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## Phthalocyanine complexes with (4-isopropylbenzyl)oxy substituents: preparation and evaluation of anti-carbonic anhydrase, anticholinesterase enzymes and molecular docking studies

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### ABSTRACT

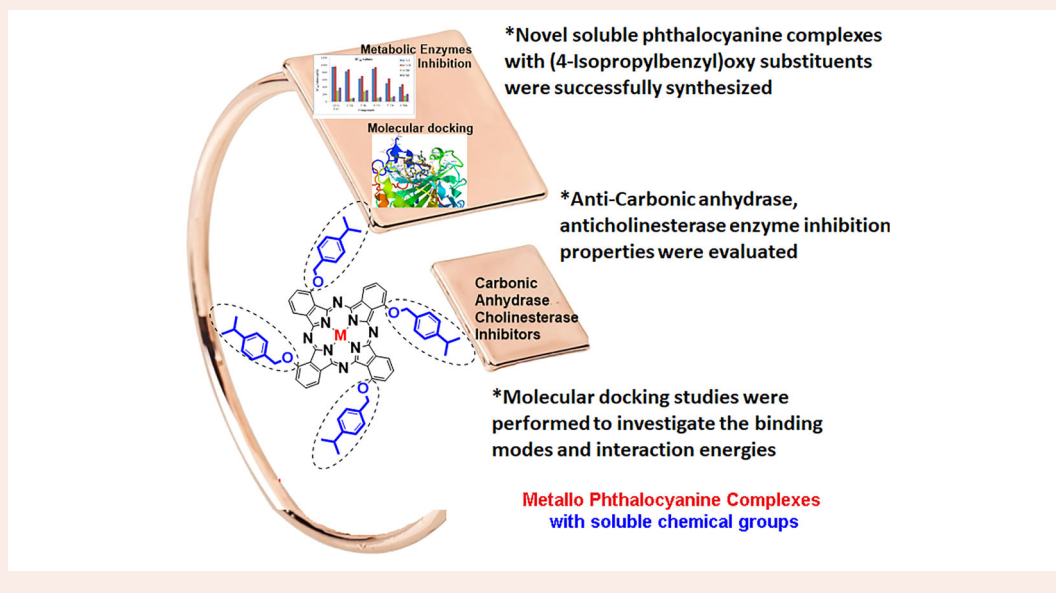
In this study, the preparation, aggregation behavior and investigation of carbonic anhydrase and cholinesterase enzyme inhibition features of non-peripherally (4-isopropylbenzyl)oxy-substituted phthalocyanines (**4–6**) are reported for the first time. The chemical structures of these new phthalocyanines were elucidated by UV-Vis (ultraviolet-visible), FT-IR (Fourier transform infrared spectrometry), NMR (nuclear magnetic resonance) and MALDI-TOF (matrix-assisted laser desorption/ionization time-of-flight) mass spectrometry. The substitution of 4-isopropylbenzyl)oxy groups benefits a remarkable solubility and redshift of the phthalocyanines Q-band. Also, these complexes were tested against some enzymes such as butyrylcholinesterase enzyme, human carbonic anhydrase I and II isoforms and acetylcholinesterase enzyme. The phthalocyanine complexes showed  $K_i$  values of in the range of  $478.13 \pm 57.25$ – $887.25 \pm 101.20 \mu\text{M}$  against hCA I,  $525.16 \pm 45.87$ – $921.14 \pm 81.25 \mu\text{M}$  against hCA II,  $68.33 \pm 9.13$ – $201.15 \pm 35.86 \mu\text{M}$  against AChE and  $86.25 \pm 13.65$ – $237.54 \pm 24.7 \mu\text{M}$  against BChE. Molecular docking studies were performed to investigate the binding modes and interaction energies of the (**2–6**) complexes with the hCA I (PDB ID:1BMZ), hCA II (PDB ID:2ABE), AChE (PDB ID:4EY6) and BChE (PDB ID:2PM8).

### ARTICLE HISTORY


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### KEYWORDS

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## Introduction

Phthalocyanines (Pcs), an important family of porphyrinoid complexes, have many applications such as gas sensors (Çimen et al., 2014), solar cells (Yıldız et al., 2019), liquid crystals phases (Chino et al., 2017; Korkut et al., 2017), electrochromic materials (Güzel, Orman, et al., 2019) and photosensitizer (Atilla et al., 2007; Güzel, 2019; Güzel et al., 2013; Sobotta et al., 2019; van de Winckel et al., 2017; Wierchowski et al., 2020) in photodynamic therapy (PDT). Soluble functional chemical groups that can be substituted to the phthalocyanine ring, the interaction between the rings is reduced and the complexes can be highly dissolved in organic or aqueous solvents. Also, in this way, the wavelength of the absorption band of the phthalocyanine complexes can be adjusted. The introduction of 4-isopropylbenzyl)oxy to the non-peripheral positions of the complexes eliminates the solubility problems of these phthalocyanines and also contributes absorption at longer wavelengths when compared to unsubstituted phthalocyanines (Gürek & Bekaroğlu, 1994; Güzel, Atmaca, et al., 2017; Önal et al., 2018; Ozoemena & Nyokong, 2002). So, these phthalocyanines are beneficial to optoelectronic, near IR devices and biological applications.

Carbonic anhydrase (CA) isoenzymes are metalloenzymes that catalyze a very easy reaction: the hydration of CO<sub>2</sub> to bicarbonate and H<sup>+</sup> (Mamedova et al., 2019; Taslimi et al., 2019). This key reaction plays a significant role in more pathological and physiological mechanisms associated with ion transport, pH control and fluid secretion (Erdemir et al., 2019). Their inhibition of these isoenzymes is the main goal correlated with the treatment of diverse diseases like obesity, glaucoma and epilepsy. More recently, CA inhibition was validated as a novel approach to fighting metastases and tumors (Boztas et al., 2019; Erdemir et al., 2019; Turkan et al., 2019a, 2019b).

