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Substituents modulate biphenyl penetration into lipid membranes

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Abstract

Electrochemical impedance techniques and fluorescence spectroscopic methods have been applied to the study of the interaction of ortho (o)-, meta (m)- and para (p)-Cl-, o-, m- and p-HO-, p-H$_3$CO-, p-H$_3$C-, p-NC- and p-O$_3$S- substituted biphenyls (BPs) with Hg supported dioleoyl phosphatidylcholine (DOPC) monolayers and DOPC vesicles. Non-planar o-substituted BPs exhibit the weakest interactions whereas planar p-substituted BPs interact to the greatest extent with the DOPC layers. The substituted BP/DOPC monolayer and bilayer interaction depends on the effect of the substituent on the aromatic electron density, which is related to the substituents’ mesomeric Hammett constants. Substituted BPs with increased ring electron density do not increase the DOPC monolayer thickness on Hg and penetrate the DOPC vesicle membranes to the greatest extent. Substituted BPs with lower ring electron density can cause an increase in the monolayer’s thickness on Hg depending on their location and they remain in the interfacial and superficial layer of the free standing DOPC membranes. Quantum mechanical calculations correlate the binding energy between the substituted BP rings and methyl acetate, as a model for the –CH$_2$-(CO)O-CH$_2$- fragment of a DOPC molecule, with the location of BPs within the DOPC monolayer.

Keywords: Electrochemical impedance, Fluorescence quenching, Phospholipid monolayers/bilayers, Monosubstituted biphenyls, Hammett constants.
1. Introduction

Biphenyl (BP) and its derivatives are well known for their thermal stability, electrical insulation and resistance to redox processes and have been widely used in the past in transformers and capacitors as dielectric fluids [1,2]. Moreover, these substance have also been employed in the preparation of pesticides, optical brighteners [3], and as fungicides in waxing many fruits. However substituted BPs are established environmental toxins [4]. Chlorobiphenyls (Cl-BPs) have been a significant issue for their toxicity, bioaccumulation and environmental persistence because of their stability [5]. BPs and Cl-BPs are metabolised to hydroxy biphenyls (HO-BPs) via formation of the arene oxide intermediate [6] with para (p)-hydroxybiphenyl (p-HO-BP) as a major product [5,7,8]. In addition to the monohydroxy metabolites, dihydroxy products can also be produced as a result of hydroxylation of BP. The hydroxy metabolites of Cl-BPs have been shown to be retained in the blood of many animals and humans [9-12]. The structures of these metabolites resemble thyroid hormones in possessing two aromatic rings and are reported as endocrine disrupters because they compete with thyroxin for its active site [13,14].

In ortho (o-) substituted BPs, the two phenyl rings have been reported to exhibit a twisted conformation with an increase in the torsional angle (\(\Pi\)) between the phenyl rings from 42.5° to 63.2° depending on the size of the substituent (Br>Cl>F). This can be explained by the destabilization effect based on the hydrogen substituent repulsion in the o- position. Changes in the molecular structure caused by substitution at the p- position does not affect the structure of BP derivatives compared to the unsubstituted BP in both ground and excited states [15]. The value of \(\Pi\) for m- and p- substituted BPs is around 42-45° but for o-substituted BPs, \(\Pi\) is around 63°. m-substituted BPs have a higher barrier height for rotation between 0°-90° than p- substituted BPs which is responsible for slightly higher values of \(\Pi\) [16]. Hence coplanarity of the two phenyl rings
decreases in the following order depending on the position of substitution: BP \approx p\text{-substituted BP} > m\text{-substituted BP} > o\text{-substituted BP}. There have been many studies on the potential toxicity of substituted BPs in vivo [2,17,18] but very little work has been carried out on the evaluation of the molecular characteristics responsible for their toxicity and the mechanism of their interaction at a cellular and plasma membrane level. According to some studies, non-planar o-substituted BPs increased the cell membrane leakiness and decreased the membrane integrity compared to the planar BPs [17-19]. While other studies have shown the non-planar o-substituted molecules to be less active compared to the planar p-substituted molecules because of the steric hindrance of the o-substituted species which influences their penetration into the phospholipid membrane [20-22]. Indeed reference [22] used electrochemical methods in a preliminary study of the relation between the biphenyl's structure and their interaction with lipid layers however no understanding was developed concerning a systematic structure-activity relationship.

The present study is an attempt to widen and deepen the work begun in reference [22] by extending the electrochemical methods and supporting them with additional techniques. An important objective of this work was to obtain an understanding of the fundamental molecular properties of aromatic and associated molecular species which enhance their interaction with phospholipid membranes as a model for biological membranes. Of interest in particular was how the substituted biphenyl affected the membrane structure, function and biomolecular organization and how this related to membrane surface interactions with the biphenyl. The final location of the substituted biphenyl in the membrane following interaction was also sought. An understanding of the factors promoting aromatic group interaction with phospholipid membranes is extremely relevant to biological membrane structure and function. A significant reason for this is that the conformation of membrane proteins and peptides is strongly affected by their aromatic residue interaction with the phospholipid skeleton [23-25].
Hg supported phospholipid monolayers and free standing bilayer vesicles have been used to investigate: (1) structure-activity relationships and (2) mechanism of interaction of BP derivatives, with phospholipid layers, using electrochemical and fluorescence spectroscopic techniques. The rationale for employing these techniques was to obtain a generic and consistent understanding of the interaction mechanism. DOPC was used as the test phospholipid in these studies. The model system of DOPC monolayer on Hg is a well established membrane model which has been well characterized internationally [26-35]. Moreover it has been shown to be particularly stable and reproducible in structure. Although Hg supports monolayers of palmitoyloleyl PC (POPC) and POPC/POPE mixtures, it was felt appropriate to carry out this study with DOPC on Hg in the first instance. The particular defect-free nature and fluidity of DOPC monolayers/bilayers enabled very precise experiments to be carried out so that the factors promoting the interaction of aromatic compounds with phospholipid membranes can be better understood. Later studies would use more complex and more relevant model membrane systems.

