SYNTHESIS, CHARACTERIZATION, NUCLEOLYTIC, ANTIBACTERIAL AND ANTIPROLIFERATIVE PROPERTIES OF VANADIUM, COPPER AND MANGANESE COMPLEXES

LIM ENG KHOON

Thesis submitted in fulfillment of the requirements

for the degree of

Doctor of Philosophy

July 2011

ACKNOWLEDGEMENTS

I would like to express profound gratitude to my supervisor, Professor Dr. Teoh Siang Guan, for his invaluable support, encouragement, enthusiasm, supervision and useful suggestions throughout this research work. His moral support and continuous guidance enabled me to complete my work successfully. I am also highly thankful to Associate Professor Dr. Ng Chew Hee, Faculty of Engineering and Science, Universiti Tunku Abdul Rahman, Professor Dr. Mohd Nazalan Mohd Najimudin, School of Biological Sciences, Universiti Sains Malaysia and Professor Dr. Tengku Sifzizul Tengku Muhammad, School of Biological Sciences, Universiti Sains Malaysia, for their valuable suggestions throughout this study.

I would also like to express my deep appreciation to Professor Dr. Fun Hoong Kun and Mr. Mohd Mustaqim Rosli, X-ray Crystallography Unit, School of Physics, Universiti Sains Malaysia, for their professional guidance, excellent help and valuable advice for running and analysis all my X-ray Crystallography samples.

Special thanks go to Miss Beh Hooi Kheng and Miss Seow Lay Jing, School of Pharmaceutical Sciences, Universiti Sains Malaysia; Mr. Cheah Yew Hoong, Bioassay Unit, Herbal Medicine Research Center, Institute Medical Centre; Miss Chew Guat Siew, Mr. Yam Hok Chai, Mr. Yuen Chee Wah and Mr. Emmanuel Jairaj Moses, School of Biological Sciences, Universiti Sains Malaysia, for their guidance and help in the biological tests.

I want to give my thanks to the staff of the School of Chemical Science, Universiti Sains Malaysia, especially to Mr. Aw Yeong Choek Hoe, Mr. Ong Chin Hwie, Mr. Ong Ching Hin, Mr. Razli Effendy bin Khalidee, Mr. Marimuthu Ayeroo, Pn. Hjh. Zali Zaiton Hj. Hussin, Pn. Yeoh Chooi Ling, Mr. Ahmad Azrulhisham Abdul Rahim, Mr. Mohd Kassim Abd Razak, Mr. Yee Chin Leng and Mr. Clement D' Silva, for their constant help and support throughout my study.

I am also sincerely grateful to USM Fellowship award from Institute of Postgraduate Studies (IPS), Universiti Sains Malaysia, for the financial support and RU Grant and Science Fund from Universiti Sains Malaysia, and FRGS Grant from Ministry of Higher Education (MOHE), Malaysia, for funding this research.

I also like to express my warmest gratitude to my friends, especially to Sharon Fatinathan, Yasodha Sivasothy, Ong Chin Hin, Oo Chuan Wei, Vijay, Yoga, Wendy, Mandy, Cynn Dee, William Yip, Karen Onn and many more, for their moral support and kindness.

Finally, I would like to thank my family members, especially my wife, Belle, for supporting and encouraging me to pursue this degree. Without my family encouragement, I would not have finished the degree.

To all of you, I say *Ribuan Terima Kasih*

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lane 9, 20 mM NaN_3; lane 10, 30 mM NaN_3; lane 11, 50 mM NaN_3; lane 12, 100 mM NaN_3.

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LIST OF ABBREVIATIONS AND SYMBOLS

Cu-2-F-6-FBZO	tetrakis(μ-2,6-diflourobenzoato-κ ^{2}*O,O'*)bis[aquacopper(II)]
Cu-2-BrBZO	tetrakis(μ-2-bromobenzoato-κ ^{2}*O,O'*)bis[aquacopper(II)]
Cu-2-Cl-4-NO_2BZO	tetrakis(μ-2-chloro-4-nitrobenzoato-κ ^{2}*O,O'*) bis[aquacopper(II)]
Cu-2-Cl-6-FBZO	tetrakis(μ-2-chloro-6-fluorobenzoato-κ^2*O,O'*) bis[aquacopper(II)]
Cu-2-ClBZO	tetrakis(μ-2-chloro-benzoato-κ^2*O,O'*)bis[aquacopper(II)]
Cu-4-Cl-2-NO_2BZO	tetrakis(μ-4-chloro-2-nitrobenzoato-κ ^{2}*O,O'*) bis[aquacopper(II)]
CuP-5-Cl-2-NO_2BZO	catena-poly[[bis(5-chloro-2-nitrobenzoato)copper(II)] bis(μ-5-chloro-2- nitrobenzoato)]
Hela	Human cervical cancer cell line
HepG2	Human liver cancer cell line
IC_{50}	A measure of the concentration of particular drug that inhibit the biological process of cells by half
bp	base pair
LC_{50}	A measure of the concentration of particular drug that kill 50 % of the cells
LMCT	Ligand to Metal Charge Transfer
MCF-7	Human breast cancer cell line
MDA-MB-231	Human invasive breast cancer cell line
MIC	minimal inhibitory concentration
Mn-4-NO_2BZO	tetraaquabis(4-nitrobenzoato)manganese(II) hydrate

LIST OF ABBREVIATIONS AND SYMBOLS

MnP-4-ClBZO	catena-poly[[diaquabis(4-chlorobenzoato) manganese(II)]bis(μ-4-chlorobenzoato)]
MnP-4-Cl-2-NO_2BZO	catena-poly[[diaquabis(4-chloro-2-nitrobenzoato) manganese(II)]bis(μ-4-chloro-2-nitrobenzoato)]
MnP-4-FBZO	catena-poly[[diaquabis(4-flourobenzoato) manganese(II)]bis(μ-4-flourobenzoato)]
MnP-4-NH_2BZO	catena-poly[[bis(4-aminobenzoato)manganese(II)] tri(μ-4-aminobenzoato)]
MnPGLY	catena-poly[[aqua(diglycolato)manganese(II) hydrate)] bis(μ-diglycolato)]
MnPyr	diaquabis(pyridine-2-carboxylato)manganese(II) hydrate
MTT	(3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide)
NMR	Nuclear Magnetic Resonance
ROS	Reactive oxygen species
SRB	Sulforhodamine B
TGI	Cytostatic activity = A measure of the concentration of particular drug that inhibit total growth of cell lines.
UV-Vis	Ultraviolet-Visible
VO_2GLY	sodium (diglycolato)dioxovanadium(V) dihydrate
VO_2HPYDC	Sodium (4-hydroxypyridine-2,6-dicarboxylato) dioxovanadium(V) dihydrate
VO_2PP	(pyridine-2-carboxylato)(pyridinium-2-carboxylato) dioxovanadium(V) monohydrate

LIST OF ABBREVIATIONS AND SYMBOLS

VODMPhen	aqua(2,9-dimethyl-1,10-phenanthroline) sulfatooxovanadium(IV)
VODMPPhen	aqua(2,9-dimethyl-4,7-diphenyl-1,10-phenanthroline) sulfatooxovanadium(IV) hydrate
VODPPhen	aqua(4,7-diphenyl-1,10-phenanthroline) sulfatooxovanadium(IV) hydrate
VOMAL	disodium diaquabis(malonato)oxovanadium(IV)
VOPhen	aqua(1,10-phenanthroline)sulfatooxovanadium(IV)
VOPYDC	diaqua(pyridine-2,6-carboxylato)oxovanadium(IV)ethanol solvate
ΔEp	| Epa – Epc |
$\mathbf{V^{V}}$	Vanadium V
$\mathbf{V^{IV}}$	Vanadium IV
ε	Molar absorptivity

SINTESIS, PENCIRIAN, SIFAT-SIFAT NUCLEOLITIK, ANTIBAKTERIA DAN ANTIPOLIFERATIF BAGI KOMPLEKS VANADIUM, KUPRUM DAN MANGAN

ABSTRAK

Siri kompleks bagi vanadium karboksilato, kuprum karboksilato, mangan karboksilato dan terbitan vanadium fenantrolina telah disintesis dan dicirikan. Maklumat penuh tentang kompleks ditunjukkan dalam Jadual 2.2 (muka surat 37). Kompleks ini telah dicirikan dengan menggunakan kristalografi X-ray, analisis unsur, FT-IR, spektroskopi UV–Vis dan siklik voltammetrik. Dalam kajian elektrokimia, kompleks vanadium karboksilato telah menunjukkan aktif redok dengan mempamerkan pasangan redok kuasi-berbalik yang selaras dengan V^{V}/V^{IV} proses redok, manakala kompleks kuprum karboksilato telah menunjukkan aktif redok dengan mempamerkan dua pasangan redok kuasi-berbalik yang selaras dengan $Cu^{II}Cu^{II}/Cu^{II}Cu^{I}$ dan $Cu^{II}Cu^{I}/Cu^{I}Cu^{I}$ proses redok dan mangan karboksilato telah menunjukkan aktif redok dengan mempamerkan pasangan redok kuasi-berbalik yang selaras dengan Mn^{II}/Mn^{III} proses redok. Dalam kajian nukleolitik, semua kompleks didapati boleh mengakibatkan pengoksidaan belahan DNA tetapi dengan cara yang berbeza. Kompleks kuprum karboksilato dan mangan karboksilato boleh mengakibatkan belahan DNA dalam kehadiran H_2O_2. Sementara itu,

