



Cite this: *Org. Biomol. Chem.*, 2021, **19**, 2312

## Reductive stability evaluation of 6-azopurine photoswitches for the regulation of CKI $\alpha$ activity and circadian rhythms†

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Photopharmacology develops bioactive compounds whose pharmacological potency can be regulated by light. The concept relies on the introduction of molecular photoswitches, such as azobenzenes, into the structure of bioactive compounds, such as known enzyme inhibitors. Until now, the development of photo-controlled protein kinase inhibitors proved to be challenging for photopharmacology. Here, we describe a new class of heterocyclic azobenzenes based on the longdaysin scaffold, which were designed to photo-modulate the activity of casein kinase I $\alpha$  (CKI $\alpha$ ) in the context of photo-regulation of circadian rhythms. Evaluation of a set of photoswitchable longdaysin derivatives allowed for better insight into the relationship between substituents and thermal stability of the *cis*-isomer. Furthermore, our studies on the chemical stability of the azo group in this type of heterocyclic azobenzenes showed that they undergo a fast reduction to the corresponding hydrazines in the presence of different reducing agents. Finally, we attempted light-dependent modulation of CKI $\alpha$  activity together with the accompanying modulation of cellular circadian rhythms in which CKI $\alpha$  is directly involved. Detailed structure–activity relationship (SAR) analysis revealed a new potent reduced azopurine with a circadian period lengthening effect more pronounced than that of its parent molecule, longdaysin. Altogether, the results presented here highlight the challenges in the development of light-controlled kinase inhibitors for the photomodulation of circadian rhythms and reveal key stability issues for using the emerging class of heteroaryl azobenzenes in biological applications.

Received 4th January 2021,  
Accepted 16th February 2021

DOI: 10.1039/d1ob00014d

rsc.li/obc

## Introduction

Protein kinases constitute a major target class for drug development.<sup>1</sup> The human kinome consists of ~500 kinases with various key cellular functions.<sup>2</sup> Due to this vastness of the kinome, the general similarities in the ATP-binding domain and a uniform catalytic mechanism, the development of specific kinase- or kinase type-selective inhibitors remains challenging.<sup>3</sup> This limitation could be partially overcome by considering, besides the biochemical target selectivity, the site

specificity – *e.g.* by limiting not the enzymes with which the compound interacts, but the parts of organs of the body in which the compound is active, thereby preventing unwanted off-site activities in the future.

Such external control over inhibitor potency in time and space could be enabled using light to locally and reversibly activate the bioactive compound only at the desired site of action. This concept forms the basis for the photopharmacological approach: photopharmacology is an emerging field of chemical biology that utilizes photo-responsive molecules to enable control over the activity of a drug, using light.<sup>4–7</sup> The main tools of photopharmacology are molecular photoswitches,<sup>8</sup> which can be reversibly switched between two or more isomeric forms using light of different wavelengths. Such control has been established over bacterial communication and resistance buildup,<sup>9,10</sup> ion channels,<sup>11</sup> G protein-coupled receptors,<sup>12</sup> lipid membranes,<sup>13</sup> and nucleic acids.<sup>14,15</sup> However, up to this point, the photocontrol of kinase activity has proven to be highly challenging for photopharmacology, leading often to chemically unstable compounds, undesired photochemical side-reactions or photoswitchable kinase

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†Electronic supplementary information (ESI) available. See DOI: 10.1039/d1ob00014d



inhibitors that show little difference in potency in their photoisomeric states. Recent examples of photoswitchable kinase inhibitors include photomodulation of RET,<sup>16</sup> PKC,<sup>17</sup> MEK1,<sup>18</sup> VEGFR2<sup>19</sup> and BRAF<sup>V600E</sup> (ref. 20) enzymes.

Here we report the evaluation of a library of green-light switchable 6-azopurines for the photocontrolled inhibition of casein kinase I $\alpha$  (CKI $\alpha$ ) in the context of our efforts towards the light regulation of circadian rhythms. Circadian rhythms are based on endogenous biochemical oscillations with a  $\sim$ 24 h cycle observed in almost every cell in the human body.<sup>21,22</sup> Various studies have linked the disruption of these rhythms to a wide variety of diseases and disorders.<sup>23–25</sup> In mammals, at the molecular level, circadian oscillations are driven by negative feedback loops of clock genes. Casein kinase I (CKI, isoforms  $\delta$ ,  $\epsilon$ , and  $\alpha$ ) family proteins play a key role in these loops (Fig. 1A) by phosphorylating the PER protein and thus promoting its proteasomal degradation and maintaining the circadian rhythms on the  $\sim$ 24 h base. Control of circadian cycles by employing small molecules has recently been investigated, opening fascinating opportunities in disease control.<sup>26–29</sup> High-throughput screening supported by rational synthetic design led to the discovery of small-molecule modulators of the circadian

period, mostly showing period lengthening,<sup>30–36</sup> but some also exhibiting period shortening effect.<sup>37–39</sup> The Kay group identified a potent modulator of cellular circadian rhythms, dubbed 'longdaysin' (Fig. 1B), which showed IC<sub>50</sub> values of 8.8 and 5.6  $\mu$ M for CKI $\alpha$  and CKI $\delta$ , respectively.<sup>30</sup> However, since the core clock mechanism resides throughout the mammalian body,<sup>22,40</sup> circadian period modifiers, such as longdaysin, display a non-selective period modulation of all peripheral clocks as well as of the master clock, suprachiasmatic nucleus (SCN). Establishing photocontrol over their activity might enable site-selective activation and pave the way towards chronopharmacology.

Towards enabling the photocontrol by switchable circadian rhythm regulators, we have recognized the *N*-benzyl-arylamine scaffold of longdaysin as the candidate for the azologization approach,<sup>41</sup> in which azobenzene photoswitches are introduced into the structure of bioactive compounds with only a minimal overall structural change. Azobenzenes are a highly versatile and widely applied class of photoswitches<sup>42,43</sup> that undergo light-induced isomerization between the thermally stable *trans*-isomer and the unstable *cis*-isomer (Fig. 1B). They have been widely exploited in photopharmacology as the most versatile and well-understood photocontrolled tools.<sup>5,6</sup>



**Fig. 1** (A) Schematic representation of the circadian feedback loop and its modulation with longdaysin. (B) Structure of longdaysin and photo-isomerization of its azo-analogue.



