SAXS measurements of azobenzene lipid vesicles reveal buffer-dependent photoswitching and quantitative Z→E isomerisation by X-rays

Abstract: Photoresponsive materials feature properties that can be adjusted by light near-instantaneously, reversibly, and with high spatiotemporal precision. There is considerable interest in maximising the degree of photoswitching, and in measuring this degree during illumination in complex environments. We study the switching of photoresponsive lipid membranes that allow for precise and reversible manipulation of membrane shape, permeability, and fluidity. Though these macroscopic responses are clear, it is unclear how large the changes of trans/cis ratio are, and whether they can be improved. Here, we used small-angle X-ray scattering to measure the thickness of photoswitchable lipid membranes, and we correlate lipid bilayer thickness to trans/cis ratios. This reveals an unexpected dependency of photoswitching ratio upon aqueous phase composition. In buffer with ionic strength, we observe thickness variations twice as large as previously observed. Furthermore, soft X-rays can quantitatively isomerise photolipid membranes to the all-trans state; enabling X-ray-based membrane control. High energy X-rays do not influence the state of the photoswitches, presumably because they deposit less dose in the sample.

Keywords: azobenzene; high-energy X-ray; isomerization; lipid bilayer; photoswitch; SAXS.

1 Introduction

Azobenzene photoswitches are molecular units, which can be switched between trans and cis isomer states at the single-molecule level by light. By embedding photoswitches into polymers or supramolecular assemblies containing many individual photoswitch units, materials with mechanical, electrical or optical properties that depend on the trans/cis ratio of the photoswitch population can be created: i.e., photoresponsive materials where these properties are adjustable by light [1–6]. Supramolecular assemblies can also provide further degrees of complexity, for example the populations of the two isomers may phase separate [7], or a highly ordered assembly process may select for or stabilise one of the isomer states [8].

As a photoresponsive material’s properties depend on the isomer ratio, there is considerable interest in measuring and maximising the degree of switching between mostly-trans and mostly-cis populations [9]. Typically, it is not possible to perform quantitative photoisomerisations to all-cis or to all-trans azobenzene populations, due to absorption overlaps [8, 10–13]. Hence, illuminations lead to mixed photostationary state (PSS) populations, with trans/cis ratios that depend on the absorption coefficients of the two isomers and the quantum yields of their photoisomerisation. However, a rare example of quantitative switching in one direction (to all-trans) was pioneered by Hecht et al. using electrocatalytic pathways that transiently oxidise or reduce the azobenzenes [14–17]. Here we study the switching of photoresponsive lipid membranes assembled from azobenzene-containing
phosphatidylcholine (azo-PC), a synthetic lipid with a light-responsive azobenzene group in one of its two hydrophobic tails. Optical control of membranes constructed from pure azo-PC allows precise and reversible manipulation of many mechanical properties including membrane shape, permeability, fluidity, and domain formation, and influence membrane protein function [4, 18–21]. So far, it remains unclear how large the changes of trans/cis ratio are that cause these property changes; and whether these ratio are close to the theoretical maxima of pure states, or whether they could be substantially improved by tailored conditions and switching stimuli – with associated improvements to switching of biophysical properties [22]. This partly derives from a technical challenge: while it is straightforward to measure trans/cis ratios in molecular solutions by a variety of methods, it is not straightforward to measure them in the pure azo-PC membranes. The reason for doubt is that membranes are dense anisotropic assemblies. These give rise to a variety of optical shifts depending on the detailed nature of intermolecular interaction, which are still under investigation [19, 23]. Furthermore, it is widely demonstrated that the azobenzene photoisomerization yield is dramatically improved by embedding amphiphilic azobenzenes in the hydrophobic environment of a lipid membrane [24]. In this study, the synthetic photolipids themselves form the vesicle. Whether the azo-PC membrane’s dielectric environment will yield the same protection from the aqueous phase needs to be tested by experiments. For example, ultrastructure studies based on X-ray experiments have shown photostimulated membrane thickness changes of ca. 4 Å for ca. 42 Å azo-PC membranes [4]: but it was not known if this 10% change is already maximal, nor was it known what population-level of trans/cis ratios were responsible for this change. Clearly, larger thickness changes would lead to more pronounced biophysically relevant effects.