untuk kompleks vanadium karboksilato, kompleks vanadium(V) karboksilato memerlukan H_2O_2 untuk mengakibatkan belahan DNA manakala kompleks vanadium(IV) karboksilato tidak memerlukan H_2O_2 untuk mengakibatkan belahan DNA. Bagi kompleks terbitan vanadium fenantrolina pula, mereka boleh mengakibatkan belahan DNA dalam kehadiran H_2O_2 dan dalam keadaan tanpa H_2O_2. Walaubagaimanapun, aktiviti belahan DNA meningkat secara mendadak dalam kehadiran H_2O_2. Agen perencet reaktif spesis oksigen (ROS) juga telah digunakan untuk menentukan spesis reaktif yang bertanggunjawab dalam belahan DNA. Hidroksil radikal dan oksigen tunggal adalah ROS yang bertanggunjawab dalam belahan DNA. Dalam penyaringan antibakteria, semua kompleks kecuali kompleks mangan karboksilato mempamerkan aktiviti antibakteria terhadap spesis bakteria Gram positif atau Gram negatif. Kompleks kuprum karboksilato menunjukkan aktiviti antibakteria yang sangat selektif di mana mereka hanya menunjukkan aktiviti antibakteria terhadap *Enterobacter aerogenes* berGram negatif bakteria. Aktiviti antibakteria kompleks terbitan vanadium fenantrolina telah menunjukan bahawa kumpulan metil yang terikat pada kedudukan 2 dan 9 pada gelang fenantrolina dapat meningkatkan aktiviti antibakteria. Dalam penyaringan antipoliferatif, secara umum kompleks vanadium karboksilato mempamerkan aktiviti antipoliferatif yang lebih tinggi berbanding dengan kompleks kuprum karboksilato dan mangan karboksilato. Kompleks kuprum karboksilato dan mangan karboksilato menunjukkan selektiviti sitotoksik terhadap garisan sel kanser HepG2 apabila dibandingkan dengan garisan sel kanser MCF-7 dan Hela. Eksperimen nukleolitik telah menyarankan bahawa belahan atau fragmentasi DNA oleh ROS yang dihasilkan oleh kompleks berkemungkinan besar bertangungjawab terhadap aktiviti antipoliferatif yang dipamerkan oleh kompleks.

Synthesis, Characterization, Nucleolytic, Antibacterial and Antiproliferative Properties of Vanadium, Copper and Manganese Complexes

ABSTRACT

Series of vanadium carboxylato complexes, copper carboxylato complexes, manganese carboxylato complexes and vanadium phenanthroline derivative complexes have been synthesized and characterized. The details of all the complexes are tabulated in Table 2.2 (page 37). The complexes have been characterized by X-ray crystallography, elemental analysis, FT-IR, UV–Vis spectroscopy and cyclic voltammetry. In electrochemistry studies, vanadium carboxylato complexes show redox active by displaying a quasi-reversible redox couple corresponding to V^{V}/V^{IV} redox process while copper carboxylato complexes show redox active by displaying two quasi-reversible redox couples corresponding to $Cu^{II}Cu^{II}/Cu^{II}Cu^{I}$ and $Cu^{II}Cu^{I}/Cu^{I}Cu^{I}$ redox processes and manganese carboxylato complexes show redox active by displaying a quasi-reversible redox couple corresponding to Mn^{II}/Mn^{III} redox process. In nucleolytic studies, all the complexes can induce oxidative DNA cleavage but in different manner. The copper carboxylato complexes and manganese carboxylato complexes can induce DNA cleavage in the presence of H_2O_2. Meanwhile for vanadium carboxylato complexes, vanadium(V) carboxylato complexes require H_2O_2 to induce DNA cleavage while vanadium(IV) carboxylato complexes do not require H_2O_2 to induce DNA cleavage. As

for vanadium phenanthroline derivative complexes, they can induce DNA cleavage in the presence and in the absence of H_2O_2. However, the DNA cleavage activity of the vanadium phenanthroline derivative complexes is greatly enhanced in the presence of H_2O_2. Reactive oxygen species (ROS) scavengers have also been used to ascertain the reactive species responsible for DNA cleavage. The hydroxyl radical and singlet oxygen species have been found to be the ROS that are responsible in the DNA cleavage reaction. In the antibacterial screening, all the complexes except manganese carboxylato complexes exhibit antibacterial activity against certain Gram negative or Gram positive bacteria species. Copper carboxylato complexes show a very selective antibacterial activity whereby they only exhibit antibacterial activity against Gram negative bacteria Enterobacter aerogenes. The antibacterial activity of vanadium phenanthroline derivative complexes reveals that methyl groups attached at the position 2 and 9 in phenanthroline ring may increase the complexes antibacterial activity. In antiproliferative screening, vanadium carboxylato complexes in general exhibit higher antiproliferative activity when compared to copper carboxylato complexes and manganese carboxylato complexes. Copper carboxylato complexes and manganese carboxylato complexes exhibit cytotoxic selectivity against HepG2 cancer cell line when compared to MCF-7 and Hela cancer cell lines. The nucleolytic experiments suggest that the cleavage or fragmentation of DNA by ROS generated by the complexes maybe responsible for the antiproliferative activity exhibited by the complexes.

Keywords: Vanadium Complexes; Copper Complexes; Manganese Complexes; DNA Cleavage; Antibacterial; Antiproliferative

CHAPTER 1

INTRODUCTION

1.1 Biological roles and medicinal applications of metal complexes and metal ions

Studies on biological activities of metal complexes have been one of our long-term interests. Metal complexes are well known to exhibit antibacterial, antiproliferative, antiapoptotic, anti-inflammatory and insulin mimetic properties [1-28]. Several of the metal complexes have entered clinical trials and few have been registered for clinical use [29-31]. Platinum based complexes such as cisplatin, carboplatin and oxaliplatin are widely used as chemotherapeutic agents against ovarian, lung, head, neck and colorectal cancers, and have greatly improving the survival rates of patients worldwide. Schematic structures of cisplatin, carboplatin and oxaliplatin are depicted in Figure 1.1. These platinum complexes react *in vivo*, crosslink the DNA in several different ways and subsequently interfering the cell division by mitosis. The damaged DNA elicits DNA repair mechanisms, which in turn activate apoptosis when repair proves impossible. Meanwhile, Aurum(I) thiolate drugs such as aurothiomalate (Myocrisin^{R}), aurothioglucose (Solganol^{R}), aurothiopropanol sulfonate (Allochrysin^{R}), and the oral drug auranofin (Ridaura^{R}), are widely used for the treatment of difficult cases of rheumatoid arthritis. Bismuth(III) compounds such as bismuth subcitrate and subsalicylate are widely used for the treatment of diarrhoea, dyspepsia and gastric and duodenal ulcers. Bismuth(III) compounds are found to be antibacterial active against

bacteria Helicobacter pylori which is associated with the mucus layer of ulcers. Sodium nitroprusside ($Na_2[Fe^{II}(CN)_5(NO)]\cdot 2H_2O$ or ($Nipride^R$) is used to treat hypotensive while Cu-salicylate ($Alcusal^R$) is used to treat inflammatory. The success of metal complexes in medicinal applications has aroused great interest in the development of new metal complexes to diagnose and treat diseases including cancers, bacteria and virus infection related diseases, inflammatory and diabetes.

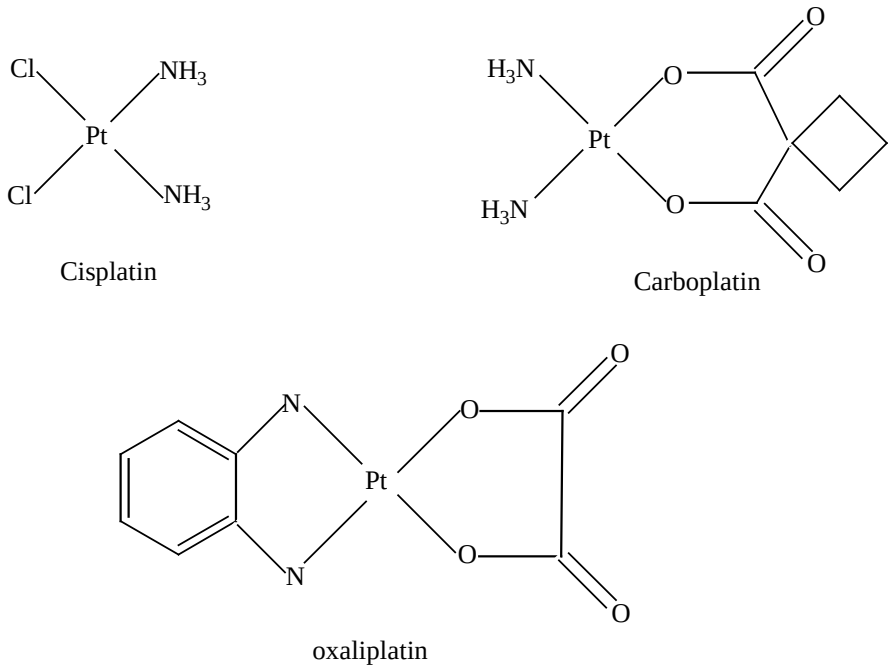


Figure 1.1: Schematic structures of cisplatin, carboplatin and oxaliplatin

Apart from metal complexes, metal ion or inorganic elements play essential roles in biological and biomedical processes in human health and disease as metalloenzymes and metalloproteins [29]. As metalloproteins, metal ions perform as catalyst or stabilizer to stabilize the protein tertiary or quaternary structure. In addition, many proteins need to bind one or more metal ions to perform their functions. Complex zinc ion is one of the

most important components of metalloproteins in the human body, which functions as DNA transcription and regulation as well as oxidation and hydrolysis, cleavage of peptide bonds as well as formation of phosphodiester bonds. Meanwhile, copper ion is presence in some of the most important metalloenzymes in the human body and functions as superoxide dismutase to neutralize free radical generated from various human biological systems. Metal ions are also very important for the structure and function (in the case of RNA) of nucleic acids. Besides, many organic compounds used in medicine do not have a purely organic mode of action; some are activated or biotransformed by metal ions including metalloenzymes, others have a direct or indirect effect on metal ion metabolism. Some of the metal ions have also been registered for clinical used as therapy and diagnosis agents, as listed in Table 1.1.

