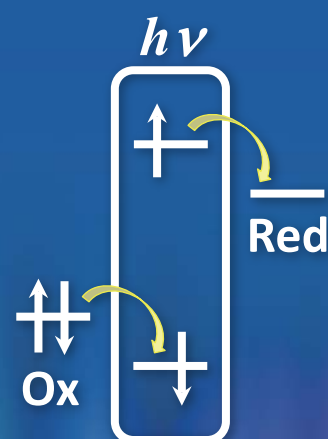
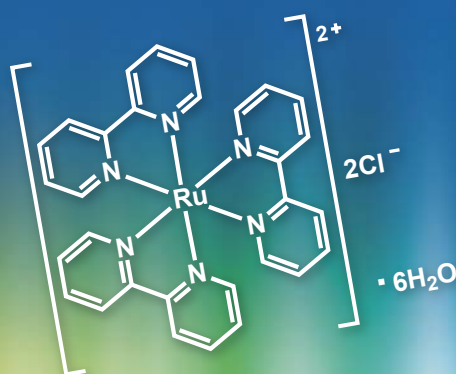
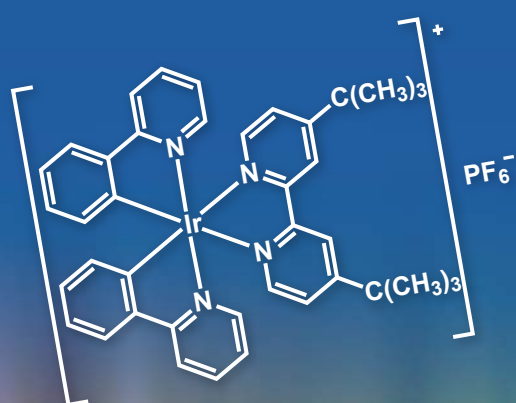




Visible Light Photoredox Catalysts



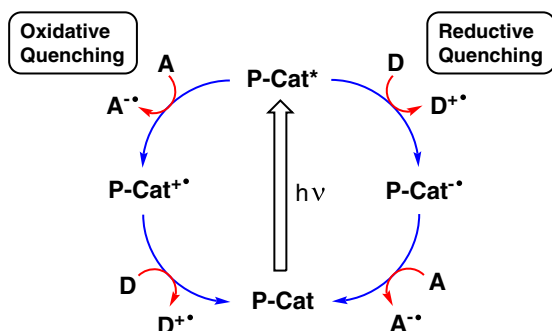
Metal Complex Catalysts

Organic Catalysts

Visible Light Photoredox Catalysts

A photoreaction undergoes a particular molecular conversion to form a product that is not obtained by a thermal reaction. Recently, visible light induced photoreactions were intensively developed. A photoreaction under visible light does not require high energy UV light and the reactions occur under mild conditions to avoid forming unexpected by-products.

A photocatalyst working for both one-electron oxidation and reduction under visible light irradiation, the so-called 'visible light photoredox catalyst', receives much attention, because there is possibility for use as a solar energy source.¹⁾ A reaction mediated by a photoredox catalyst particularly works simply, whereas a thermal reaction under coexistence of oxidizing and reducing reagents is normally hard to make work. The reaction cycle using a photoredox catalyst involves both oxidative and reductive pathways, thus it shows a 'redox-neutral' mechanism overall.