The acetylcholine (ACh) molecule in the brain can be hydrolyzed by two kinds of cholinesterases including acetylcholinesterase (AChE) and butyrylcholinesterase (BChE) (El-Sayed et al., 2019; Genç Bilgiçli et al., 2019). AChE enzyme is more active than BChE enzyme and counting for approximately 80% ACh hydrolysis. Indeed, the inhibition act of AChE enzyme provides an impressive approach for the treatment of Alzheimer's disease (AD). Also, four AChE inhibitors (AChEIs) like donepezil, tacrine, galantamine and rivastigmine compounds have been approved by the FDA to improve cognitive functions and memorial of AD patients in the clinic. AChE enzyme is a protein involved in the termination of impulse transmission via rapid hydrolysis of the important neurotransmitter ACh, an advanced reduction of which had a main role in the pathogenesis of AD (Genç Bilgiçli et al., 2019; Turkan et al., 2019a, 2019b). Various drugs utilized in the therapy of AD are based on the established cholinergic hypothesis, where the objective is to raise the concentration of ACh molecule in the synaptic cleft by reducing of cholinesterase activity (Bursal et al., 2019; Turkan et al., 2019a, 2019b).

In this study, to investigate some metabolic enzymes inhibition properties, non-peripherally 4-isopropylbenzyl)oxy group containing phthalocyanines have been prepared. The synthesized phthalocyanine complexes showed good

solubility and also recorded as anticholinergic and antiepileptic potentials. Besides, we investigate the activity with the mentioned metabolic enzymes of studied complexes by molecular docking studies.

## Materials and methods

### Instruments and chemicals

Shimadzu UV 2600 spectrophotometer system was used for UV-Vis studies. Infrared spectra were performed on a PerkinElmer FT-IR spectrophotometer. NMR spectra were performed on a Bruker 300 MHz spectrometer. The clusters of peaks that corresponded to the calculated isotope composition of the molecular ion were observed by MALDI-TOF mass spectrometry (Bruker). 2,5-dihydroxybenzoic acid (DHB) was used as matrix for MALDI-TOF mass spectrometry. The elemental analysis was performed on CHNS-932 (LECO). All starting materials and solvents were purchased from Merck and Aldrich Chemical Company. Silica gel (200–400 mesh, F254, Kieselgel 60, Merck) was utilized in the purification of molecules by column chromatography assay. 3-((4-isopropylbenzyl)oxy)phthalonitrile (**1**) and its soluble zinc and indium phthalocyanine derivatives (**2** and **3**) were prepared according to the reported method (Güzel, Arslan, et al., 2019).

### General procedure for the synthesis of the phthalocyanines (2–6)

A mixture of 3-((4-isopropylbenzyl)oxy)phthalonitrile (**1**) (0.100 g, 0.36 mmol), anhydrous ZnCl<sub>2</sub>, InCl<sub>3</sub>, CuCl<sub>2</sub>, CoCl<sub>2</sub>, CuCl<sub>2</sub>, MnCl<sub>3</sub> ~ (0.030 g, excess) in hexanol and 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU, 0.05 mL) were heated to reflux temperature in sealed tube under an argon atmosphere for 16 h. After completion of the reaction, the resulting green mixture was cooled to room temperature. The crude products were precipitated by methanol–water, filtered off and then washed with the methanol–water mixture. The crude products were further purified by column chromatography using a mixture of THF:EtOH (10:1 v/v) as an eluent. Obtained Pcs **2–6** were highly soluble in ethylacetate, CH<sub>2</sub>Cl<sub>2</sub>, CHCl<sub>3</sub>, DMF and DMSO.

Yield of **2**: 42 mg (40.1%). M.p. > 200 °C. FT-IR ( $\nu_{\max}/\text{cm}^{-1}$ ): 3054 (m, Ar. C-H), 2957 (s, Aliph. C-H), 1590 (m, Ar. -C=C), 1487 (m, Aliph. C-H), 1334 (w, Ar.-C-N), 1271 (s, -R-O-Ar). <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>)  $\delta$  7.56–7.48 (m, 4H), 7.45–7.05 (m, 24H), 5.55–5.46 (m, 8H), 3.09–2.94 (m, 4H), 1.30–1.24 (m, 24H). UV-Vis  $\lambda_{\max}$  (nm) THF: 705, 633, 377. Anal. Calc. For C<sub>72</sub>H<sub>64</sub>N<sub>8</sub>O<sub>4</sub>Zn: C, 73.87; H, 5.51; N, 9.57; O, 5.47; Zn, 5.58; Found: C, 73.18; H, 5.34; N, 9.68. MS (MALDI-TOF):  $m/z$  1169.75 [M + H]<sup>+</sup>.

Yield of **3**: 34 mg (32.1%). M.p. > 200 °C. FT-IR ( $\nu_{\max}/\text{cm}^{-1}$ ): 3057 (m, Ar. C-H), 2957s, Aliph. C-H), 1587 (m, Ar. -C=C), 1487 (m, Aliph. C-H), 1334 (w, Ar.-C-N), 1271 (s, -R-O-Ar). <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>)  $\delta$  7.80–7.59 (m, 4H), 7.36–7.03 (m, 24H), 5.49–5.41 (m, 8H), 2.92–2.79 (m, 4H), 1.21–1.16 (m, 24H). UV-Vis  $\lambda_{\max}$  (nm) THF: 721, 647, 366. Anal. Calc. For C<sub>72</sub>H<sub>64</sub>ClInN<sub>8</sub>O<sub>4</sub>: C, 68.87; H, 5.14; Cl, 2.82; In, 9.14; N, 8.92; O,