Monolayers of phospholipids on the Hg surface act as a sensor element for aromatic compounds and many other species, including biomembrane-active peptides and nanoparticle dispersions [22, 26-35]. At potentials around -0.4 V which is close to the position of zero charge (PZC) of Hg, the Hg supported DOPC monolayer is completely intact and impermeable to inorganic ions. At more negative potentials, capacitance peaks appear due to underlying field induced phase transitions occurring in the monolayer [36-39]. Alterations in the nature of these capacitance peaks are very sensitive to any changes in the monolayer structure. The capacitance peak occurring at around -0.94 V (capacitance peak-1) is associated with electrolyte penetration of the monolayer. The capacitance peak at around -1.0 V (capacitance peak-2) represents a phase transition involving a nucleation and growth process that results in the formation of a bilayer in equilibrium with electrolyte on the Hg surface. The described profile is specific to the capacitance-potential curves for DOPC monolayers. Different capacitance-potential profiles are obtained with different phospholipid
molecules [40]. The capacitance-potential profile is itself a fingerprint for a specified phospholipid monolayer. Interaction of organic compounds with the monolayer significantly influences the capacitance-potential profile of the layer in a selective and systematic manner. It has been established by direct imaging that binding of SiO$_2$ nanoparticles [34] to the DOPC polar groups elicits significant depression of the capacitance current peaks in proportion to the extent of binding. In addition it is clearly stated in other work [28-30] that the decrease in the DOPC capacitance peak-1 may be due to an interaction of the peptide with the DOPC polar heads and to an intercalation of the peptide between the hydrocarbon tails or to a possible combination of the two effects.

Phospholipid vesicles have been used extensively as a biological membrane mimic [41] to study the interactions of membrane active compounds [42,43] with membranes. The BPs have fluorescent properties which allow the use of fluorescence spectroscopy to estimate their fractions which have not penetrated into biomimetic assemblies such as phospholipid vesicles and are thus available for quenching by a quencher (e.g. I$^-$) [44-47]. This is defined as the accessible fraction in this study. These experiments were carried out as a corollary to the electrochemical experiments to substantiate the nature of the BP-DOPC interactions. Table 1 summarizes the BP derivatives used in this study and their physical properties including log P values [48] and Hammetts constants [49].

2. Experimental

2.1. Materials

o-chlorobiphenyl (o-Cl-BP), m-chlorobiphenyl (m-Cl-BP), p-chlorobiphenyl (p-Cl-BP), o-hydroxybiphenyl (o-HO -BP), m-hydroxybiphenyl (m-HO-BP), p-hydroxybiphenyl (p-HO-BP), p-methoxybiphenyl (p-H$_3$CO-BP), p-methylbiphenyl (p-H$_3$C-BP), p-cyanobiphenyl (p-NC-BP) and p-sulphonic acid biphenyl (p-HO$_3$S-BP) were purchased from Sigma Aldrich and their stock solutions (1000 mol dm$^{-3}$) were prepared in acetone and ethanol for electrochemical and fluorescence
studies respectively. p-HO$_3$S-BP exists in ionised form at the solution pH of 7.4 because its pK$_a$ value is near to that of toluene sulfonic acid [50] (~2.8) and is represented as p-O$_3$S-BP throughout the text. The electrolyte, 0.1 mol dm$^{-3}$ KCl was prepared from KCl (Fisher Chemicals Ltd.) calcined at 600 °C in a muffle furnace and dissolved in 18.2 M MilliQ water containing 0.001 mol dm$^{-3}$ phosphate buffer at pH 7.4. A 2.54 mmol dm$^{-3}$ DOPC (Avantilipids) solution in pentane (HPLC grade, Fisher Scientific Chemicals Ltd.) was prepared for electrochemical experiments and a 12.7 mmol dm$^{-3}$ stock DOPC dispersion in 0.1 mol dm$^{-3}$ KCl containing 0.001 mol dm$^{-3}$ phosphate buffer was used for preparation of vesicles in the fluorescence studies. An 8.0 mol dm$^{-3}$ stock solution of KI (Fisher Scientific Ltd.) was prepared with 0.2 mmol dm$^{-3}$ sodium thiosulphate (Na$_2$S$_2$O$_3$) in 18.2 M MilliQ water for use in quenching experiments.

2.2. Electrochemical methods

2.2.1. Apparatus and procedure

The experiments were performed in a standard three electrode electrochemical cell containing; a Ag/AgCl, 3.5 mol dm$^{-3}$ KCl as reference electrode with porous sintered ceramic frit separating the 3.5 mol dm$^{-3}$ KCl solution from the electrolyte solution, a platinum rod as a counter electrode and a DOPC coated MFE as working electrode. The electrochemical apparatus was contained in a Faraday cage. The Hg film electrode (MFE) was prepared using a micro-fabricated Pt electrode (Tyndall National Institute, Ireland) having a 1 mm Pt disc area and a contact pad [24,31,32]. Fabricated electrodes were rinsed with acetone and then MilliQ water, placed in hot Piranha solution for 10-15 minutes, subsequently rinsed with MilliQ water and dried under N$_2$ gas. Hg was deposited using an Eppendorf pipette on the Pt disc electrode to cover the Pt completely and the fabricated electrode inserted into electrochemical cell. A potential excursion from -0.2 to -3.0 V at 40V s$^{-1}$ was used to remove organic material from the surface. The area (~0.009 mm$^2$) of the Hg was determined from the capacitance-potential plot of the clean Hg surface as described previously. This electrode is stable and operational for up to three months [51].
The electrochemical cell, electrodes and all other glass apparatus were washed with Piranha solution to remove organic contamination. Subsequently, the apparatus and the electrodes were rinsed with MilliQ water to remove traces of Piranha residue. A PGSTAT 30 Autolab potentiostat (Ecochemie, Utrecht, Netherlands) interfaced to PowerLab 4/25 signal generator (AD Instruments Ltd.) controlled by Scope software was used to record rapid cyclic voltammetry (rcv) scans. The Autolab systems GPES (general purpose electrochemical studies) and FRA (frequency response analyser) with PGSTAT 30 and controlled with Autolab software were used for the alternating current voltammetry (acv) and impedance measurements respectively. The electrolyte solution was purged with argon gas for about 15-20 minutes followed by a redirection of argon gas as a blanket on top of electrolyte during the experiment. DOPC monolayers were prepared by spreading about 15 $\mu$m$^3$ of 2.54 mmol dm$^{-3}$ DOPC in pentane at the argon-electrolyte interface in the electrochemical cell. A period of 5-10 minutes was required for the pentane to evaporate. The MFE electrode was vertically lowered into the solution slowly through the interface to allow the DOPC monolayer to self-assemble on the Hg surface. Current-potential and impedance scans were recorded using electrochemical techniques. The MFE was cleaned by applying a cycling potential from -0.2 to -3.0 V to remove the DOPC monolayer and a new monolayer was deposited prior to each series of experiments. For electrochemical studies of substituted BP interactions, calculated amounts of their respective stock solution were injected below the DOPC monolayer into the electrolyte followed by gentle stirring of the solution for 5 minutes. Subsequent electrochemical analysis was then performed on the modified lipid coated electrode.