In this work, we use orthogonal solvent measurements to relate photolipid bilayer thickness to the population-level of trans/cis ratios. Comparison of illumination in chloroform and in water reveals an unexpected sensitivity of the photoswitches inside the hydrophobic part of the lipid membrane to the aqueous buffer conditions outside. In water, we observe a smaller-than-expected photoswitching yield under low ionic strength. Furthermore, by mixing pre-switched monomers from chloroform, we show that membrane variations of minimum 8 Å are experimentally possible when photoswitching efficiency is high. In water with high ionic strength, we observe large thickness variations close to illumination in chloroform. These thickness changes are twice as large as thickness variation observed in previous vesicle photoswitching experiments [4], and much larger than the membrane thickness variations achievable within typical physiologically relevant conditions in natural lipids by change of temperature [25–28].

We also discovered that while hard X-rays do not switch the membranes under study, soft X-rays (8 keV) efficiently and quantitatively isomerise photolipid membranes to the all-trans state within seconds, which we attribute to radical redox reactions following X-ray dose deposition in the medium. This enables soft X-rays to enforce a higher degree of membrane property control than photoswitching alone can achieve, while emphasising the role of high energy X-rays as low dose probes in soft matter experiments.

2 Materials and methods

2.1 Synthesis of azo-PC and reference photoswitch FAzoM

Azo-PC was purchased from Avanti Polar Lipids, Inc. (Alabama, United States). Novel reference photoswitch FAzoM was synthesised by standard reactions (see Supporting Information).

2.2 Synthesis of azo-PC and reference photoswitches for benchmarking

In our hands, standard techniques to measure trans/cis ratios in molecular solutions (H-NMR, HPLC) were not reproducible when applied in situ to lipid membranes (ordered assemblies). Destructive readouts (e.g. adding cosolvents to homogenise membrane/water mixtures before HPLC measurement) were also tested but were also not reproducible, which we attributed to difficulties arising from the surfactant nature of the AzoPC. Finally, we developed a method to relate photolipid membrane trans/cis isomer ratio to the thickness determined by SAXS by using a calibration series for lipid membranes of known trans/cis composition. This series was created by mixing preconditioned all-trans and mostly-cis stocks (see below). The trans/cis ratios in the preconditioned AzoPC stocks were in turn determined by comparison of their UV–Vis spectra with that of the isoelectronic reference photoswitch FAzoM in the same conditions, since the trans/cis ratio of the apolar FAzoM can be reliably quantified by HPLC (see Supporting Information section S1 for details).

2.3 Preconditioning and mixing of azo-PC

Azo-PC was dissolved in chloroform (25 mg/mL) and stored at −20 °C until further use. Azo-PC stocks have been stored in dark for several days to reach the all-trans state. After illumination of molecularly-dissolved azo-PC by UV-A LED (Roschwege Star-UV365-03-00-00, λ = 365 nm, 9 nm FWHM, Conrad Electronic SE, Germany) the photostationary state (PSS(τ)) has a cis fraction of 83%
(see Supporting Notes for further detail). To adjust the cis fraction in the assembly, we mix azo-PC in the all-trans state with azo-PCs with cis fraction of 83%, in appropriate proportions. After mixing, we follow the protocol for vesicle preparation. All preparation steps were made in the dark.

2.4 Small unilamellar vesicle (SUV) preparation

The Azo-PC chloroform stock solution was evaporated under a nitrogen stream and stored under vacuum for 12 h. The resulting dry lipid film was dissolved in cyclohexane, and exposed to a vacuum of $6 \times 10^{-3}$ mbar at a temperature of $-60 \degree C$ yielding a fluffy lipid powder. Immediately after lyophilisation, the azo-PC powder was hydrated with deionized (DI) water (Milli-Q, Reptile Bioscience Ltd., Boston, MA), or with PBS buffer (pH 7.5), or with 1× TE buffer (10 mM Tris, 1 mM EDTA, pH 8), to a final concentration of 30 mg/mL. The suspension was gently vortexed and subjected to several freeze/thaw cycles. Finally, the sample solution was extruded ca. 25 times through a polycarbonate membrane with a pore diameter of 50–80 nm using a Mini Extruder (Avanti Polar Lipids, Inc., Alabama, United States). The small vesicle size promotes unilamellar membranes [29].