Table 1.1: Some of metal ions in clinical use

Compounds	Function/Treat
Li_2CO_3	Prophylaxis for bipolar disorders
$CaCO_3$, $Mg(OH)_2$	Antacid
$La_2^{III}(CO_3)_3$	Chronic renal failure
$MgSO_4$	Hypomagnesemia
Potassium citrate	Kidney stones
Magnesium citrate	Saline laxative

1.2 Antibacterial and antiproliferative activities of transition metal complexes

Transition metal complexes such as vanadium, copper and manganese complexes are known to exhibit excellent antibacterial and antiproliferative activities. The summary of antibacterial and antiproliferative activities of selected few vanadium, copper and manganese complexes that have already described in the literatures is tabulated in Table 1.2. Referring to Table 1.2, it can be seen that transition metal complexes are rich in antibacterial and antiproliferative activities, being active against a wide spectrum of bacteria species or cancer cells. Transition metal complexes may induce cell death through disruption of the cell cycle or by DNA strand scission [32]. There are evidences to indicate that metal complexes can induce DNA strand scission not directly reacting with DNA components but acting mainly through the production of highly reactive oxygen species, especially hydroxyl radicals generated in cells. These reactive oxygen species actually cause the DNA strand scission. Transition metal complexes through Fenton-like reactions and/or during the intracellular reduction can generate reactive oxygen species. Besides, some transition metal complexes, which are photoactivatable, can induce DNA cleavage upon UV irradiation and singlet oxygen is the common reactive oxygen species that is generated in this process. Rich diversity of antibacterial and antiproliferative activities by transition metal complexes provides exciting prospects for the design of novel therapeutic agents with unique mechanisms of action to act against certain bacteria or cancers, as different metal complexes can produce different therapeutic effect. Therefore, detailed investigations could be helpful in designing more potent antibacterial and anticancer agents for the therapeutic use.

Table 1.2: The antibacterial and antiproliferative activities of some vanadium, copper and manganese complexes

Complexes	Antibacterial/ Antiproliferative activity	Bacteria species/Cancer cells	Ref:
$[Cu^{II}$(4-(2-pyridylmethyl)-1,7-dimethyl-1,4,7-triazonane-2,6-dione)$(CH_3CN)_2](ClO_4)_2$	Antibacterial	Escherichia coli (T7), Staphylococus aureus, Pseudomonas aeruginosa	13
Cu^{II}(2-furancarbaldehyde thiosemicarbazone) $0.5H_2O$	Antibacterial	Bacillus subtilis, Staphylococus aureus	18
$[Cu_2{}^{II}$(N,N'-bis(3-aminopropyl)oxamide)(2,2'-bipyridine)(2,4,6-trinitrophenol)$(H_2O)]$(2,4,6-trinitrophenol)	Antibacterial	Escherichia coli, Bacillus subtilis, Staphylococus aureus	16
$Cu^{II}{}_2$(N,N'-bis(N-hydroxyethylaminoethyl) oxamide)$(2,4,6\text{-trinitrophenol})_2$	Antiproliferative	SMMC-7721 human hepatocellular carcinoma cells, A549 human lung adenocarcinoma cells	8
Cu^{II} (ethyl 2-bis(2-pyridylmethyl)aminopropionate)Cl_2	Antiproliferative	Eca-109 human esophageal cancer cells, A549 human lung adenocarcinoma cells	10
Cu^{II} (norfloxacinato)(2,2'-bipyridine)Cl_2	Antiproliferative	HL-60 and K562 human leukemia cells	19

Table 1.2: Continued

Complexes	Antibacterial/ Antiproliferative activity	Bacteria species/Cancer cells	Ref:
Mn^{II}(tetraphenyl porphyrin), (ebselen–porphyrin conjugate)	Antibacterial	Staphylococus aureus	15
Mn^{II}(tetraamide macrocyclic)NO_3	Antibacterial	Pseudomonas cepacicola, Klebsella aerogenous	25
Mn^{II}(6,7-dicycanodipyrido[2,2-*d*:29,39-*f*]quinoxaline) $(NO_3)(H_2O)]NO_3.CH_3OH$	Antiproliferative	BGC-823 human stomach cancer cells, HL-60 human leukemia cells	27
$V^V{}_4O_{10}(\mu\text{-}O)_2[VO(H\text{-}ciprofloxacin)_2)]_2.13H_2O$	Antibacterial	Staphylococus aureus, Escherichia coli, Pseudomonas aeruginosa	21
V^V (2-methyl-3H-5-hydroxy-6-carboxy-4-pyrimidinone ethyl ester)	Antiproliferative	Hela human cervical cancer cells	6
V^VO_2(salicylaldehydesemicarbazone)	Antiproliferative	MC3T3-E1 osteoblastic mouse calvaria-derived cells, UMR106 rat osteosarcoma-derived cells	12
$V^{IV}O$(3-amino-6(7)-chloroquinoxaline-2-carbonitrile N^1, N^4-dioxide)$_2$	Antiproliferative	V79 chinese hamster lung fibroblasts cells	24

1.3 Background of nucleolytic activity of metal complexes

Recently, research on nucleolytic activity of metal complexes has blossomed leading to the discovery of the capacity of metal complexes to interact with DNA and further to induce DNA cleavage in the presence of co-factor. Transition metal complexes such as ruthenium, copper, cobalt, manganese and vanadium complexes have been reported to promote DNA cleavage in the presence of co-factor [33-55]. The DNA cleavage by metal complexes can occur via oxidative, photolytic and hydrolytic cleavage.

Double helical DNA consists of two complementary, antiparallel polydeoxyribonucleotide strands associated by specific hydrogen bonding interactions between nucleotide bases, Figure 1.2. The backbone of the DNA strand is made from alternating phosphate and sugar residues. The sugar in DNA is 2-deoxyribose, which is a pentose (five-carbon) sugar. The sugars are joined together by phosphate groups that form phosphodiester bonds between the third and fifth carbon atoms of adjacent sugar rings. The sugar phosphate backbone of paired strands defines the helical grooves, within which the edges of the heterocyclic bases are exposed. The biologically relevant B-form structure of the DNA double helix is characterized by a shallow, wide major groove and a deep, narrow minor groove. The major and minor grooves provide a lot of hydrogen binding sites. The DNA double helix is stabilized by hydrogen bonds between the nucleotide bases attached to the two strands. The four bases found in DNA are adenine, cytosine, guanine and thymine, Figure 1.3. These four bases are attached to the sugar/phosphate to form the complete nucleotide.