2.2.2. Rapid cyclic voltammetry (rcv)

rcv was used to acquire rapid scans and examine the properties of the monolayer at a scan rate ($\gamma$) of 40 Vs$^{-1}$ with a potential excursion between -0.2 and -1.2 V on the DOPC coated Hg electrode.
The specific capacitance defined as the capacitance per unit area is related to the current (I) from rcv scans by the following equation:

$$C_{sp} = \frac{I}{A}$$  

(1)

Suppression of the capacitance current peaks in the pure electrolyte solution may indicate either imperfections in the monolayer or the presence of contamination in the electrolyte solution. The electrode area, A, was estimated from the capacitance of the uncoated Hg [40].

### 2.2.3. Electrochemical impedance

Alternate current voltammograms (acv) were recorded by measuring the out-of-phase current in response to application of an ac sine wave of 0.005 V amplitude and 75 Hz frequency (f) added to a negatively increasing (ramp rate 5 mVs$^{-1}$) DC potential (E) from -0.2 to -1.2 V. $C_{sp}$ was calculated using the current from the acv scans by the following equation:

$$C_{sp} = \frac{I}{A} \times \frac{\Delta V}{2\pi f}$$  

(2)

where $\frac{1}{2\pi f}$ is the angular frequency and $\Delta V$ is the amplitude of the sine wave. At least three $C_{sp}$-E curves were recorded following fresh DOPC depositions on the MFE for each experiment. Measurements of the real and imaginary impedance of the electrode system using frequencies logarithmically distributed between 65000 to 0.1 Hz, with ac amplitude of 0.005 V and at potential -0.4 V (~PZC of Hg [52]) were carried out both in the absence and presence of BPs in the electrolyte solution. The real and imaginary impedance was normalized at respective frequencies and transformed to the complex capacitance plane as “Cole-Cole” plots using Microsoft EXCEL spreadsheet. Complex plane axes were expressed as $Y'/\omega A$ vs. $Y''/\omega A$ where $Y'/\omega A$ and $Y''/\omega A$ represent real and imaginary normalised admittances respectively divided by electrode area (A) as defined previously [53]. Incidentally, the series combination of a capacitance, C, and a resistance, R, which is simulated by the Hg-supported DOPC monolayer, is such that $Y''/\omega A$ equals $C/(\omega^2R^2C^2+1)A$, while $Y'/\omega A$ equals $\omega RC(Y''/\omega)/A$; moreover, the $Y'/\omega A$ vs. $Y''/\omega A$ plot yields a semicircle of diameter $C/A$ and centre of coordinates $(C/2A)$. This value $C/A$ is frequency
independent and its value is extracted from measurements carried out over the whole frequency range from $10^5$ to 0.1 Hz. This frequency independent capacitance (C/A) approximates to the monolayer capacitance divided by electrode area which is directly related to the monolayer dielectric permittivity and inversely related to the monolayer thickness. In this study the C/A was graphically determined from the complex capacitance plots and expressed as specific frequency independent capacitance ($FIC_{sp}$). In the case of the phospholipid coated Hg electrode, any additional circuit element (second semicircle and "tail") at lower frequencies outside of RC is reflective of ionic penetration [54] and physical modifications of the phospholipid monolayer respectively [53].

2.3. Fluorescence Spectroscopy

2.3.1. Principle

The basic principle in fluorescence spectroscopy involves the emission of light as a result of singlet-singlet electronic relaxation subsequent to absorbing light energy. The fluorescence has a typical life time of nanoseconds. In fluorescence spectroscopy, an incident light beam (usually UV) excites the electrons in the fluorophore molecule to the excited electronic states. These excited molecules then lose vibrational energy to return to the lowest vibrational state of the excited electronic state. As a result, the emission spectrum is recorded keeping the excitation wavelength constant and measuring the fluorescent light at different wavelengths. In this study, the substituted BP acted as an effective fluorophore [55]. The penetration of the BP into the vesicle bilayer was followed by recording the extent of quenching of the BP's fluorescence by KI (i.e. I⁻ as active quencher) which itself at lower solution KI concentrations did not penetrate the phospholipid bilayer. BPs which remained in the polar region of the bilayer were accessible to quenching by KI. BPs which penetrated the bilayer have their fluorescence quenched to varying degrees depending on their extent of penetration.