2.5 UV-A/blue light illumination

For photoswitching of azo-PC membranes during SAXS, we built a dual UV-A and blue light LED setup shown in the Supporting Information section S2. For UV-A illumination, we focus a high-power LED (Roschwege Star-UV365-03-00-00, $A = 365 \text{ nm}$, 9 nm FWHM, Conrad Electronic SE, Germany) on the sample capillary. The total maximum optical power of 170 mW and a focal spot size of 4 mm$^2$ yield an irradiance of 4.25 W cm$^{-2}$. For blue light illumination, fed in by a dichroic mirror, a high-power LED (Roschwege LSC-B, $A = 465 \text{ nm}$, 18 nm FWHM, Conrad Electronic SE, Germany) was used. The blue light is focused with the same focal spot size and a total maximum optical power of 120 mW, resulting in an irradiance of 3.0 W cm$^{-2}$. The LEDs and the X-ray detector were remote controlled by TTL signals from an Arduino microprocessor (Reichelt electronics GmbH & Co. KG, Germany). For both LEDs, photostationary states are achieved within a few seconds and multiple switching cycles yield identical SAXS intensities.

2.6 SAXS measurements at 17.4 keV

X-ray data from azo-PC SUVs with preconditioned trans/cis ratios were recorded at a Mo-sourced small angle X-ray scattering (SAXS) setup [30]. The Mo anode delivers a beam with an energy of 17.4 keV, a beam size of 1.0 mm$^2$, and a flux of $2 \times 10^9 \text{ cnts} \cdot s^{-1} \text{ mm}^{-2}$. X-ray data were recorded by a Dectris Pilatus 3R Detector with 516 pixels of size 172 × 172 µm$^2$. All in-house SAXS measurements were performed in darkness.

2.7 SAXS measurements at 8 keV

SAXS data from azo-PC SUVs were recorded at the Austrian SAXS beamline at ELETTRA synchrotron using a beam energy of 8 keV [31] and a beam size of $0.5 \times 2.0 \text{ mm}^2$. The sample solution was loaded in 1.5–2 mm diameter quartz glass capillaries by flow-through and placed in our UV-A/blue LED setup. A Pilatus detector from Dectris Ltd., Switzerland with 981 × 1043 pixels of size 172 × 172 µm$^2$ served as detector.

2.8 SAXS measurements at 54 keV

The high energy SAXS data from azo-PC SUVs were recorded at beamline P21.1 at the PETRA III ring at DESY. We measured with a beam energy and size of 54 keV and $1 \times 1 \text{ mm}^2$, respectively. For the high energy SAXS experiments, the sample solution was first switched optically and then loaded in Kapton tubes of 40 mm in length and 2.5 mm in diameter. A Lambda detector (X-Spectrum GmbH, Germany) with 772 × 516 pixels with $55 \times 55 \text{ µm}^2$ pixel size was used.

3 Results and discussion

Azo-PC monomers dissolved in low-dielectric chloroform are driven towards cis-rich photostationary state by UV-A light at 365 nm, and towards trans-rich state by blue light of 465 nm (Figure 1a). The absorption spectra of these mixed states, either in molecular solution or in assemblies, have been reported by us and others [4, 18, 19] and substantial differences between them have been noted [19]. Previously, we have used SAXS to determine the membrane thickness for blue or UV light photostationary states reached in lipid vesicles in DI water [4]. Here, we prepared calibration series of azo-PC vesicles of known trans/cis ratios by mixing stocks of dark-adapted all-trans azo-PC with 83% cis azo-PC (see methods and SI). We use calibrants of 0%, 10%, 19%, 39%, 58% and 83% cis isomers to cover the full range of ratios accessible during membrane photoswitching, with an estimated error margin of ±5% of the given value to account for mixing precision as well as cis fraction uncertainties in the preconditioned samples. The referencing method developed to determine the stock cis percentage is novel in this field. This method can prove generally useful for analysis of systems which lack reliable bulk references, i.e. which cannot be interpreted with help of UV–Vis spectra of monomer solution, see Methods and Supporting Information for details.

