

DOCTORAL THESIS

Development of red light-activated porphycene-based photosensitizers for hypoxic anti-tumor photodynamic therapy

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ABSTRACT

This work focuses on the development of red light-activated porphycene-based photosensitizers for anti-tumor photodynamic therapy (PDT) under both normoxic and hypoxic conditions. A total of seven water-soluble porphycenes have been designed, synthesized and evaluated as potential PDT agents in terms of their photophysical and photobiological properties using principally the human nasopharyngeal carcinoma (HK-1) cells. Among the porphycenes synthesized, two were neutral amphiphilic aryl porphycenes, TDEGPPo and Zn(II) TDEGPPo, with relatively weak photo-cytotoxic activities even under normoxic condition. Two cationic porphycenes, TPyBPo and TriPyPPo, exhibited strong photo-cytotoxic activities, with LD₅₀ of 0.3 μM at a light dose of 3 J/cm², under normoxic condition. However, much lower photo-cytotoxicity was observed under hypoxic condition for TPyBPo and TriPyPPo, with LD₅₀ of 3 μM and 3.5 μM, respectively, obtained at high light doses (>10 J/cm²). Two alkyl porphycenes with one and two sulfonamide diglycol functionalities, TBPoS-OH and TBPoS-2OH, were synthesized and shown to exhibit very potent photo-cytotoxic activities, with respective LD₅₀ of 53 nM and 20 nM (light dose 8 J/cm²) under normoxic conditions. Most importantly, comparably potent photo-cytotoxicity was also observed for these porphycenes under hypoxic conditions, with respective LD₅₀ of 65 nM and 50 nM (light dose 8 J/cm²). In addition, these porphycenes were taken up by the HK-1 cells very rapidly, with >90% accumulated inside the cells after only 1 h of incubation. Confocal microscopy revealed that these porphycenes were localized at the lysosomes, mitochondria as well as endoplasmic reticulum. Furthermore, the predominant mode of cell death caused

by the PDT action of these porphycenes was shown to be apoptosis. In an attempt to effect mitochondria localization to enhance apoptotic cell death for these porphycenes, TBPoS-OH was conjugated with rhodamine B to produce the TBPoS-Rh B conjugate. This porphycene-Rh B conjugate also displayed very potent photo-cytotoxicity under both normoxic and hypoxic conditions, with LD₅₀ of 52 nM and 85 nM, respectively, at a light dose of 8 J/cm². However, confocal microscopy revealed its principal subcellular localization was at the lysosomes, not the mitochondria. The PDT activities of these porphycenes were compared to a well-known patented PDT agent, EtNBS, which is active under both normoxic and hypoxic conditions, with LD₅₀ of 58 nM and large than 1000 nM, respectively, towards the HK-1 cells. This comparison clearly shows that our sulfonoamido-porphycenes, TBPoS-OH, TBPoS-2OH and TBPoS-Rh B conjugate, display a 15- to 25-fold stronger hypoxic PDT activity relative to EtNBS, thus making these porphycenes excellent candidates for hypoxic anti-tumor photodynamic therapy.

Table of Contents

Declaration	i
Abstract	iii
Acknowledgements	v
Table of Content	vi
List of Figures	x
List of Schemes	xv
List of Tables	xvii
List of Abbreviations and Symbols	xix
Chapter 1 Introduction	1
1.1 Photodynamic Therapy (PDT).....	3
1.2 Synthesis of Porphycenes.....	13
1.3 Photophysical Properties of Porphycenes.....	24
1.4 Biological Applications of Porphycene Derivatives	29
1.5 Scopes of This Thesis.....	31
1.6 Reference.....	33
Chapter 2 Aryl Porphycene-Based Photosensitizers for Anti-Cancer Photo	

-dynamic Therapy	43
2.1 Introduction	45
2.2 Synthesis of Aryl-Porphycenes.....	47
2.3 Results and Discussion	68
2.4 Conclusion	75
2.5 Reference.....	76
 Chapter 3 Novel Red-Absorbing Porphycene-Based Photosensitizers for Hypoxic Anti-Tumor Photodynamic Therapy	 79
3.1 Introduction	81
3.2 Synthesis Procedures.....	84
3.3 Results and Discussion	88
3.4 Conclusion	104
3.5 Reference.....	105
 Chapter 4 Alkyl-Porphycene Conjugate in vitro Studies for Photodynamic Therapy against Hypoxic Tumors.....	 107
4.1 Introduction	109

4.2 Synthesis Procedures.....	111
4.3 Results and Discussion	114
4.4 Conclusion	124
4.5 Reference.....	125
Chapter 5 Experimental Details.....	127
5.1 Synthesis and Characterization of <i>t</i> -Bu ₄ Po.....	129
5.2 Photophysical Measurement.....	133
5.3 Evaluation of Protocol and Setup for Hypoxic PDT.....	135
5.4 <i>In Vitro</i> Studies	137
Chapter 6 Conclusion and Future Work	141
6.1 Conclusion	142
6.2 Future works.....	144
Appendix.....	149