2.3.2. Preparation of liposomes
A DOPC stock dispersion of 12.7 mmol dm\(^{-3}\) was prepared by directly dispersing DOPC in 0.1 mol dm\(^{-3}\) KCl with added 0.001 mol dm\(^{-3}\) phosphate buffer. Unilamellar vesicles containing 3.9% ethanol and 0.249 mmol dm\(^{-3}\) DOPC were prepared by a modification of the solvent exchange procedure [56]. This involved adding 2 cm\(^3\) of the DOPC stock dispersion to 4 cm\(^3\) ethanol in a measuring flask followed by the addition of 96 cm\(^3\) of 0.1 mol dm\(^{-3}\) KCl with added 0.001 mol dm\(^{-3}\) phosphate buffer gently along the wall of flask with constant swirling. The resulting vesicles formed a unidisperse dispersion of ~220 nm diameter unilamellar vesicles as characterised by dynamic light scattering (DLS) and confocal microscopy (not shown) and were found to be stable for up to 24 hours (Figure 1(a)). No attempt was made to remove the 3.9% ethanol from the dispersion since the ethanol facilitated the stability of the added biphenyl in solution for the subsequent fluorescence measurements. It has been shown that 4% ethanol does not significantly influence dimyristoyl PC (DMPC) conformation in vesicle dispersions [57] and shows no significant interaction with DOPC monolayers supported on Hg (data not shown).

2.3.3. Instrumentation and procedure

A FluoroMax-3 Fluorimeter from Horiba Scientific with xenon lamp was used to obtain fluorescence spectra. Fluorescence spectra of DOPC vesicles at the excitation wavelengths specific to different BPs were recorded as controls. Substituted BP solutions were added to the DOPC vesicle dispersions to a concentration of 1 mol dm\(^{-3}\) with an incubation period of 15 minutes before measurement. KI was used as a quencher. Secondary stock solutions of different concentrations of KI (1.0-7.0 mol dm\(^{-3}\)) were prepared from the primary stock solution of 8.0 mol dm\(^{-3}\) KI and 0.2 mmol dm\(^{-3}\) Na\(_2\)S\(_2\)O\(_3\) in 18.2 M\(^{\circ}\) MilliQ water. Na\(_2\)S\(_2\)O\(_3\) was used as a reducing agent to prevent the oxidation of iodide to iodine. 0.15 cm\(^3\) portions of the KI secondary stock solutions were added to the 3 cm\(^3\) of vesicle dispersion containing substituted BPs in the cuvette to give 0.047 to 0.333 mole dm\(^{-3}\) KI concentrations and fluorescence spectra were recorded at appropriate excitation and emission wavelengths. The cuvette was dried from inside using
compressed air before taking all the measurements to eliminate the error from small dilutions. Maximum fluorescence intensities were measured from the fluorescence spectra. All fluorescence spectra were corrected for the fluorescence of a vesicle dispersion with no biphenyl and no quencher. An example of a corrected fluorescence spectrum is shown in Figure 1(b). Fluorescence data were plotted versus the inverse of the KI concentration according to the modified Stern-Volmer equation [44].

\[
\frac{F^0}{\Delta F} = \left( \frac{1}{f_a K_{c_{KI}}} \right) + \frac{1}{f_a} \quad (3)
\]

Where \( F^0 \) is the intensity of emitted light in the absence of quencher and \( \Delta F \) is the difference in the intensities of emitted light in the absence and presence of quencher respectively, \( K \) is the Stern-Volmer quenching constant, \( c_{KI} \) is the KI concentration and \( f_a \) is the accessible fraction. The accessible fraction is defined as the fraction (expressed in text as %) of the fluorophore present in the polar region of the DOPC bilayer and hence available for quenching. The accessible fraction therefore reports on the location of the BP fluorophore in the membrane. The accessible fractions \( (f_a) \) for the quenching of the BP derivative fluorescence were calculated from the intercept values (intercept = \( 1/f_a \)) with an intercept value of 1.0 indicating 100% accessible fraction of the BP derivatives to quenching by I\(^-\). The slope of these plots gives the effective quenching constant (K) for accessible fluorophores which is similar to the binding constant for the quencher-acceptor systems [58] and represents the quenching efficiency of the BP derivatives by I\(^-\). It is not a significant parameter in terms of their interaction with the membrane. 0.04 mmol dm\(^-3\) I\(^-\) has been shown to interact with the polar but not penetrate the apolar region in Hg supported DOPC monolayers [54] and is known to associate with the choline group of DOPC [59]. Indeed it was shown from I\(^-\) quenching of Laurdan (6-dodecanoyl-2-dimethylaminonaphthalene) in the DOPC phospholipid bilayer that I\(^-\), from solution concentrations above 0.15 mol dm\(^-3\), penetrates down to the glycerol level of the phospholipid bilayer [59].
In this study, all experiments were carried out under the same conditions; thus it is unlikely that the
I/DOPC interaction will affect the relative BP penetration into the membrane. This assumption is
supported by the characteristic Stern-Volmer plot (Figure 1(c)) where the slope of the plot is linear
indicating that the increase in I concentration is not affecting the BP accessible fraction and hence
BP penetration. However, at higher solution concentrations of I, a discrete increase in the slope of
the modified Stern-Volmer plots was observed. This increase in slope is due to I penetrating
the apolar region of the monolayer and quenching penetrated BP. This data is therefore excluded
from the plot used to estimate the intercept. All the BP derivatives were found to be stable on
exposure to light (constant intensity of re-emitted light on repetitive exposure to light) except o-Cl-
BP and were studied using the method described above. For o-Cl-BP, the spectra were recorded in
kinetic acquisition mode with an interval of 0.1 s and 6 s total time, at an excitation wavelength of
260 nm and an emission wavelength of 316 nm to eliminate the effects from photo-degradation.
The intercept from the linear regression of these intensity values versus time yielded the intensity of
re-emitted light at zero time (where there is no photo degradation) and was used to plot the
modified Stern-Volmer plot for I quenching of o-Cl-BP.

