Upon the photostability of 8-nitro-cGMP and its caging as a 7-dimethylaminocoumarinyl ester†

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Light is a very attractive trigger for biological systems since it allows for the precise control over space, time and dosage. It is important that the caged compound does not bind to the target of the unleashed molecule prior to activation and is biologically fully inactive. This then allows us to equilibrate biological systems with the caged compound at high concentrations and to study fast processes without the need for considering slow pharmacokinetics.

A very advanced and even commercially available photocaging group for cGMP and cAMP is the 7-dimethylamino-coumarinyl moiety (DMACM) that is attached to the cyclic nucleotide via a phosphoric acid ester bond. The respective triester is hydrolysed upon irradiation between 330 and 440 nm under fast release of the desired cyclic nucleotides (within 1 nano second) and coumarinyl alcohol (9). Due to the good absorption coefficient (λ\(_{\text{max}}\) = 395 nm, \(\varepsilon = 16 \text{ mM}^{-1} \text{ cm}^{-1}\)) and high quantum yield (ϕ = 0.3) the uncaging is of excellent efficiency. Under physiological salt and pH conditions the caged compound is sufficiently soluble (10 \(\mu\)M and hydrolytically stable in the dark, making these compounds highly applicable. It has also been demonstrated that DMACM-caged cyclic nucleotides penetrate the mammalian cell membrane to get effective inside the cell.

In order to study the role of 8-nitro-cGMP in signalling and to explore the S-guanylation of the proteome we decided to develop a photoactivatable 8-nitro-cGMP precursor carrying the DMACM group. However, in contrast to cGMP or 8-bromo-cGMP, 8-nitro-cGMP was known from the literature to absorb in the visible spectrum (λ\(_{\text{max}}\) = 400 nm, \(\varepsilon = 9.1 \text{ mM}^{-1} \text{ cm}^{-1}\)). Since the \(N,C\)-glycosidic bond in 8-nitro-cGMP was described to be unusually long and prone to hydrolysis we expected 8-nitro-cGMP to possibly depurinate upon irradiation at 330–440 nm, the light energy that is required for the release of the coumarinyl group. However, nothing was known about the photostability of 8-nitro-cGMP to our knowledge. To test this, we synthesised 8-nitro-cGMP from commercially available 8-bromo-cGMP by nucleophilic substitution, close to a microwave-assisted heating we were able to accelerate the reaction from 5 days to 30 h and to improve the yield from 20% to 27%. To test the photostability, 8-nitro-cGMP (27 \(\mu\)M in degassed phosphate buffer was incubated under conditions that are typical for cGMP signalling experiments (10 mM Tris-HCl buffer, pH 7.4 and 1 mM MgCl\(_2\)) at room temperature. To test the conjugation efficiency we synthesized a fluorescein derivative of 8-nitro-cGMP and used this for UV–vis and fluorescence measurements. To test the uncaging efficiency, we synthesized the 8-nitro-cGMP(DMACM) hydrochloride (81\% yield).

Cyclic nucleotides are highly abundant messenger molecules that are found in all kingdoms of life. In humans they signal and regulate a whole set of physiologically relevant processes including ion channel conductance, blood vessel relaxation, metabolism or apoptosis for instance. Cyclic nucleotides are formed as a second messenger in response to extracellular signals like peptide hormones or nitric oxide (NO). Even though cyclic nucleotides have been studied for decades, the research field is still vivid. In particular, the development of new methods for sensing cyclic nucleotides inside living tissues and animals drives the progress. Particularly, the development of new methods for sensing cyclic nucleotides inside living tissues and animals drives the progress.3

Recently, Takaaki Akaike and co-workers reported 8-nitroguanosine 3',5'-cyclic monophosphate 1 (8-nitro-cGMP) as a novel second messenger that is formed inside the cell in a NO-dependent manner. Interestingly, 8-nitro-cGMP can also lead to protein S-guanylation by reacting with active cysteine residues. Later, the antagonisation of 8-nitro-cGMP by hydrogen sulphide was shown, demonstrating a complex interaction of 8-nitro-cGMP with various signalling molecules. More recently, the involvement of 8-nitro-cGMP in autophagy and guard cell signalling in plants was reported.

Although the cyclic nucleotide concentration inside a cell is tightly regulated, a local burst in the concentration may occur in response to extracellular signals. To study nucleotide signalling researchers have developed tools for the flash-like activation of cyclic nucleotides by light, so-called photocaged compounds.

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buffer, pH 7.0) was irradiated in a fluorescence spectrometer with precise control over irradiation wavelength (400 ± 16 nm) and time. At various time points, small aliquots were taken out and analysed by UV-spectroscopy and HPLC. As expected, 8-nitro-cGMP was unstable upon irradiation, and a new compound was formed (Fig. 1A) with a bathochromic shift of ≈30 nm in the UV-absorbance (λmax = 435 nm, Fig. S1, ESI†). The half-life was determined to be 67.4 ± 1.8 min (Fig. 1B) for the decomposition of the starting material. By HPLC-assisted high-resolution electrospray ionisation mass spectrometry (HPLC-HRMS) we clearly identified the constitution of the products to be C8H4N6O2 and C5H9O8P (ESI,† supplementary spectra) that formally fits to 8-nitrosoguanine (Scheme 1A, 3), or the corresponding oxime-isomer, and to an open form (4) of ribonolactone 7 (Scheme 1A).