Especially the recent interest in heterocyclic azobenzenes is inspired by the increasing number of methods for their synthesis<sup>5,14,44</sup> and the common presence of heterocyclic moieties in drugs.<sup>45-47</sup> Despite the fact that SAR analysis of longdaysin was conducted by extensive structural modifications of its C(2) and N(9) positions,<sup>35</sup> to the best of our knowledge, modification of the benzene ring (*o*, *m*, *p*, Fig. 2) has never been investigated. Thus, a synthetic methodology and a library of photoswitchable compounds developed previously by our group<sup>48</sup> were evaluated for enabling reversible circadian clock modulation (Fig. 1).

## Results and discussion

Microwave-assisted nucleophilic aromatic substitution of 9-chloro purines (2 and 3) with substituted aryl hydrazines (1a-q), followed by oxidation (Fig. 2), provided in good to high yields the desired photoswitches based on the purine core ( $R_2 = \text{H}$ , 4a-q;  $R_2 = \text{NH}_2$ , 5a-f). The choice of substituents was primarily driven by investigation of their influence on the biological activity, but was also aimed at obtaining the optimal photochemical properties, such as visible-light photo-isomerization and longer thermal half-lives of the metastable *cis*-isomers. Since the ultimate application of this work is to control the biological activity, substituents were chosen to electronically resemble the electron-withdrawing  $\text{CF}_3$  group present originally in longdaysin. These substituents ranged from strongly electron-withdrawing to weakly electron-donating groups, with

the exception of compound 4c, the only one with a strong electron-donating methoxy substituent.

For the application of azobenzenes in biological systems, their photochemical properties must be adjusted for a particular purpose. The most common parameters to be optimized are the photostationary state distribution (PSD) of isomers under irradiation with light of different wavelengths, thermal half-life of the metastable isomer, wavelength used for photo-isomerization, and (photo)chemical stability. Given that the cellular circadian assay takes several days, thermal *cis*-to-*trans* isomerization had to be minimized to permit a pronounced biological effect of the *cis*-isomer, with the least background activity of the *trans*-isomer. Thus, by modifying the benzene ring, next to increased biological activity (*vide infra*), we also aimed for longer half-lives. If this condition is not met, continuous or pulse irradiation with cell-nontoxic light (>500 nm)<sup>49-51</sup> would be required during the measurement to keep the content of the *cis*-isomer as high as possible for a prolonged period. However, this is not possible in circadian rhythm assays due to the nature of the experimental setup in which the luminescence of the cells is being measured in a plate reader under carefully maintained (dark) conditions, precluding continuous or pulsed irradiation.<sup>37</sup>

### Photo-isomerization

The wavelength used for isomerization is one of the key features for utilization of azobenzene in biological systems.<sup>51,52</sup> Generally, UV-light is needed for *trans*-to-*cis* isomerization, while back-isomerization can be achieved by either thermal



Fig. 2 Synthesis of longdaysin azolones 4a-q and 5a-f.



relaxation or visible light irradiation.<sup>43</sup> As UV-light is strongly absorbed and scattered by the skin, visible-light mediated photo-isomerization is highly desirable due to increased tissue penetration and reduced phototoxicity.<sup>53,54</sup> Heterocyclic azobenzenes, presented in this work, allow for using green light ( $\lambda_{\text{max}} = 530 \text{ nm}$ ) for *trans*-to-*cis* photo-isomerization (Fig. S4†). To quantitatively determine *trans*-to-*cis* conversion with green light, PSDs of two switches (**4f** and **5e**) were measured by <sup>1</sup>H-NMR (DMSO-*d*<sub>6</sub>, 1 mg mL<sup>-1</sup>, 25 °C, Fig. S4†). The photostationary state was achieved after 2 h of irradiation with green light. For azopurine **4f**, the PSS distribution (*cis*:*trans*) was determined to be 61 : 39 while for **5e** it was 71 : 29.

### Thermal half-life of compounds *cis*-4a–q and *cis*-5a–f

After the photostationary state was reached, thermal back-isomerization was studied in DMSO, CKI assay buffer and the medium used in the cellular assay (Table 1). The relationship between the structure and thermal stability in DMSO revealed that back-isomerization was highly dependent on the electronic nature of the substituents, as well as their position on the aromatic ring. Increased electron density of the benzene ring created a push–pull system with an electron-poor adenine core, causing shorter half-lives.<sup>55</sup> Thus, *para*-methoxy substi-

tuted azopurine **4c** exhibited the shortest half-life of 17 s. On the other hand, increasing the electron density of the purine core by incorporation of the C(2)-amino group (**5a–f**) had the opposite effect. Further analysis has shown that *para*- and *meta*-substituents on the benzene ring had a very limited influence on the thermal stability of the *cis*-isomer. Conversely, incorporation of *ortho*-substituents had a significant impact on the half-lives. Interestingly, the half-life of di-*ortho*-methyl compound **4f** was shorter than of mono-*ortho*-trifluoromethyl compound **4m**, indicating the importance of not only the steric effects but also the electronic effects of the *ortho*-substituents. Inspired by the work of Woolley<sup>51,52</sup> and Hecht<sup>56</sup> and our recent studies,<sup>57</sup> we expected to enhance the thermal stability of the *cis*-isomer by introducing *ortho* fluoro- or chloro-substituents. Incorporation of only one chlorine atom (**4h**) led to a significantly longer half-life of >1 h. The thermal half-life for the relaxation of di-*ortho*-fluoro compound **4q** was 1.4 h, while di-*ortho*-chloro compound **4n** displayed the longest half-life among 2-H-azopurines of almost 3 h. Remarkably, di-*ortho*-fluoro compound **5f** exhibited the slowest thermal relaxation, with a half-life of almost 12 h. In summary, for longdaysin azologs **4** and **5**, a very broad range of half-lives in DMSO (from 17 s to 12 h) were obtained. Altogether, the stability of the *cis*-isomer could be fine-tuned by controlling the electronic and steric effects of the substituents.