3. Results and discussion
The effect of the interaction of the substituted mono HO-BPs with the DOPC monolayer on the
capacitance-potential plot is shown in Figure 2(a). Suppression of the peak-1 capacitance with an
increase in solution concentration of o-, m- and p-HO- BPs is clear and the extent of suppression
follows the order of o-<m-<p- HO-BP/DOPC interaction (Figure 2(b)). This suppression of the
capacitance peak is directly related to the extent of interactions of these BPs with the DOPC
monolayer [22,27] indicating that the p-HO-BP showed the strongest interactions with the DOPC
monolayer compared to that of m- and o-HO-BP. Notably the potentials (-E) characterising
capacitance peaks-1 and -2 do not markedly change on penetration of the HO-BP into the
monolayer. The accessible HO-BP fraction for I quenching in the presence of vesicles prepared
from a 0.249 mmol dm\(^{-3}\) DOPC dispersion indicates that this fraction is lowest for p-HO-BP and increases in the following order: p-HO-BP (17.21 \%) < m-HO-BP (29.03 \%) < o-HO-BP (90.63 \%) which indicates the penetration extent of HO-BP into the apolar core is in the order: p-HO-BP (82.79 \%) > m-HO-BP (70.97\%) > o-HO-BP (9.37\%). This order arises from: (i) the symmetrically linear and planar structure of the p-isomer enabling its ready penetration into the apolar membrane core; (ii) the lack of two-fold symmetry of the m-isomer rendering it less able to penetrate the phospholipid membrane; and, (iii) the twisted geometry of the o-isomer arising from the increased torsion angle between the two phenyl rings further impeding its penetration.

The capacitance peak-1 height correlates with the accessible fraction of HO-BPs for quenching (Figure 2(c)) indicating that the fluorescence results are commensurate with and support the electrochemical results and together relate qualitatively to the interaction of the HO-BP with the DOPC layer. This suggests that the HO-BPs behave in a comparable manner at the Hg supported monolayer/electrolyte interface and at the vesicle/electrolyte interface. Figure 2(d) shows the impedance data represented as a complex capacitance plot of a DOPC coated Hg electrode in the presence of o-, m- and p-HO-BPs. In this plot the interaction of these species with the DOPC monolayer is represented by the presence of a low frequency "tail" at the right hand side of the plot. The frequencies defining this tail are lower than ~22 radians s\(^{-1}\) and are usually representative of imperfections in the layer and species adsorbed within the DOPC [53]. It is interesting that the monolayer capacitance does not decrease following the interaction of DOPC with the HO-BP isomers and the interaction of p-HO-BP with DOPC and its penetration introduces an additional very small irregular semicircular element defined by frequencies 1900 to 13 radians s\(^{-1}\) between the RC element and the "tail". The inset to Figure 2(d) shows this intermediate circuit element more clearly. Current work investigating the interaction of small molecular weight organic compounds with DOPC layers show that phenolic compounds tend to have a disruptive effect on DOPC monolayers compared to non-polar aromatics. This intermediate circuit element could relate to the
disruptive effect of the p-HO-BP leading to ionic penetration within the polar head region [54]. Further more focused work would need to be carried out to establish this.

Interaction of Cl-BPs with DOPC monolayers on Hg had a significant effect on the capacitance-potential plot (see Figure 3 (a)). Suppression of the peak-1 capacitance with an increase in solution concentration of o-, m- and p-Cl-BPs is clear (Figure 3(b)) and following interaction of the Cl-BPs with the DOPC monolayer, the capacitance peak suppression was more pronounced compared to that following HO-BP/DOPC interactions (Figure 3(a) and (b) compared to Figures 2(a) and (b)). This observation indicates that Cl-BP interacts more strongly than HO-BP with the DOPC polar heads. This shows a deeper penetration of HO-BP than Cl-BP in the DOPC layer. In addition, capacitance peak-1 and more extensively peak-2 (Figures 3 (a)) were shifted to more negative potentials in the presence of m- and p-Cl-BPs. The accessible Cl-BP fraction for I\(^-\) quenching in the presence of vesicles prepared from a 0.249 mmol dm\(^{-3}\) DOPC dispersion showed that this fraction increased in the following order: p-Cl-BP (76.47 %) < m-Cl-BP (93.14 %) < o-Cl-BP (96.08 %). This indicates that the extent of penetration of Cl-BP into the monolayer apolar core is in the order: p-Cl-BP (23.53 %) > m-Cl-BP (6.86 %) > o-Cl-BP (3.92%). The isomers of Cl-BPs exhibited a similar order of incorporation/penetration into the bilayer to that of HO-BPs but the extent of the penetration is decreased which is commensurate with the capacitance peak suppression results above. Cl-BPs have a comparable 3D structure to their HO- analogues but there is an essential difference in distribution of electron density among the ring atoms in these substituted BPs. It is well established that the HO- group acts as an electron donor but the Cl- group acts as an electron acceptor when attached to aromatic rings. This is described by Hammetts substitution constant (\(\xi\)) [49] (see Table 1) of the functional group attached to an aromatic ring where negative and positive values indicates the substituent is electron donating and withdrawing respectively relative to H-. Accordingly HO-BP has an increased ring electron density than Cl-BP and penetrates the DOPC bilayer to a greater extent than Cl-BP. In Figure 3(c) it can be seen that the capacitance peak-1
height from the capacitance-potential plot of a DOPC monolayer on Hg correlates with the Cl-BP accessible fraction for quenching in DOPC vesicle dispersions. This indicates that the fluorescence results are commensurate with and support the electrochemical results together relating to the extent of interaction of the Cl-BP with the DOPC layer.