5.10. Found: C, 68.16; H, 5.02; N, 8.46. MS (MALDI-TOF):  $m/z$  1219.98 [M-Cl + H]<sup>+</sup>.

Yield of **4**: 49 mg (46.8%). M.p. > 200 °C. FT-IR ( $\nu_{\max}$ /cm<sup>-1</sup>): 3062 (m, Ar. C-H), 2957 (s, Aliph. C-H), 1603 (m, Ar. -C=C), 1477 (m, Aliph. C-H), 1348 (w, Ar.-C-N), 1271 (s, -R-O-Ar). UV-Vis  $\lambda_{\max}$  (nm) THF: 696, 334. Anal. Calc. For C<sub>72</sub>H<sub>64</sub>CoN<sub>8</sub>O<sub>4</sub>: C, 74.28; H, 5.54; Co, 5.06; N, 9.62; O, 5.50 Found: C, 73.66; H, 5.55; N, 9.76. MS (MALDI-TOF):  $m/z$  1164.33 [M + H]<sup>+</sup>.

Yield of **5**: 52 mg (49.7%). M.p. > 200 °C. FT-IR ( $\nu_{\max}$ /cm<sup>-1</sup>): 3056 (m, Ar. C-H), 2960 (s, Aliph. C-H), 1607 (m, Ar. -C=C), 1486 (m, Aliph. C-H), 1350 (w, Ar.-C-N), 1228 (s, -R-O-Ar). UV-Vis  $\lambda_{\max}$  (nm) THF: 700, 350, 320. Anal. Calc. For C<sub>72</sub>H<sub>64</sub>CuN<sub>8</sub>O<sub>4</sub>: C, 73.98; H, 5.52; Cu, 5.44; N, 9.59; O, 5.47 Found: C, 73.16; H, 5.39; N, 9.69. MS (MALDI-TOF):  $m/z$  1168.72 [M + H]<sup>+</sup>.

Yield of **6**: 31 mg (31.1%). M.p. > 200 °C. FT-IR ( $\nu_{\max}$ /cm<sup>-1</sup>): 3060 (m, Ar. C-H), 2957 (s, Aliph. C-H), 1578 (m, Ar. -C=C), 1488 (m, Aliph. C-H), 1338 (w, Ar.-C-N), 1246 (s, -R-O-Ar). UV-Vis  $\lambda_{\max}$  (nm) THF: 751, 523, 321. Anal. Calc. For C<sub>72</sub>H<sub>64</sub>ClMnN<sub>8</sub>O<sub>4</sub>: C, 72.32; H, 5.40; Cl, 2.96; Mn, 4.59; N, 9.37; O, 5.35 Found: C, 73.12; H, 5.23; N, 9.46. MS (MALDI-TOF):  $m/z$  1160.86 [M-Cl + H]<sup>+</sup>.

## Biological investigation

### Carbonic anhydrase and cholinesterase activities assays

The inhibitory efficacy of new complexes on BChE/AChE activities was obtained conforming to the spectrophotometric procedure of Ellman et al. (Bursal et al., 2019) and also performed according to our previous study (Huseynova et al., 2019). Briefly, red blood cells that were extracted from fresh blood were hemolyzed and the pH of the hemolysate obtained was modified at 4 °C to 8.7 with solid Tris (Ekinci et al., 2007). The Sepharose 4BL-tyrosine-sulfanilamide affinity column was coated with Human erythrocyte hemolysate. For 1.0 M NaCl/25 mM Na<sub>2</sub>HPO<sub>4</sub> (pH 6.3) and 0.1 M NaCH<sub>3</sub>COO/0.5 M NaClO<sub>4</sub> (pH 5.6), the human carbonic anhydrase isozymes (hCA I and hCA II) were eluted. Sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) was applied and obtained in a single band for the purity of both isoenzymes. The protein determination in effluents showing activity was evaluated at 595 nm using the Bradford method spectrophotometrically (Bradford, 1976). Activities of CA isoenzymes were obtained following the Verpoorte et al. (1967) procedure. The spectrophotometric increase in the absorbance of the reaction medium was observed at 348 nm. The Lineweaver-Burk procedure (Lineweaver & Burk, 1934) CA activity (percent) versus inhibitory concentration was used to assess the inhibitory agents, and 1/V versus 1/[S] graphs were drawn. Bovine serum albumin has been used in detail as standard protein as previously given (Bayram et al., 2008; Burmaoğlu et al., 2016; Caglar et al., 2016). A percentage of activity versus inhibitor concentration graph was drawn for the designation of the inhibition efficacy of each phthalocyanine complexes with (4-isopropylbenzyl)oxy substituents on both hCA isoenzymes. Those graphs obtained the IC<sub>50</sub> values. Three different novel concentrations of

phthalocyanine complexes were used to calculate Ki values. In this study, graphs were given which are exhibited for the most effective inhibition of the studied compounds. The esterase activity determination method was used to determine these effects. According to this method, measurements were made every 5 min at a wavelength of 412 nm. Related graphs were created for the inhibition values of each molecule (%).