Before evaluating photoswitchable azo-longdaysin derivatives in biological assays, the half-lives in the corresponding aqueous media were measured (CKI assay buffer and cellular assay medium, Table 1). In contrast to the results obtained in DMSO, and despite the incorporation of different substituents, a very fast thermal back-isomerization of all photoswitches was observed. The obtained half-lives varied from a few seconds (**4c**) to slightly more than half an hour (**4n**), and the observed trend was the same as in DMSO (Table 1). Di-*ortho*-fluoro substituents in **4q** and **5f**, di-*ortho*-chloro in **4n**, and C(2)-amino substituents (**5a–f**) slowed down the thermal relaxation of the *cis*-isomer, but considering the length of the *in vitro* assay (3 h) and cellular assay (5 d), the back-isomerization was too fast for the photomodulation of the circadian period without constant or prolonged irradiation during the assay.

Despite the photophysical properties of azo-switches having been thoroughly investigated in the past,<sup>58–60</sup> their chemical stability in biologically relevant media is still underexplored. However, to enable photocontrol in cells, tissues or living organisms, the chemical and metabolic stability of azobenzenes must be established. It is known that enzyme<sup>61,62</sup> or thiol-mediated<sup>63</sup> azobenzene reduction can occur in cells, particularly by glutathione (GSH) that is present in cells at a concentration of up to 10 mM.<sup>64,65</sup> In addition, a recent seminal work by Peifer, Herges and co-workers<sup>66</sup> showed that photo-switchable CKI $\delta$  inhibitors based on imidazole- and thiazole-type heterocyclic azobenzenes undergo reduction to the corresponding hydrazines when exposed to thiols such as dithiothreitol (DTT) or GSH. Therefore, following the previous work and led by the finding that photoswitches **4h**, **4l**, **4n** and **5b**

**Table 1** Thermal *cis*-to-*trans* isomerization in three different media: DMSO (40  $\mu\text{M}$ , 25 °C), CKI assay buffer (40  $\mu\text{M}$ , 30 °C) containing ~100  $\mu\text{M}$  dithiothreitol (DTT) and cellular assay medium (40  $\mu\text{M}$ , 35 °C). Half-life of the *trans*-azobenzene (40  $\mu\text{M}$ ) reduction in CKI buffer containing 500  $\mu\text{M}$  DTT (30 °C)

Compound	$t_{1/2}$ , <i>cis</i> -to- <i>trans</i> (min)			$t_{1/2}$ , reduction (min)
	DMSO	CKI buffer	Cellular assay medium	CKI buffer
<b>4a</b>	3.5	0.55	0.35	120
<b>4b</b>	1.7	0.28	0.18	54
<b>4c</b>	0.28	0.12	0.033	21
<b>4d</b>	18	0.65	0.82	33
<b>4e</b>	15	0.87	0.70	19
<b>4f</b>	43	1.2	0.93	25
<b>4g</b>	18	0.82	0.70	28
<b>4h</b>	68	ND <sup>a</sup>	ND <sup>a</sup>	5.9
<b>4i</b>	8.5	0.80	0.57	52
<b>4j</b>	10	0.70	0.58	55
<b>4k</b>	9.7	1.1	0.37	14
<b>4l</b>	36	ND <sup>a</sup>	0.73	9.1
<b>4m</b>	54	1.9	1.5	63
<b>4n</b>	180	ND <sup>a</sup>	36	6.7
<b>4o</b>	2.4	0.42	0.25	95
<b>4p</b>	4.1	0.77	0.45	79
<b>4q</b>	85	9.3	4.6	20
<b>5a</b>	78	3.1	2.0	13
<b>5b</b>	66 <sup>b</sup>	ND <sup>b</sup>	ND <sup>b</sup>	92
<b>5c</b>	81	3.5	1.1	2.4
<b>5d</b>	46	1.8	1.0	83
<b>5e</b>	91	0.73	1.0	8.0
<b>5f</b>	710	18	7.3	29

<sup>a</sup>The half-life was not determined due to a constant absorption decrease upon dissolving the azopurine in CKI buffer or cellular assay medium. <sup>b</sup>Compound **5b** showed slow decomposition in DMSO.



showed a continuous absorption decrease when dissolved in CKI assay buffer (Table 1), we tested all photoswitches for their reduction by DTT present in this buffer solution (Fig. 3A). Initially, the process was analyzed by  $^1\text{H-NMR}$  in  $\text{DMSO-}d_6$  solution of **4k**, **4n** and **5e** (9 mM) and different quantities of DTT (0 eq., 1 eq., 2 eq., and 4 eq., Fig. 3B). The NMR spectrum of azopurine **4k** did not change over time in the DTT-free sample. On the other hand, all three samples with DTT showed a second set of signals appearing after 10 min. In the presence of four equivalents of DTT, the starting azopurine **4k** was almost completely (89%) converted to the corresponding hydrazine after 4 h. Additional analysis by UPLC-MS confirmed that the newly formed compound shows a mass increase of 2 units, consistent with the reduction product (Fig. S3†). Interestingly, when compound **4n** was subjected to 1 eq. of DTT in  $\text{DMSO-}d_6$ , bleaching due to azo bond reduction occurred instantaneously. This indicates that di-*ortho*-chloro substituents do not only play a stabilization role but also increase the susceptibility to reduction of the reported azobenzene system. This highlights the need for a systematic screening and better understanding of the chemical stability of heterocyclic azobenzenes under reductive conditions present in biological systems.<sup>66</sup>

Azopurines with a C(2)-amino group (**5a-f**) feature higher electron density on the aromatic rings. Therefore, we expected to see a slower reduction rate than for the corresponding C(2) unsubstituted azobenzenes. Testing the reduction of **4k** and **5e** with DTT confirmed our prediction (Fig. 3C). On the other hand, in an aqueous solution containing DTT (CKI buffer), both compounds are reduced so quickly that the difference is indistinguishable (Table 1).