Figure 4(a) displays the impedance plots of the DOPC coated Hg electrode in the presence of Cl-BPs in the electrolyte solution. It is clear from the presence of the low frequency "tail" on the right hand side of the plot, that the Cl-BPs interact with the DOPC monolayer. Figure 4(b) displays the FIC$_{sp}$ versus Cl-BP concentration plots and it can be seen that the monolayer capacitance decreases in the presence of Cl-BPs in solution. In fact the following order of FIC$_{sp}$ decrease subsequent to interaction of the following Cl-BPs with the DOPC monolayer is: p-Cl-BP > m-Cl-BP > o-Cl-BP commensurate with the degree of interaction of these compounds with the DOPC monolayer. If it is assumed that the relative permittivity of Cl-BPs (~5-6 [60,61]) is about two times higher than that of the hydrocarbon layer of the monolayer, a decrease in monolayer capacitance clearly indicates the formation of a thicker monolayer. Indeed, the increase in monolayer thickness must be the overriding factor leading to a decrease in FIC$_{sp}$. Furthermore, the negative shift of the capacitance peaks associated with Cl-BP/DOPC interaction can be related to the increasing thickness of the low dielectric region. This increases the applied potential required for the instigation of the phase transition underlying the capacitance peak (Figure 3 (a)) enabling the phase transition to take place at the critical field value [62]. A further explanation for the peak shift is that the polarizability of the aromatic rings (assumedly located in the polar group region) lessens the steepness of the potential drop across the layer [62]. This leads to a greater applied potential to be used to initiate the phase transitions.

Figure 5(a) plots the BP substituent $\int_1$ values against the substituted BP accessible fraction in the membrane for all substituted BP/DOPC membrane interactions. It is significant that p-H$_3$C-BP and
p-O₃S-BP are significant outliers on this plot. The \( t \) values employed have been calculated for substituted benzenes and include both long range mesomeric (\( t_m \)) and short range inductive (\( t_r - t_m \)) effects of substitution [63]. The benzene molecule is small and the inductive effect is significant in changing its properties but in the case of BP and other larger molecules, the inductive effect does not extend to the unsubstituted ring compared to the mesomeric effect [64]. In order to estimate the precise and realistic electron density on the unsubstituted BP ring, mesomeric Hammetts constants \( t_m \) [49] were used. Figure 5(b) shows the correlation between the substituted BP accessible fraction and \( t_m \) of the substituent. The substituent value of \( t_m \) generally showed a good correlation with the accessible fraction of BPs subject to \( \Gamma \) quenching. p-O₃S-BP is a marginal outlier to this trend but it is a negatively charged molecule with a large polar O₃S- group which will additionally favour its preference for the polar region. Figure 6 shows a plot of the FIC\( _{sp} \) extracted from the complex capacitance plots of BP penetrated DOPC on Hg versus the accessible fraction of substituted BPs subject to \( \Gamma \) quenching in the presence of DOPC vesicles. The V shaped plot shows a minimum FIC\( _{sp} \) representing a maximum thickness of the DOPC corresponding to the intermediate value of the accessible fraction. An outlier to the trend is the point representing the o-HO-BP/DOPC interaction which could be due to a variety of factors and requires further investigation. The accessible fraction is in inverse proportion to the penetration depth of the substituted BP in the bilayer membrane and, assuming that the penetration of BPs in the monolayer and bilayer are the same or similar, the mid-location of the Cl-BP in the monolayer is related to a maximum thickness of the monolayer. By the same token, when the substituted BP is located outside this region either nearer to the surface of the monolayer or deeper in the apolar core, the increase in thickness is not so pronounced.

4. Model

The experimental observations support a model whereby NC-BP is located within the DOPC polar head region, Cl-BP is located at the DOPC polar-apolar interface and HO-BP is located towards the...
DOPC apolar core. The model is commensurate with the existence of CH-□ associations in which a significant contribution comes from dispersion interactions [65] favoured by the increased electron density or polarizability of the aromatic ring [65,66]. This is facilitated by electron donating substituents (eg for HO-, $\int_r \int_m = -0.37$ and $\int_m = -0.735$) on the biphenyl aromatic rings which will enable deeper insertion of the biphenyl rings into the apolar core. The model is also substantiated by the existence of H-bonding between the substituted BPs and the DOPC molecules. In the case of aromatic rings acting as electron donors, this interaction favours CH-□ association where the acyl chain $>\text{CH}$- group is the electron acceptor [67,68]. On the other hand the aromatic ring can act as acceptor in the presence of electron withdrawing substituents [69]. A particular example of this is pentafluorobenzene which forms H-bonds due to the electron withdrawing effect of the F- substituents [70]. These effects would favour interactions between electron withdrawing (eg for NC-, $\int_r \int_m = 0.66$ and $\int_m = 0.15$ [49]) substituted biphenyl aromatic rings and the DOPC polar group region.

The Cl- substituent of p-Cl-BP has a strong inductive electron accepting effect ($\int_r \int_m = 0.42$ [49]) and a weak mesomeric electron donating effect ($\int_m = -0.19$) which renders the substituted BP ring relatively more electron deficient than the unsubstituted ring. This is because the inductive effect only operates on the substituted ring [64]. This difference in electron density of the substituted and unsubstituted rings respectively leads to a directional location of p-Cl-BP in the DOPC layer. The unsubstituted ring remains in the apolar region through CH-□ interactions whereas the substituted ring tends towards the apolar/polar interface. Such a structure can be stabilized by a halogen bond between Cl- of p-Cl-BP and the phosphate group oxygen [71,72] of PC. The head groups of DOPC in the liquid phase of the monolayer are oriented parallel to the layer plane [73,74] whereas the apolar tails of DOPC molecule are in a plane normal to the monolayer. As a result, the positioning of the rigid p-Cl-BP in the right-angled interfacial region will increase the angle between the polar
groups and the alkyl chains and cause a thickening of the layer as observed from the capacitance decrease in Figure 4(b).