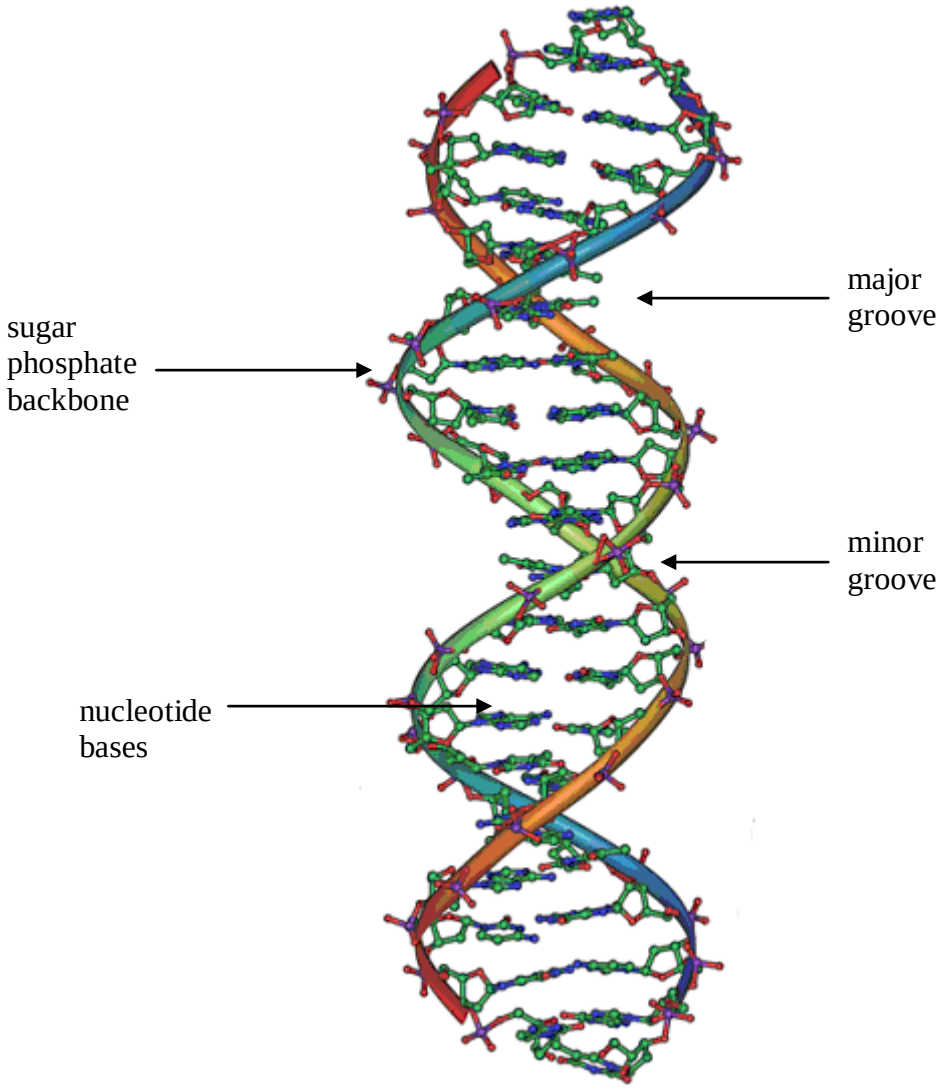


Figure 1.2: The structure of part of a DNA double helix

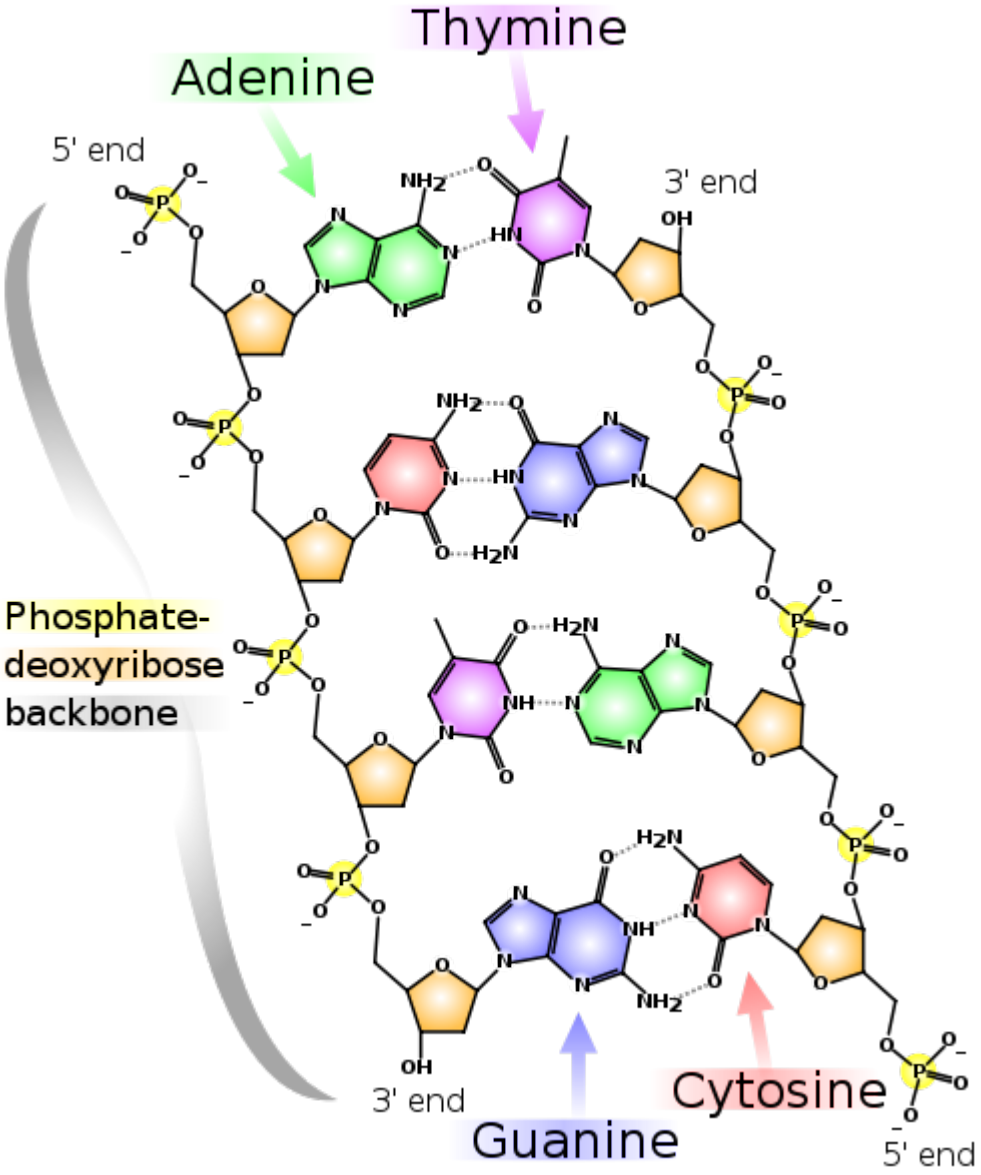


Figure 1.3: The chemical structure of DNA. Hydrogen bonds are shown as dotted lines

The DNA cleavage by metal complexes can be monitored by agarose gel electrophoresis. The DNA strain that was used in DNA cleavage studies is pBR322 DNA. The pBR322 DNA is a double helix DNA and it exists in supercoil form. In general, if scission or cleavage occurs on one strand of the supercoil DNA, the supercoil (Form I) will relax and convert to nicked form (Form II) while if scission occurs on both strands, a linear form (Form III) will be generated (Figure 1.4). These three forms of DNA will migrate in different rate in gel electrophoresis (Figure 1.5) with supercoil form migrates the fastest while nicked form migrates the slowest and linear form migrates in between supercoil and nicked forms. DNA cleavage by metal complexes is varied among the complexes, with some metal complexes can induce both single and double strand scissions while some metal complexes can only induce single scission.

In oxidative and photolytic DNA cleavage, metal complexes cannot induce DNA cleavage directly but indirectly through generating reactive oxygen species (ROS) such as hydroxyl radical and singlet oxygen. These ROS are actually responsible in DNA cleavage reaction. In order to study the DNA cleavage mechanism by metal complexes, various inhibiting agents have been used such as DMSO, t-butanol, mannitol and sodium azide. DMSO, t-butanol and mannitol are used as hydroxyl radical inhibitors while sodium azide is used as singlet oxygen inhibitor. Meanwhile in hydrolytic DNA cleavage, metal complexes can induce DNA cleavage directly by cleaving the P–O bonds in the phosphodiester of DNA.