Next, the reduction rate of all azobenzenes was measured in CKI buffer containing DTT ( $\sim 500 \mu\text{M}$ , Table 1). No clear correlation between the reduction rate and structure was observed. At the end of the measurement, all compounds were fully reduced, with reduction  $t_{1/2}$  ranging from  $\sim 2$  min (**5c**) to  $\sim 2$  h (**4a**). In that respect, measuring the thermal half-lives of the *cis*-isomers in the buffer was possible for almost all compounds except for **4h**, **4l**, **4n** and **5b** which underwent fast reduction competing with thermal back-isomerization. Since bleaching was also observed in cellular assay medium, we proceeded to the experimental testing of the stability of azopurine **4e** in the presence of other reducing agents, such as GSH, cysteine and ascorbic acid and compared their reducing efficiency to DTT (Fig. 3D). Interestingly, the reduction was found to proceed with all reducing agents. The fastest



**Fig. 3** Reductive stability of azopurines. (A) The reduction process. (B) Reduction of **4k** by different amounts of DTT followed by  $^1\text{H-NMR}$  in  $\text{DMSO-}d_6$  solution (25 °C, 9 mM **4k**). (C) Comparison of the reduction rate of *trans*-**4k** and *trans*-**5e** by DTT (1 eq.). The reduction is followed by  $^1\text{H-NMR}$  in  $\text{DMSO-}d_6$  solution (25 °C, 9 mM). (D) Reduction of *trans*-**4e** followed by UV-Vis spectroscopy at 340 nm in buffer solution (100 mM HEPES, pH 7.4) without (green) or with different reducing agents (500  $\mu\text{M}$ ) – DTT (black), GSH (red), ascorbic acid (purple), and cysteine (blue).



reduction occurred in the case of DTT and the slowest with GSH, while ascorbic acid and cysteine showed similar, intermediate rates. The obtained results show the importance of testing every photoswitchable drug for the possible reduction with intracellular GSH, or reductive components present in *in vitro* buffer (such as DTT) that are used for preventing the oxidation of peptides and proteins.

### Photo-modulation of the CKI activity

Selected azopurines were tested *in vitro* in recombinant CKI $\alpha$  inhibition assays to assess their potency and the extent to which this potency can be photomodulated through azobenzene switching before reduction to the corresponding light-nonresponsive hydrazines occurs. Nine azopurines were selected for the screening of their activity in the dark (black bars, Fig. 4) and upon irradiation with green ( $\lambda_{\max} = 530$  nm) light (green bars, Fig. 4). All the compounds were initially tested at 20  $\mu\text{M}$  concentration, assuming that after azologization the potency of longdaysin (IC<sub>50</sub> value of 5.6  $\mu\text{M}$ ) was retained.<sup>30</sup> Among the tested compounds (Fig. 4), azopurines **4b** and **4c** showed almost no inhibition of CKI $\alpha$ , similarly to *di-ortho*-methyl **4f** and *meta*-carboxylic **4p** azopurines. Irradiation with green light during the assay increased the activity of **4d**, while **5e** proved to be less potent after irradiation. The induced alteration in inhibition might be attributed to the difference in affinity between the two isomers. The relatively small difference in activity between the *cis*- and *trans*-isomer of **4d** or **5e** can be explained by the relatively fast reduction to the corresponding hydrazine by DTT present in the assay buffer, yielding the same light-nonresponsive product and preventing a significant difference in inhibition to be observed during the assay.

In order to test this hypothesis, the inhibition assay was performed starting with azobenzenes **4d** and **5e**, and their corresponding hydrazines (Fig. S5<sup>†</sup>). We observed that the inhibitory potency of azobenzenes and their reduced analogues was the same, confirming that reduction of the azo

bond under the assay conditions occurs fast enough so that the difference in inhibition cannot be efficiently regulated with light. As a consequence, we attribute the biological effects observed in Fig. 4 and 5 to the reduced forms of photoswitches (**4<sup>red</sup>** and **5<sup>red</sup>**).

### Modulation of the circadian period

Despite the observed reduction, we decided to test the azopurines for the modulation of the circadian period (Fig. 5). The assay employs human U2OS cells with a *Bmal1-dLuc* reporter (established at the University of California San Diego, USA) in DTT-free cellular medium.<sup>37</sup> The experimental readout is based on chemiluminescence produced during the oxidative decarboxylation of luciferin. Luciferin is present in a high concentration (0.2 mM) in the cellular assay medium, and it strongly absorbs in the UV region. This prevents photoisomerization during the cellular assay with UV-light, which is generally used for the photoisomerization of azobenzene photoswitches.<sup>5</sup> Gratifyingly, the possibility to isomerize the azopurines reported here with green light gives them an advantage for the application even in the presence of luciferin. Additionally, in contrast to UV-light, visible light is less cytotoxic, enabling longer irradiation prior to the performance of the cellular assay. Prolonged cellular irradiation (continuous or pulse) would maintain the presence of the *cis*-isomer for a longer time despite having a short half-life; however, again, it was not feasible in the current experimental setup (*vide infra*).

For the initial screening of the activity of the thermally stable *trans*-isomers, azopurines were dissolved in DMSO and applied to the cells in 3-fold dilution series. The chemiluminescence signal was measured every 100 min over 5 d and the concentrations required for 4 h circadian period lengthening from the DMSO control were plotted (Fig. 5). Compound **4a** showed significantly reduced potency in comparison with longdaysin (Fig. 5A) and reduced reporter intensity (Fig. S6<sup>†</sup>), which likely arises from toxicity at higher concentrations. Similarly, azopurines substituted in the *para*-position (**4b** and