## Molecular docking

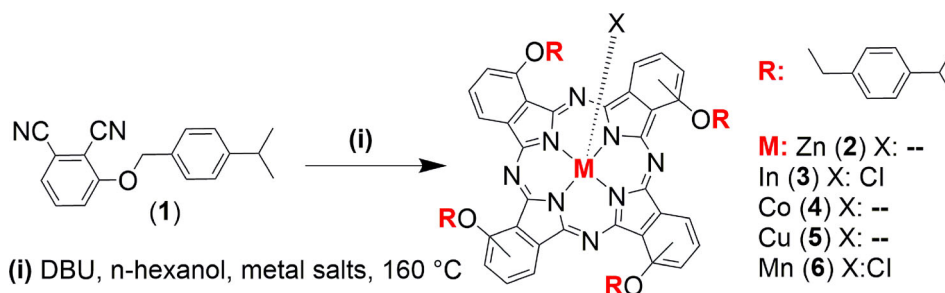
Docking studies were performed against hCA I and II, cholinesterase enzymes for non-peripheral metallophthalocyanine complexes (**2-6**). As a result of the literature review, target proteins representing enzymes were procurement from the protein data bank. The 1BMZ target protein represents human carbonic anhydrase isoenzyme I, a major member of the metalloenzymes (Karthick & Tandon, 2016). 2ABE is a protein representing the L-histidine structure situated as the most abundant activator in human carbonic anhydrase isoenzyme II (Temperini et al., 2006). 4EY6 represents the catalytic anionic region to which the quaternary trimethylammonium choline moiety of ACh is attached and hydrolyzed at cholinergic chemical synapses in the central nervous system and the bottom of the binding site of the enzyme AChE found in neuromuscular junctions (da Costa et al., 2019; Malik et al., 2017). 2PM8 represents the N-terminal residue and toxic esters and an amino acid chain on the protein surface of butyrylcholine esterase, an enzyme involved in insulin resistance, which can hydrolyze many medical (Boyko et al., 2019). The molecular docking data were obtained by using the Development Company's Docking Server service and Virtua Drug research (Sayin et al., 2018).

## Results

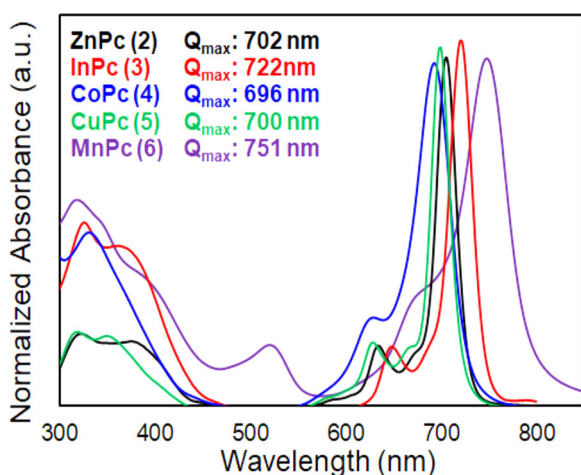
### Synthesis and characterization

Figure 1 represents the synthetic route of novel soluble metallophthalocyanine complexes (**2-6**).

Phthalocyanine precursor (**1**) and its soluble zinc and indium phthalocyanines (**2** and **3**) were prepared according to the reported procedure (Güzel, Arslan, et al., 2019). Other targeted non-peripherally substituted phthalocyanines (**4-6**) were prepared by utilizing the anhydrous metal salts in DBU, the reaction colors turned dark green and the reaction was stopped. The reactions were continued using DBU and the anhydrous metal salts (CoCl<sub>2</sub> for **4**, CuCl<sub>2</sub> for **5** and MnCl<sub>3</sub> for **6**) and appropriate solvents (*n*-hexanol) at 165 °C. The crude complexes were obtained after washing several times with the alcohol/water mixture. The yields obtained for the cobalt, copper and manganese phthalocyanines (**4**, **5** and **6**) are 47%, 50% and 31%, respectively. Elucidation of the novel phthalocyanine complexes was performed with UV-Vis spectrophotometry, elemental analysis, FT-IR, <sup>1</sup>H-NMR spectroscopy and MALDI-TOF mass spectrometry. The FT-IR spectrum of phthalocyanines (**2-6**), the peak of the C≡N vibration of compound (**1**) at 2230 cm<sup>-1</sup> disappeared as expected.



**Figure 1.** The synthesis of soluble metallophthalocyanine complexes (2–6).



**Figure 2.** Normalized absorption spectra of the soluble non-peripherally substituted phthalocyanine complexes (2–6) in THF.

Stretching vibrations of aliphatic C–H bonds of phthalocyanines (2–6), as in the phthalonitrile derivative, were detected at around  $2950\text{ cm}^{-1}$ . Similarly, stretching vibrations of aromatic C–H bonds of phthalocyanines (2–6) were observed at around  $3050\text{ cm}^{-1}$ .  $^1\text{H-NMR}$  investigations of phthalocyanine complexes 2 and 3 provided the expected chemical changes for the structure (Güzel, Orman, et al., 2019).  $^1\text{H-NMR}$  spectra of (4, 5 and 6) were precluded owing to their paramagnetic nature (Nas et al., 2017). The wide range of derivatives obtained by MALDI-TOF techniques confirmed the expected structures. Also, it has been found that the complexes synthesized in the structures observed without significant fragmentation are stable under MALDI-MS conditions. The molecular ion peaks were observed at  $m/z$ : 1169.75  $[\text{M} + \text{H}]^+$  for 2, 1219.98  $[\text{M} - \text{Cl} + \text{H}]^+$  for 3, 1164.33  $[\text{M} + \text{H}]^+$  for 4, 1168.72  $[\text{M} + \text{H}]^+$  for 5 and 1160.86  $[\text{M} - \text{Cl} + \text{H}]^+$  for 6.