The SAXS measurements for these calibrant ratios are summarized in Figure 1b. The X-ray intensity distributions are typical for lipid bilayer samples [32, 33] and are stable over several hours (see Figure S3). The distributions vary strongly with the trans-to-cis ratio, e.g. the intensity dip around $q = 0.05 \text{ Å}^{-1}$ shifts consecutively to higher $q$-values with increasing cis content. This indicates that the membrane thickness decreases with increasing cis isomer fraction. To extract the head-to-head distances $d_{HH}$, we model the SAXS intensities in Figure 1b by an established lipid
bilayer electron density profile, using the software SasView (see Supporting Information section S6) [34]. The results of this analysis are shown in Figure 1c. The dark-adapted state shows a head-to-head-distance of $d_{HH} (0 \pm 5\% \text{cis}) = 40.9 \pm 0.6$ Å. The analysis reveals that the thickness of the azo-PC membrane depends almost linearly on the trans-to-cis ratio over a broad range. Increasing the percentage of cis azo-PC thins the membrane down to $d_{HH} (83 \pm 5\% \text{cis}) = 33.0 \pm 0.5$ Å for 83% cis. Thus, the membrane thickness changes by 8 Å.

In the following measurements, we use this relation between membrane thickness and cis fraction to infer the cis fraction of various photostationary states in response to illumination and buffer conditions (Figure 1). Our previous photoswitching experiments have been performed on vesicles in deionized water (DI water): a common choice for lipids since DI water facilitates unilamellar membrane formation by increasing lipid vesicle stability. The maximal membrane thickness change obtained by optical switching under these conditions was only 4 to 5 Å. The new experiments with premixed lipids reported in Figure 1 now reveal that this optical control window covers only about half of the thickness change effect which could be achieved in ideal photoswitching conditions. We therefore studied two widely used buffer systems, phosphate-buffered saline (PBS) and 1×TE buffer. The dark-adapted state, several photostationary states induced via UV-A and blue light are labelled accordingly. Horizontal lines indicate mean values of $d_{HH}$ for the dark-adapted, UV-A, and blue light photostationary states, shown as solid, dashed, and dotted line, respectively. The exposure time of UV-A light while approaching the photostationary state is indicated in seconds. The optical control window for the different buffer conditions is indicated by a red double-headed arrow.
PBS), and a mixture of Tris with EDTA (1×TE), for their influence on the optical control window. The analysis of the SAXS data is condensed in Figure 2, the full data set is shown in Figures S4 and S5a.

The dark-adapted states (all-trans) yielded mean membrane thicknesses of \( d_{\text{HH}} \) (dark, buffers) = 42.9 ± 0.2 Å and \( d_{\text{HH}} \) (dark, deionized water) = 41.9 ± 0.9 Å (Figure 2a and b, circles, solid line). Hence, all conditions allow for the formation of dense, all-trans membranes in the dark. To photoswitch towards high cis contents, UV-A illumination was used until a photostationary state was achieved. Independent of the usage of buffer or not, the UV photostationary states do not reach a cis fraction of 83% as obtained by preconditioning monomers in chloroform. Instead, the observed membrane thicknesses of \( d_{\text{UV}} \) = 34.8 ± 0.6 Å (see Figure 2b, dashed line and Figure S5) indicates cis contents around 64%. This is not unexpected, given the blueshifting of azobenzene absorption spectra seen upon going from molecular solutions to assembled systems [19].

However, when using blue light to photoswitch from a cis- to a trans-enriched state, the outcomes depend greatly on whether the system is buffered or not. In DI water, the maximal blue light photostationary state membrane thickness is \( d_{\text{blue}} \) = 39.0 ± 0.3 Å (Figure 2b, squares, dotted line), i.e. 30% cis isomer still remains. Instead, in PBS or 1×TE buffer, the membrane thickness increases to \( d_{\text{blue}} \) = 42.3 ± 0.4 Å (Figure 2a, squares, dotted line), i.e. only 3% cis isomer remains. Optical control of vesicles is therefore highly efficient in these buffered aqueous solutions. Follow up experiments with NaCl solutions (see Figure S9), indicate that ionic strength rather than pH buffering is key to efficient photoswitching towards trans-enriched states in lipid vesicles. This effect is not observed for molecular monomers in aqueous solutions, where azobenzenes lose their ability to efficiently isomerize anyway [24]. Increasing ionic strength was reported to remove water molecules from inside the membrane promoting tighter packing of lipids [35, 36]. This should enhance the protection of the azobenzene photoswitch from water.