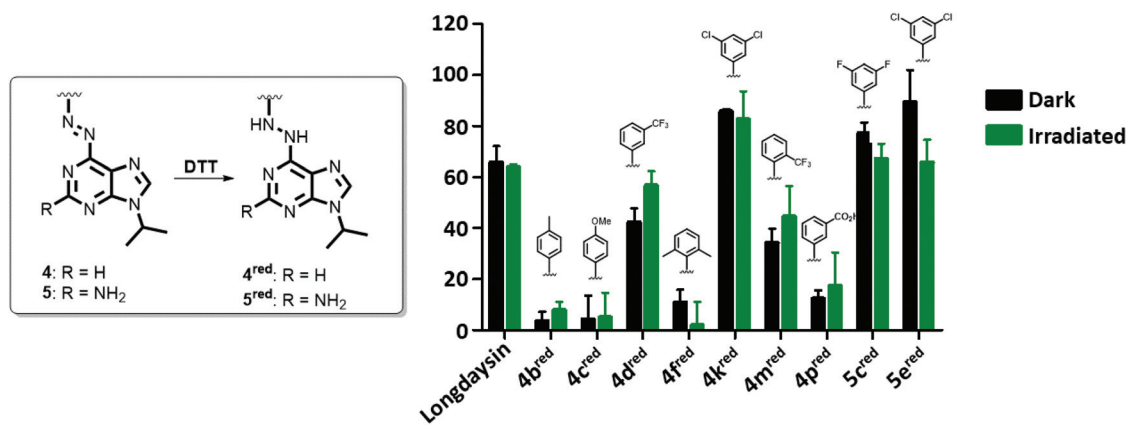
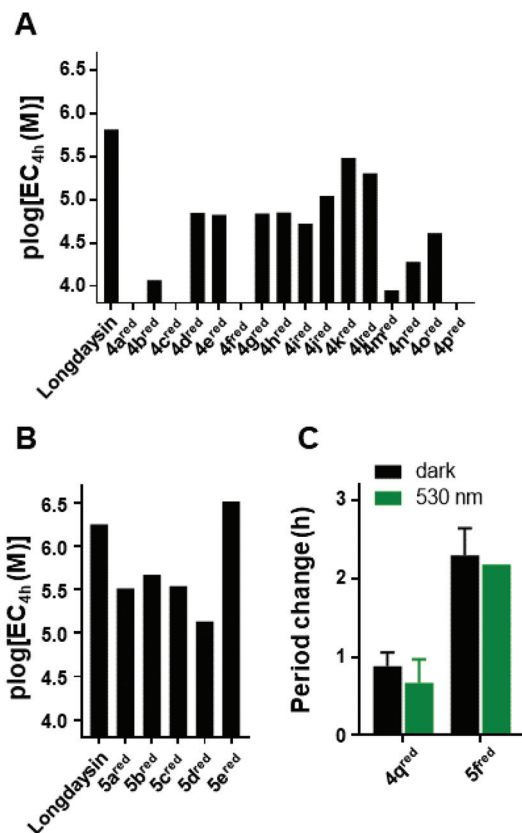


Fig. 4 Degree of CKI $\alpha$  inhibition in the enzymatic assay starting with different azopurines (**4** and **5**, 20  $\mu\text{M}$ , 30  $^{\circ}\text{C}$ ) that get reduced to the corresponding hydrazines (**4<sup>red</sup>** and **5<sup>red</sup>**) under the assay conditions. Samples kept in the dark are shown in black, and samples irradiated during the assay in green. The results (of the assays are mean  $\pm$  SD of at least two independent measurements.





**Fig. 5** The effect of azopurines 4 and 5, which undergo reduction of the azo bond under the assay conditions to hydrazines **4<sup>red</sup>** and **5<sup>red</sup>**, on the circadian period in human U2OS cells. Luminescence rhythms were monitored in the presence of various concentrations of compounds (12 points of 3-fold dilution series in DMSO; final 0.7% DMSO) in the dark (A and B) or upon irradiation with green light ( $\lambda_{\max} = 530$  nm) (C). In (A) and (B),  $[EC_{4h}]$  (M) is the concentration required for a 4 h period change and it is plotted to compare the potency.  $[EC_{4h}]$  values are calculated by fitting the 12-point dilution ( $n = 2$ ) data with the exponential growth curve, and for **4a<sup>red</sup>**, **4c<sup>red</sup>**, **4f<sup>red</sup>** and **4p<sup>red</sup>** they could not be obtained due to significant toxicity of these compounds. In (C), the period change relative to DMSO (at 7.1  $\mu$ M) is plotted.

**4c**) showed a very small period change and caused reduction of the reporter intensity, indicating that introduction of a *para*-substituent is not beneficial for the biological activity and renders these compounds toxic. The *meta*-substituted azopurines exhibited strong period lengthening, with compound **5e** having an even stronger effect than that of longdaysin (Fig. 5A and B). Incorporation of substituents at the *ortho*-position led to a decrease of period lengthening, but in contrast to the *para*-substituted azopurines, these compounds did not exhibit reduced reporter intensity (with the exception of **4f**). Due to the beneficial effect of the *meta*-substituents, particular emphasis was put on introducing groups in this position. SAR analysis revealed that a polar carboxylic group in the *meta*-position fully suppressed the activity of **4p**. The methyl group in **4o** and the fluorine substituent in **4i** slightly increased the activity, but the period change remained low. Moreover, **4o** and **4p** showed reduction of the reporter intensity.

Introduction of  $CF_3$  or CN groups at the *meta*-position led to an almost equal increase of activity, while the chloro-substituent yielded the best circadian period modulator based on mono-*meta*-substituted azopurines. Interestingly, the cytotoxic effect was suppressed by introduction of these *meta*-substituents. The addition of the second *meta*-substituent contributed to a higher activity in comparison with the corresponding mono-*meta*-substituted compound.

In summary, comparing the circadian period lengthening effect of longdaysin and its direct azo-analogue **4d**, it is evident that azologization of longdaysin decreased the potency, probably caused by the loss of an important interaction of the benzylic 6-NH group with  $CKI\alpha$ .<sup>67</sup> The hydrogen bond is possibly restored by reduction of the azo-group to the hydrazine **4d<sup>red</sup>**. Thus, a reduced form of the **4d** azo-analogue of longdaysin, at a concentration of 7.9  $\mu$ M, exhibited only 2.2 h period lengthening in comparison with 11.5 h for longdaysin. However, the potency can be recovered by varying substituents on the benzene and purine moiety, and this strategy provided the reduced azopurine **5e<sup>red</sup>** with an activity even higher than that of longdaysin.

