


 Cite this: *RSC Adv.*, 2021, **11**, 6188

Synthesizing and evaluating the photodynamic efficacy of asymmetric heteroleptic A₇B type novel lanthanide bis-phthalocyanine complexes†

 Emel Önal, ^{ab} Özge Tüncel, ^c Mohamad Albakour, ^a Gizem Gümüşgöz Çelik, ^a Ayşe Gül Gürek ^{*,a} and Serdar Özçelik ^{*,c}

In this study heteroleptic A₇B type novel Lu(III) and Eu(III) lanthanide phthalocyanines (LnPc(Pox)[Pc'(AB₃SH)]) with high extinction coefficients have been synthesized as candidate photosensitizers with reaction yields higher than 33%. The singlet oxygen quantum yields of LuPc(Pox)[Pc'(AB₃SH)] and EuPc(Pox)[Pc'(AB₃SH)], respectively, were measured 17% and 1.4% by the direct method in THF. The singlet oxygen quantum yield of LuPc(Pox)[Pc'(AB₃SH)] in THF is the highest among lutetium(III) bis-phthalocyanine complexes to date. The photodynamic efficacy of the heteroleptic lanthanide phthalocyanines was evaluated by measuring cell viabilities of A549 and BEAS-2B lung cells, selected to representing *in vitro* models for testing cancer and normal cells against potential drugs. The cell viabilities demonstrated concentration dependent behavior and were varied by the type of phthalocyanines complexes. Irradiation of the cells for 30 minutes with LED array at 660 nm producing flux of 0.036 J cm⁻² s⁻¹ increased cell death for LuPcPox-OAc, LuPc(Pox)[Pc'(AB₃SH)] and ZnPc. The IC₅₀ concentrations of LuPc(Pox)[Pc'(AB₃SH)] and ZnPc were determined to be below 10 nM for both cell lines, agreeing very well with the singlet oxygen quantum yield measurements. These findings suggest that LuPc(Pox)[Pc'(AB₃SH)] and particularly LuPcPox-OAc are promising drug candidates enabling lowered dose and shorter irradiation time for photodynamic therapy.

 Received 9th January 2021
 Accepted 20th January 2021

DOI: 10.1039/d1ra00197c

rsc.li/rsc-advances

Introduction

Sandwich-type phthalocyanine complexes of rare-earth elements (Ln(III)bisPcs) are well-studied classes of coordinating compounds with tetrapyrrole ligands.^{1–4} Owing to characteristic overlapping of ligand with π-orbitals,⁵ which is supplemented with a specific interaction of two f-electronic systems in the case of triple-decker complexes,^{6,7} these compounds possess a wide range of unique properties determining a variety of applications.^{8–11} The structural modifications in sandwich-type phthalocyanines referring synthesis of mixed-ligand and heteronuclear derivatives,¹² effectively control the intramolecular interactions and tailor materials' properties.^{13,14}

Homoleptic Ln(III)bisPcs are made of two identical phthalocyanine rings whereas heteroleptic complexes are composed of two different phthalocyanine rings. Bis-phthalocyanines are

naturally neutral organic radicals in which one macrocycle is negatively charged and the other macrocycle is deprotonated π-radical while the oxidation state of the central metal ion is (+3). In heteroleptic Ln(III)Pcs family, synthesis of asymmetric heteroleptic Ln(III)bisPcs functionalized in only one of the eight isoindole subunits of the bis- are rarely reported because it is a highly challenging in terms of chemical synthesis. In general, there are four alternative synthetic approaches for the preparation of heteroleptic bis-complexes depending on the type of reactants involved in the reactions: (1) two different metal-free Pc ligands reacting with a rare-earth salt, (2) reaction of the mono rare-earth complexes with phthalonitrile or its analogs, (3) reaction of two different phthalonitrile in the presence of a rare-earth salt and (4) mono rare-earth complexes and metal-free or alkali metal macrocycles reaction.

Photodynamic therapy (PDT) is a cancer treatment method receiving increased attention recently. PDT is a binary therapy (hopefully curative) that requires combination of preferably near-infrared light and a photosensitizer (PS). Each component is harmless by itself, but, when they are combined that they lead to the generation of reactive oxygen species (ROS) causing oxidative cell damage and eventually kill cells. Phthalocyanines (Pcs) are among promising second-generation photosensitizers having extended four benzo rings on the pyrrole units resulting in enhanced absorption in the far-red/infrared region.

^aDepartment of Chemistry, Gebze Technical University, Gebze, 41400 Kocaeli, Turkey. E-mail: gurek@gtu.edu.tr

^bFaculty of Engineering, Doğuş University, Ümraniye, 34775 Istanbul, Turkey

^cDepartment of Chemistry, Izmir Institute of Technology, Urla, 35430 Izmir, Turkey. E-mail: serdarozcelik@iyte.edu.tr

† Electronic supplementary information (ESI) available. See DOI: 10.1039/d1ra00197c



Phthalocyanines are emerging as PS-candidates with a strong light absorption ability and capable of generating higher numbers of singlet oxygen initiating death of cancer cells triggered by light. The optical properties of candidate PSs can be finely tuned in many ways through modifying type of coordinating metal^{15,16} and structure of ligand moieties.¹⁷ These capabilities make synthesis of novel Pc metal complexes including the lanthanides attractive.

Among the new generation Pc complexes, lanthanide(III) phthalocyanines are of a high interest because of possible coordination of two or more Pc macrocyclic units per metal atom forming LnPc₂, or Ln₂Pc₃.^{18–20} Lutetium ion (Lu³⁺) has high interest due to its ability to coordinate two or more Pc-molecules per one Lu ion.^{21,22} These lanthanide bis-phthalocyanine derivatives exhibit high intrinsic electrical conductivity with exciting electrochemical and electrochromic properties.⁴ The optical properties of LnPc₂ such as higher extinction coefficient in the far-red region (>670 nm) with an optimal singlet oxygen quantum yield (>0.3) and red-shifted fluorescence (>680 nm) are favorable for PDT applications. The presence of the heavy lanthanide central metal^{23,24} enhances the rate of intersystem crossing (ISC) to the triplet state.²⁵ Therefore, LnPc₂ complexes are expected to have higher excited state absorption in the triplet state compared to that in the singlet state due to enhanced intersystem crossing (ISC). The well-known lutetium complexes used in PDT are texaphyrin derivatives which are clinically well accepted for radiation and photodynamic therapy in oncology.^{26,27} The second metal ion for formation of LnPc₂ complex, which is explored in the present study, is europium(III). Similar to Lu, Eu ions tend to form bis-structures having a high yield of singlet oxygen (¹O₂) generation.^{28,29}

To the best of our knowledge there is no report on synthesis of the A₇B target asymmetric heteroleptic Eu(III) and Lu(III)

bisPcs with a polar thiol group for the “B” part, and for A part hydrophobic polyoxyethylene chains in the PDT application. In this study, we present synthesis methods allowing expeditious access to these target derivatives, schematically shown in Fig. 1, which can be described as A₇B complexes. The significance of this study is thus devoted to developing synthesis methods for A₇B targeted asymmetric heteroleptic Pc complexes, and evaluating biological responses of cells towards these novel phthalocyanine complexes as candidate photosensitizers in PDT. The Pc rings are substituted with polyoxyethylene substituents at the peripheral positions to enhance the solubility. Introduction of mercaptohexanol on the eighth isoindole subunit of the Pc ring is to ease potential conjugation of these Pcs to noble metal nanoparticles.

Photosensitizers can be activated with specific light irradiation and produce reactive oxygen species (ROS) such as singlet oxygen, oxygen free radicals that may be produced only in the irradiated area. Therefore, it is crucial to evaluate toxic influence of PSs by measuring cell viability and to determine IC₅₀ values *in vitro* conditions. We demonstrated that **LuPcPox(OAc)** among the Pcs tested offers higher photodynamic efficacy for A549 and BEAS-2B cells *in vitro* conditions.

Experimental section

Materials

All solvents and reagents were of reagent grade quality, and obtained commercially from Aldrich, Fluka or Merck. Comprehensive synthesis methods and characterizations of phthalodinitrile **FN-Pox**, **FN-OH** and mono-Ln-phthalocyanine derivatives (**H₂PcAB₃OH**, **LnPcPox(OAc)**; Ln: Lu(III) or Eu(III)) as starting materials were provided in the ESI (Scheme S1 and Fig. S1–S24).†

Synthesis of asymmetric heteroleptic A₇B type bis-phthalocyanines

Hydroxylated asymmetric heteroleptic bis-phthalocyanines [LnPc(Pox)[Pc'(AB₃OH)] (Ln = Lu(III) or Eu(III))

General procedure. In a reaction flask 1 eq. of **H₂PcAB₃OH** and 1 eq. (0.19 g, 0.087 mmol) for **LuPcPox**, or 1 eq. (0.16 g, 0.087 mmol) for **EuPcPox** were dissolved in 5 mL dry *o*-DCB/octanol mixture and 2–3 drops of organic base DBU was added at 190 °C under argon atmosphere and achieved for 6 h. The reaction was cooled to room poured in 50 mL hexane solution. For purification silica gel column chromatography was used by dichloromethane/ethanol (15 : 1) as an eluent mixture. **LuPc(Pox)[Pc'(AB₃OH)]** and **EuPc(Pox)[Pc'(AB₃OH)]** molecular formula C₁₆₈H₂₄₀LuN₁₆O₄₃S₁₅ and C₁₆₈H₂₄₀EuN₁₆O₄₃S₁₅, molecular weight was calculated 3827.70 and 3804.70 g mol⁻¹ respectively. Yield: 0.15 g (45%) and 0.21 g (55%), respectively.

LuPc(Pox)[Pc'(AB₃OH)]. FT-IR (ATR): ν_{\max} (cm⁻¹), 3478.2 (OH), 2913–2869.7 (CH₂, CH₃), 1593 (C–N), 1324, 1281, 1197, 1101–1028 (C–O–C). MALDI-TOF (matrix: DIT): calculated C₁₆₈H₂₄₀LuN₁₆O₄₃S₁₅ [M]⁺ 3827.70 *m/z*, founded [M]⁺ 3827.69 *m/z*. UV-Vis-NIR (CHCl₃), λ_{\max} /nm: 315, 535, 631, 699, 949, 1356–1973.