### Ground state electronic absorption spectra and aggregation behavior

Absorption spectroscopy is the first and most widely used method to determine the absorption properties of phthalocyanine complexes. These complexes show typical electronic spectra with two absorption regions. One of them is the UV region at about 300–350 nm (B band) and the other in the visible area at 600–700 nm (Q-band). UV-Vis spectra of the obtained phthalocyanines, the Q-bands, which are one of the characteristic bands for phthalocyanines and observed in the visible part,

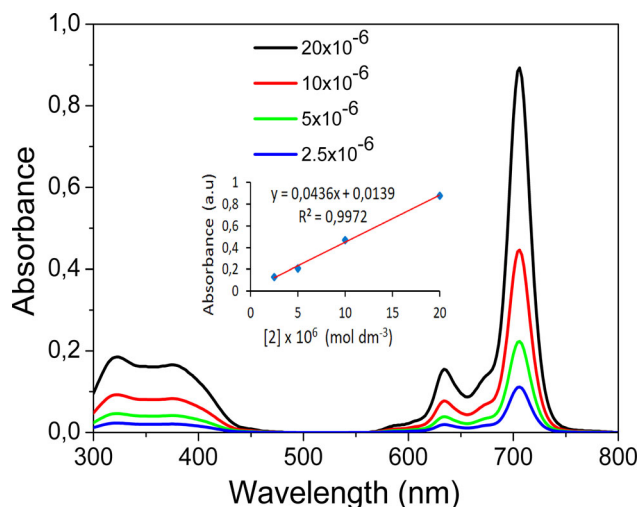
were seen at 702 nm for 2, 722 nm for 3, 696 nm for 4, 700 nm for 5 and 751 nm for 6 (Figure 2).

In metallophthalocyanine complexes (2–6), the single (narrow) Q-bands shown that these phthalocyanines are in monomeric form in the solvent medium. All complexes demonstrated intense single Q-absorption bands which are indicative of the metallophthalocyanines with the  $D_{4h}$  symmetry. Also, indium and manganese phthalocyanine complexes (3 and 6) showed redshift when compared to other metallophthalocyanine complexes in this study. This could be explained by the fact that the larger indium atom is located out of the plane of the Pc cavity for complex 3. The Q-band maximum of complex 6 shows a significant redshift concerning 2, 4 and 5, respectively, due to the oxidation state of Mn in the Pc ligand is +3, whereas Zn, Co and Cu are in oxidation state +2. Also, for manganese phthalocyanine complex (6) shows absorption at 524 nm, which was interpreted as a charge transfer absorption (phthalocyanine to metal, LMCT). Aggregation can often be explained as a very tight stacking of monomers, dimers and rings in the studied solvent. There are many factors in phthalocyanines that affect aggregation, such as concentration, nature of the solvent, peripheral and non-peripheral substituents, the metal ion in the macrocyclic nucleus and operating temperature (Atmaca et al., 2019; Güzel, Günsel, et al., 2017; Kurt et al., 2014).

To better observe the solubility and aggregation features of the phthalocyanines, dilution studies were carried out in THF solvent. As an illustrative example, non-peripherally substituted zinc phthalocyanine complex (2) was carried out using concentrations ranging from  $2.5\ \mu\text{M}$  to  $20\ \mu\text{M}$  (Figure 3).

### Metabolic enzyme inhibition results

The hCA I and II isozymes are widely distributed throughout the body. CA I isoform is involved in cerebral and retinal edema, while CA II isoforms are involved in epilepsy and glaucoma. For example, inhibitory compounds that target hCA I are involved in cerebral and retinal edema, inhibitory compounds targeting hCA IV, II, XIV and XII as diuretics, in management edema, as drugs Antiepileptic, anticoagulant and also used for the treatment of some diseases. Carbonic anhydrase inhibitors (CAIs) have been developed to treat a variety of diseases since CA is involved in several important physiological functions such as lipogenesis,  $\text{CO}_2$  and bicarbonate transfer, calcification, gluconeogenesis, tumorigenesis



**Figure 3.** Electronic absorption spectra of **2** in THF at different concentrations. The inset shows the calibration plot for the Q-band maximum.

and bone resorption. Both hCA I and II isoenzymes, and also BChE and AChE were evaluated in this study. Enzymes results of this study:

1. For hCA I isoform, the  $K_i$  values were determined in the range of  $478.13 \pm 57.25$ – $887.25 \pm 101.20$   $\mu\text{M}$ . In comparison, the  $K_i$  for the standard CA inhibitor AZA (acetazolamide), a definitive hCA I inhibitor, was  $851.53 \pm 87.33$   $\mu\text{mol/L}$  against hCA I (Table 1). All metallophthalocyanine complexes (**2–6**) had effective inhibition effects than that of AZA. Also, between these compounds, manganese metallophthalocyanine complex (**6**), was the best hCA I inhibitor ( $K_i$   $478.13 \pm 57.25$   $\mu\text{M}$ ). As shown in Table 1,  $IC_{50}$  values are in the range of  $412.46$ – $903.74$   $\mu\text{M}$  toward hCA I, for hCA II are in the range of  $481.04$ – $945.88$   $\mu\text{M}$ . Metallophthalocyanine complexes (**2–6**) synthesized in this paper significantly inhibited hCA II with  $K_i$  in the low nanomolar range.  $K_i$  values were calculated between  $525.16 \pm 45.87$  and  $921.14 \pm 81.25$   $\mu\text{M}$  (Table 1 and Figure 4). On the other hand, manganese metallophthalocyanine complex (**6**) ( $K_i$ :  $525.16 \pm 45.87$   $\mu\text{M}$ ) is, in fact, the best inhibitor in these complexes, and also was better hCA II inhibitor compared to the clinical candidate drug ( $K_i$  of AZA:  $1060.77 \pm 98.0$   $\mu\text{M}$ ).
2. AChE and BChE enzymes play an important role in the regulation of blood enzymes in the brain as important enzymes that differ in the kinetics, specificity of velocity and activity in several parts of brain cells. In the past decade, efforts to treat AD have focused on improving cholinergic neurotransmission in brain cells. This mechanism is largely based on the cholinergic hypothesis, which is evaluating cholinesterase enzyme inhibitors (ChEIs) that inhibit central cholinergic neurotransmission by inhibiting ACh degradation. In fact, ChEIs are currently the most promising treatment for AD because they reintroduce cholinergic function in human cells by increasing the amount of carbon in the central nervous system. As seen in Table 1,  $IC_{50}$  values are in the range of  $87.63$ – $284.92$   $\mu\text{M}$  toward AChE enzyme, for BChE