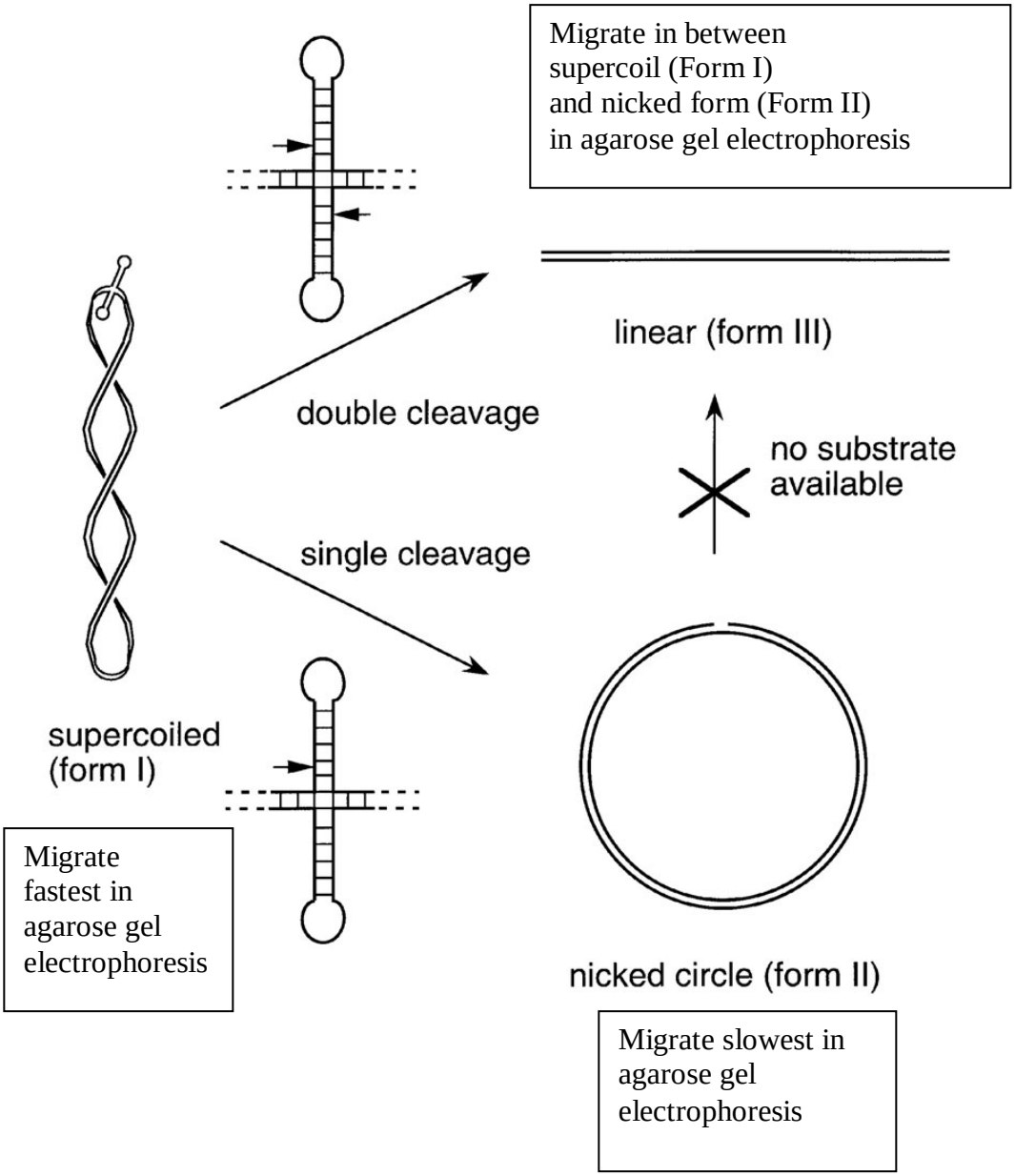


Figure 1.4: Supercoiled, nicked and linear DNA

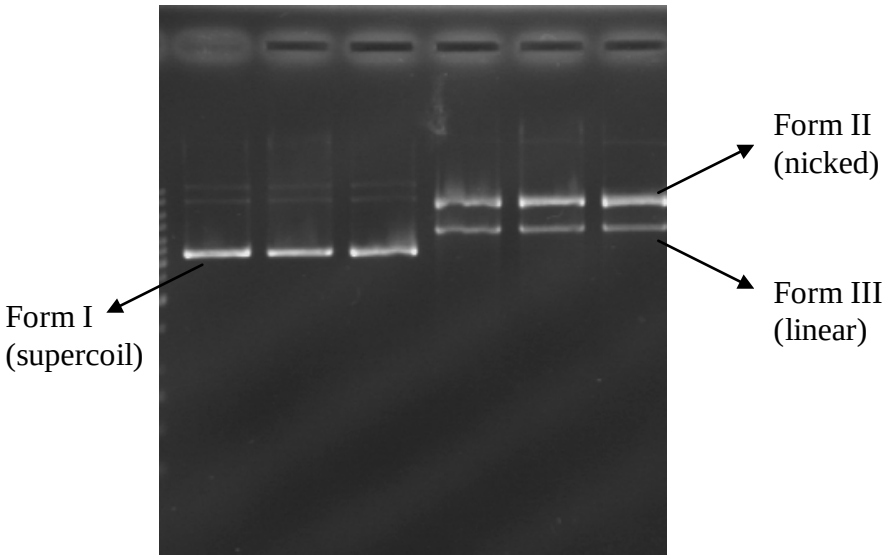
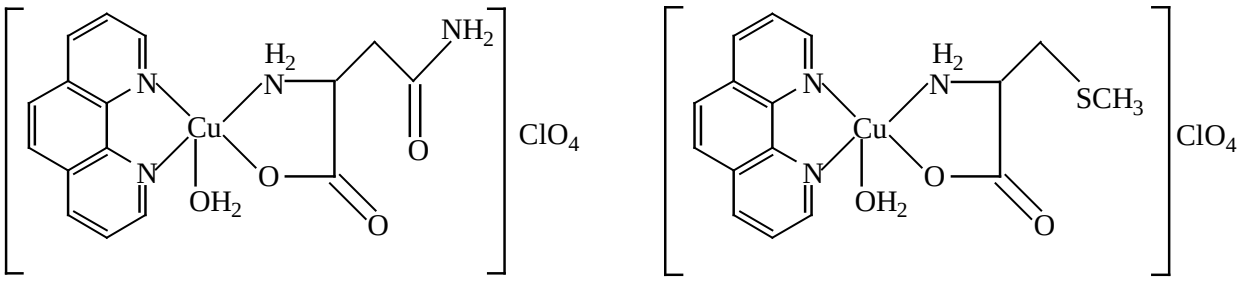


Figure 1.5: Supercoiled, nicked and linear DNA bands in gel electrophoresis diagram

1.3.1 Oxidative DNA cleavage by metal complexes in the presence of 3-mercaptopropionic acid (MPA)

Complexes [Cu^{II}(ternary-L-glutamine)(1,10-phenanthroline)(H_2O)](ClO_4) and [Cu^{II}(ternary-S-methyl-L-cysteine)(1,10-phenanthroline)(H_2O)](ClO_4) (Figure 1.6) can exhibit oxidative DNA cleavage in the presence of 3-mercaptopropionic acid (MPA) [33, 34]. MPA plays as reduction agent in the DNA cleavage reaction. Both of the complexes can only induce single DNA scission by converting supercoil DNA to nicked form. The mechanistic aspects of the DNA cleavage reactions have been investigated with various inhibiting agents and the results show that hydroxyl radical scavenger DMSO can inhibit the DNA cleavage induced by both of the complexes. This indicates the involvement of

hydroxyl radical in the cleavage reaction. The proposed DNA cleavage mechanism of metal complex in the presence of MPA is illustrated in Figure 1.7.



[a] [b]

Figure 1.6: The schematic structures;
a) [Cu^{II}(ternary-L-glutamine)(1,10-phenanthroline)(H_2O)](ClO_4)
b) [Cu^{II}(ternary-S-methyl-L-cysteine)(1,10-phenanthroline)(H_2O)](ClO_4)

$$[\text{M complex}]^{n} + \text{H-MPA} \longrightarrow [\text{M complex}]^{n-1} + \bullet\text{MPA} + H^{+}$$

$$[\text{M complex}]^{n-1} + O_2 \longrightarrow [\text{M complex}]^{n} + \bullet O_2^{-}$$

$$2 \bullet O_2^{-} + 2H^{+} \longrightarrow H_2O_2 + O_2$$

$$[\text{M complex}]^{n-1} + H_2O_2 \longrightarrow [\text{M complex}]^{n} + OH^{-} + \bullet OH$$

$$[\text{Supercoil DNA}] + \bullet OH \longrightarrow \text{DNA cleavage}$$

Figure 1.7: The proposed DNA cleavage mechanism of metal complex in the presence of 3-mercaptopropionic acid (MPA)

1.3.2 Oxidative DNA cleavage by metal complexes in the presence of ascorbic acid

Complexes [Ru^{II}(imidazo[4,5-f][1,10]phenanthroline)$(NH_3)_4$]$(PF_6)_2$ and [Cu^{II}(L-threonine)(1,10-phenanthroline)(H_2O)](ClO_4) (Figure 1.8) can induce oxidative DNA cleavage in the presence of ascorbic acid [35, 36]. Similar to MPA, ascorbic acid also acts as the reduction agent in the DNA cleavage reaction. Complex [Ru^{II}(imidazo[4,5-f][1,10]phenanthroline)$(NH_3)_4$]$(PF_6)_2$ can only induce single DNA scission by converting supercoil DNA to nicked form while complex [Cu^{II}(L-threonine)(1,10-phenanthroline)(H_2O)](ClO_4) can induce both single and double DNA scissions by converting supercoil DNA to nicked and linear forms. In comparison, complex [Cu^{II}(L-threonine)(1,10-phenanthroline)(H_2O)](ClO_4) appears to be a better DNA cleaver when compared to complex [Ru^{II}(imidazo[4,5-f][1,10]phenanthroline)$(NH_3)_4$]$(PF_6)_2$ in the presence of ascorbic acid. In mechanistic studies, it is evident that the hydroxyl radical scavenger DMSO diminish significantly the nuclease activity of complex [Cu^{II}(L-threonine)(1,10-phenanthroline)(H_2O)](ClO_4)], which is indicative of the involvement of the hydroxyl radical in the cleavage process. The proposed DNA cleavage mechanism of metal complex in the presence of ascorbic acid is illustrated in Figure 1.9.