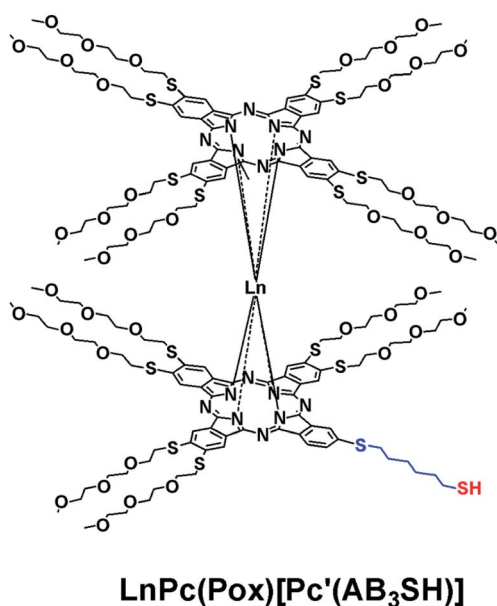


Fig. 1 Structures of the target heteroleptic thiol functionalized lanthanide bis-phthalocyanine complexes: Ln = Eu(III) or Lu(III).



EuPc(Pox)[Pc'(AB₃OH)]. FT-IR (ATR): ν_{\max} (cm⁻¹), 3502.5 (OH), 2914–2864.8 (CH₂, CH₃), 1592.4 (C–N), 1314.3, 1281.3, 1198.3, 1103.3–1027.8 (C–O–C). MALDI-TOF (matrix: DIT): calculated C₁₆₈H₂₄₀EuN₁₆O₄₃S₁₅ [M]⁺ 3804.70 *m/z*, founded [M + 7H]⁺ 3813.97 *m/z*. UV-Vis-NIR (CHCl₃), λ_{\max} /nm: 317, 573, 638, 710, 942, 1402–1896.

Mesylated asymmetric heteroleptic bis-phthalocyanines (LnPc(Pox)[Pc'(AB₃Mes)], Ln: Lu(III) or Eu(III))

General procedure. In an ice bath ~0 °C, LnPc(Pox)[Pc'(AB₃-OH)] (0.15 g, 0.04 mmol for Lu complex or 0.25 g, 0.05 mmol for Eu complex) was dissolved in dichloromethane (150 mL) in the presence of trimethylamine (2.5 mL, excess). 2.5 mL (excess of) methane sulfonyl chloride was added dropwise to the solution during 1 h. The mixture was allowed to warm to room temperature and stirred for 4 h. The reaction was followed by thin-layer chromatography (TLC) using a 15 : 1 dichloromethane : ethanol solvent mixture as an eluent. When the starting complexes were completely consumed, first 200 mL saturated NaHCO₃ solution was added to the solution and stirred vigorously for 5 min, then 200 mL diluted HCl solution was added. The two phases resulting mixture was extracted with more dichloromethane, organic phase was dried with anhydrous sodium sulphate and evaporated dichloromethane. The crude products mesylated Ln-complexes were purified by preparative TLC (silica gel) using a 15 : 1 dichloromethane : ethanol solvent system as an eluent. Yield: Lu(Pox)Pc(AB₃Mes)Pc' 0.11 g (75%), Eu(Pox)Pc(AB₃Mes)Pc' 0.16 g (72%) respectively.

LuPc(Pox)[Pc'(AB₃Mes)]. FT-IR (ATR): ν_{\max} (cm⁻¹), 2870.6 (CH₂, CH₃), 1590 (C–N), 1330.9–1194.2 (–S=O), 1248, 1071 (C–O–C), 942.5. MALDI-TOF (matrix: DIT): calculated C₁₆₈H₂₄₂LuN₁₆O₄₅S₁₆ [M]⁺ 3905 *m/z*, founded [M]⁺ 3905.79 *m/z*. ***EuPc(Pox)[Pc'(AB₃Mes)]***: FT-IR (ATR): ν_{\max} (cm⁻¹), 2860.7–2929.6 (CH₂, CH₃), 1590.7 (C–N), 1313.3–1197.7 (–S=O), 1280.1, 1097.3, 1024.8 (C–O–C). MALDI-TOF (matrix: DIT): calculated C₁₆₉H₂₄₂EuN₁₆O₄₅S₁₆ [M]⁺ 3882.79 *m/z*, founded [M + H]⁺ 3883.87 *m/z*.

Thiol substituted asymmetric heteroleptic bis-phthalocyanines (LnPc(Pox)Pc'(AB₃SH)], Ln: Lu(III) or Eu(III))

General procedure. In a reaction flask LnPc(Pox)[Pc'(AB₃Mes)] (0.11 g, 0.028 mmol) for Lu or 0.16 g, 0.04 mmol for Eu complex was dissolved in deoxygenated (sonicated for 30 min using an ultrasonic bath) tetrahydrofuran (THF)/ethanol (EtOH) mixture (75 : 25 mL). The solution was refluxed under an argon atmosphere in the dark. Then, thiourea (17 mg, excess) was added during reflux over the night. After the starting Ln complexes were consumed, argon was bubbled through the reaction mixture. Aqueous sodium hydroxide solution (20%, 12 mL) which was previously deoxygenated by sonication for 30 min was added and the mixture was refluxed for 2 h. When the reaction was complete (monitoring with TLC), the resulting mixture was poured into a mixture of dilute hydrochloric acid and ice and extracted with dichloromethane. The organic phase was extracted and dried with anhydrous sodium sulphate. The crude products were purified by preparative TLC (silica gel) using a 15 : 1 dichloromethane : ethanol solvent system as an eluent. Yield: 0.052 g (33%) for LuPc(Pox)[Pc'(AB₃SH)] or 0.058 g (38%) for EuPc(Pox)[Pc'(AB₃SH)].

LuPc(Pox)[Pc'(AB₃SH)]. FT-IR (ATR): ν_{\max} (cm⁻¹), 2867.1 (CH₂, CH₃), 2577.7 (S–H), 1591.5 (C–N), 1283.1, 1100–1028.3 (C–O–C). MALDI-TOF (matrix: DIT): calculated C₁₆₈H₂₄₀LuN₁₆O₄₂S₁₆ [M]⁺ 3843.76 *m/z*, founded [M]⁺ 3843.58 *m/z*. UV-Vis-NIR (CHCl₃), λ_{\max} /nm: 315, 533, 631, 700, 753, 953, 1467–1993.

EuPc(Pox)[Pc'(AB₃SH)]. FT-IR (ATR): ν_{\max} (cm⁻¹), 3502.5 (OH), 2914–2864.8 (CH₂, CH₃), 1592.4 (C–N), 1314.3, 1281.3, 1198.3, 1103.3–1027.8 (C–O–C). MALDI-TOF (matrix: DIT): calculated C₁₆₉H₂₄₀EuN₁₆O₄₆S₁₆ [M]⁺ 3820.76 *m/z*, founded [M – H]⁺ 3819.68 *m/z*. UV-Vis-NIR (CHCl₃), λ_{\max} /nm: 317, 574, 637, 703, 942, 1652–1890.

Cell culture

A549 cells (human non-small lung carcinoma cell lines) and BEAS-2B cells (non-tumorigenic human bronchial epithelium cells) were purchased from American Type Culture Collection. A549 cells were maintained in DMEM/F-12 (Biological Industries), supplemented with 10% FBS (fetal bovine serum) (GIBCO), 1% penicillin (Biological Industries) and 1% L-glutamine (Biological Industries). BEAS-2B cells were grown in bronchial epithelial cell growth medium (BEGM) bullet kit from Lonza. Both cell cultures were seeded in 96-well plates and kept at 37 °C in a 5% CO₂ humidified incubator.

Irradiation of cells with LED array emitting at 660 nm

The cells seeded in the 96-wells were irradiated with red light at 660 nm using a light emitting diode array (LED). The distance between the LED source and the 96-well plate was set to 15 cm. The number of incoming red photons was 2000 $\mu\text{mol m}^{-2} \text{s}^{-1}$, measured with a calibrated PAR meter (Apogee SQ-520). Light irradiation flux of 0.036 J cm⁻² s⁻¹ was used throughout the experiments. The flux and the temperature were not varied during the irradiation and continuously monitored. Variations in flux and temperature were determined to be about $\pm 1\%$, ensuring unchanged conditions for the cells during the irradiation.³⁰

The irradiation procedure is as follow: the 96-well plates were seeded with 1500 cells per well and incubated for 24 hours before adding the phthalocyanine complexes to the wells. The cells were incubated for 24 and 48 hours with various amounts (0.01 to 50 μM) of the phthalocyanine complexes. Before the LED irradiation, the cell medium was replaced with a fresh medium, eliminating the phthalocyanines that were not internalized. The irradiation time was set to 30 min.

Cell viability analysis

Cell viabilities in the dark and light conditions were evaluated by the MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide) (purchased from Sigma Aldrich) assay. The Pcs were dissolved in DMSO. All Pcs were transferred to cell culture media DMEM + FBS 10% or RPMI + FBS 10%. The final DMSO concentration was <0.001% (v/v). The control group of cells (A549 or BEAS-2B) had DMEM without DMSO in the viability test (MTT). The final concentration of the DMSO was considered negligible. The cells were incubated for further 24 and 48 hours following



the treatment (light irradiation and dark conditions). MTT ($0.5 \mu\text{g mL}^{-1}$) was added into each well and the plates were additionally incubated for 4 hours at 37°C . The formazan crystals were solubilized by addition of DMSO at the end of the incubation period. The change in MTT absorbance was measured by a Variskan spectrophotometer at both 570 nm and 720 nm, to determine the cell viability.

IC_{50} values were determined by applying curve fitting procedure to the cell viability data. We fitted a sigmoidal equation (reported in the ESI†) to obtain a dose–response curve reporting the IC_{50} value by using Origin software.

Results and discussion

Design and synthesis

For synthesis of asymmetric heteroleptic bis-Pcs it is a challenging work to control the number and location of the functional groups on the phthalocyanine ring which requires the synthesis of asymmetrically substituted derivatives. However, the increasing scientific interest of such mono-functional heteroleptic derivatives is evidenced by new research reports. For this special design most easy but less selective method belongs to reaction of two different phthalonitrile (PNs) in boiling solvent results with formation of single-decker, triple-decker, and open-chain oligomeric products, and separation of such reaction mixture seems to be very challenging.^{31–36} Recently, promising alternative one-step method was reported by Alpugan and co-workers³⁷ to access mono functionalized heteroleptic lanthanide bis complexes. In their report, a mixture of different phthalonitrile in different proportions was used and the maximum yield (37%) for A₇B heteroleptic bis-phthalocyanines and minimum for other complexes were achieved in the ratio of 10 : 1 in the presence of $\text{Eu}(\text{OAc})_3$ in refluxing dry pentanol. A reaction between rare-earth mono-Pc and PN represents better selective approach for the preparation of asymmetric heteroleptic bis-complexes.³⁸ To reduce formation of not desired mono-Pcs, another approach for the preparation of asymmetric heteroleptic complex requires synthesis of A₃B type macrocycle using a statistical template condensation of two different PNs and then its reaction with lanthanide salts and unsubstituted phthalocyanine.^{39–43} However, not surprisingly, overall yield of the target complexes are quite low.