enzyme, are in the range of  $104.83$ – $316.54$   $\mu\text{M}$ . In this work, AChE enzyme was also significantly inhibited by metallophthalocyanine complexes (**2–6**) at the low micromolar inhibition with  $K_i$  values in the range of  $68.33 \pm 9.13$ – $201.15 \pm 15.86$   $\mu\text{M}$  (Table 1). Indeed, the most powerful AChE inhibition was recorded by zinc phthalocyanine complex (**2**) with a  $K_i$  value of  $68.33 \pm 9.13$   $\mu\text{M}$ . Finally, metallophthalocyanine complexes (**2–6**) inhibited BChE with  $K_i$  values in the range of  $86.25 \pm 13.65$ – $237.54 \pm 24.7$   $\mu\text{M}$  (Table 1 and Figure 4). However, the most powerful BChE inhibition was obtained by zinc metallophthalocyanine (**2**) with a  $K_i$  value of  $86.25 \pm 13.65$   $\mu\text{M}$ .

### Docking study

Docking studies were performed to investigate the inhibitory activity of newly synthesized metallophthalocyanine complexes (**2–6**) by central atom exchange against hCA I, hCA II, AChE and BChE enzymes. It was also detailed the secondary chemical interactions between the complexes studied with the amino acid residues of the target proteins. In docking studies, the estimated free energy of binding values of target proteins and metal complexes are given in Table 2.

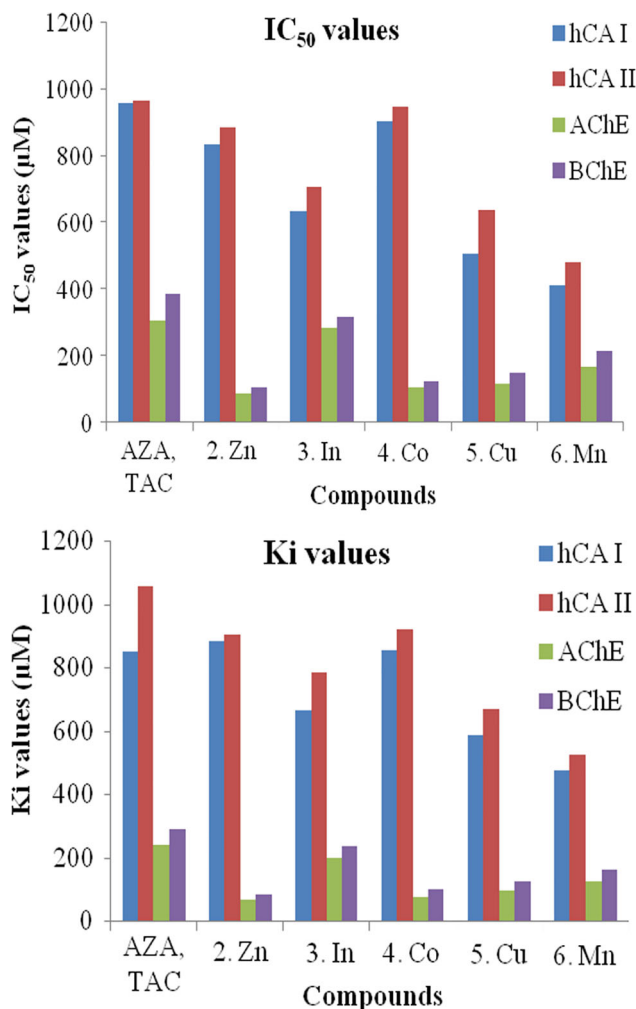
The binding energies obtained from docking results for complexes and target proteins are given in Table 2. For a good physiological protein–ligand interaction, the binding energy should increase at negative values. The results show that the experimental  $IC_{50}$  values and the inhibition efficiencies determined by docking results are almost compatible. However, the predicted inhibition efficiency of complexes with the hCA I enzyme showed more affirmative results than the experimental  $IC_{50}$  values. This may be explained depending on the target protein structure selected. The selected target protein represents an active site of the enzyme. When the interactions between 1BMZ target protein and ZnPc (**2**) are examined, the oxygen atom formed a hydrogen bond with the nitrogen of GLN92. The oxygen, nitrogen and metal atom contain polar interaction with the Thr307 and GLU417 amino acid residues. The carbon atoms exhibit hydrophobic interactions with TRP5, PHE91, ALA121, LEU131, ALA135 and  $\pi$ – $\pi$  interactions with TRP5, HIS67 and HIS200. In interactions of InPc (**3**) with 1BMZ, the oxygen and nitrogen atoms formed H-bonds with HIS64 and HIS67. Carbon atoms were performed hydrophobic interactions with ILE60, VAL62, ALA121, LEU141, VAL143, LEU198, TRP209 and  $\pi$ – $\pi$  interactions with HIS64, HIS67, PHE91, HIS200, HIS119. As in the metal complexes with ZnPc (**2**) and InPc (**3**), the between the metal complex and 1BMZ target protein are present H-bond with GLN92, HIS96, THR199, polar interactions with GLN92, HIS94, hydrophobic interactions with HIS67, PHE91, ALA121, LEU131, PRO202, LEU198, VAL143, LEU141, LEU31,  $\pi$ – $\pi$  interactions with HIS200, HIS94, PHE91. There are polar interactions with ASN69, hydrophobic interactions with PRO3, TRP5, VAL62, HIS67, PHE9, HIS94, ALA121, LEU131, PRO202 and  $\pi$ – $\pi$  interactions with PHE91, HIS94 and HIS200 between the complex with Mn metal in MnPc (**6**) and 1BMZ without H-bonds. The absence of H-bonds from the

**Table 1.** The enzyme inhibition results of phthalocyanine complexes (2–6) against human carbonic anhydrase isoenzymes I and II (hCA I and II), acetylcholinesterase (AChE) and butyrylcholinesterase (BChE) enzymes.