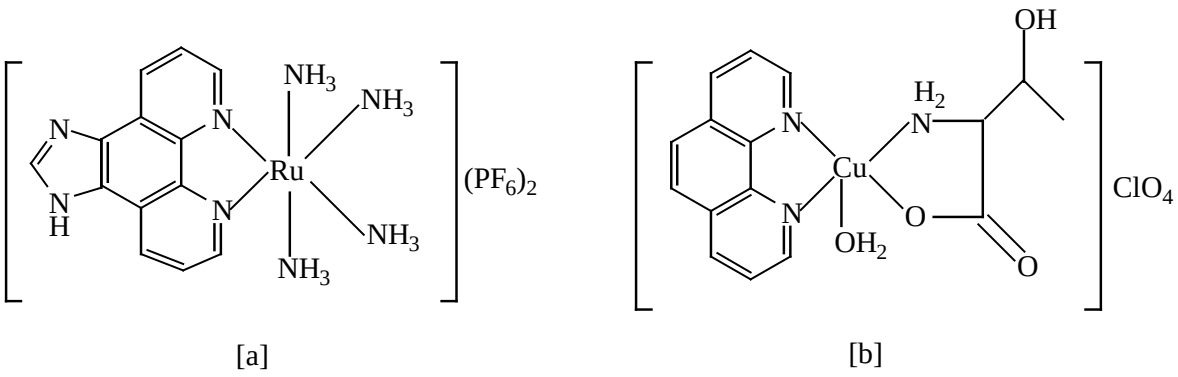


Figure 1.8: The schematic structures;
a) $[Ru^{II}(imidazo[4,5\text{-}f][1,10]phenanthroline)(NH_3)_4](PF_6)_2$
b) $[Cu^{II}(L\text{-}threonine)(1,10\text{-}phenanthroline)(H_2O)](ClO_4)$

$$[M\ complex]^{n} + H\text{-}Asc^{-} \longrightarrow [M\ complex]^{n-1} + \bullet Asc^{-} + H^{+}$$

$$[M\ complex]^{n-1} + O_2 \longrightarrow [M\ complex]^{n} + \bullet O_2^{-}$$

$$2 \bullet O_2^{-} + 2H^{+} \longrightarrow H_2O_2 + O_2$$

$$[M\ complex]^{n-1} + H_2O_2 \longrightarrow [M\ complex]^{n} + OH^{-} + \bullet OH$$

$$[Supercoil\ DNA] + \bullet OH \longrightarrow DNA\ cleavage$$

Figure 1.9: The proposed DNA cleavage mechanism of metal complex in the presence of ascorbic acid

1.3.3 Oxidative DNA cleavage by metal complexes in the presence of H_2O_2

Complexes $[Co^{II}(imidazole\text{-}terpyridine)_2](ClO_4)_2$ and $[Cu^{II}(imidazole\ terpyridine)_2](ClO_4)_2$ (Figure 1.10) can promote oxidative DNA cleavage in the presence of H_2O_2 [37, 38]. In contrast to MPA and ascorbic acid, H_2O_2 acts as oxidation agent in the DNA cleavage reaction. Complex $[Co^{II}(imidazole\text{-}terpyridine)_2](ClO_4)_2$ can only induce single DNA scission by converting supercoil DNA to nicked form while complex $[Cu^{II}(imidazole\ terpyridine)_2](ClO_4)_2$ can induce both single and double DNA scissions by converting supercoil DNA to nicked and linear forms. This indicates that the DNA cleavage efficiency of complex $[Cu^{II}(imidazole\ terpyridine)_2](ClO_4)_2$ is higher than the DNA cleavage efficiency of complex $[Co^{II}(imidazole\text{-}terpyridine)_2](ClO_4)_2$ in the presence of H_2O_2. From the mechanistic studies, it is shown that the hydroxyl radical scavenger DMSO can reduce significantly the nuclease activity of complex $[Co^{II}(imidazole\text{-}terpyridine)_2](ClO_4)_2$ while the hydroxyl radical scavenger ethanol can reduce significantly the nuclease activity of complex $[Cu^{II}(imidazole\ terpyridine)_2](ClO_4)_2$. This results reflect that the participation of hydroxyl radical in the cleavage process. The proposed DNA cleavage mechanism of metal complex in the presence of H_2O_2 is illustrated in Figure 1.11.

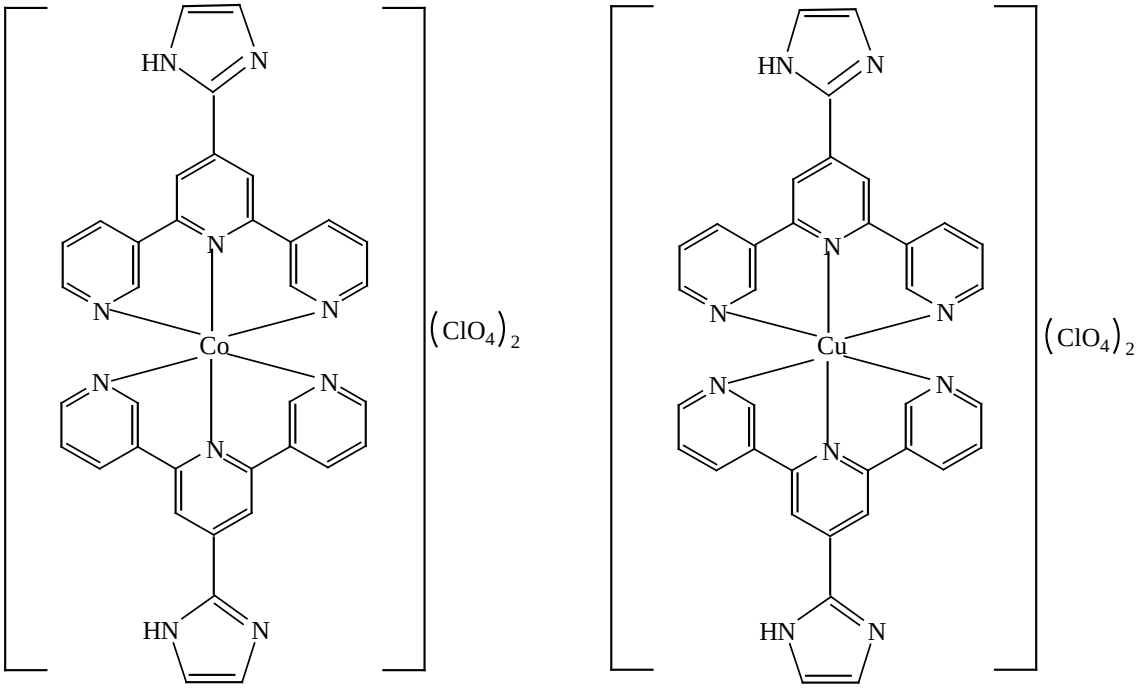


Figure 1.10: The schematic structures;
a) $[Co^{II}(imidazole\text{-}terpyridine)_2](ClO_4)_2$
b) $[Cu^{II}(imidazole\ terpyridine)_2](ClO_4)_2$

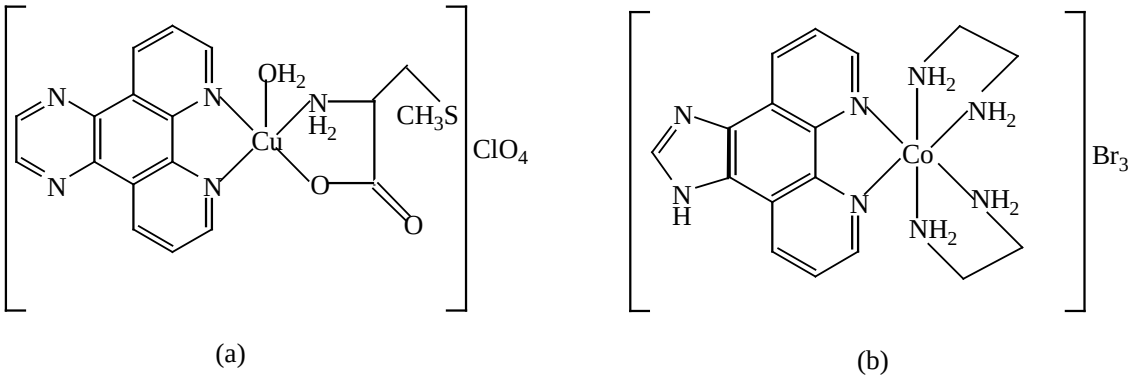
$$[M\ complex]^n + H_2O_2 \longrightarrow [M\ complex]^{n-1} + \bullet O_2^- + 2H^+$$
$$[M\ complex]^{n-1} + O_2 \longrightarrow [M\ complex]^n + \bullet O_2^-$$
$$[M\ complex]^n + \bullet O_2^- \longrightarrow [M\ complex]^{n-1} + O_2$$
$$[M\ complex]^{n-1} + H_2O_2 \longrightarrow [M\ complex]^n + OH^- + \bullet OH$$
$$[Supercoil\ DNA] + \bullet OH \longrightarrow DNA\ cleavage$$

Figure 1.11: The proposed DNA cleavage mechanism of metal complex in the presence of H_2O_2