In our study, the preparation of A₇B type rare-earth asymmetric heteroleptic bis-Pc complexes operates interactions between lanthanide (Lu and Eu) mono-Pc complex and a metal-free asymmetric Pc macrocycle in a protic or an aprotic solvent⁴⁴ which is a method 4 known as the “raise-by-one-story” method⁴⁵ that is mentioned in the introduction part. Starting synthon 4-(6-mercaptohexan-1-ol)phthalonitrile (**FN-OH**) was synthesized according to literature reports,⁴⁶ for production of 4,5-bis(4,7,10-trioxaundecan-1-sulfonyl)phthalonitrile (**FN-Pox**) simple solvent modifications⁴⁷ was done. Metal free phthalocyanines **H₂PcAB₃OH** and **H₂PcPox** were obtained by cyclic tetramerization of the phthalonitrile (**FN-Pox** and **FN-OH**) in refluxing solvent. For symmetric **H₂PcPox** synthesis, two methods were applied. In method A; the first step was

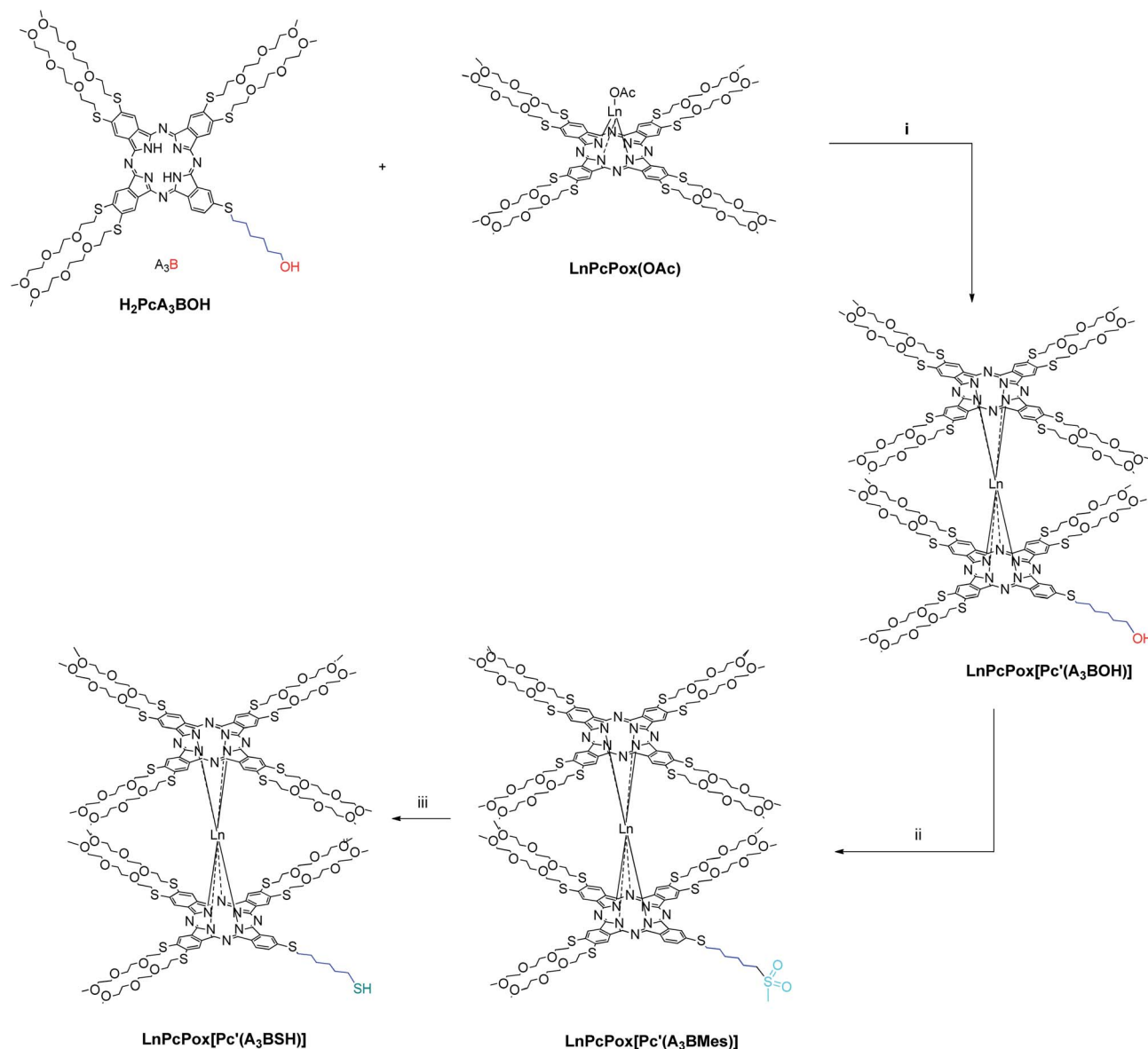
production of **ZnPc** first then second step by removing Zn metal to produce free Pc with reaction yield 72%. Even the reaction yield was higher than method B, time consuming two steps reactions and difficulty of purification of **ZnPcPox** directed us following method B. In method B which is based on *in situ* two reactions; Li-template assisted tetramerization, and *in situ* removal of the lithium ions by acidic treatments. In method B **H₂PcPox** was obtained as a byproduct during synthesis of asymmetric phthalocyanine **H₂PcAB₃OH**. Our advantage in this method B is that the A₄ metal free phthalocyanine derivative **H₂PcPox** (yield 63%) which was obtained during the synthesis of asymmetric A₃B type **H₂PcAB₃OH** (yield 35%) (Scheme S1†), it was reused to form mono-Ln(III)Pc metal complexes. Thus, two different symmetric and asymmetric phthalocyanine products that formed as a result of a single reaction were used again for formation of Lu(III) and Eu(III) bis-phthalocyanine complexes (Scheme 1). The substituted groups on asymmetric A₇B type Ln(III)bisPcs are crucial for design of PDT active asymmetric heteroleptic bisPcs as a PS candidate.

In this context, our aim was not only herein to combine different substituted mono-Pc complex and a metal-free Pc macrocycle by using multiple reaction steps. Also a new synthetic approach have been developed to obtain targeted A₇B type asymmetric heteroleptic Pcs containing only one thiol group (Scheme S1†). Ln(III)bisPcs containing SH group were prepared from **LnPc(Pox)[Pc'(AB₃OH)]** in two steps: mesylation and nucleophilic substitution by thiourea, with an overall yield of practically max. 38% (Scheme 1).^{48,49} The structures of novel compounds were characterized by a combination of IR, UV-Vis, MS and if possible ¹H NMR, ¹³C NMR spectral data (Fig. S1–S42†). All results of the complexes are consistent with the assigned formulations (Schemes 1 and S1†). To mention, although satisfactory results could not be obtained, ¹H NMR spectral experiments were carried out with reduced form of Lu and Eu bis-complexes.⁵⁰

Ground state electronic absorption spectra

In accordance with their electronic structure, Pcs present intense π – π bands in the visible (Q band) and UV (B or Soret band) spectral regions.⁵¹ Absorption spectrum with typical spectral pattern of metallo phthalocyanines shows sharp and intense band in lowest energy region (674 nm) corresponding a π – π^* transition, referred to as the Q band. The broad band in the 300–400 nm region consists of more than one component, including B₁ and B₂ bands, and referred to as Soret region.⁵² Several components observed at the foot of the Q band are considered to be of vibronic origin.⁵³ The 500–600 nm area of the spectrum is called an optical window, where there is almost no light absorption. When additional bands are observed in the window region, for example due to charge transfer transitions between the central metal ion and the π ring, the color of the complex changes.⁵⁴ The spectral properties of the lanthanide complexes are more complicated than for the metal phthalocyanines with only one Pc-ring. Furthermore, oxidation and reduction of the Pc ring produces π -cationic or π -anionic species, which have both of them characteristic spectroscopic





Scheme 1 Synthesis of asymmetric Lu(III) and Eu(III) bis-phthalocyanine complexes (Ln = Lu, Eu) (i) *o*-DCB/octanol, DBU, 190 °C, under argon; (ii) methane sulfonyl chloride, DCM, 0 °C; (iii) thiourea, THF/EtOH.

properties.⁵⁴ In our study, the Lambert–Beer law indicating linear relationship between absorbance and concentrations were observed for all lanthanide monomer and double decker complexes in DMSO and CHCl_3 in the whole concentration measurement range (Fig. S37–S42[†]). The UV–NIR spectrum of neutral Ln(III)bisPcs show a relatively intense absorption at 631 nm and 662 nm for $\text{LuPc(Pox)[Pc'(AB}_3\text{SH)]}$ and $\text{EuPc(Pox)[Pc'(AB}_3\text{SH)]}$ complexes, respectively due to the phthalocyanine Q band. The broad band at 380 and 378 nm is attributed to the phthalocyanine Soret band. In addition, the intramolecular charge transfer (ICT) between two Pc rings of neutral Ln(III) bisPcs show broad and intense absorption regions in the infrared region appearing between 1400 and 1900 nm. The two *n*-radical anion bands appeared at 535(Lu), 571(Eu) nm (BV) and 943(Lu), 936(Eu) nm (RV). The position of the ICT band for the

LnPc_2 is dependent on the size of central rare earths.⁵⁵ These results indicate that the interaction between the two rings becomes stronger as the ring-to-ring separation decreases along with the lanthanide contraction. More accurately, the blue-shift is due to the decreased distance between the Pc rings. The ionic radii of the lanthanides are in the order $\text{Lu} - 2.65 \text{ pm} < \text{Eu} - 2.84 \text{ pm}$.⁵⁶ Consequently, the blue-shift is due to the decreased distance between the Pc rings. It can be correlated with the radius of the lanthanide(III). Our results are consistent with the reports showing that for Ln(Pc)_2 .^{10,57} The spectral changes due to chemical oxidation in the UV–Vis–NIR spectra of Ln bis-phthalocyanines are illustrated in Fig. 2 and 3. The oxidation of Ln(III)bisPcs was exploited by adding iodine in CHCl_3 (40 μL , 0.1 M) to 10 mL of a chloroform solution of the Pc (1×10^{-5} M). For Lu complex $\text{LuPc(Pox)[Pc'(AB}_3\text{SH)]}$, the strongest