This was a surprise since we had expected 8-nitro-cGMP to decay via hydrolysis under release of 8-nitro-guanine. However, based on the previous reports on photodecomposition of nitrated nucleosides, we propose a mechanism (Scheme 1A) which involves abstraction of the C1′–H by the excited triplet state of the nitro-group to generate a C1′-radical (5) followed by the collapse of this di-radical intermediate to a five-membered cyclic structure (6) and its decomposition to ribonolactone 7. Since the analogous 2′-deoxyribonolactone is also reported to be unstable, undergoing various reactions including β-eliminations, the apparent decomposition of 7 to other products was not a surprise and may explain the failure of its direct detection. However, a hydrolysed, open form of 7 (4) could be detected with high accuracy by mass spectrometry. Due to the presence of the cyclic phosphate in these compounds, β-elimination products of 4 or 7 cannot be discriminated by mass spectrometry and thus we are unable to clearly define the configuration of 4 or to exclude that it is a mixture of related products. Taking the stoichiometric formation of 8-nitrosoguanine (3) into account that is also supported by the appearance of two isosbestic points (410 and 310 nm) in the UV spectra (Fig. S1, ESI†) we exclude the possibility of the formation of 4 by direct oxidation of 5 with triplet oxygen.

Even though 8-nitro-cGMP is not photostable we continued to synthesise the caged compound because the determined half-life for the photodecomposition was relatively long and it could be expected that DMACM photo-removal could possibly compete. The DMACM group was installed similarly as reported for the synthesis of DMACM-caged 8-bromo-cGMP by incubation of 8-nitro-cGMP with a diazo precursor (8). The product was obtained in an overall yield of 22% (Scheme 1B) as a nearly 1-to-1 mixture of the axial and equatorial isomers that could be resolved during HPLC purification. First, we tested the photoactivation with the equatorial isomer of the caged compound (10 μM) in degassed buffer was irradiated at 400 nm for up to 300 s. Aliquots at various time points were subjected to HPLC analysis to monitor the formation of the 8-nitro-cGMP. (C) Selected HPLC-traces (260 nm) of the irradiated reaction mixture; (D) plot of the HPLC peak area values (traces at 370 nm) of liberated 1 versus irradiation time revealed a first-order half-life of ~35 s.

Scheme 1 (A) Proposed photodecomposition mechanism of 8-nitro-cGMP; (B) synthesis of DMACM-caged 8-nitro-cGMP.
half-life of 35 ± 3.5 seconds for the formation of 8-nitro-cGMP (Fig. 1D). For the axial isomer we found similar first-order kinetics with a half-life of 36 ± 2.1 seconds (Fig. S3, ESI†). Since photoactivation of the caged compound is nearly 116-fold faster than photodecomposition of the released product, more than 95% of the caged compound can be photoactivated within 5 minutes without significant decomposition of the released cyclic nucleotide (<5%). This is in accordance with the HPLC-data that show no detectable amounts of degraded nucleotide within the first few minutes of irradiation (Fig. 1C). Thus, photoactivation is clearly fast enough to successfully compete with product degradation. Furthermore, we determined the photochemical quantum yield (ϕ) at 400 nm using relative methods. For this we compared the photostability of both caged 8-bromo-cGMP (Fig. S4, ESI†). Taking an extinction coefficient of ε = 25 mM⁻¹ cm⁻¹ into account, we determined ϕ to be 7.5 ± 0.4% for the axial and 7.8 ± 0.7% for the equatorial isomer of 2. As also observed for other DMACM-caged cyclic nucleotides ϕ is very similar for both isomers. In contrast, ϕ was calculated to be ~0.18% for the photodecomposition of 8-nitro-cGMP at 400 nm. Since longer-wavelength light is more suitable for application in biology we also characterised the photoactivation at 430 ± 16 nm. Due to the reduced absorption at this wavelength slightly slower first-order kinetics were observed with 53 ± 1.3 seconds and 59 ± 1.2 seconds for the axial and equatorial isomers, respectively (Fig. S5 and S6, ESI†). Compared to the uncaging kinetics of 8-nitro-cGMP, its photodegradation at 430 nm was stronger attenuated resulting in a half-life of ~8 hours (Fig. S7, ESI†) making this wavelength even more suitable for photoactivation experiments.

The spontaneous hydrolysis of the DMACM-caged 8-nitro-cGMP (2) in the dark was tested at neutral pH and was found to be negligible, as also reported for other DMACM-caged cyclic nucleotides (Fig. S8, ESI†). With respect to their photochemical and hydrolytic properties, the two isomers of 2 are fully interchangeable – thus their resolution is dispensable for usage in biology.

For later applications it is essential to demonstrate that the caged precursor is biologically entirely inactive. To test this for DMACM-caged 8-nitro-cGMP (2), we applied an in vitro phosphorylation assay in cell extracts. This assay depends on the phosphorylation of vasodilator-stimulated phosphoprotein (VASP) by cGMP-dependent protein kinase type I (cGKI). It has been demonstrated that activation of cGKI by cGMP or 8-nitro-cGMP at 400 nm. Since longer-wavelength light is more suitable for application in biology we also characterised the photoactivation at 430 ± 16 nm. Due to the reduced absorption at this wavelength slightly slower first-order kinetics were observed with 53 ± 1.3 seconds and 59 ± 1.2 seconds for the axial and equatorial isomers, respectively (Fig. S5 and S6, ESI†). Compared to the uncaging kinetics of 8-nitro-cGMP, its photodegradation at 430 nm was stronger attenuated resulting in a half-life of ~8 hours (Fig. S7, ESI†) making this wavelength even more suitable for photoactivation experiments.

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