The propensity for the substituted BPs to form H-bonds with the polar DOPC moieties has been explored. This is a semi-quantitative exercise which has been carried out to substantiate the conclusions from the preceding experiments. It is acknowledged that a more complete treatment, outside the scope of this paper, would take account of the water molecular environment. H-bonding energies of substituted BPs with the glycerol-dicarboxylic ester moiety of DOPC have been estimated. Methyl acetate \((\text{CH}_3-(\text{CO})\text{O-CH}_3)\) was chosen as a molecule to model the glycerol-dicarboxylic ester grouping of DOPC and its binding with the substituted BPs. The choice of methyl acetate is justified since its structure is analogous to each of the two parts of the glycerol-dicarboxylic ester of DOPC namely R-(CO)O-CH< and R-(CO)O-CH< where R- is the oleoyl chain. It is to be noted that the binding energy between p-X-BP (where X is the substituent) to methyl acetate is half that of the binding energy of two p-X-BP molecules with each of the two ester groups of one DOPC molecule. The critical H-bonding moieties in both the methyl acetate molecule and the glycerol-dicarboxylic ester fragment are C-O-C and >C=O. Binding energies were also estimated between the >C=O (methyl acetate) and HO- (p-HO-BP) groups and between the -PO2- (DOPC) and Cl- (p-Cl-BP) groups to substantiate the conclusions. All energies were calculated using the program package Jaguar from Schrodinger, Inc. (Maestro Version 10.2.011, MMshare Version 3.0.011, Release 2015-2, Platform Windows-x64) and are displayed in Tables 2 and 3. The binding energy between two water molecules was calculated with the same program in order to compare it with the other estimated binding energies.

Table 2 shows complexes of unsubstituted and substituted BPs with CH3-(CO)O-CH3. Complex A exhibits two hydrogen bonds between the m- and p- hydrogens of the unsubstituted BP ring and the C-O-C and >C=O oxygens respectively of CH3-(CO)O-CH3. Complex B exhibits: (a) two
hydrogen bonds between the o- hydrogens of the substituted and unsubstituted BP rings and the >C=O oxygen of CH$_3$-(CO)O-CH$_3$ and, (b) a hydrogen bond between the m-hydrogen of the substituted BP and the C-O-C oxygen of CH$_3$-(CO)O-. Table 2 lists the binding energies calculated for complexes between the BPs and CH$_3$-(CO)O-CH$_3$. Complexes A and B with unsubstituted BP show similar binding energies. The Complex B of p-Cl-BP shows a higher binding energy of 2.27 compared to 1.8 kcal mol$^{-1}$ for Complex A. The preferable formation of Complex B would result in the positioning of the unsubstituted phenyl ring inside the apolar region of DOPC layer and the substituted phenyl ring in the interfacial layer of DOPC. The bonding of Cl- with the DOPC -PO$_2^-$ grouping with an energy of 2.08 kcal mol$^{-1}$ (see Table 3) would re-enforce this structure. Binding energies of p-HO-BP with CH$_3$-(CO)O-CH$_3$ are 1.31 and 1.03 kcal mol$^{-1}$ for Complexes A and B respectively. This is more than offset by the relatively high binding energy of HO- with the methyl acetate, >C=O group, of 7.43 kcal mol$^{-1}$ (see Table 3) which has a higher energy than the HOH-HOH bond energy (5.05 kcal mol$^{-1}$). This favours the positioning of both p-HO-BP rings within the apolar core of DOPC. The binding of p-NC-BP rings with CH$_3$-(CO)O-CH$_3$ favours Complex A (2.86 kcal mol$^{-1}$). The relatively high binding energy in this case is enabled by the electron withdrawing NC- group. The bonding of the unsubstituted ring and the high polarity of the NC- group [75] will promote the positioning of the p-NC-BP substituted ring within the polar head region of the DOPC layer.

5. Conclusions
The interaction of monosubstituted BPs with DOPC bilayers and monolayers is strongly dependent on the structural and electronic configuration of the molecules and specifically:-.

1. p-substituted BPs compared to o- and m-substituted BPs exhibit stronger interactions with DOPC bilayers and monolayers because of retained co-planarity of the two aromatic rings irrespective of the nature of substituent.
2. BPs with electron donor substituents penetrate the apolar core of the bilayer whereas BPs with electron acceptor substituents are incorporated in the polar group region of the DOPC bilayer. BPs located at the interface between the polar and apolar region increase the DOPC monolayer thickness.

3. Experimental results and binding energy calculations indicate that the three structural and electronic features influencing the location of substituted BPs in DOPC bilayers and monolayers are respectively: (i) The position of the substituent, (ii) the electron density or polarizability of the BP rings and, (iii) the hydrogen bonding capability of the aromatic rings.

4. Both electrochemical and fluorescence studies exhibit a consistent pattern of interactions between substituted BPs and DOPC bilayers and monolayers respectively suggesting that the Hg electrode support has an insignificant influence on the interactions of the substituted biphenyl compounds with DOPC.

These findings are directly relevant to the initial objective of the work in that they relate the molecular properties of substituent position and ring polarizability to the extent of interaction of the substituted biphenyl with phospholipid monolayers/bilayers as a model for biological membranes. This has important relevance to biological membrane function not only with respect to the interaction of aromatic pharmaceuticals with biomembranes but also in relation to the conformation of proteins containing aromatic rings within biomembranes. In both cases the biomembrane structure, function and biomolecular organization is strongly affected by the nature of the aromatic ring/biomembrane associations.

6. Acknowledgments

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7. References


[33] F Neville, D Gidalevitz, G Kale and A Nelson, Electrochemical screening of anti-microbial peptide LL-37 interaction with phospholipids, Bioelectrochemistry 70 (2) 205.


Captions

Figure 1.
(a) DLS measurement of prepared vesicle dispersion. (b) Fluorescence of 1 μmol dm⁻³; o-HO-BP at excitation = 295 nm (blue), m-HO-BP at excitation= 290 nm (black) and p-HO-BP at excitation = 288 nm (red), in the presence of 0.249 DOPC mmol dm⁻³ and 0.12 mol dm⁻³ of KI. All fluorescence values corrected for fluorescence in absence of BP and KI. (c) Modified Stern-Volmer plots for I⁻ quenching of 1 μmol dm⁻³ p- (filled circle), m- (triangle) and o- (square) HO-BP in the presence of 0.249 mmol dm⁻³ DOPC vesicles in 0.1 mol dm⁻³ KCl with 0.001 mol dm⁻³ of phosphate.