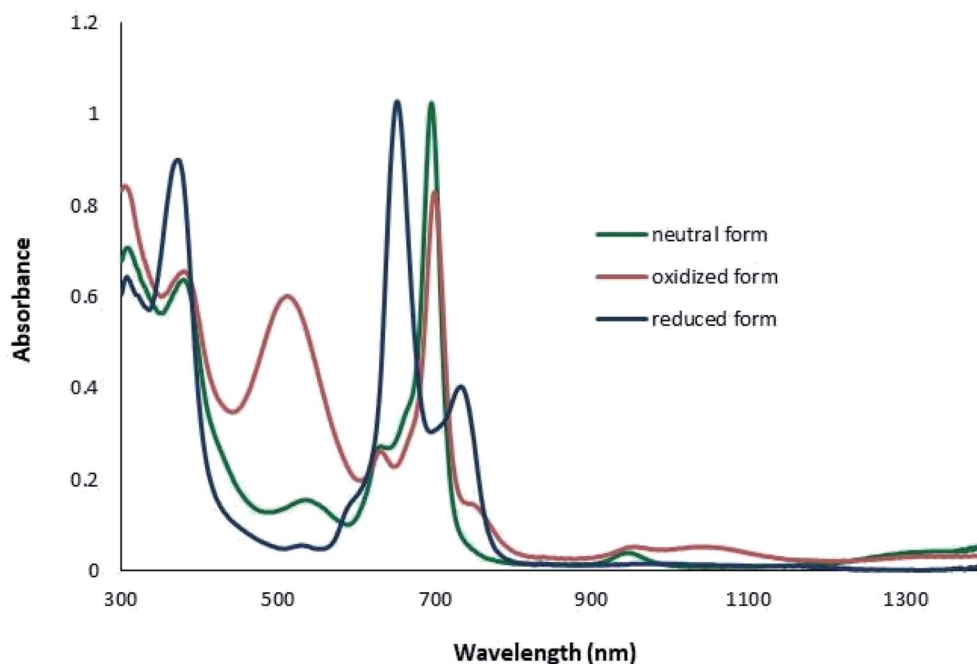


Fig. 2 UV-Vis-NIR spectra of neutral, oxidized and reduced form of LuPc(Pox)[Pc'(AB₃SH)] (1×10^{-5} M) in CHCl₃.

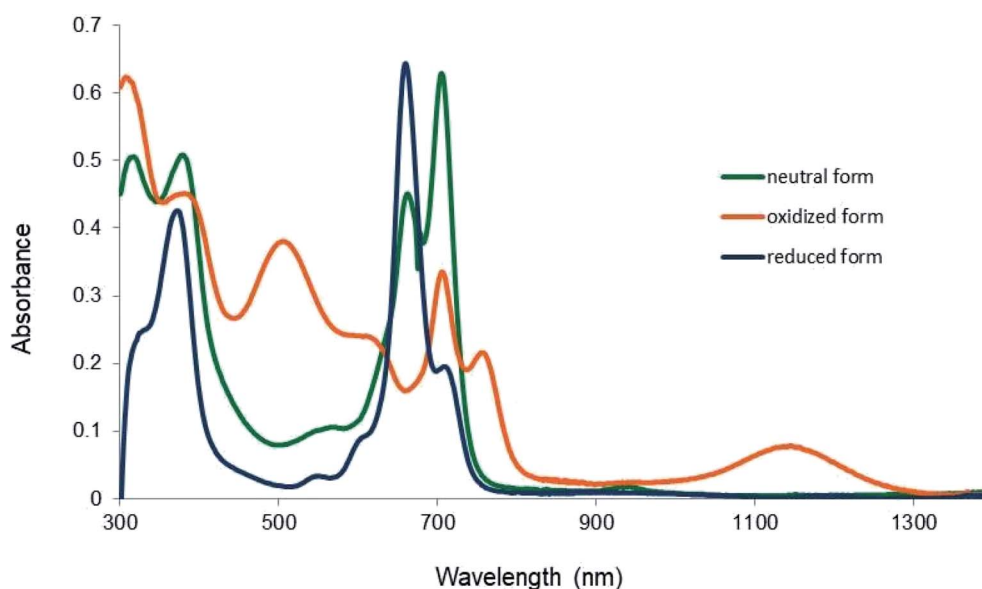


Fig. 3 UV-Vis-NIR spectra of neutral, oxidized and reduced form of EuPc(Pox)[Pc'(AB₃SH)] (1×10^{-5} M) in CHCl₃, reduced in THF, oxidation in CHCl₃.

absorption Q band at 695 nm loses intensity and exhibits slightly red shift to 699 nm, also shows that intensity of band at 511 nm and 1045 nm, corresponding to radical $\pi-\pi^*$ transition, significantly increase. In addition, one electron which causes the intermolecular transition that was prominent in the neutral form complexes, is removed from the complex, all bands corresponding to those transitions disappeared. Same results were observed for Eu complex **EuPc(Pox)[Pc'(AB₃SH)]** as well which are summarized in Table 1. The reduced forms of the

compounds were obtained by adding 4 mg of NaBH₄ to 10 mL of a solution of the Pc (1×10^{-5} M) in THF under argon atmosphere. The anionic complex (Pc₂Lu⁻) displays slightly blue shifted a splitted Q band absorbing at 651 nm and 732 nm. The intervalence transition is absent due to addition of the second electron by reduction. Moreover, the band at 1045 nm, which is linked to the radical part of the complex, also disappeared by reduction of complex. The electronic absorption spectra of the reduced form of **EuPc(Pox)[Pc'(AB₃SH)]** complex are similar to



Table 1 UV-NIR spectral data of neutral, oxidized and reduced forms of Lu and Eu bis-phthalocyanines (1×10^{-5} M). B: Soret band, BV and RV: radical transition bands, ICT: intramolecular charge transfer band

Compound	Form	λ_{\max} (nm)				
		B	BV	Q	RV	ICT
LuPc(Pox)[Pc'(AB₃SH)]	Neutral	380	535	631, 695	943	1467–1993
	Oxidized	380	511	630, 699, 760	1045	—
	Reduced	372	530	651, 732	—	—
EuPc(Pox)[Pc'(AB₃SH)]	Neutral	378	571	662, 705	936	1652–1890
	Oxidized	381	506	706, 755	1146	—
	Reduced	373	550	659, 708	—	—

reduced form of **LuPc(Pox)[Pc'(AB₃SH)]** derivative and the electronic spectral data are summarized in Table 1.

Photophysics and photochemistry

The photophysical (fluorescence quantum yield) and photochemical (singlet oxygen generation) properties of Bis-Ln-Pc₂ were investigated in comparison with unsubstituted zinc(II) phthalocyanine (**ZnPc**) in DMSO. DMSO was used as a solvent due to nontoxic property of DMSO for biological media. All of the data discussed below are summarized in Table 2.

Photophysical properties

The fluorescence quantum yield (Φ_F) is the ratio of photons emitted through fluorescence to number of photons absorbed. The fluorescence quantum yields were determined in DMSO, using a comparative method. The values were all lower than 0.01 for **LuPc(Pox)[Pc'(AB₃SH)]** and **EuPc(Pox)[Pc'(AB₃SH)]** complexes. Even monomer complex derivatives did not show any fluorescence emission in DMSO (Table 2). The absorption, excitation and emission spectra for all complexes are shown in Fig. S47–S50,† respectively. The emission signals for complexes are weak and broad and not comparable to the excitation and absorption spectrum. The low Φ_F values for all complexes could be attributed to the high atomic number of lutetium and europium which are coordinating atoms in the cavity of these phthalocyanines may facilitate the intersystem crossing to its excited triplet state after excitation.

Photochemical properties

The singlet oxygen quantum yield (Φ_Δ) is described as the number of molecules of singlet oxygen generated per number of

photons absorbed by the sensitizer. A number of different techniques for the determination of Φ_Δ measurement of efficiency have been developed over the five decades.^{58–60} In our work, we had applied a direct method in THF. Singlet-oxygen phosphorescence spectra for all compounds excited with a xenon-arc source at their respective absorption maxima were recorded by a near-IR-sensitive detector. Singlet oxygen phosphorescence spectra of Pc derivatives in THF at equal absorbances (0.23) were obtained to directly determine Φ_Δ . Φ_Δ for lutetium(III) mono **LuPcPox(OAc)** and bis phthalocyanines **LuPc(Pox)[Pc'(AB₃SH)]** derivatives respectively are 0.27 and 0.17 (Table 2). We determined the singlet oxygen quantum yield 0.17 which is the greatest value in THF for A₇B type heteroleptic lutetium(III) bis-phthalocyanine complex to date. Table 2 shows Φ_Δ values of lutetium complexes in THF along with mono and bis **EuPcPox(OAc)** and **EuPc(Pox)[Pc'(AB₃SH)]** derivatives (Fig. 4). The lower Φ_Δ values observed for europium complexes were showing inefficient molecular oxygen energy transfer.

