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Abstract

The interaction of light with photosensitizing drugs lead to formation of highly reactive oxygen species exerting oxidative damage the cells. This so called photodynamic effect represents a promising method for selective destruction of tumor cells loaded with a photosensitizer. Wider clinical use of this approach requires development of new photosensitizing drugs with higher affinity to tumor cells and further optimization of the irradiation procedure. Here, we report a simple approach allowing screening of the