Finally, we investigated the possibility of light-induced circadian period modulation using selected compounds **4q** and **5f**. These two azopurines have a different C(2)-substituent, but the same benzene-ring substituents (di-*ortho*-fluoro). Both compounds were applied to cells in the dark (black bars, Fig. 5C) or irradiated with green light ( $\lambda_{\max} = 530$  nm) prior to the assay (green bars, Fig. 5C). As expected, **5f<sup>red</sup>** showed a stronger effect on the circadian period modulation than that of **4q<sup>red</sup>**. Nevertheless, a clear light-induced effect was not observed. This can be explained by both a short *cis*-isomer half-life in the cellular assay medium and a fast reduction to the corresponding hydrazines **4q<sup>red</sup>** and **5f<sup>red</sup>** that are not responsive to light.

As a result of the assay's length (5 d) and a possible fast reduction by intracellular GSH, we assume that the observed period lengthening effect mostly arises from the corresponding hydrazine. Due to, *inter alia*, unavoidable endogenous GSH production by the cells, it was not possible to prove if the circadian period can be additionally modulated by light. Nevertheless, a comprehensive library of compounds revealed a molecule with stronger period lengthening than that of longdaysin.

## Conclusion

A library of azopurines was synthesized and photochemically characterized and their chemical stability in the biological environment was investigated. Furthermore, their biological activity as  $CKI\alpha$  kinase inhibitors was tested in enzyme activity and circadian period modulation assays.

The two-step one-pot synthesis provided a quick and efficient access to the library of differently substituted azopurines. Photochemical characterization revealed that green light



can be utilized for photo-isomerization, yielding moderate PSS distributions. The thermal stability of the *cis*-isomer was measured in DMSO and aqueous media, showing that the rate of the *cis*-to-*trans* isomerization can be efficiently manipulated by varying substituents on the benzene ring or at the C(2)-position. While a broad range of half-lives was obtained in DMSO, thermal isomerization in aqueous media occurred on shorter timescales, rendering these photoswitches unsuitable for light modulation in long-term circadian rhythm experiments. Furthermore, azopurines were tested for their sensitivity to reduction with reducing agents commonly present in cells (GSH) or used for stabilization of proteins (*e.g.*, DTT). Both in DMSO and aqueous media, the reduction took place, and the rate was similar for all the compounds. The reduction process was clean, yielding pure light-nonresponsive hydrazines. Despite the observed reduction, all the compounds were tested for CKI $\alpha$  inhibition and circadian period modulation. Photo-modulation of the CKI $\alpha$  inhibition activity was observed, but due to the rapid DTT-mediated reduction of the azo functional group, the effect was not significantly pronounced and probably can be attributed to a different reduction rate of the photo-isomers. Interestingly, the cellular assay screening yielded compound **5e<sup>red</sup>** with stronger period lengthening than that of longdaysin. At a concentration of 7.9  $\mu$ M, azopurine **5e<sup>red</sup>** displayed a 12 h period-lengthening effect, in comparison with 9.2 h of longdaysin. However, the reduction rate and the nature of the circadian cellular assay prevented photo-modulation of the circadian period and CKI $\alpha$  inhibition.

In summary, a comprehensive SAR analysis of the ability of reduced azopurines to modulate the circadian period was performed. The reduction study and screening of reducing agents revealed DTT as the most potent reducing agent for the azo group. These results support previous findings by Peifer, Herges and co-workers,<sup>66</sup> confirming that DTT is a crucial reagent to be used in photopharmacology to test the biological stability of heteroazoarenes towards reduction. This work reveals challenges associated with azobenzene-type photomodulators of circadian rhythms in the broader context of photopharmacology. It also emphasizes the importance of achieving a long half-life, low fatigue, high photostationary state distributions (PSD) and a chemically stable photoswitch when long-term biological light modulation is required.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

We thank Natsuko Ono and Dr Kaori Goto for technical assistance, and Carla Miró Viñals for the help with the graphical design of Fig. 1. We gratefully acknowledge the generous support from The Netherlands Organization for Scientific Research (NWO-CW, Top grant to B. L. F., and VIDI Grant No.

723.014.001 for W. S.), the Royal Netherlands Academy of Arts and Sciences (KNAW), the Ministry of Education, Culture and Science (Gravitation program 024.001.035), the European Research Council (Advanced Investigator Grant No. 227897 to B. L. F.), Grant-in-Aid for Scientific Research (B) 18H02402 and Challenging Research (Exploratory) 20K21269 from JSPS (T. H.), Uehara Memorial Foundation (T. H.), and Takeda Science Foundation (T. H.). We would also like to acknowledge Konstantin Hoffer for the great technical assistance and the Deutsche Forschungsgemeinschaft DFG for grant no. PE1605\_2\_2.