Figure 2.
(a) Specific capacitance (Csp)-potential (-E) plot of DOPC coated Hg electrode in 0.1 mol dm⁻³ KCl containing 0.001 mol dm⁻³ phosphate buffer as control (black) in the presence of 1.0 mol dm⁻³ o- (blue), m- (green) and p- (red) HO-BP; (b) Peak-1 specific capacitance (from (a)) versus BP concentration (cBP) plots of o- (filled triangle), m- (filled inverted triangle) and p-(open square) HO-BP; (c) % peak-1 capacitance (relative to control from (a)) against accessible fraction from fluorescence measurements in 0.249 mmol dm⁻³ DOPC vesicle dispersion with 1.0 mol dm⁻³ of HO-BP; (d) Complex capacitance plane plots of DOPC coated Hg electrode in 0.1 mol dm⁻³ KCl containing 0.001 mol dm⁻³ phosphate buffer (black) in the presence of 1.0 mol dm⁻³ o- (blue), and p- (red) HO-BP.

Figure 3.
(a) Specific capacitance (Csp)-potential (-E) plot of DOPC coated Hg electrode in 0.1 mol dm⁻³ KCl containing 0.001 mol dm⁻³ phosphate buffer as control (black) in the presence of 1.0 mol dm o- (blue), m- (green) and p- (red) Cl-BP; (b) Peak-1 specific capacitance (from (a)) versus concentration ((cBP) plots of o- (filled triangle), m- (open triangle) and p-(filled square) Cl-BP; (c)
% peak-1 capacitance (relative to control from (a)) against accessible fraction from fluorescence measurements of DOPC vesicle dispersion with 1.0 \( \mu \text{mol dm}^{-3} \) Cl-BP.

**Figure 4.**

(a) Complex capacitance plane plot of representative impedance data of DOPC coated Hg chip electrode in 0.1 mol dm\(^{-3}\) KCl with 0.001 mol dm\(^{-3}\) phosphate buffer (black) containing 1.0 \( \mu \text{mol dm}^{-3} \) of o- (blue) and p- (red) Cl-BP acquired using EIS at -0.4 V applied potential; (b) FIC\(_{sp}\) from (a), versus BP concentration (c\(_{BP}\)) plot of o- (filled triangle), m- (open triangle) and p- (filled square) Cl-BP. Error bars are within the symbol size.

**Figure 5.**

Plot of BP substituent's Hammetts constants: (a) \( \sigma_t \) and, (b) \( \sigma_m \) [35] against BPs accessible fraction to I\(^-\) quenching on the DOPC bilayer with 1 \( \mu \text{mol dm}^{-3} \) BP in solution. Error bars within symbol size.

**Figure 6.**

Plot of FIC\(_{sp}\) (closed triangle) of DOPC coated Hg against accessible fraction to I\(^-\) quenching of substituted BP on the DOPC bilayer with 1 \( \mu \text{mol dm}^{-3} \) BP in solution. Horizontal red dashed line corresponds to FIC\(_{sp}\) of pure DOPC. Error bars are within the symbol size.
Figure 1

(a) Graph showing intensity (%) vs. diameter (nm) with two curves labeled time 0 and time 24hr.

(b) Graph showing fluorescence (c.p.s.) vs. wavelength (nm) with three curves.

(c) Graph showing $F_0/\Delta F$ vs. $(1/c_{KI})$ (mmol$^{-1}$dm$^3$) with data points and error bars.
Figure 2

(a) C_{sp}/\mu F\ cm^{-2} vs. -E/V

(b) C_{sp} of peak-1/\mu F\ cm^{-2} vs. c_{BP}/\mu mol\ dm^{-3}

(c) Relative peak-1 height /% vs. Accessible fraction /%

(d) Y_{\omega A^{-1}} / \mu F\ cm^{-2} vs. Y_{\omega^{-1} A^{-1}} / \mu F\ cm^{-2}
Figure 3

(a) $C_{sp}/\mu F \text{ cm}^{-2}$ vs. $-E/V$ for different compounds.

(b) $C_{sp}$ of Peak-1/µF cm$^{-2}$ vs. $c_{BP}$ / µmol dm$^{-3}$.

(c) Relative peak-1 height (%) vs. Accessible fraction/%

- $o$-Cl-BP
- $m$-Cl-BP
- $p$-Cl-BP
Figure 4
Figure 5
Figure 6

![Graph showing the relationship between FICsp (µF cm⁻²) and the accessible fraction (%) for various groups such as p-HO-, m-HO-, p-H₂CO-, o-HO-, o-Cl-, p-Cl-, p-NC-, and p-O₃⁻S.](image-url)
Table 1.

Log P [48], Hammetts constant (\( \sigma_t \) and \( \sigma_m \)) [49] and torsional angle [16] \((\varphi)\) values.

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Table 2.

Binding energies of BP, Cl-BP, HO-BP and NC-BP molecules with CH$_3$-(CO)O-CH$_3$ (calculated with Schrodinger Jaguar software).

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Table 3.

Binding energies of HO- (p-HO-BP) with CH₃-(CO)O-CH₃, Cl- (Cl-BP) with –PO₂⁻ of DOPC molecule and HOH with HOH; normalised per single bond (calculated with Schrodinger Jaguar software).

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Graphical abstract
Highlights

Substituents modulate biphenyl penetration into lipid membranes

- Substituted biphenyls (BP) interact with DOPC membranes in the order p- > m- > o-
- Substituents modulate the penetration of BPs into DOPC membranes
- BP substituent position and ring polarizability influence BP/membrane interaction
- Monolayer/bilayer experiments give consistent results for BP/membrane interaction