NO-independent stimulation or activation of soluble guanylyl cyclase during early reperfusion limits infarct size

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Aims

Guanylyl cyclase-cyclic guanosine monophosphate signalling plays an important role in endogenous cardioprotective signalling. The aim was to assess the potential of direct pharmacological activation and stimulation of soluble guanylyl cyclase, targeting different redox states of the enzyme, to limit myocardial necrosis during early reperfusion.

Methods and results

Rat isolated hearts were subjected to reversible left coronary artery occlusion (ischaemia-reperfusion) and infarct size was assessed by the tetrazolium staining technique. Administration during early reperfusion of BAY 41-2272, an NO-independent, haem-dependent stimulator of soluble guanylyl cyclase targeting the reduced state, or BAY 60-2770, an NO-independent, haem-independent activator targeting the oxidized state, significantly limited infarct size. Inhibition of NO synthesis did not abrogate this protection, but exogenous perfusion of NO with BAY 41-2272 produced a synergistic effect. The haem site oxidiser, ODQ abrogated the protection afforded by BAY 41-2272 but potentiated the protection afforded by BAY 60-2770. Targeting both the reduced and oxidized forms of sGC together did not afford additive protection.

Conclusions

Targeting either reduced or oxidized forms of sGC during early reperfusion affords cardioprotection, providing support for the concept that direct sGC manipulation at reperfusion has therapeutic potential for the management of acute myocardial infarction.

Keywords

Ischaemia-reperfusion • cGMP • NO • sGC

1. Introduction

Guanosine-3’5’-cyclic monophosphate (cyclic guanosine monophosphate, cGMP) is a second messenger generated by soluble and particulate guanylyl cyclases (sGC and pGC). Manipulating cGMP signalling through natriuretic peptide receptors/pGC and NO/sGC can limit the development of myocardial ischaemia-reperfusion injury.¹ Most pertinently, activation of these signalling pathways during the early phase of reperfusion is associated with marked limitation of infarct size.²–⁵ an effect mediated through elevation of intracellular cGMP levels and activation of protein kinase G (PKG).²

sGC is a heterodimeric protein which incorporates a prosthetic haem group, required for stimulation by NO, its major biological activator.⁶,⁷ It is known that the balance in the redox state of the haem moiety is shifted from a reduced (Fe²⁺) state to an NO-insensitive (Fe³⁺) state under conditions of oxidative stress.⁸–¹⁰ Since the bioavailability of NO may be severely compromised in disease states, especially those associated with oxidative stress,¹¹ a range of directly acting, NO-independent activators and stimulators have been developed for the management of vascular diseases. These compounds include direct NO-independent stimulators of sGC, such as BAY 41-2272, which bind to the haem moiety in its normal Fe²⁺ state and stimulate GTP catalytic activity independently of NO. Synthetic haem-independent activators of sGC in the oxidized Fe³⁺ haem state include BAY 60-2770.³,⁸,¹²,¹³ It is assumed that the redox balance of sGC under conditions of ischaemia-reperfusion is shifted towards the oxidized form which is insensitive to NO stimulation.

Since there is evidence that cGMP signalling is a tractable target for limiting the injurious consequences of reperfusion, we set out to investigate the infarct-limiting properties of this unique class of compounds...
whose mechanism of action is to stimulate/activate sGC, thereby elevating cGMP levels independently of NO.\(^{14–19}\) The primary objective of the study was to evaluate the effects of agents that target either the reduced or oxidized/haem-free states of sGC. It was hypothesized that perfusion of a sGC stimulator, BAY 41-2272 or activator, BAY 60-2770 during early reperfusion would limit infarct size and this protection would be in part as a result of elevated cGMP concentrations in the myocardium.

2. Methods

2.1 Animals

Male Sprague-Dawley rats (300 – 350 g; from Charles River, UK; B&K Universal Ltd, UK; or Harlan UK Ltd, UK) were used for these studies. Their care and use were in accordance with the UK Home Office guidelines on the Animals (Scientific Procedures) Act 1986 (The Stationary Office, London, UK) and approved by the Cardiff University ethics review board. Rats were housed in the institutional animal house under 12 h on/12 h off light cycles and allowed to acclimatize for at least 7 days with the standard chow containing 4% fat and 18% protein and water available ad libitum.

2.2 Experimental infarction and infarct size measurement

Rats were anaesthetized by pentobarbital sodium (175 mg/kg) with heparin (200 units) given concomitantly by i.p. injection. Hearts were excised and Langendorff perfused with modified Krebs–Henseleit buffer at constant pressure (74 mmHg) as previously described\(^1\) (see Supplementary material online for full details). Using a fluid filled balloon, left ventricular end-diastolic pressure (LVEDP) was set between 5 and 10 mmHg and isovolumic developed pressure continuously recorded (Chart, Powerlab data acquisition software, ADInstruments, UK). Coronary flow rate (CFR) was measured by collecting the coronary effluent from the apex of the heart. Before commencing any experimental protocol, hearts were left to equilibrate at 37°C for 10 min using a heated circulator.