## References

- 1 D. Fabbro, *Mol. Pharmacol.*, 2015, **87**, 766–775.
- 2 G. Manning, D. B. Whyte, R. Martinez, T. Hunter and S. Sudarsanam, *Science*, 2002, **298**, 1912–1934.
- 3 M. I. Davis, J. P. Hunt, S. Herrgard, P. Ciceri, L. M. Wodicka, G. Pallares, M. Hocker, D. K. Treiber and P. P. Zarrinkar, *Nat. Biotechnol.*, 2011, **29**, 1046–1051.
- 4 M. W. H. Hoorens and W. Szymanski, *Trends Biochem. Sci.*, 2018, **43**, 567–575.
- 5 W. A. Velema, W. Szymanski and B. L. Feringa, *J. Am. Chem. Soc.*, 2014, **136**, 2178–2191.
- 6 J. Broichhagen, J. A. Frank and D. Trauner, *Acc. Chem. Res.*, 2015, **48**, 1947–1960.
- 7 M. M. Lerch, M. J. Hansen, G. M. van Dam, W. Szymanski and B. L. Feringa, *Angew. Chem., Int. Ed.*, 2016, **55**, 10978–10999.
- 8 I. M. Welleman, M. W. H. Hoorens, B. L. Feringa, H. H. Boersma and W. Szymański, *Chem. Sci.*, 2020, **11**, 11672–11691.
- 9 W. A. Velema, J. P. van der Berg, M. J. Hansen, W. Szymanski, A. J. M. Driessen and B. L. Feringa, *Nat. Chem.*, 2013, **5**, 924–928.
- 10 M. J. Hansen, J. I. C. Hille, W. Szymanski, A. J. M. Driessen and B. L. Feringa, *Chem*, 2019, **5**, 1293–1301.
- 11 M. Stein, S. J. Middendorp, V. Carta, E. Pejo, D. E. Raines, S. A. Forman, E. Sigel and D. Trauner, *Angew. Chem., Int. Ed.*, 2012, **51**, 10500–10504.
- 12 J. A. Frank, D. A. Yushchenko, N. H. F. Fine, M. Duca, M. Citir, J. Broichhagen, D. J. Hodson, C. Schultz and D. Trauner, *Chem. Sci.*, 2017, **8**, 7604–7610.
- 13 J. A. Frank, D. A. Yushchenko, D. J. Hodson, N. Lipstein, J. Nagpal, G. A. Rutter, J. S. Rhee, A. Gottschalk, N. Brose, C. Schultz and D. Trauner, *Nat. Chem. Biol.*, 2016, **12**, 755–762.
- 14 K. Hüll, J. Morstein and D. Trauner, *Chem. Rev.*, 2018, **118**, 10710–10747.
- 15 A. S. Lubbe, W. Szymanski and B. L. Feringa, *Chem. Soc. Rev.*, 2017, **46**, 1052–1079.
- 16 R. Ferreira, J. R. Nilsson, C. Solano, J. Andréasson and M. Grötli, *Sci. Rep.*, 2015, **5**, 9769.
- 17 D. Wilson, J. W. Li and N. R. Branda, *ChemMedChem*, 2017, **12**, 284–287.