Next, we explored possible influence of X-ray on photostationary states. For this study, we used X-ray exposures to deposit a certain X-ray dose into the vesicle solution, i.e. a certain energy of absorbed X-rays per mass of the exposed sample [37]. The X-ray absorption drops drastically for increasing energies and in turn high X-ray energies enable low-dose SAXS experiments and vice versa (cf. Figure S7 and Note S7). Thus, the dosing experiment was conducted as follows. First, we prepared the vesicles in a high cis fraction by UV-A illumination. Next, we exposed the sample to a rather high X-ray energy of 54 keV for 60 s. Five such consecutive exposures yield an identical SAXS signal. There is no sign of any hard X-ray induced effects after 5 min in total, even though this experiment was performed at a high brilliance Petra III synchrotron beamline with full beam on the sample, cf. Figure 3a. This finding is in agreement with the fact that the cross section for photoabsorption is dramatically reduced for high X-ray energies.
Depositing high doses in water requires soft X-rays (discussion in the Supporting Information section S7). Such soft X-rays give rise to radiolysis of water, and oxidising and reducing radicals and reactive species [38, 39], which may provide pathways for catalytic redox-based unidirectional switching towards the thermodynamic groundstate (all-trans) [15]. Therefore, we tested also the change of membrane thickness during six consecutive 8 keV X-ray exposures each of 2 s (Figure 3a). After each X-ray exposure, a shift of the SAXS pattern was visible. To quantify the thickness change of the photomembrane, we modelled to extract the head-to-head distances, which increased dramatically after each soft X-ray exposure, depending on the total delivered X-ray dose. Remarkably, the final mean membrane thickness of $d_{\text{HH}}^{\text{xray}} = 42.5 \pm 0.3$ Å (Figure 3b, star and square datapoints at high dose, dashed line; Figure S5b) matches the thickness of the dark-adapted all-trans state $d_{\text{HH}}^{\text{dark}} = 41.9 \pm 0.9$ Å (Figure 2b, circular data points from repeated measurements, solid line). This is a direct indication of quantitative switching obtained by 8 keV X-ray exposure.

These X-ray induced changes of the isomerisation state of the azobenzene photoswitch are fully reversible and should not be mistaken as irreversible radiation damage. They can however be employed as a reversible X-ray dose readout. To demonstrate this, we plot the membrane thickness as observed by hard X-ray SAXS in response to soft X-ray exposure with dose expressed in kGray [kGy] in Figure 3b. The calculation of the dose, a routine calculation in radiation protection, is explained in the Supporting Information. Here, quantitative switching starts above 100 kGy, and saturates for an X-ray dose above 700 kGy. We propose that the measurement of azo-PC cis to trans transition for SUVs in DI water can be used to read out the effective X-ray dose in a regime of up to 700 kGy. This range may help to calibrate critical doses for biological SAXS experiments which can range from 51 kGy [40], to 400 kGy [41], to 284–7056 kGy [38].

The SAXS signal is almost independent of the X-ray energy (Figure S7, inset). Therefore, the minimum of the X-ray dose for samples with high water fraction at 36 keV (Figure S7) should be used to prevent radiation effects. Our experiments demonstrate that high quality SAXS data may be obtained even for weakly scattering biological samples.

4 Conclusions and summary

SAXS with hard X-rays provides a direct read out of the membrane bilayer thickness. We find that membrane thickness has a linear dependency of photolipid isomerization fraction. The 8 Å thickness change accessible by photoswitching in buffered solutions (20% of the membrane thickness) is massive, and apparently only possible because the membrane itself seals the azobenzene from the unfavourable polar solvent. Hydrophobic matching of membrane thickness with trans-membrane proteins is a wide research field that could benefit enormously from the large thickness change effects observed here [42]. Due to the rapid development of high power UV-A LEDs, and the availability of pulsed UV-A lasers at synchrotron sources [43], we expect that future experiments with pulsed illumination may allow rapid membrane thickness transition times of few ms and below. At these short timescales, the hydrodynamic coupling of the intercalated water to the photolipids may give rise to further interesting transient phenomena [25]. We also highlight the versatility of high energy X-rays as low-dose probes.

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Author contribution: MFO performed vesicle preparation, and SAXS experiments and X-ray data analysis, and wrote the manuscript; AMD performed synthesis of azo-PC and FAzoM, and spectroscopic and HPLC analyses of cis/trans isomer ratios; AB developed a protocol for photolipid SUV preparation; BA performed the SAXS experiments in NaCl solution; HA enabled the SAXS experiments at ELETTRA and provided in depth expertise on SAXS calibration and data conversion; OTS designed cis/trans ratio measurements and wrote the manuscript; BN designed the study, and joined the X-ray experiments, and wrote the manuscript.

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