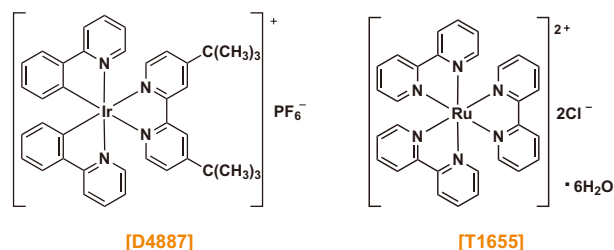


Photoredox catalysis by oxidative and reductive pathways

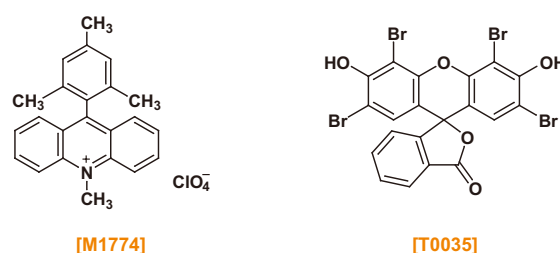
The formation of radical species normally requires a photoirradiation, redox reactions, and thermal activation by supplying high energy. The research field of photocatalysis developed transition metal complex catalysts and organic catalysts to form radical species under mild conditions, such as irradiating with visible light.

Some ruthenium(II) polypyridyl complexes and iridium(III) phenylpyridyl complexes work as photoredox catalysts under irradiation of visible light.²⁾ These transition metal complexes are useful photocatalysts, because they can form a long-lived triplet-excited state under photoirradiation. A chemical modification of the coordinating ligands controls the redox potentials of the transition metal complexes.³⁾ Metal-free organic catalysts have also been developed. Some acridinium compounds with a donor-acceptor structure can be photoredox catalysts, because the excited state exhibits a long-lived charge separation by irradiating with visible light.⁴⁾ In addition, one reported that eosin and xanthene dyes are photoredox catalysts as well.⁵⁾

Transition metal Photoredox catalysts

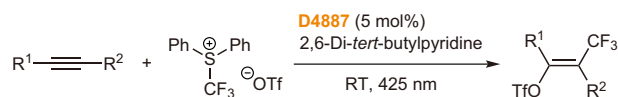


Transition metal-free photoredox catalysts

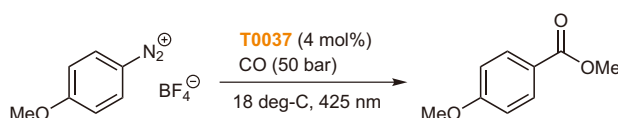


Reaction examples mediated by visible light photoredox catalysts

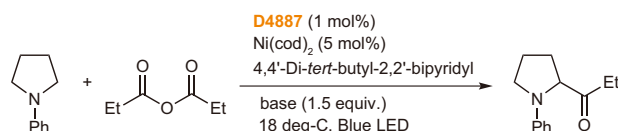
(1) Trifluoromethylation⁶⁾



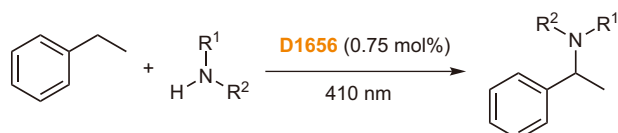
(2) Carbonylation⁷⁾

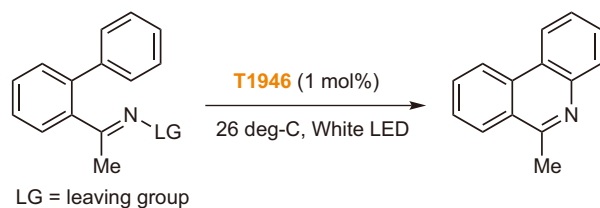
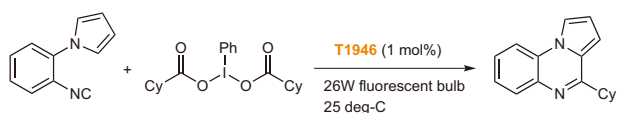
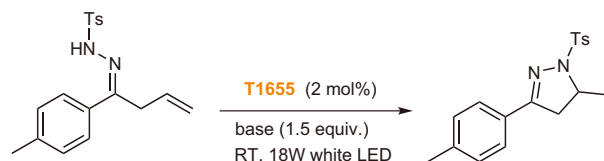