- 18 Y. H. Tsai, S. Essig, J. R. James, K. Lang and J. W. Chin, *Nat. Chem.*, 2015, **7**, 554–561.
- 19 D. Schmidt, T. Rodat, L. Heintze, J. Weber, R. Horbert, U. Girreser, T. Raeker, L. Bußmann, M. Kriegs, B. Hartke and C. Peifer, *ChemMedChem*, 2018, **13**, 2415–2426.
- 20 M. W. H. Hoorens, M. E. Ourailidou, T. Rodat, P. E. van der Wouden, P. Kobauri, M. Kriegs, C. Peifer, B. L. Feringa, F. J. Dekker and W. Szymanski, *Eur. J. Med. Chem.*, 2019, **179**, 133–146.
- 21 M. H. Hastings, A. B. Reddy and E. S. Maywood, *Nat. Rev. Neurosci.*, 2003, **4**, 649–661.
- 22 C. L. Partch, C. B. Green and J. S. Takahashi, *Trends Cell Biol.*, 2014, **24**, 90–99.
- 23 J. Bass and M. A. Lazar, *Science*, 2016, **354**, 994–999.
- 24 R. W. Logan and C. A. McClung, *Nat. Rev. Neurosci.*, 2019, **20**, 49–65.
- 25 S. M. Abbott, R. G. Malkani and P. C. Zee, *Eur. J. Neurosci.*, 2018, **51**, 567–583.
- 26 T. Wallach and A. Kramer, *FEBS Lett.*, 2015, **589**, 1530–1538.
- 27 T. Hirota and S. A. Kay, *Methods Enzymol.*, 2015, **551**, 267–282.
- 28 Z. Chen, S. h. Yoo and J. S. Takahashi, *Annu. Rev. Pharmacol. Toxicol.*, 2018, **58**, 231–252.
- 29 S. Miller and T. Hirota, *J. Mol. Biol.*, 2020, **432**, 3498–3514.
- 30 T. Hirota, J. W. Lee, W. G. Lewis, E. E. Zhang, G. Breton, X. Liu, M. Garcia, E. C. Peters, J.-P. Etchegaray, D. Traver, P. G. Schultz and S. A. Kay, *PLoS Biol.*, 2010, **8**, e1000559.
- 31 T. Oshima, Y. Niwa, K. Kuwata, A. Srivastava, T. Hyoda, Y. Tsuchiya, M. Kumagai, M. Tsuyuguchi, T. Tamaru, A. Sugiyama, N. Ono, N. Zolboot, Y. Aikawa, S. Oishi, A. Nonami, F. Arai, S. Hagihara, J. Yamaguchi, F. Tama, Y. Kunisaki, K. Yagita, M. Ikeda, T. Kinoshita, S. A. Kay, K. Itami and T. Hirota, *Sci. Adv.*, 2019, **5**, eaau9060.
- 32 T. Hirota, J. W. Lee, P. C. St. John, M. Sawa, K. Iwaisako, T. Noguchi, P. Y. Pongsawakul, T. Sonntag, D. K. Welsh, D. A. Brenner, F. J. Doyle, P. G. Schultz and S. A. Kay, *Science*, 2012, **337**, 1094–1097.
- 33 S. Miller, Y. L. Son, Y. Aikawa, E. Makino, Y. Nagai, A. Srivastava, T. Oshima, A. Sugiyama, A. Hara, K. Abe, K. Hirata, S. Oishi, S. Hagihara, A. Sato, F. Tama, K. Itami, S. A. Kay, M. Hatori and T. Hirota, *Nat. Chem. Biol.*, 2020, **16**, 676–685.
- 34 S. Miller, Y. Aikawa, A. Sugiyama, Y. Nagai, A. Hara, T. Oshima, K. Amaiike, S. A. Kay, K. Itami and T. Hirota, *Cell Chem. Biol.*, 2020, **27**, 1192–1198.
- 35 J. W. Lee, T. Hirota, D. Ono, S. Honma, K. I. Honma, K. Park and S. A. Kay, *J. Med. Chem.*, 2019, **62**, 1989–1998.
- 36 Y. Isojima, M. Nakajima, H. Ukai, H. Fujishima, R. G. Yamada, K. Masumoto, R. Kiuchi, M. Ishida, M. Ukai-Tadenuma, Y. Minami, R. Kito, K. Nakao, W. Kishimoto, S. h. Yoo, K. Shimomura, T. Takao, A. Takano, T. Kojima, K. Nagai, Y. Sakaki, J. S. Takahashi and H. R. Ueda, *Proc. Natl. Acad. Sci. U. S. A.*, 2009, **106**, 15744–15749.
- 37 T. Hirota, W. G. Lewis, A. C. Liu, J. W. Lee, P. G. Schultz and S. A. Kay, *Proc. Natl. Acad. Sci. U. S. A.*, 2008, **105**, 20746–20751.
- 38 T. K. Tamai, Y. Nakane, W. Ota, A. Kobayashi, M. Ishiguro, N. Kadofusa, K. Ikegami, K. Yagita, Y. Shigeyoshi, M. Sudo, T. Nishiwaki-Ohkawa, A. Sato and T. Yoshimura, *EMBO Mol. Med.*, 2018, **10**, e8724.
- 39 Z. Chen, S. h. Yoo, Y.-S. Park, K. h. Kim, S. Wei, E. Buhr, Z.-Y. Ye, H.-L. Pan and J. S. Takahashi, *Proc. Natl. Acad. Sci. U. S. A.*, 2012, **109**, 101–106.
- 40 J. Gaucher, E. Montellier and P. Sassone-Corsi, *Trends Cell Biol.*, 2018, **28**, 368–379.
- 41 J. Morstein, M. Awale, J. L. Reymond and D. Trauner, *ACS Cent. Sci.*, 2019, **5**, 607–618.
- 42 A. A. Beharry, G. A. Woolley, M. M. Nass, N. H. Wassermann, B. F. Erlanger, M. Takagi, M. Komiyama, M. Kokkinidis, A. Rompp, B. Spengler, A. Pingoud and W. Zinth, *Chem. Soc. Rev.*, 2011, **40**, 4422–4437.
- 43 *Molecular Switches*, ed. B. L. Feringa and W. R. Browne, WILEY-VCH, 2nd edn, 2011.
- 44 S. Crespi, N. A. Simeth and B. König, *Nat. Rev. Chem.*, 2019, **3**, 133–146.
- 45 A. P. Taylor, R. P. Robinson, Y. M. Fobian, D. C. Blakemore, L. H. Jones and O. Fadeyi, *Org. Biomol. Chem.*, 2016, **14**, 6611–6637.
- 46 A. Gomtsyan, *Chem. Heterocycl. Compd.*, 2012, **48**, 7–10.
- 47 Y. J. Wu, in *Progress in Heterocyclic Chemistry*, Elsevier Ltd, 2012, vol. 24, pp. 1–53.
- 48 D. Kolarski, W. Szymanski and B. L. Feringa, *Org. Lett.*, 2017, **19**, 5090–5093.
- 49 J. D. Stoen and R. J. Wang, *Proc. Natl. Acad. Sci. U. S. A.*, 1974, **71**, 3961–3965.
- 50 M. Hori, K. Shibuya, M. Sato and Y. Saito, *Sci. Rep.*, 2014, **4**, 7383.
- 51 M. Dong, A. Babalhavaeji, S. Samanta, A. A. Beharry and G. A. Woolley, *Acc. Chem. Res.*, 2015, **48**, 2662–2670.
- 52 S. Samanta, A. A. Beharry, O. Sadovski, T. M. McCormick, A. Babalhavaeji, V. Tropepe and G. A. Woolley, *J. Am. Chem. Soc.*, 2013, **135**, 9777–9784.
- 53 R. Weissleder and V. Ntziachristos, *Nat. Med.*, 2003, **9**, 123–128.
- 54 J. L. Sandell and T. C. Zhu, *J. Biophotonics*, 2011, **4**, 773–787.
- 55 N. Nishimura, T. Sueyoshi, H. Yamanaka, E. Imai, S. Yamamoto and S. Hasegawa, *Bull. Chem. Soc. Jpn.*, 1976, **49**, 1381–1387.
- 56 C. Knie, M. Utecht, F. Zhao, H. Kulla, S. Kovalenko, A. M. Brouwer, P. Saalfrank, S. Hecht and D. Bléger, *Chem. – Eur. J.*, 2014, **20**, 16492–16501.
- 57 L. N. Lameijer, S. Budzak, N. A. Simeth, M. J. Hansen, B. L. Feringa, D. Jacquemin and W. Szymanski, *Angew. Chem., Int. Ed.*, 2020, **59**, 21663–21670.
- 58 J. García-Amorós and D. Velasco, *Beilstein J. Org. Chem.*, 2012, **8**, 1003–1017.
- 59 H. M. D. Bandara and S. C. Burdette, *Chem. Soc. Rev.*, 2012, **41**, 1809–1825.
- 60 H. Rau, in *Photoreactive Organic Thin Films*, Elsevier, 2002, pp. 3–47.



- 61 H. R. López-Mirabal and J. R. Winther, *Biochim. Biophys. Acta, Mol. Cell Res.*, 2008, **1783**, 629–640.
- 62 N. S. Kosower and E. M. Kosower, *Int. Rev. Cytol.*, 1978, **54**, 109–160.
- 63 C. Boulègue, M. Löweneck, C. Renner and L. Moroder, *ChemBioChem*, 2007, **8**, 591–594.
- 64 A. Meister and M. E. Anderson, *Annu. Rev. Biochem.*, 1983, **52**, 711–760.
- 65 H. Østergaard, C. Tachibana and J. R. Winther, *J. Cell Biol.*, 2004, **166**, 337–345.
- 66 M. Schehr, C. Ianes, J. Weisner, L. Heintze, M. P. Müller, C. Pichlo, J. Charl, E. Brunstein, J. Ewert, M. Lehr, U. Baumann, D. Rauh, U. Knippschild, C. Peifer and R. Herges, *Photochem. Photobiol. Sci.*, 2019, **18**, 1398–1407.
- 67 D. Kolarski, A. Sugiyama, G. Breton, C. Rakers, D. Ono, A. Schulte, F. Tama, K. Itami, W. Szymanski, T. Hirota and B. L. Feringa, *J. Am. Chem. Soc.*, 2019, **141**, 15784–15791.

