Cytotoxicity and lipophilicity of ruthenium(II) polypyridyl complexes as anti-cancer drugs

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The anti-cancer drugs based on the other metals than platinum have been attracting much attention, because the platinum-based drugs such as cisplatin show the heavy side effects and/or the drug resistance of tumor cells. Especially, the cytotoxicity of the ruthenium complexes have been actively reserched.¹ These complexes show the effects of the photo-dynamic theray (PDT) and the photo-activated chemotherapy (PACT) when they are irradiated. We have systematically synthesized [Ru(p-Cym)(L)]+ and $[Ru(L1)_2(L2)]^{2+}$ -type complexes with various 2,2'-bipyridine and 1,10-phenanthroline derivatives (Fig. 1). We have found that some of the ruthenium complexes show the strong cytotoxicity for tumor cells such as A549 and MDAMB231.² We have estimated the hydrophobicity by using reverse-phase HPLC, and discuss the relationship between the cytotoxicity and the hydrophobisity.

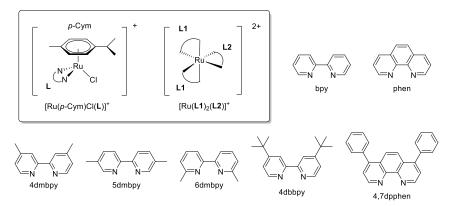


Figure 1. Ruthenium complexes synthesized in this work.

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¹Bashir, M.; Yousuf, I. et al., Coord. Chem. Rev., 2023, 487, 215169.

² Ishida, H.; Hara, M.; Hirata, Y. Jpn Pat. Appl., 2023-086639.



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