(3) Direct acylation to C-H bond⁸⁾



(4) Direct amination to C-H bond⁹⁾

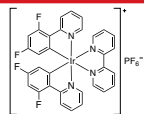
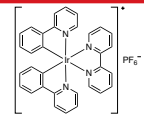
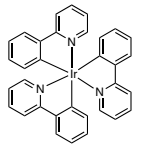
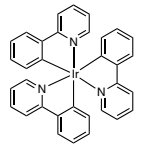
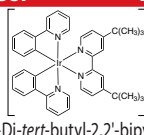
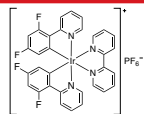
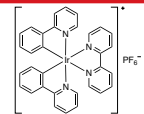
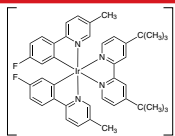
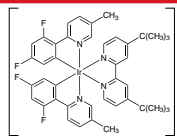
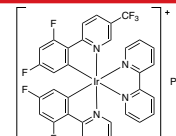
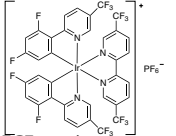
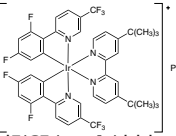
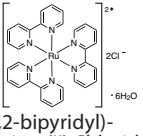
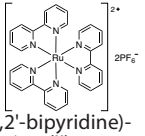
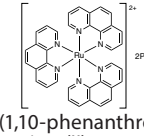
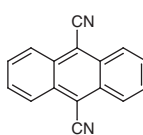
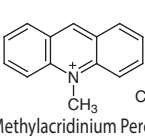
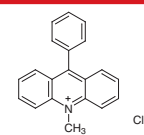
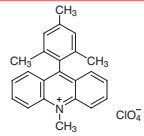
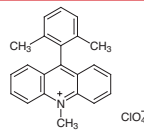
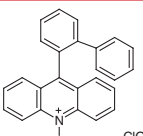
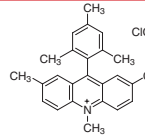
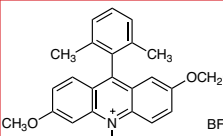
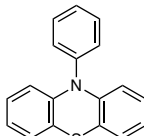
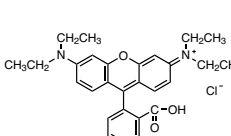
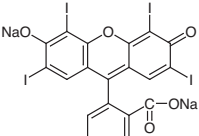
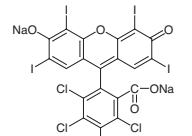
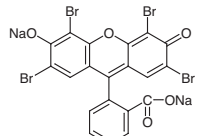
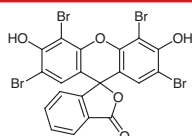
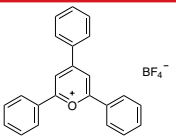
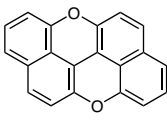


(5) Formation of iminyl radical¹⁰⁾(6) Synthesis of quinoxaline derivative¹¹⁾(7) Formation of oxazole by [3 + 2] cycloaddition¹²⁾(8) Formation of hydrazonyl radical¹³⁾