Regional ischaemia was induced by placing a reversible suture and snare occluder around the left main coronary artery close to its origin. Ischaemia was confirmed by a CFR reduction of >30%. Hearts were subjected to 35 min coronary artery occlusion followed by 120 min reperfusion at 37°C. Following reperfusion the coronary artery was ligated and hearts were perfused with Evans’ Blue dye to stain the non-ischaemic tissue and thereby delineate the ischaemic risk zone by dye exclusion. Hearts were then frozen at −20°C, sectioned transversely from the apex into 2 mm thick sections and incubated in 1% triphenyltetrazolium chloride in phosphate buffer (pH 7.4) at 37°C to determine the unstained necrotic region within the ischaemic risk zone. After fixing for 48 h in 4% formaldehyde, sections were imaged from both sides (Image version 1.45q, NIH, USA) and a graphics tablet (Toon International B.V., Netherlands) was used to measure the total tissue area, risk zone, and necrotic zone of each section. These values were converted into volumes and combined to give total risk zone volume (R) and infarct size as a percentage of the risk zone (R/IR %).

Hearts were included for analysis only if they met strict inclusion criteria during the experimental protocols. Hearts were excluded if they failed to develop sinus rhythm during stabilization, if heart rate was < 200 b.p.m., baseline CFR >20 mL/min or if left ventricular developed pressure was <50 mmHg during stabilization. Regional ischaemia was confirmed by a CFR reduction >30% during coronary artery occlusion. Following release of the coronary artery snare, successful reperfusion was confirmed by a return towards baseline of CFR.

2.3 Myocardial cGMP assay

cGMP measurements were made in LV and RV myocardial tissue samples, harvested from hearts perfused in separate experiments as described below. Tissue samples were snap frozen, crushed, and immediately added to 200 μL lysis buffer containing 1 mM 3-isobutyl-1-methylxanthine. Samples were centrifuged for 60 min at 10,000 g. Supernatants were re-suspended in 400 μL 10% trichloroacetic acid and left for 30 min followed by a further centrifugation at 2500 g for 10 min. The supernatant was used for the cGMP measurements using the commercially available cyclic nucleotide radiomunnoassay kit (IBL International GMBH, Germany).

2.4 Treatment protocols

The following pharmacological agents were used in this study. BAY 41-2272 (Sigma-Aldrich, UK) is a sGC stimulator; BAY 60-2770 (gift from Bayer Pharma AG, Germany) is a sGC activator; NG-nitro-L-arginine methyl ester (L-NAME, Tocris Bioscience, UK) was used to inhibit NO synthase activity; 1H-[1,2,4]oxadiazolo[4,3-a]quinoxalin-1-one (ODQ, Tocris Bioscience) was used to selectively inhibit sGC; 2-(4-carboxyphenyl)-4,4,5,5-tetramethyl ylmidazoline-1-oxyl-3-oxide potassium salt (C-PTIO, Tocris Bioscience) was used as a scavenger of free NO; 1-hydroxy-2-oxo-3-[(N-methyl-6-aminohexyl)-3-methyl-1-triazene (NOC-9, Sigma-Aldrich) was used as NO donor. ODQ and BAY 41-2272 were dissolved in DMSO; the final maximum concentration of DMSO; in modified Krebs–Henseleit buffer was 0.05% v/v.

Following stabilization, all hearts were randomized to one of the experimental protocols, illustrated in Figure 1 (see Supplementary material online for details). All drug perfusions were started 5 min before release of the coronary artery snare until 10 min after reperfusion apart from L-NAME, C-PTIO, and ODQ which were perfused from 7 min before release of the snare. In Series 1, the infarct-limiting effects of BAY 41-2272 given at early reperfusion were examined and the relationship to NO and sGC redox state was interrogated by co-administration of L-NAME, C-PTIO, and ODQ. In Series 2, the contribution of exogenous NO was explored using NOC-9. In Series 3, the effects of BAY 60-2770 were examined to explore the targeting of the oxidized Fe\(^{3+}\) or haem-free state of sGC. Allied to each series, separate groups of hearts were prepared, without infarct size determination, for biochemical analysis of tissue cGMP concentration as described in Section 2.3.

