crystals were sent for analysis.

4.2 Crystal structure of 5,5'-Bi-1,10-phenanthroline (diphen)

Bond lengths and bond angles for phen were chosen from a shallow-boat shape, all the six-membered rings are exactly planar. Therefore, phen exhibits good symmetry with the identical in above and bottom half.

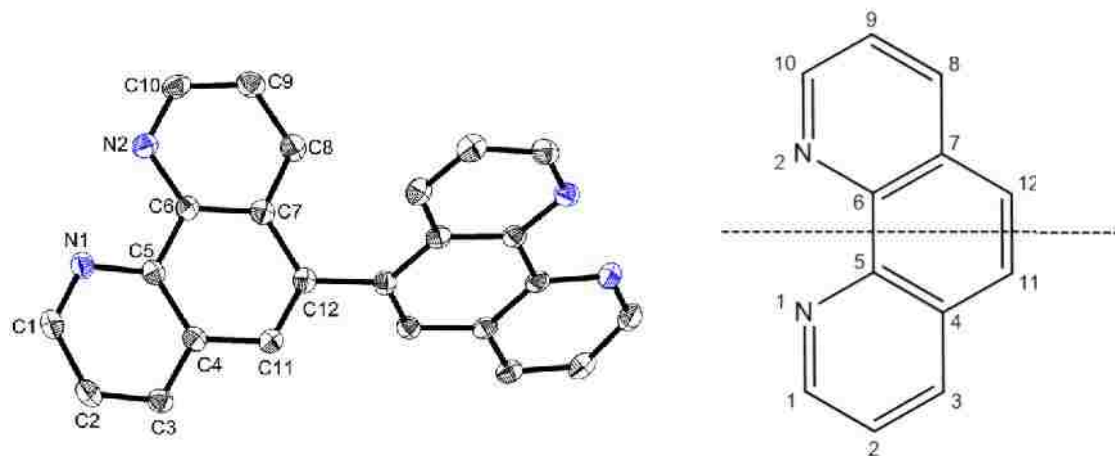
In the two columns of Table 4.1 the bond lengths and bond angles are compared with the phen literature values by Satoshi Nishigaki.⁴⁴ The agreement is fairly good, except for the bonds contain C12, whose lengths are somewhat larger than the phen. The bond length for C12-C12, which linked two phen subunits, is 1.494(3) Å, longer than the average bond lengths for N-C (1.326 Å) or C-C (1.372 Å)⁴⁵ in the literature for phen-coordinated metal complexes. This reflects that the bridging bond between two phen subunits is stretched and also caused the neighbor rings slightly stretched.

Table 4.1 Selected bond lengths (Å) and band angles (°) for diphen and phen

Bond lengths (Å)	diphen	phen
N1-C1	1.325(2)	1.317(7)
C1-C2	1.403(3)	1.402(11)
C3-C2	1.369(3)	1.352(11)
C4-C3	1.412(2)	1.410(9)
C4-C11	1.435(2)	1.427(10)
C12-C11	1.357(2)	1.330(12)
C12-C7	1.451(2)	1.426(12)
C7-C8	1.412(3)	1.414(11)
C8-C9	1.371(3)	1.348(13)
C9-C10	1.405(3)	1.394(12)
N2-C10	1.321(2)	1.312(8)
Band angles (°)		
N1-C1-C2	123.78(17)	123.8(6)
C3-C2-C1	119.39(17)	118.9(6)
C2-C3-C4	118.68(17)	119.0(7)
C3-C4-C11	122.04(16)	124.2(7)
C12-C11-C4	121.51(16)	122.5(8)
C11-C12-C7	119.79(16)	120.9(6)
C8-C7-C12	122.77(17)	124.5(6)
C9-C8-C7	120.02(18)	122.2(7)
C8-C9-C10	118.23(17)	116.9(7)
N2-C10-C9	123.86(17)	125.1(8)

Figure 4.1 Left: The model of the diphen single-crystal structure. Anions are omitted for clarity.

Atom color code: carbon, grey; nitrogen, blue. Right: The structure for phen



4.3 Synthesis of $[(\text{phen})_2\text{Ru-diphen-Ru}(\text{phen})_2](\text{PF}_6)_4$ complexes.

Synthesis of this dimer utilized a commonly used polymeric ruthenium carbonyl compound as the entry point. This entry point was chosen because the polymeric material readily reacts with one chelating ligand such as 1,10-phenanthroline under very mild conditions with no further reaction. Reaction with diphen yields a bridged dimer. Further reaction with additional ligands requires the remaining carbonyl groups be removed with trimethylamine-N-oxide. The product was fully characterized as described earlier. As expected the dimer shares many common features with $\text{Ru}(\text{phen})_3^{2+}$. For example the maximum absorbance in the visible region of the spectrum occurs at 446 nm with the dimer and at 448 nm with the monomer. Emission maxima are identical at room temperature.