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Metal Complex Catalysts

<p>B4944 200mg</p>  <p>(2,2'-Bipyridine)bis[2-(2,4-difluorophenyl)pyridine]iridium(III) Hexafluorophosphate CAS RN: 864163-80-4</p>	<p>B4893 200mg</p>  <p>(2,2'-Bipyridine)bis(2-phenylpyridinato)iridium(III) Hexafluorophosphate CAS RN: 106294-60-4</p>	<p>T3716 200mg 1g</p>  <p>Ir(ppy)₃ CAS RN: 94928-86-6</p>	<p>T1946 200mg</p>  <p>Ir(ppy)₃ (purified by sublimation) CAS RN: 94928-86-6</p>	<p>D4887 200mg</p>  <p>(4,4'-Di-tert-butyl-2,2'-bipyridine)-bis[(2-pyridinyl)phenyl]iridium(III) Hexafluorophosphate CAS RN: 676525-77-2</p>
<p>B4944 200mg</p>  <p>(2,2'-Bipyridine)bis[2-(2,4-difluorophenyl)pyridine]iridium(III) Hexafluorophosphate CAS RN: 864163-80-4</p>	<p>B4893 200mg</p>  <p>(2,2'-Bipyridine)bis(2-phenylpyridinato)iridium(III) Hexafluorophosphate CAS RN: 106294-60-4</p>	<p>B6258 200mg 1g</p>  <p>[Ir(p-F(Me)ppy)₂-(4,4'-dtbbpy)]PF₆ CAS RN: 808142-88-3</p>	<p>B6254 200mg 1g</p>  <p>[Ir(dF(Me)ppy)₂-(dtbbpy)]PF₆ CAS RN: 1335047-34-1</p>	<p>B6161 200mg 1g</p>  <p>Ir[dF(CF₃)ppy]₂(bpy)PF₆ CAS RN: 1092775-62-6</p>
<p>B6451 100mg 500mg</p>  <p>[Ir(dFCF₃ppy)₂-(5,5'-dCF₃bpy)]PF₆ CAS RN: 1973375-72-2</p>	<p>D5817 200mg 1g</p>  <p>[Ir(dF(CF₃)ppy)₂(dtbbpy)]PF₆ CAS RN: 870987-63-6</p>	<p>T1655 1g 5g</p>  <p>Tris(2,2'-bipyridyl)-ruthenium(II) Chloride Hexahydrate CAS RN: 50525-27-4</p>	<p>T3435 1g</p>  <p>Tris(2,2'-bipyridine)-ruthenium(II) Bis(hexafluorophosphate) CAS RN: 60804-74-2</p>	<p>T3208 200mg 1g</p>  <p>Tris(1,10-phenanthroline)-ruthenium(II) Bis(hexafluorophosphate) CAS RN: 60804-75-3</p>
<h2>Organic Catalysts</h2>		<p>D1656 1g 5g</p>  <p>9,10-Dicyanoanthracene CAS RN: 1217-45-4</p>	<p>M1787 250mg 1g</p>  <p>10-Methylacridinium Perchlorate CAS RN: 26456-05-3</p>	<p>M1775 1g 5g</p>  <p>10-Methyl-9-phenylacridinium Perchlorate CAS RN: 36519-61-6</p>
<p>M1774 1g 5g</p>  <p>9-Mesityl-10-methylacridinium Perchlorate CAS RN: 674783-97-2</p>	<p>D3429 1g</p>  <p>9-(2,6-Dimethylphenyl)-10-methylacridinium Perchlorate CAS RN: 1276539-32-2</p>	<p>B2897 1g 5g</p>  <p>9-(2-Biphenyl)-10-methylacridinium Perchlorate</p>	<p>M2072 1g</p>  <p>9-Mesityl-2,7,10-trimethylacridinium Perchlorate CAS RN: 1216909-33-9</p>	<p>D5983 200mg 1g</p>  <p>Acr-450 CAS RN: 2771238-32-3</p>
<p>P2470 1g 5g</p>  <p>10-Phenylphenothiazine CAS RN: 7152-42-3</p>	<p>R0040 25g 250g</p>  <p>Rhodamine B CAS RN: 81-88-9</p>	<p>T0557 25g</p>  <p>Erythrosine B CAS RN: 16423-68-0</p>	<p>R0041 25g</p>  <p>Rose Bengal CAS RN: 632-69-9</p>	<p>T0037 25g</p>  <p>Eosine CAS RN: 17372-87-1</p>
<p>T0035 25g</p>  <p>Eosin Y CAS RN: 15086-94-9</p>	<p>T3968 1g</p>  <p>2,4,6-Triphenylpyrylium Tetrafluoroborate CAS RN: 448-61-3</p>	<p>X0083 1g</p>  <p>peri-Xanthenoxanthene CAS RN: 191-28-6</p>		

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