The Φ_Δ values were also evaluated by indirect method in DMSO, a solvent in which the phthalocyanine is not aggregated. No aqueous system can entirely mimic biological media, whose complexity is likely to affect the solubility/aggregation of phthalocyanines. As indirect methods, chemical trapping and O₂ consumption methods exist which can be suited for use in both organic and aqueous media.^{61,62} In our study, the measurements were performed by examining the disappearance of the absorption band of 1,3-diphenylisobenzofuran (DPBF) upon irradiation and subsequent quenching by the singlet oxygen generated was monitored by UV-Vis spectroscopy (Fig. S43–S46†). Φ_Δ are higher for mono-lutetium **LuPcPox(OAc)** and mono-europium **EuPcPox(OAc)** complexes in DMSO. The higher values for mono complexes could be due to red shifts of

Table 2 Photophysical and photochemical parameters of mono and bis Ln(III) phthalocyanine complexes

Compound	λ_{\max} (nm)	$\log \epsilon$	λ_{ex} (nm)	$\lambda_{\text{max}}^{\text{em}}$ (nm)	$\lambda_{\text{max}}^{\text{ex}}$ (nm)	$\Delta\lambda_{\text{ST}}$ (nm)	Φ_F	Indirect Φ_Δ in DMSO	Direct Φ_Δ in THF
LuPcPox(OAc)	709	5.01	640	717	715	2	0.004	0.300	0.270
LuPc(Pox)[Pc'(AB₃SH)]	695	5.02	640	716	707	9	0.002	0.120	0.170
EuPcPox(OAc)	709	5.22	673	738	736	2	0.010	0.050	0.090
EuPc(Pox)[Pc'(AB₃SH)]	705	5.18	665	699	697	2	0.005	0.007	0.014
ZnPc	672	5.14 (ref. 63)	643	682	673	9	0.180 (ref. 64)	0.670 (ref. 65)	0.530 (ref. 66)



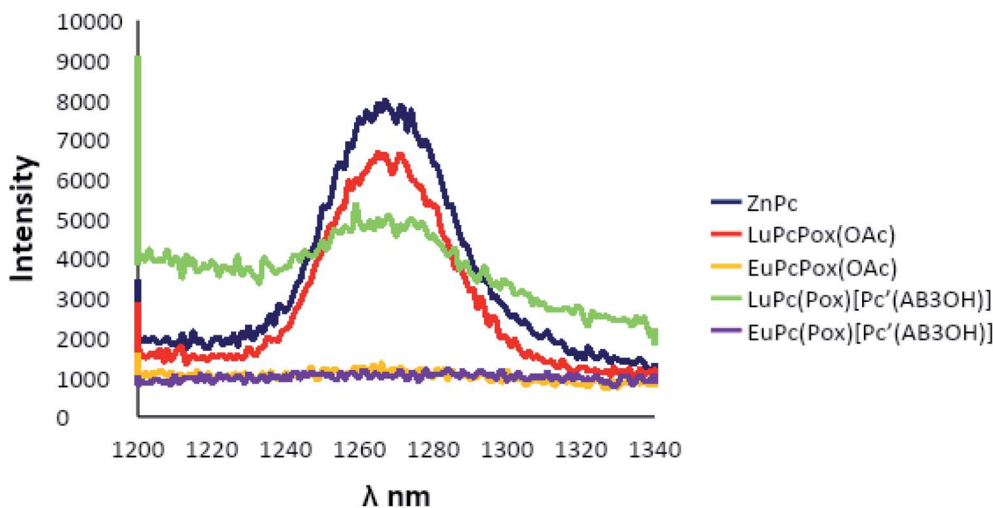


Fig. 4 Singlet oxygen phosphorescence spectra of the Pc derivatives in THF at equal absorbances (0.23) at the maximum absorption wavelength.

the Q-bands of these complexes. There were no changes in the DPBF spectra (Fig. S45 and S46[†]) even after long period of light exposure in the presence of europium complexes. It indicated that DPBF was not degraded because of lack of singlet oxygen production by these complexes.

Electrochemical determination

Although both complexes, Ln(III) mono and bis-Pcs has the same metal atom but their molecular structures are different. They exhibit different electronic interactions which can affect their voltametric behaviors. In our study, electrochemical characterizations of lanthanide series bis-Pcs having Lu³⁺ (Bis-LuPc₂), Eu³⁺ (Bis-EuPc₂) metal centers have been carried out to investigate the effects of metal center on the electrochemical responses of LnPc₂ type complexes. Fig. 5, 6, S51 and S52[†] represent the CV and SWV responses of mono and their

sandwich complexes LuPc(Pox)[Pc'(AB₃SH)] and EuPc(Pox)[Pc'(AB₃SH)] in DCM/TBAP respectively.

In Fig. 5 mono phthalocyanine LuPcPox(OAc) gives one oxidation redox couple, O₁ at 0.64 V, and two reduction couples, R₁ at -0.91 V, R₂ at -1.25 V. In Fig. 6, mono EuPcPox(OAc) showed similar electrochemical responses with only one oxidation-reduction couple O₁ at 0.77 V, and R₁ at -0.86 V. $\Delta E_{1/2}$ values of mono Lu and Eu complexes (1.55 and 1.62 V) reflect the energy band gap which define the energy necessary for the transition of an electron from the highest occupied molecular orbital (HOMO) to the lowest unoccupied molecular orbital (LUMO), are in harmony with those of MPc having redox inactive metal center and mono Lu and Eu Pc complexes.^{67,68}

The Ln(III) bis-Pc₂ complexes studied here, their radical structures are well reflected by their electrochemical behaviors since radical complexes are easily reduced or oxidized⁶⁹ Bis-Lu phthalocyanine LuPc(Pox)[Pc'(AB₃SH)] gives two oxidation

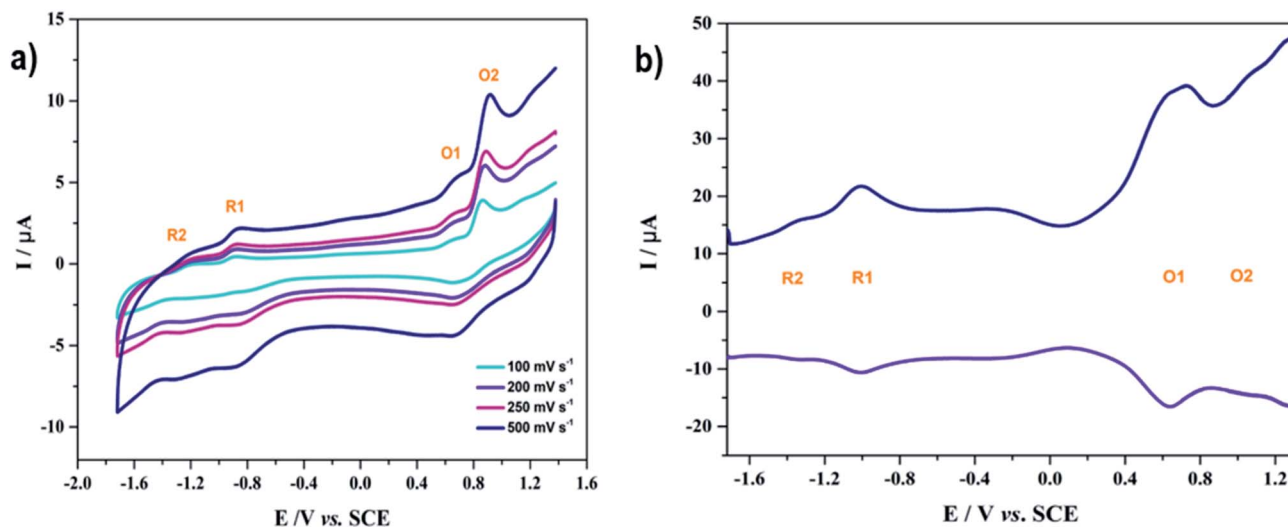


Fig. 5 (a) CVs of LuPcPox(OAc) at various scan rates on Pt in DCM/TBAP. (b) SWV of LuPcPox(OAc), SWV parameters: pulse size = 100 mV; step size = 5 mV; frequency = 25 Hz.



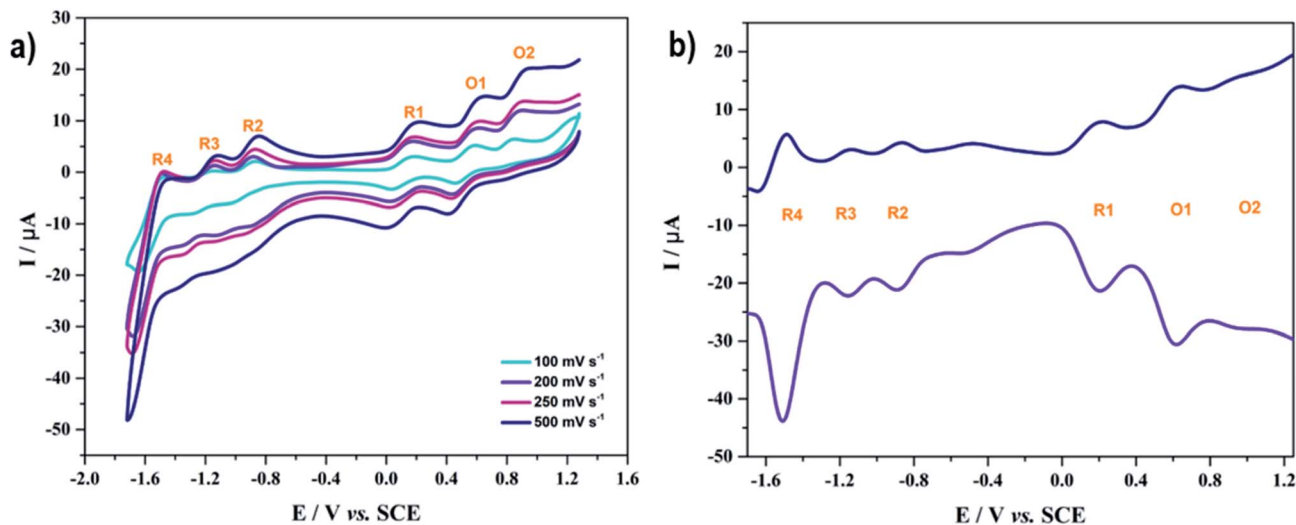


Fig. 6 (a) CVs of LuPc(Pox)[Pc'(AB₃SH)] at various scan rates on Pt in DCM/TBAP. (b) SWV of LuPc(Pox)[Pc'(AB₃SH)], SWV parameters: pulse size = 100 mV; step size: 5 mV; frequency: 25 Hz.

redox couples, O₂ at 0.71 V, O₁ at 0.44 V, and four reduction couples, R₁ at 0.03 V, R₂ at -1.02 V, R₃ at -1.31 V and R₄ at -1.65 V. Also, Bis-Eu phthalocyanine **EuPc(Pox)[Pc'(AB₃SH)]** gives two oxidation redox couples, O₂ at 0.80 V, O₁ at 0.62 V, and four reduction couples, R₁ at 0.16 V, R₂ at -0.95 V, R₃ at -1.18 V and R₄ at -1.50. Due to the π - π interaction of phthalocyanine rings around the lutetium and europium core in Ln(III) bis-Pc₂ complexes, oxidation-reduction peak separation ($\Delta E_{1/2}$) decreased to 0.42 and 0.46 V that easier electron transition reactions between two Pc rings closed to each other.