1.3.4 Photolytic DNA cleavage by metal complexes

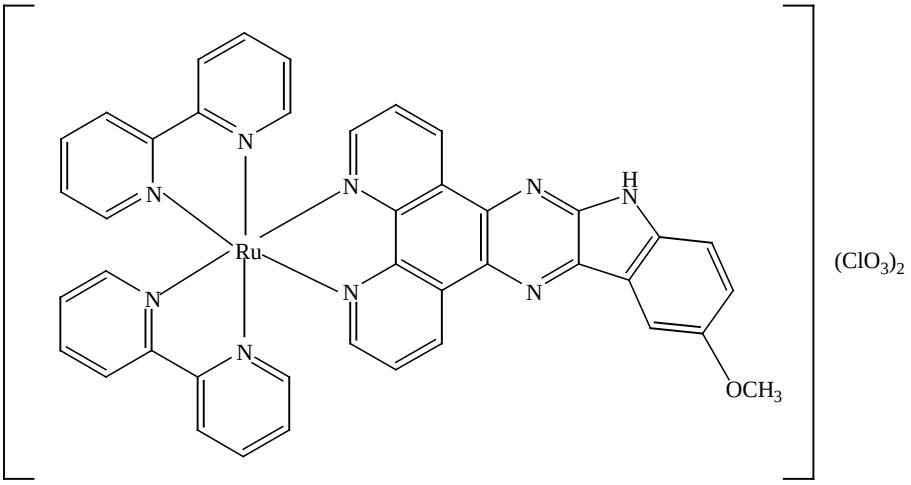
Complexes $[Cu^{II}$(ternary-S-methyl-L-cysteine)(dipyridoquinoxaline)$(H_2O)](ClO_4)$, $[Co^{III}$ (ethylenediamine$)_2$(imidazo[4,5-f][1,10]-phenanthroline)$]Br_3$, $[Ru^{II}$(2,2'-bipyridine$)_2$(5-methoxy-isatino-[1,2-b]-1,4,8,9-tetraazatriphenylene$)](ClO_4)_2$ and $[Ni^{II}$(naptho[2,3-*a*] dipyrido[3,2-*h*:2',3'-*f*]phenazine-5,18-dione)(1,10-phenanthroline$)](PF_6)_2$ (Figure 1.12) can trigger photolytic DNA cleavage upon irradiation [34, 39, 40, 41]. Complexes $[Cu^{II}$(ternary-S-methyl-L-cysteine)(dipyridoquinoxaline)$(H_2O)](ClO_4)$ and $[Ru^{II}$(2,2'-bipyridine$)_2$(5-methoxy-isatino-[1,2-b]-1,4,8,9-tetraaza triphenylene$)](ClO_4)_2$ can induce both single and double DNA scissions by converting supercoil DNA to nicked and linear forms while complexes $[Co^{III}$(ethylenediamine$)_2$(imidazo[4,5-f][1,10]-phenanthroline$)]Br_3$ and $[Ni^{II}$(naptho[2,3-*a*]dipyrido[3,2-*h*:2',3'-*f*]phenazine-5,18-dione)(1,10-phenanthroline$)](PF_6)_2$ can only induce single DNA scission by converting supercoil DNA to nicked form. In comparison, DNA cleavage efficiency of complexes $[Ru^{II}$(2,2'-bipyridine$)_2$(5-methoxy-isatino-[1,2-b]-1,4,8,9-tetraazatriphenylene$)](ClO_4)_2$ and $[Cu^{II}$(ternary-S-methyl-L-cysteine)(dipyridoquinoxaline)$(H_2O)](ClO_4)$ is higher than the DNA cleavage efficiency of complexes $[Co^{III}$(ethylenediamine$)_2$(imidazo)[4,5-f][1,10]-phenanthroline$)]Br_3$ and $[Ni^{II}$(naptho[2,3-*a*]dipyrido[3,2-*h*:2',3'-*f*]phenazine-5,18-dione)(1,10-phenanthroline$)](PF_6)_2$ under photolytic DNA cleavage. In mechanistic studies, DNA cleavage activity of complexes $[Ru^{II}$(2,2'-bipyridine$)_2$(5-methoxy-isatino-[1,2-b]-1,4,8,9-tetraazatriphenylene$)](ClO_4)_2$ and $[Cu^{II}$(ternary-S-methyl-L-cysteine) (dipyridoquinoxaline)$(H_2O)](ClO_4)$ can be inhibited by singlet oxygen inhibitor sodium azide which indicate the contribution of singlet oxygen in the cleavage process. The

DNA cleavage mechanism of the complexes Co^{III}(ethylenediamine)$_2$(imidazo[4,5-f][1,10]-phenanthroline)]Br_3 and [Ni^{II}(naptho[2,3-*a*]dipyrido[3,2-*h*:2',3'-*f*]phenazine-5,18-dione)(1,10-phenanthroline)]$(PF_6)_2$ is still under investigation. It is proposed that photon from the excitation source excites the metal complexes, which then transfers the energy to the ground state oxygen molecule ($^3O_2$) and excites it to the $^1\Delta g$ state ($^1O_2$). The proposed DNA cleavage mechanism of metal complex under irradiation is illustrated in Figure 1.13.

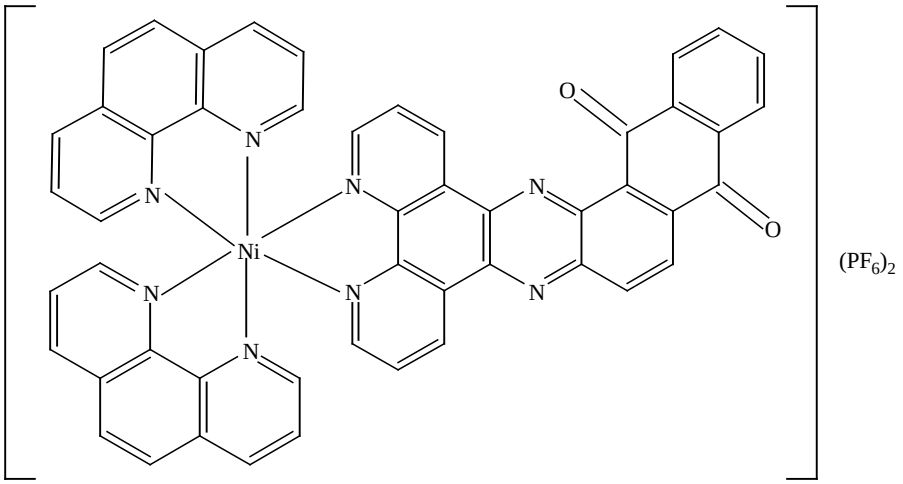


(a) (b)

Figure1.12: The schematic structures;
a) [Cu^{II}(ternary-S-methyl-L-cysteine)(dipyridoquinoxaline)(H_2O)](ClO_4)
b) [Co^{III}(ethylenediamine)$_2$(imidazo[4,5-f][1,10]-phenanthroline)]Br_3



(c)



(d)

Figure 1.12: Continued

c) [Ru^{II}(2,2'-bipyridine)$_2$(5-methoxy-isatino-[1,2-b]-1,4,8,9-tetraaza triphenylene)](ClO_4)$_2$

d) [Ni^{II}(naptho[2,3-*a*]dipyrido[3,2-*h*:2',3'-*f*]phenazine-5,18-dione)(1,10-phenanthroline)](PF_6)

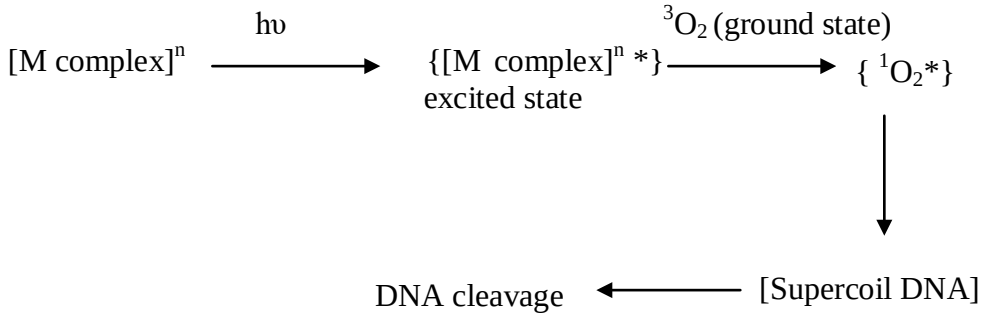
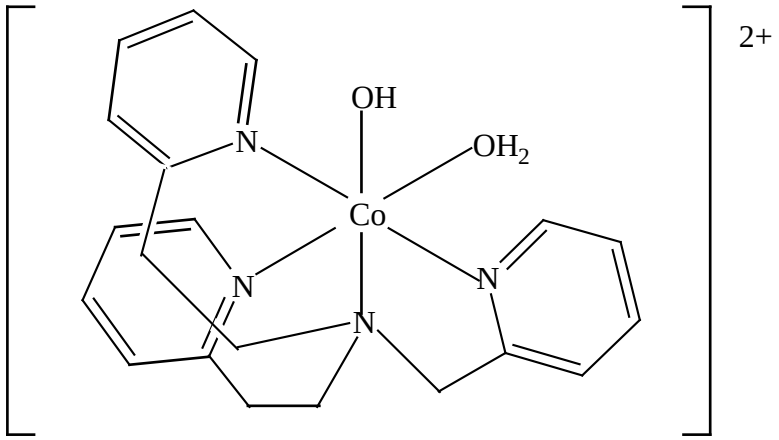


Figure 1.13: The proposed DNA cleavage mechanism of metal complex upon irradiation