4.4 Quenching of $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2]\text{Cl}_4$ by Peroxydisulfate

The photoredox reactions of $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2]\text{Cl}_4$ were studied through a series of quenching reactions. The analogous monomeric complexes $\text{Ru}(\text{bpy})_3^{2+}$ and $\text{Ru}(\text{phen})_3^{2+}$

were used as comparisons. In previous research, Fe^{3+} was used as an oxidative quencher in 0.5 M H_2SO_4 . Laser flash photolysis measurements under conditions where more than 50 % of $\text{Ru}(\text{bpy})_3^{2+}$ was quenched to yield $\text{Ru}(\text{bpy})_3^{3+}$ consistently indicated the yield of oxidized $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2]^{4+}$ was limited to less than 50 %. It appears at this time that oxidation of one half of the dimer prevents the quenching reaction by the other half.

Quenching of $\text{Ru}(\text{bpy})_3^{2+}$ or $\text{Ru}(\text{phen})_3^{2+}$ by peroxydisulfate ion is known to proceed with a quantum yield of 2. The standard reduction potential for the two electron reduction of peroxydisulfate anion to the sulfate anion:

$$E^0(\text{S}_2\text{O}_8^{2-}/2\text{SO}_4^{2-}) = 1.94 \text{ V}$$

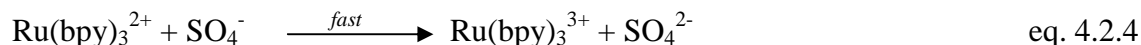
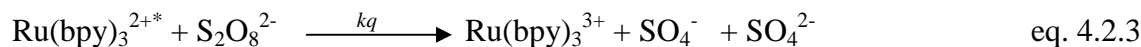
The standard reduction potential for the reduction of $\text{S}_2\text{O}_8^{2-}$ to SO_4^- occurs at a lower potential:

$$E^0(\text{S}_2\text{O}_8^{2-}/\text{SO}_4^-, \text{SO}_4^{2-}) = 1.45 \text{ V}$$

The sulfate radical anion generated from one-electron reduction of $\text{S}_2\text{O}_8^{2-}$ is a stronger oxidant with an oxidation potential of 2.44 V:

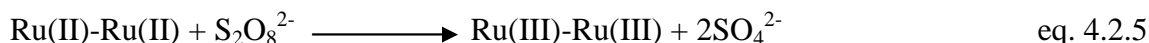
$$E^0(\text{SO}_4^-/\text{SO}_4^{2-}) = 2.44 \text{ V}$$

The reaction of excited state of $\text{Ru}(\text{bpy})_3^{2+}$ with $\text{S}_2\text{O}_8^{2-}$ to produce Ru(III) species has been investigated by Bolletta.⁴⁶ The following photoinduced oxidation scheme appears to describe the high yield



A strong oxidizing intermediate, SO_4^- , is produced during the reaction (eq 4.2.3) which generates a second Ru(III) (eq 4) through a thermal reaction. Previous reports show that the initial photoinduced electron transfer occurs in a collision between excited $\text{Ru}(\text{bpy})_3^{2+*}$ and $\text{S}_2\text{O}_8^{2-}$ in the $\text{Ru}(\text{bpy})_3^{2+}$ $\text{S}_2\text{O}_8^{2-}$ system⁴⁶⁻⁴⁷.

Unlike the mononuclear ruthenium (II) complex, $\text{Ru}(\text{bpy})_3^{2+}$, the symmetric dimer contains two reducing sites. These two sites are potentially capable of reducing $\text{S}_2\text{O}_8^{2-}$ through a single two-electron reaction, or sequence of two one-electron reactions. The reactions of interest in this system are shown below.



A Stern-Volmer investigation of the quenching reaction indicated that the dimer had a higher Stern-Volmer quenching constant, k_q , than $\text{Ru}(\text{phen})_3^{2+}$. One possible explanation is simply the difference in charge of the complexes. Since the quencher has a negative charge it is reasonable to assume that electrostatics will play a significant role in the quenching reaction.

The most interesting result that may have a bearing on the mechanism is the laser flash photolysis transients. Figure 3.18 shows a clear step function with the dimer. On the other hand the transient absorbance changes with $\text{Ru}(\text{bpy})_3^{2+}$ clearly show a biphasic reaction. The first is very fast and is probably the initial quenching reaction. The second slower phase can be interpreted as the diffusion of the sulfate radical ion to another reduced complex. Since the dimer has a second ruthenium in close proximity it is reasonable to assume that the diffusion to the adjacent ruthenium center will be very fast and appear as a single step function. Unfortunately, these observations were made during the data analysis and it was not possible to

examine the reactions on a shorter time scale. Figure 3.17, however does give a small indication that a similar slow phase is present in $\text{Ru}(\text{phen})_3^{2+}$.

4.5 Quenching of $[(\text{phen})_2\text{Ru}(\text{diphen})\text{Ru}(\text{phen})_2]\text{Cl}_4$ by Ascorbate

A Stern-Volmer investigation of the quenching reaction indicated that the dimer had a similar Stern-Volmer quenching constant k_q with $\text{Ru}(\text{phen})_3^{2+}$. In the laser flash photolysis transients, there is no big difference between the absorbance changes of the dimer and $\text{Ru}(\text{phen})_3^{2+}$. The possible explanation is ascorbate is different from $\text{S}_2\text{O}_8^{2-}$, one ascorbate molecular only can donate one electron to ruthenium complexes. Here so after the half of the dimer was reduced, the other half had no advantage to be reduced when compare with monomers.

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