From CV and SWV data represented on Table 3, it is seen that the effect of the central metal on the redox reactions is reflected by the peak potentials for second and first oxidations and also first reduction peak potentials which are size-sensitive. The highest negative potential shift was observed with 0.03 V for **LuPc(Pox)[Pc'(AB₃SH)]** first reduction. We can conclude,

EuPc(Pox)[Pc'(AB₃SH)] complex reduction easier than **LuPc(Pox)[Pc'(AB₃SH)]** derivative which is in good agreement with literature and can be explained by the decreasing $\Delta E_{1/2}$ value of Ln(Pc)₂ with decreasing rare earth ionic radius.⁶⁹⁻⁷¹

To be able to decide that a redox reaction is reversible, peak potential should not change with scan rate and the ratio of anodic and cathodic peak currents should be equal to 1. The reaction is not reversible if these conditions are not fit. We have more expansive peaks on both anodic and cathodic directions at 500 mV s⁻¹ scan rates (Fig. 5a, 6a, S51a and S52a†). The potential shifts at higher scan rates are indicated that redox reactions have quasi-reversible systems. In the forward scan, formed species occur in the oxidation reactions, and in the backward scan, formed species are observed in the reduction reaction. The reversibility of the complex was also checked with the square wave voltammogram (SWV) given in Fig. 5b, 6b, S51b

Table 3 The electrochemical data for mono and bis Ln(III) phthalocyanines. All electrochemical data were given versus SCE in DCM. HOMO and LUMO energy level values determined from experimental cyclic voltammetry (CV)

Complex		O ₂	O ₁	R ₁	R ₂	R ₃	R ₄	^d $\Delta E_{1/2}$ (V)	^f E_{HOMO} (eV)	^g E_{LUMO} (eV)
LuPcPox(OAc)	^a $E_{1/2}$ (V)	—	0.64	-0.91	-1.25	—	—	1.55	-4.97	-3.42
	^b ΔE_p (mV)	—	48	88	31	—	—	—	—	—
	^c $I_{\text{pa}}/I_{\text{pc}}$	—	0.95	0.73	0.69	—	—	—	—	—
LuPc(Pox)[Pc'(AB₃SH)]	^a $E_{1/2}$ (V)	0.71	0.44	0.03	-1.02	-1.31	-1.65 ^e	0.42	-4.76	-4.35
	^b ΔE_p (mV)	—	130	158	150	161	—	—	—	—
	^c $I_{\text{pa}}/I_{\text{pc}}$	—	0.74	0.94	0.98	0.96	—	—	—	—
EuPcPox(OAc)	^a $E_{1/2}$ (V)	—	0.77	-0.86	—	—	—	1.62	-5.10	-3.48
	^b ΔE_p (mV)	—	102	57	—	—	—	—	—	—
	^c $I_{\text{pa}}/I_{\text{pc}}$	—	0.90	0.16	—	—	—	—	—	—
EuPc(Pox)[Pc'(AB₃SH)]	^a $E_{1/2}$ (V)	0.80	0.62	0.16 ^e	-0.95	-1.18	-1.50	0.46	-4.95	-4.49
	^b ΔE_p (mV)	—	136	38 ^e	65	63	119	—	—	—
	^c $I_{\text{pa}}/I_{\text{pc}}$	—	0.88	0.89 ^e	0.83	0.96	0.77	—	—	—

^a $E_{1/2} = (E_{\text{pa}} + E_{\text{pc}})/2$ at 0.100 V s⁻¹. ^b $\Delta E_p = |E_{\text{pa}} - E_{\text{pc}}|$. ^c $I_{\text{pa}}/I_{\text{pc}}$ for reduction, $I_{\text{pc}}/I_{\text{pa}}$ for oxidation processes at 0.100 V s⁻¹ scan rate. ^d $\Delta E_{1/2} = E_{1/2}$ (first oxidation) - $E_{1/2}$ (first reduction) = HOMO-LUMO gap for MPC having an electro inactive metal center. ^e This value is derived from SWV. ^f $E_{\text{HOMO}} = -[(E_{\text{ox}} - E_{1/2}(\text{ferrocene})) + 4.8]$. ^g $E_{\text{LUMO}} = -[(E_{\text{red}} - E_{1/2}(\text{ferrocene})) + 4.8]$.



and S52b.† While the first reduction and the first oxidation couples of all monomeric LnPc complexes are reversible, the second oxidation and reductions are quasi-reversible with respect to ΔE_p and $I_{p,a}/I_{p,c}$ values. Peak analyses of the complexes could not be performed very well for mono LnPc complexes as expected. We believe that broadening of the reduction processes may be due to the aggregation of the complexes in DCM which is non-polar and non-coordinating solvent. For Ln(III) bis-Pc₂ complexes, as shown in Fig. 5a and 6a, not only the oxidation couple O₁ and the first reduction couple R₁ but also oxidation couple O₂ and the second reduction couple R₂ are reversible with respect to ΔE_p , $\Delta E_p - E_{p/2}$, I_{pa}/I_{pc} values. All couples were monitored clearly with CV measurements (Fig. 6a and S52a†) and SWV measurements as well, except R₄ couples could not be analyzed properly and thus, were ill-defined with SWV measurements (Fig. S52b†) for **EuPc(Pox)[Pc'(AB₃SH)]** complex.

Evaluating cytotoxicity and determining IC₅₀ values

The cytotoxicity of the A549 and BEAS-2B cell lines incubated with **EuPc(Pox)[Pc'(AB₃SH)]**, **LuPc(Pox)[Pc'(AB₃SH)]** and **EuPcPox(OAc)** with a concentration range of 0.1–50.0 μM were evaluated for a time period of 24 and 48 hours. **LuPcPox(OAc)** and **ZnPc** complexes in the same incubation conditions but varying the concentration from 0.01 to 1.00 μM were used for the assessment of cytotoxicity. Fig. 7 represents the cell viabilities of A549 cells for a period of 24 hours. The viability graphs for A549 and BEAS-2B cells after 48 hours of Pc treatment were represented in ESI Part as Fig. S53 and S54.†

The type of phthalocyanine complexes was found to be significant for the biological response of the A549 cells. The cells were mostly alive (viability > 60%) for **EuPc(Pox)[Pc'(AB₃SH)]** regardless of the Pc concentrations and light irradiation.

The light irradiation did not increase toxicity of the cells in the presence of **EuPc(Pox)[Pc'(AB₃SH)]**. When the cells were incubated with **LuPc(Pox)[Pc'(AB₃SH)]** and then irradiated the cell viability was reduced 10-fold with concentration. The effect of irradiation on the viability was insignificant (statistically same) when the cells were incubated with **EuPcPox(OAc)**. The viability of the cells incubated with **ZnPc** was gradually decreased with increased concentrations and the light irradiation caused high toxicity in the sub- μM range. The most dramatic effect was observed for **LuPcPox(OAc)**: the cells were severely killed when they were exposed to the light in the presence of **LuPcPox(OAc)**. At 10 nM almost all of the cells were killed. Interestingly, the cells remained alive under the irradiation (the total dose rate of $0.036 \text{ J cm}^{-2} \text{ s}^{-1}$) when they were not treated with **LuPcPox(OAc)** in the concentration range. This finding suggests that the red-light irradiation alone up to 65 J cm^{-2} is not capable of inducing toxicity in the cells. The cytotoxicity response of the A549 cells incubated with **LuPc(Pox)[Pc'(AB₃SH)]**, **EuPcPox(OAc)** and **ZnPc** are concentration dependent.

By using these viability data, IC₅₀ values were determined by a nonlinear curve fitting. We used a dose–response sigmoidal fitting (provided in the ESI Fig. S55†) to obtain the IC₅₀ values. The IC₅₀ values were tabulated in Table 4. IC₅₀ graphs were provided in ESI Part as Fig. S56–S65† for both cell lines. **EuPc(Pox)[Pc'(AB₃SH)]** and **EuPcPox(OAc)** complexes have IC₅₀ values above 25 μM and not changing with the light irradiation. When **ZnPc** is used the IC₅₀ is about 0.25 μM when the cells are irradiated. The IC₅₀ is reduced to below 0.01 μM for **LuPcPox(OAc)** with the irradiation. When the A549 cells were treated with **LuPcPox(OAc)** complex, we microscopically observed that the cells remained on the plate were loosely attached to the plate surfaces (data not provided), suggesting that **LuPcPox(OAc)** is severely cytotoxic to A549 cells when irradiated with red-light,

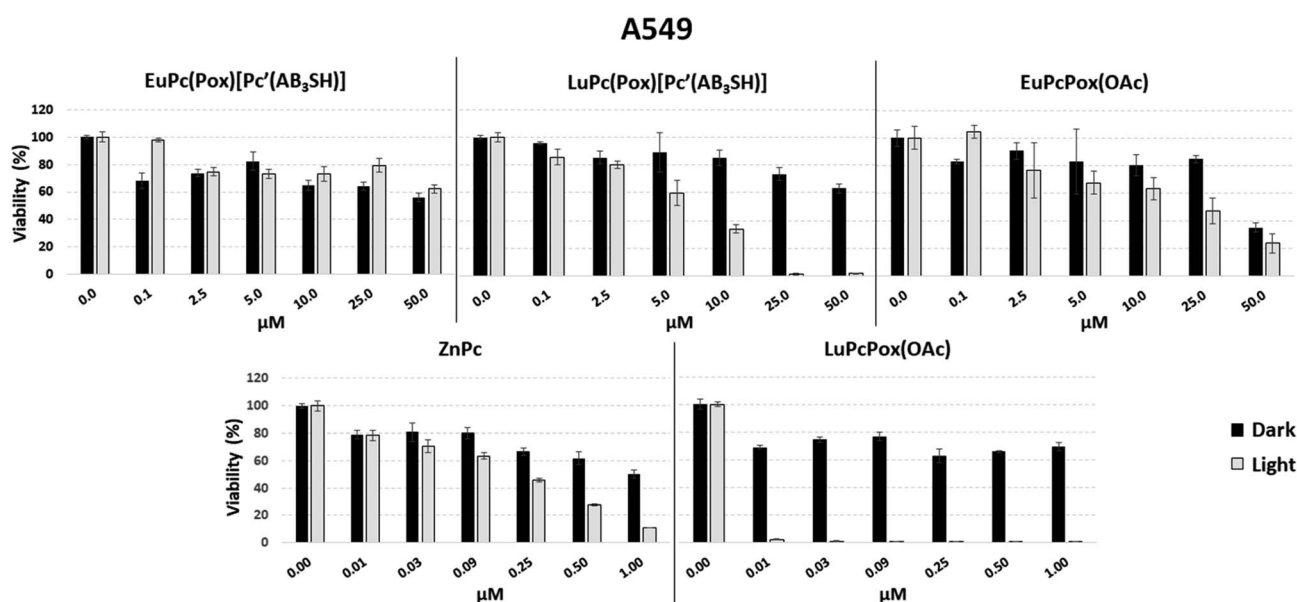


Fig. 7 Cell viabilities of A549 cells incubated with LnPcs in the dark and under the light irradiation. The light dose was $0.036 \text{ J cm}^{-2} \text{ s}^{-1}$ for 30 min of irradiation ($n = 4$).