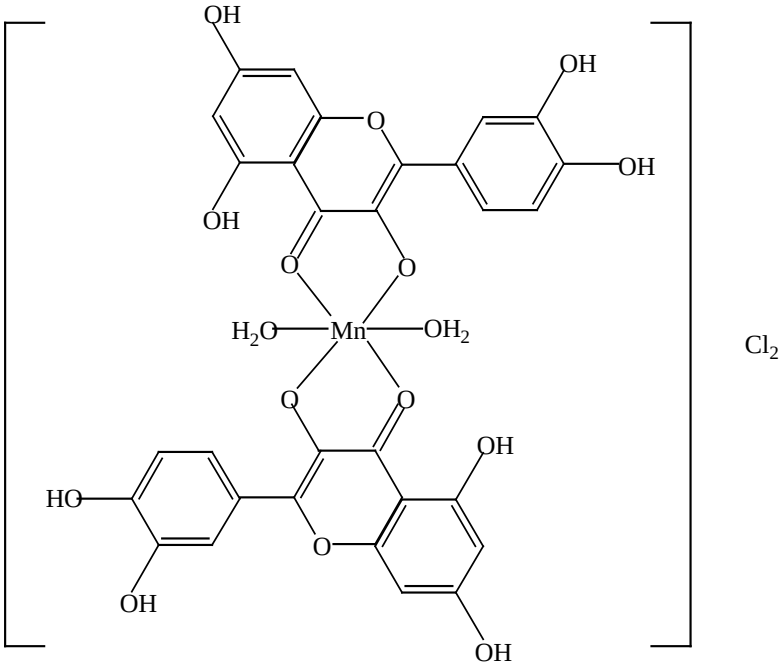
1.3.5 Hydrolytic DNA cleavage by metal complexes

Cis-aquahydroxo-tetraamine-cobalt(III) complex, $[Co^{III}$(bis[2-(2-pyridylethyl)](2-pyridylmethyl)amine)$(OH)(H_2O)]^{2+}$, generated from $[Co^{III}$(bis[2-(2-pyridylethyl)](2-pyridylmethyl)amine)$(CO_3)]ClO_4$ and complex $[Mn^{II}(quercetin)_2(H_2O)_2]Cl_2$ (Figure 1.14) can induce DNA cleavage via hydrolytic pathway [42, 43]. Both of the complexes can induce DNA cleavage in the absence of co-factor and in the dark. The mechanistic aspects of the DNA cleavage reaction have been investigated with various inhibiting agents and the results show that hydroxyl radical and singlet oxygen scavengers cannot inhibit the DNA cleavage induced by both of the complexes. These observations indicate that hydroxyl radical and singlet oxygen species are not involved in the cleavage reaction. The DNA cleavage characteristics of complexes $[Co^{III}$(bis[2-(2-pyridylethyl)](2-pyridylmethyl)amine)$(OH)(H_2O)]^{2+}$ and $[Mn^{II}(quercetin)_2(H_2O)_2]Cl_2$ support hydrolytic cleavage. Both of the complexes can induce single and double DNA scissions by converting supercoil DNA to nicked and linear forms. In hydrolytic cleavage, it is

proposed that DNA cleavage occurs at the P–O bond in the phosphodiester of DNA. The proposed hydrolytic DNA cleavage mechanism by metal complex is illustrated in Figure 1.15.



(a)



(b)

Figure 1.14: The schematic structures;
a) $[Co^{III}(bis[2\text{-}(2\text{-pyridylethyl})](2\text{-pyridylmethyl})amine)(OH)(H_2O)]^{2+}$
b) $[Mn^{II}(quercetin)_2(H_2O)_2]Cl_2$

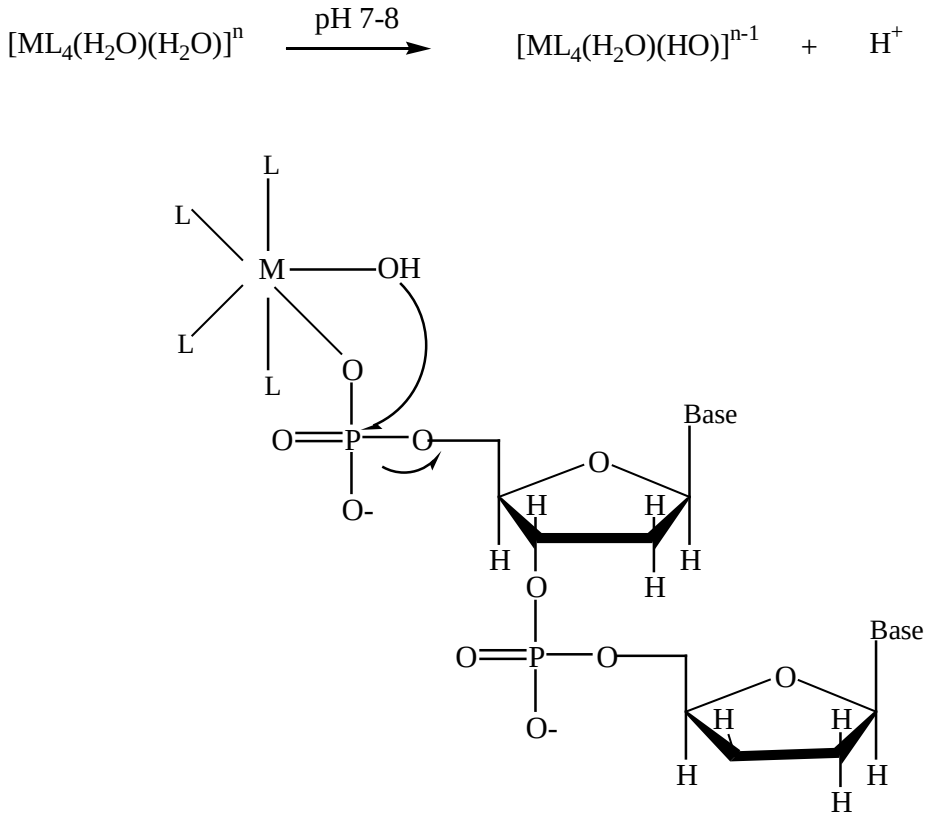
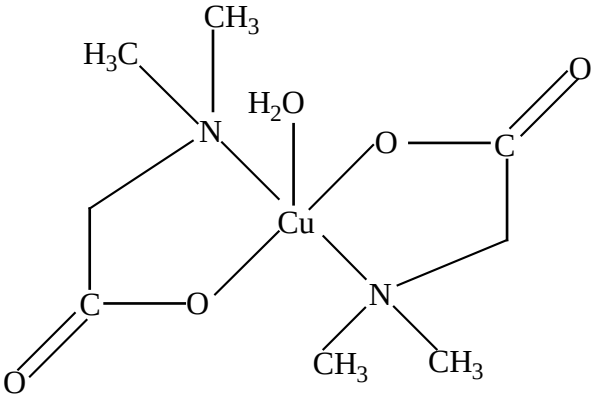


Figure 1.15: The proposed hydrolytic DNA cleavage mechanism by the metal complex

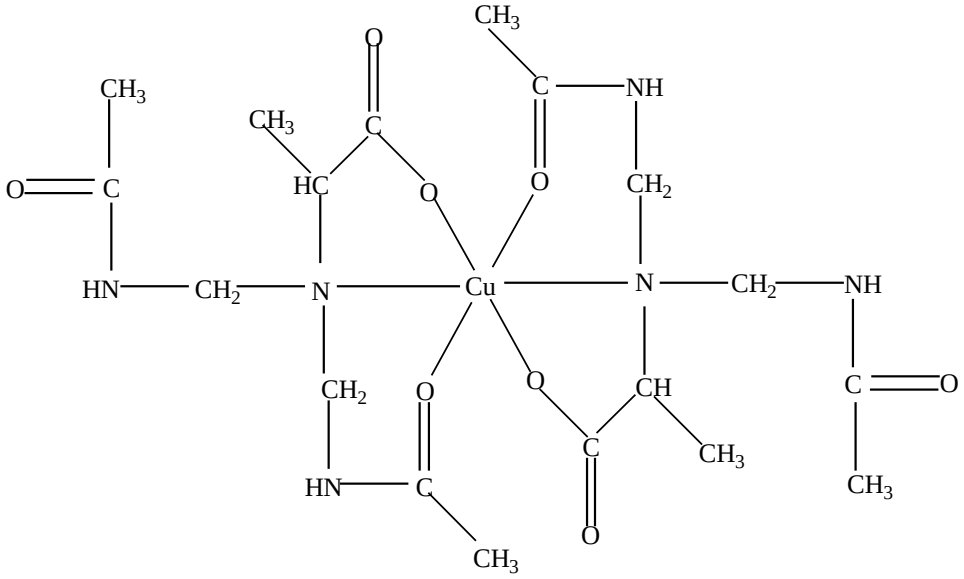
1.3.6 Oxidative DNA cleavage by copper(II) amino acid complexes in the presence of H_2O_2

Recently, Ng et al. have demonstrated that neutral Cu^{II} amino acid complexes such as Cu^{II}(*N*,*N*-di-(*N*'-methylacetamido)-L-alaninato)$_2$ and Cu^{II}(N,N'-dimethylglycinato)$_2$ (Figure 1.16) can induce oxidative cleavage of DNA in the presence of H_2O_2 [44, 45]. Both of the complexes can induce single and double DNA scissions by converting supercoil DNA to nicked and linear forms. Hydroxyl radical scavenger DMSO can inhibit significantly the cleavage reaction induced by complex Cu^{II}(N,N'-

dimethylglycinato)$_2$ which reflect the involvement of hydroxyl radical in cleavage reaction. The proposed DNA cleavage mechanism by complex Cu^{II}(N,N'-dimethylglycinato)$_2$ is similar to the proposed DNA cleavage mechanism illustrated in Figure 1.11.



(a)



(b)

Figure 1.16: The schematic structures;
a) Cu^{II}(N,N'-dimethylglycinato)$_2$
b) Cu^{II}(*N*,*N*-di-(*N*'-methylacetamido)-L-alaninato)$_2$