Compounds	IC <sub>50</sub> (μM)				K <sub>i</sub> (μM)							
	hCA I	r <sup>2</sup>	hCA II	r <sup>2</sup>	AChE	r <sup>2</sup>	BChE	r <sup>2</sup>	hCA I	hCA II	AChE	BChE
ZnPc (2)	834.62	0.9727	883.62	0.9889	87.63	0.9582	104.83	0.9881	887.25 ± 101.20	905.42 ± 104.35	68.33 ± 9.13	86.25 ± 13.65
InPc (3)	632.04	0.9715	705.83	0.9526	284.92	0.9790	316.54	0.9733	668.37 ± 89.36	786.46 ± 143.54	201.15 ± 15.86	237.54 ± 24.7
CoPc (4)	903.74	0.9543	945.88	0.9830	105.94	0.9384	124.92	0.9824	856.14 ± 85.66	921.14 ± 81.25	78.21 ± 15.78	103.45 ± 10.13
CuPc (5)	506.24	0.9618	635.17	0.9783	114.85	0.9384	150.25	0.9518	588.13 ± 78.25	670.31 ± 90.36	98.36 ± 7.88	125.11 ± 10.05
MnPc (6)	412.46	0.9722	481.04	0.9598	168.37	0.9788	212.43	0.9711	478.13 ± 57.25	525.16 ± 45.87	125.24 ± 17.25	165.38 ± 23.84
AZA <sup>a</sup>	957.44	0.9350	964.78	0.9353	–	–	–	–	851.53 ± 87.33	1060.77 ± 98.0	–	–
TAC <sup>b</sup>	–	–	–	–	305.84	0.9631	383.80	0.9614	–	–	244.37 ± 25.47	290.25 ± 31.26

<sup>a</sup>Acetazolamide (AZA) was used as a control for hCA I and II.

<sup>b</sup>Tacrine (TAC) was used as a control for AChE and BChE enzymes.

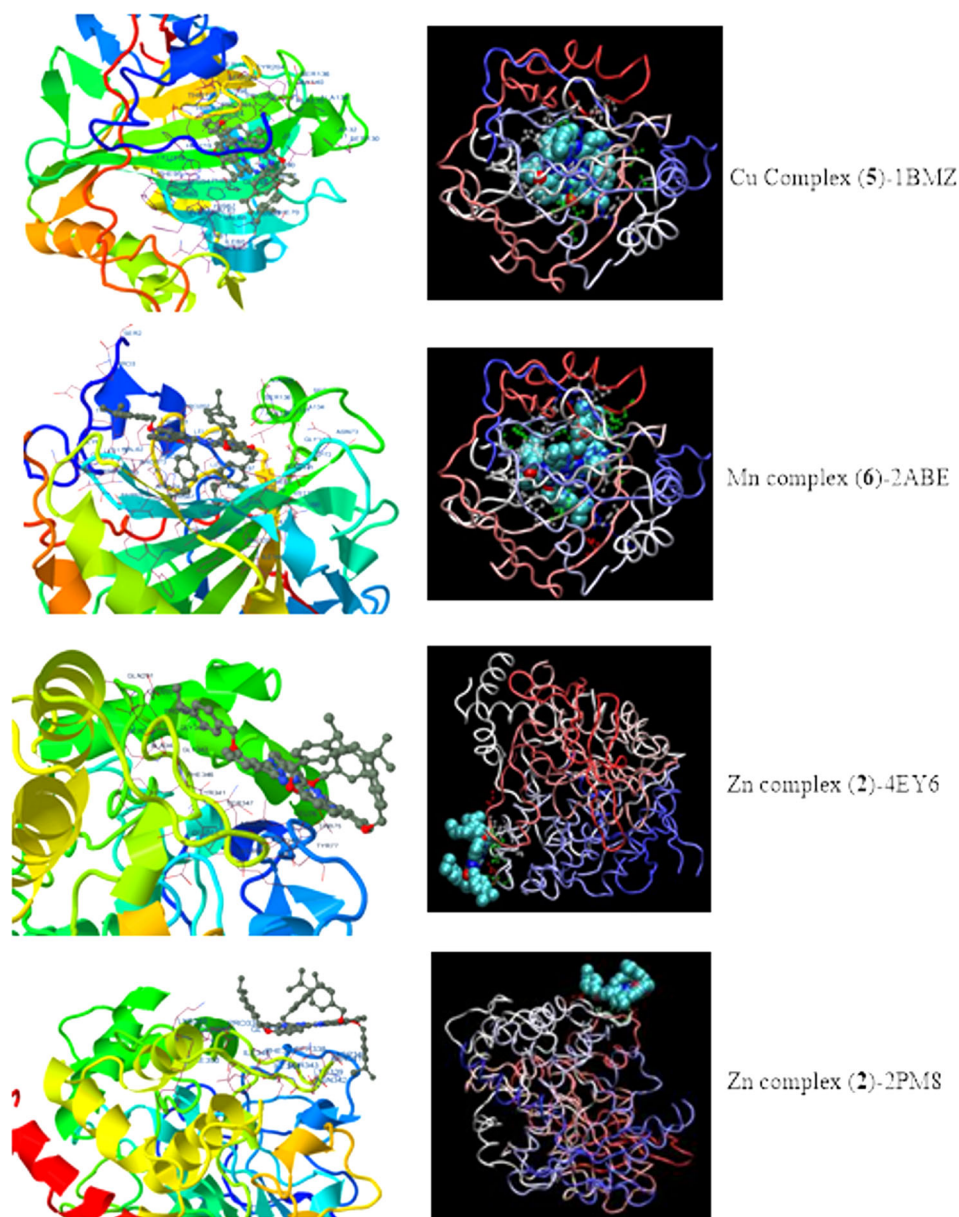
**Figure 4.** Comparative results of IC<sub>50</sub> and K<sub>i</sub> values of all compounds.**Table 2.** The estimated free energy of binding (kcal/mol) between (2–6) complexes and the target protein.