Table 4 IC₅₀ values of all Pcs used to treat A549 and BEAS-2B cells and singlet oxygen quantum yields

Compound	Φ_{Δ} (%)		A549				BEAS-2B			
	Indirect in DMSO	Direct in THF	24 hours		48 hours		24 hours		48 hours	
			Dark	Light	Dark	Light	Dark	Light	Dark	Light
LuPcPox(OAc)	30	27	1.34	<0.01	0.93	<0.01	0.35	<0.01	0.49	<0.01
LuPc(Pox)[Pc'(AB₃SH)]	12	17	59.63	7.59	44.59	8.64	51.97	3.35	44.74	3.31
EuPcPox(OAc)	5	9	44.15	27.64	58.75	47.68	33.99	39.86	19.72	16.52
EuPc(Pox)[Pc'(AB₃SH)]	0.7	1.4	136.07	60.25	39.68	42.8	32.39	42.19	42.78	45.74
ZnPc	67	53	0.95	0.20	0.88	0.38	0.11	<0.01	1.28	0.28

confirming the IC₅₀ value (<0.01 μM). This is unprecedented for lanthanide phthalocyanines to the best of our knowledge.

Fig. 8 shows the cytotoxic responses of the BEAS-2B cells in the dark and under the irradiation. The trend for the BEAS-2B cell viability has similar trend compared to the response of the A549 cells, but the BEAS-2B cells are much more sensitive. The BEAS-2B cells were not responding to the light irradiation in the presence of **EuPc(Pox)[Pc'(AB₃SH)]** with concentrations up to 50 μM . However, the viability was significantly reduced with the light when the BEAS-2B cells were incubated with **LuPc(Pox)[Pc'(AB₃SH)]** with concentration above 2.5 μM . **ZnPc** complex was found to be toxic for the concentrations below 1.0 μM . The irradiation reduced the viability of the BEAS-2B cells when **ZnPc** with 0.25 μM or above concentration was used. The BEAS-2B cells incubated with LuPc(OAc) complexes were highly sensitive to the light irradiation: the viability reduced well below 0.01 μM of LuPc(OAc) complex used.

The BEAS-2B cells are more sensitive to the light and the phthalocyanines complexes compared to the A549 cells. The IC₅₀ values of BEAS-2B cells are smaller compared to those for

the A549 cells and tabulated in the Table 4. The IC₅₀ values for **EuPcPox(OAc)** and **EuPc(Pox)[Pc'(AB₃SH)]** were determined to be about 33 μM . **LuPc(Pox)[Pc'(AB₃SH)]** and **LuPcPox(OAc)** were highly toxic compared to Eu-complexes: 0.01 μM of **LuPcPox(OAc)** with the light was enough for killing almost all the cells. The **ZnPc** was more toxic for the BEAS-2B cells compared to the A549 cells: 0.11 μM of **ZnPc** was plentiful for cell death even in the dark (Table 4). It implies that **ZnPc** may not be considered as a good candidate photosensitizer because it is just toxic in the absence of light. These findings show that the endurance of the normal cells exposed to light is much lower. These results further imply that we need to deliver these Pc-complexes to cancer cells with higher specificity exploring the difference between cancer and normal healthy cells. In general, the expression level of certain receptors such as folate and epidermal growth factor receptors is higher on cancer cell membranes compared to normal healthy cells. This difference may be explored to direct the phthalocyanines complexes conjugating with antibodies or peptides recognizing certain

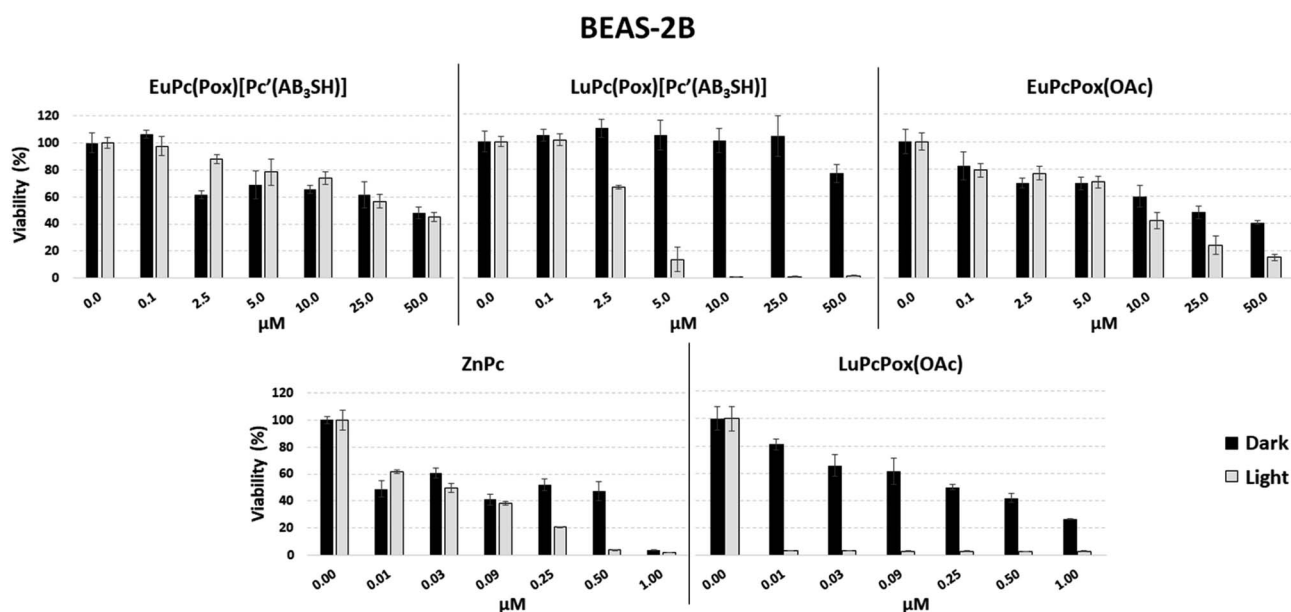


Fig. 8 Cell viabilities of BEAS-2B cells incubated with Pc complexes in the dark and light irradiations. The cells were irradiated with flux of 0.036 J cm⁻² s⁻¹ for 30 min ($n = 4$).



receptors specifically. Thus, more phthalocyanines may be accumulated in cancer cells.

The singlet oxygen generation quantum yields are in good agreement with the viability (MTT assay) findings. **LuPcPox(OAc)** and **ZnPc** generate higher amount of singlet oxygen and induce cell death at nanomolar concentrations. While the singlet oxygen generation is measured *in silico*, it is still a good indicator related to generation of radicals *in vitro* conditions. Since MTT assay measures mitochondrial activity, suggesting that these complexes producing reactive oxygen species produced by the light may interfere with mitochondrial processes. More work should be conducted to decipher cell death mechanism(s) in the dark and under the light irradiation. Nevertheless, the significance of this work is to present synthesis methods to prepare novel asymmetric heteroleptic Lu(III) complexes as promising photosensitizers that may offer lower concentration and a shorter exposure time with reduced irradiation that make faster and effective phototherapy feasible.

Conclusion

We synthesized novel bis-lanthanide Lu(III) and Eu(III) phthalocyanine complexes as candidate photosensitizers in PDT. The synthetic strategy to get A₇B type **LuPc(Pox)[Pc'(AB₃SH)]** and **EuPc(Pox)[Pc'(AB₃SH)]** involved in preparation of one polyoxo group 4,5-bis(4,7,10-trioxaundecan-1-sulfonyl) substituted Pc ring and one unsymmetrical group possessing a single functional thiol reactive substituent. Challenging multi step reactions produced **LuPc(Pox)[Pc'(AB₃SH)]** and **EuPc(Pox)[Pc'(AB₃SH)]** with reaction yields of 33% (0.052 g) and 38% (0.058 g) respectively. These reaction yields encouraged us that these newly synthesized complexes may be considered as attractive species for both fundamental studies and practical applications. The Ln(III) bis-Pc₂ sandwich complexes may be classified radicals that are well characterized by electrochemical measurements. We determined that the reduction of **EuPc(Pox)[Pc'(AB₃SH)]** is easier than **LuPc(Pox)[Pc'(AB₃SH)]** because of decreased distance between the Pc rings as a consequence of smaller radius of Eu ions. For photodynamic efficacy, the quantum yields of singlet oxygens generated by **LuPc(Pox)[Pc'(AB₃SH)]**, **LuPcPox(OAc)** and **ZnPc** are favorable. These complexes induce concentration dependent cell killing and respond to light irradiation reducing the IC₅₀ values down to nanomolar. **LuPcPox(OAc)** is determined as the best candidate photosensitizer among five phthalocyanine complexes developed in this study.

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

We are thankful to the Scientific and Technological Research Council of Turkey (TÜBİTAK) for financial support (project number: 116Z337).