	1BMZ	2ABE	4EY6	2PM8
ZnPc (2)	−3.61	−1.22	−6.04	−5.30
InPc (3)	−4.13	−1.57	−4.78	−4.27
CoPc (4)	−3.05	−1.24	−6.02	−5.25
CuPc (5)	−5.25	−2.12	−5.98	−5.18
MnPc (6)	−2.97	−4.28	−5.17	−5.02

secondary chemical interactions and the low number of interactions with amino acid residues proves why the binding energy is low. When the interactions of 2ABE target protein and ZnPc (2) were examined, nitrogen and oxygen

atoms are made polar interactions with GLN92, HIS94, HIS96, HIS107, GLU117, HIS119, TRP209, HIS300, hydrophobic interactions with carbon atoms and PRO30, ALA65, VAL121, LEU141, VAL143, LEU198 and  $\pi$ - $\pi$  interactions with HIS64, HIS94, HIS107, HIS109, TRP209, HIS300. The interactions of the InPc (3) are polar with GLN92, HIS94, GLU117, THR199, hydrophobic with PRO30, HIS64, ALA65, HIS119, VAL121, VAL121, VAL143, LEU198, VAL245 and  $\pi$ - $\pi$  interactions with PHE131, TRP209, HIS300. The CuPc (5) as well as in the ZnPc (2) had polar interactions with HIS64, HIS119, HIS94, GLU106, HIS107, hydrophobic interactions with TYR7, PRO30, ALA65, VAL121, LEU198 and  $\pi$ - $\pi$  interactions with HIS119, TRP209, HIS300 and HIS96. The complex with the highest interaction energy between 2ABE target protein and complexes is the MnPc (6). This phthalocyanine, unlike others, made H-bond with HIS64, THR200, GLU106, HIS96. Besides, the MnPc (6) is had hydrophobic interaction with ALA65 and cation- $\pi$  interaction with HIS94. The type and number of the interaction of complexes with 4EY6 and 2PM8 target proteins differ from the type and number of the interaction of other proteins. This may be due to the unsymmetrical molecular structures of proteins. Between 4EY6 and ZnPc (2), there is H-bond with SER347, ASP349 and hydrophobic interaction with LEU76. InPc (3) with the mentioned target protein exhibits polar interactions with SER347, ASP349, ASN350, hydrophobic with LEU76, TYR77, PRO344, LEU353,  $\pi$ - $\pi$  interactions with TYR77. When the CoPc (4) is examined, it is seen that it makes H-bond with ASP349. Also, carbon atoms are in polar interaction with LEU79 and PRO344 and  $\pi$ - $\pi$  interaction with TYR77. In the MnPc (6), the nitrogen atoms and the carbon atoms with ASP349 interact with H-bond and polar, respectively. Also, the carbon atoms are done hydrophobic and  $\pi$ - $\pi$  interaction with PRO78 and TYR77, respectively. Finally, the H-bond between the ZnPc (2) with the 2PM8 target is formed with SER338 and polar interactions are performed with SER338 and ASN341. In the InPc (3), nitrogen, oxygen and indium metal only interact polarly with SER 388. In the CoPc (4) and CuPc (5) complexes, oxygen and nitrogen atoms formed H-bonds with SER338, respectively. Similarly, in these metals, carbon atoms exhibit hydrophobic interaction with ILE344 and PRO335, respectively. In the MnPc (6), nitrogen atom produced the polar interaction with SER338 amino acid residue and the hydrophobic interaction with the carbon atom and PRO335.

The molecular structures of the investigated complexes are similar. However, partial charges of the terminal atoms



**Figure 5.** Docking pose of the most active inhibitor of each target protein.

vary depending on the gravitational force of the bond electrons of the central atom and exhibit different inhibitory activities and different secondary chemical interactions. Molecular simulation results can provide important information for the further development of enzyme inhibitors. While the interaction energy of CuPc (**5**) and MnPc (**6**) complexes with the 1BMZ and 2ABE target proteins was the highest, the interaction energy of 4EY6 and 2PM8 proteins with ZnPc (**2**) was the highest. According to the results given in Table 2, the mode of interaction with the most active inhibitor of each target protein is given in Figure 5.

## Discussion

In this study, a series of new soluble metallophthalocyanine complexes synthesized and investigated enzyme inhibition properties. Synthesized complexes showed good inhibition effects against BChE, AChE, hCA I and hCA II enzymes.

Especially, zinc and manganese metallophthalocyanine complexes (**2** and **6**) can be good candidate drugs, the same as AChE, BChE and CA inhibitor complexes, respectively. In this manner, these novel complexes can use for therapy of some diseases like epilepsy, mountain sickness, AD, gastric and duodenal ulcers, glaucoma, osteoporosis or neurological disorders. The binding interaction of these active compounds with hCA isoenzymes I and II (hCA I and II), AChE and BChE enzymes was supported in detail using molecular insertion.

## Author contributions

Specific author contributions are as follow: Designing the study: Emre Güzel, Ümit M. Koçyiğit, Parham Taslimi, İlhami Gülçin; Collecting and analyzing the data: Emre Güzel, Ümit M. Koçyiğit; Drafting the manuscript: Emre Güzel, Ümit M. Koçyiğit; Visualization and revising: İlkay Şişman, Mehmet



Nebioğlu, Barış S. Arslan, Emre Güzel; Theoretical calculations: Sultan Erkan; Supervision: Emre Güzel, Ümit M. Koçyiğit.

## Disclosure statement

The authors declare that there is no conflict of interest.

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