References

- 1 R. Weiss and J. Fischer, *The Porphyrin Handbook*, Elsevier Science, San Diego, USA, 2003.
- 2 K. Kadish, K. M. Smith and R. Guilard, *The porphyrin handbook: bioinorganic and bioorganic chemistry*, Academic Press, 2000.
- 3 N. Kobayashi, *Coord. Chem. Rev.*, 2002, **227**, 129–152.
- 4 Y. Bian, Y. Zhang, Z. Ou and J. Jiang, *Handbook of Porphyrin Science: With Applications to Chemistry, Physics, Materials Science, Engineering, Biology and Medicine*, 2011.
- 5 N. Ishikawa, T. Okubo and Y. Kaizu, *Inorg. Chem.*, 1999, **38**, 3173–3181.
- 6 N. Ishikawa, T. Iino and Y. Kaizu, *J. Am. Chem. Soc.*, 2002, **124**, 11440–11447.
- 7 N. Ishikawa, T. Iino and Y. Kaizu, *J. Phys. Chem. A*, 2002, **106**, 9543–9550.
- 8 M. M. Nicholson, *Phthalocyanines: Properties and Applications*, 1993.
- 9 J. Jiang, K. Kasuga and D. P. Arnold, *Supramolecular Photosensitive and Electro-active Materials*, 2001.
- 10 M. M. Ayhan, A. Singh, C. Hirel, A. G. Gürek, V. Ahsen, E. Jeanneau, I. Ledoux-Rak, J. Zyss, C. Andraud and Y. Bretonnière, *J. Am. Chem. Soc.*, 2012, **134**, 3655–3658.
- 11 H. Al-Sagur, S. Komathi, M. A. Khan, A. G. Gurek and A. Hassan, *Biosens. Bioelectron.*, 2017, **92**, 638–645.
- 12 D. K. P. Ng and J. Jiang, *Chem. Soc. Rev.*, 1997, **26**, 433–442.
- 13 W. Huang, H. Xiang, Q. Gong, Y. Huang, C. Huang and J. Jiang, *Chem. Phys. Lett.*, 2003, **374**, 639–644.
- 14 S. Casilli, M. De Luca, C. Apetrei, V. Parra, Á. A. Arrieta, L. Valli, J. Jiang, M. L. Rodríguez-Méndez and J. A. De Saja, *Appl. Surf. Sci.*, 2005, **246**, 304–312.
- 15 A. C. Tedesco, J. C. Rotta and C. N. Lunardi, *Curr. Org. Chem.*, 2003, **7**, 187–196.
- 16 M. Durmuş and T. Nyokong, *Photochem. Photobiol. Sci.*, 2007, **6**, 659–668.
- 17 A. G. Gürek and Ö. Bekaroğlu, *J. Chem. Soc., Dalton Trans.*, 1994, 1419–1423, DOI: 10.1039/DT9940001419.
- 18 A. L. Thomas, *Phthalocyanine Research and Applications*, Taylor & Francis, 1990.
- 19 T. Sokolova, T. Lomova, V. Morozov and B. Berezin, *Koord. Khim.*, 1994, **20**, 637–640.
- 20 I. Kalashnikova, S. Nefedov, L. Tomilova and N. Zefirov, *Russ. Chem. Bull.*, 2007, **56**, 2426–2432.
- 21 M. Durmuş, S. Yeşilot, B. Çoşut, A. G. Gürek, A. Kılıç and V. Ahsen, *Synth. Met.*, 2010, **160**, 436–444.
- 22 R. Zugle, C. Litwinski and T. Nyokong, *Polyhedron*, 2011, **30**, 1612–1619.
- 23 L. Chen, R. Hu, J. Xu, S. Wang, X. Li, S. Li and G. Yang, *Spectrochim. Acta, Part A*, 2013, **105**, 577–581.
- 24 R. Rousseau, R. Aroca and M. Rodríguez-Mendez, *J. Mol. Struct.*, 1995, **356**, 49–62.
- 25 J. R. Darwent, P. Douglas, A. Harriman, G. Porter and M.-C. Richoux, *Coord. Chem. Rev.*, 1982, **44**, 83–126.
- 26 J. L. Sessler and R. A. Miller, *Biochem. Pharmacol.*, 2000, **59**, 733–739.



- 27 S. W. Young, K. W. Woodburn, M. Wright, T. D. Mody, Q. Fan, J. L. Sessler, W. C. Dow and R. A. Miller, *Photochem. Photobiol.*, 1996, **63**, 892–897.
- 28 C. Hou, L. Zhao, F. Geng, D. Wang and L.-H. Guo, *Anal. Bioanal. Chem.*, 2016, **408**, 8795–8804.
- 29 P. Chen, Y.-F. Huang, G.-Y. Xu, J.-P. Xue and J.-J. Chen, *J. Porphyrins Phthalocyanines*, 2019, **23**, 619–627.
- 30 E. Dube, D. O. Oluwole, N. Nwaji and T. Nyokong, *Spectrochim. Acta, Part A*, 2018, **203**, 85–95.
- 31 L. G. Tomilova, Y. G. Gorbunova, M. L. Rodriguez-Mendez and J. A. De Saja, *Mendeleev Commun.*, 1994, **4**, 127–128.
- 32 N. Ishikawa and Y. Kaizu, *Chem. Lett.*, 1998, **27**, 183–184.
- 33 N. Ishikawa and Y. Kaizu, *J. Phys. Chem. A*, 2000, **104**, 10009–10016.
- 34 Y. Liu, K. Shigehara, M. Hara and A. Yamada, *J. Am. Chem. Soc.*, 1991, **113**, 440–443.
- 35 N. Ishikawa and Y. Kaizu, *Coord. Chem. Rev.*, 2002, **226**, 93–101.
- 36 I. Alcón, M. Gonidec, M. R. Ajayakumar, M. Mas-Torrent and J. Veciana, *Chem. Sci.*, 2016, **7**, 4940–4944.
- 37 S. Alpugan, Ü. İsci, F. Albrieux, C. Hirel, A. G. Gürek, Y. Bretonnière, V. Ahsen and F. Dumoulin, *Chem. Commun.*, 2014, **50**, 7466–7468.
- 38 N. Sheng, R. Li, C.-F. Choi, W. Su, D. K. Ng, X. Cui, K. Yoshida, N. Kobayashi and J. Jiang, *Inorg. Chem.*, 2006, **45**, 3794–3802.
- 39 D. Pernin, K. Haberstroth and J. Simon, *J. Chem. Soc., Perkin Trans. 1*, 1997, 1265–1266, DOI: 10.1039/A701305A.
- 40 S. Kyatskaya, J. R. Galán Mascarós, L. Bogani, F. Hennrich, M. Kappes, W. Wernsdorfer and M. Ruben, *J. Am. Chem. Soc.*, 2009, **131**, 15143–15151.
- 41 B. Ballesteros, G. de la Torre, A. Shearer, A. Hausmann, M. Á. Herranz, D. M. Guldi and T. Torres, *Chem.–Eur. J.*, 2010, **16**, 114–125.
- 42 M. Urdampilleta, S. Klyatskaya, J. P. Cleuziou, M. Ruben and W. Wernsdorfer, *Nat. Mater.*, 2011, **10**, 502–506.
- 43 V. E. Pushkarev, V. V. Kalashnikov, A. Y. Tolbin, S. A. Trashin, N. E. Borisova, S. V. Simonov, V. B. Rybakov, L. G. Tomilova and N. S. Zefirov, *Dalton Trans.*, 2015, **44**, 16553–16564.
- 44 V. E. Pushkarev, L. G. Tomilova and V. N. Nemykin, *Coord. Chem. Rev.*, 2016, **319**, 110–179.
- 45 N. Ishikawa and Y. Kaizu, *Chem. Phys. Lett.*, 1993, **203**, 472–476.
- 46 M. N. Yaraşir, M. Kandaz, A. Koca and B. Salih, *Polyhedron*, 2007, **26**, 1139–1147.
- 47 S. S. Erdem, I. V. Nesterova, S. A. Soper and R. P. Hammer, *J. Org. Chem.*, 2009, **74**, 9280–9286.
- 48 N. Nombona, E. Antunes, C. Litwinski and T. Nyokong, *Dalton Trans.*, 2011, **40**, 11876–11884.
- 49 T. P. Mthethwa, S. Tuncel, M. Durmuş and T. Nyokong, *Dalton Trans.*, 2013, **42**, 4922–4930.
- 50 D. O. Oluwole, A. V. Yagodin, N. C. Mkhize, K. E. Sekhosana, A. G. Martynov, Y. G. Gorbunova, A. Y. Tsivadze and T. Nyokong, *Chem.–Eur. J.*, 2017, **23**, 2820–2830.
- 51 *Handbook of Porphyrin Science*, ed. K. M. Kadish, K. M. Smith and R. Guilard, World Scientific Publishing Company, 2011.
- 52 T. Nyokong, Z. Gasyna and M. J. Stillman, *Inorg. Chem.*, 1987, **26**, 1087–1095.
- 53 A. B. P. Lever, in *Advances in Inorganic Chemistry and Radiochemistry*, ed. H. J. Emeléus and A. G. Sharpe, Academic Press, 1965, vol. 7, pp. 27–114.
- 54 M. J. Stillman and T. J. Nyokong, in *Phthalocyanines: Properties and Applications*, ed. C. C. Leznoff and A. B. P. Lever, VCH, New York, 1996, ch. 9, vol. 4.
- 55 F.-L. Lu, *Polyhedron*, 2007, **26**, 3939–3946.
- 56 S. Cotton, in *Lanthanide and Actinide Chemistry*, 2006, pp. 9–22.
- 57 J. W. Buchler, P. Hammerschmitt, I. Kaufeld and J. Löffler, *Chem. Ber.*, 1991, **124**, 2151–2159.
- 58 A. A. Gorman, I. Hamblett, C. Lambert, A. L. Prescott, M. A. J. Rodgers and H. M. Spence, *J. Am. Chem. Soc.*, 1987, **109**, 3091–3097.
- 59 M. Niedre, M. S. Patterson and B. C. Wilson, *Photochem. Photobiol.*, 2002, **75**, 382–391.
- 60 R. Schmidt, C. Tanielian, R. Dunsbach and C. Wolff, *J. Photochem. Photobiol., A*, 1994, **79**, 11–17.
- 61 W. Spiller, H. Kliesch, D. Wöhrle, S. Hackbarth, B. Röder and G. Schnurpfeil, *J. Porphyrins Phthalocyanines*, 1998, **2**, 145–158.
- 62 I. Kraljić and S. E. Mohsni, *Photochem. Photobiol.*, 1978, **28**, 577–581.
- 63 I. Gürol, M. Durmuş, V. Ahsen and T. Nyokong, *Dalton Trans.*, 2007, 3782–3791.
- 64 P. Jacques and A. M. Braun, *Helv. Chim. Acta*, 1981, **64**, 1800–1806.
- 65 N. A. Kuznetsova, N. S. Gretsova, O. A. Yuzhakova, V. M. Negrimovskii, O. L. Kaliya and E. A. Luk'yanets, *Russ. J. Gen. Chem.*, 2001, **71**, 36–41.
- 66 L. Kaestner, M. Cesson, K. Kassab, T. Christensen, P. D. Edminson, M. J. Cook, I. Chambrier and G. Jori, *Photochem. Photobiol. Sci.*, 2003, **2**, 660–667.
- 67 A. B. P. Lever and E. Milaeva, in *The Phthalocyanines, Properties and Applications*, ed. C. C. Leznoff, Wiley, 1992, p. 162.
- 68 M. Arici, C. Bozoğlu, A. Erdoğan, A. L. Uğur and A. Koca, *Electrochim. Acta*, 2013, **113**, 668–678.
- 69 P. Zhu, F. Lu, N. Pan, D. P. Arnold, S. Zhang and J. Jiang, *Eur. J. Inorg. Chem.*, 2004, 510–517.
- 70 E. B. Orman, A. Koca, A. R. Özkaya, İ. Gürol, M. Durmuş and V. Ahsen, *J. Electrochem. Soc.*, 2014, **161**, H422.
- 71 V. E. Pushkarev, A. Y. Tolbin, N. E. Borisova, S. A. Trashin and L. G. Tomilova, *Eur. J. Inorg. Chem.*, 2010, 5254–5262.