2.5 Data analysis

Data are presented as means ± SEM and analysed using GraphPad Prism 5.0 (USA). Normal distribution of data was confirmed with the Kolsmogorov–Smirnoff test. Differences in mean values for infarct size, cGMP concentrations, and baseline haemodynamic parameters were compared by one-way ANOVA and Newman–Keuls post hoc test. Correlation of cGMP concentration with infarct size was determined using Spearman’s rank correlation coefficient. A P-value of < 0.05 was considered significant.

3. Results

Two hundred and eighty-five rats were used for this study. In the infarct experiments, 19 hearts failed to meet one or more of the inclusion criteria and were excluded from further experimentation. There were no technical exclusions in the groups prepared for cGMP analysis. Thus, we report data from 205 infarct experiments and from 61 hearts prepared for cGMP analysis.

3.1 Cardioprotective effects of sGC stimulator BAY 41-2272

In Series 1, we investigated the infarct-limiting properties of the sGC stimulator BAY 41-2272 and explored its dependency on endogenous NO and its effects on myocardial cGMP concentration.

3.1.1 Infarct size

Baseline haemodynamic data are presented in Table 1. All experimental groups displayed comparable flow (CFR) and functional (HR, LVD...
RPP) parameters at the end of the pre-ischaemic stabilization period. The ischaemic area at risk of infarction for these groups was 44–55% of total myocardial volume with no statistically significant differences between groups. Under control conditions, infarct size was 31.5±2.8% of the ischaemic risk zone. Treatment at reperfusion with BAY 41-2272 affected a concentration-dependent reduction in infarct size, maximal at 3 μM the highest concentration employed (17.0±2.1%, P < 0.05) (Figure 2A). The cGMP inhibitor, ODQ 2 μM, which oxidizes the haem group of the NO binding site in sGC, abrogated the need for the haem site to be in its reduced state, although ODQ had no effect on infarct size per se (Figure 2B). However, in the presence of the NO synthase inhibitor L-NAME 100 μM, infarct limitation was still afforded by BAY 41-2272 (20.5±2.5%, P < 0.05 vs. control). In the presence of C-PTIO 30 μM, BAY 41-2272 also produced a significant limitation of infarct size (23.6±0.9%, P < 0.05 vs. control). Neither L-NAME nor C-PTIO perfused alone had any effect on infarct size (Figure 2C).

3.1.2 Myocardial cGMP concentration

cGMP measurements for Series 1 are presented in Figure 2D. In parallel groups of hearts subjected to 35 min left coronary artery occlusion and 10 min reperfusion, treatment with BAY 41-2272 (from 5 min before until 10 min after reperfusion) elicited an approximately two-fold elevation of cGMP content in both left ventricle subjected to ischaemia-reperfusion (17.76±1.87 vs. 9.44±0.61 fmol/mg tissue, P < 0.01) and normoxically perfused right ventricle (28.79±3.01 vs. 13.69±0.50 fmol/mg tissue, P < 0.01). It is of interest to note that the cGMP content of naive (non-perfused) hearts was significantly higher in right ventricle than in left ventricle samples (17.87±2.56 vs. 11.28±0.54 fmol/mg tissue, P < 0.01).

3.2 Cardioprotective effects of exogenous NO

In Series 2, we explored the effects of exogenously administered NO, from the donor compound NOC-9, on limiting infarct size when administered during early reperfusion.

3.2.1 Infarct size

The baseline haemodynamic parameters and area at risk for all groups in this series were comparable between groups (Table 1). NOC-9 perfused across the concentration range 1 nM–1 μM at early reperfusion limited infarct size in a concentration-dependent manner from 34.7±1.6% in control hearts to 20.5±1.3% (P < 0.001) in hearts treated with NOC-9 at the highest concentration (Figure 3A).

3.2.2 Myocardial cGMP concentration

To further explore the relationship between BAY 41-2272-induced cardioprotection and NO, cGMP measurements were made in hearts perfused with BAY 41-2272 and concomitantly with either the NO synthase inhibitor L-NAME or the NO scavenger C-PTIO. Tissue samples from concomitant BAY 41-2272 and L-NAME perfused LV had cGMP levels 48% lower than those perfused with BAY 41-2272 alone (11.15±0.91 vs. 17.76±1.87 fmol/mg tissue, P < 0.05). Similarly, cGMP levels in LV tissue perfused with both C-PTIO and BAY 41-2272 were 54% lower than BAY 41-2272 alone (8.29±0.52 vs. 17.76±1.87 fmol/mg tissue, P < 0.001) (Figure 3B).

3.3 Cardioprotective effects of sGC activator BAY 60-2770

In Series 3, we investigated the infarct-limiting properties of sGC activator BAY 60-2770 which targets the oxidized Fe3+ and haem-free forms of the protein.

3.3.1 Infarct size

Baseline haemodynamic parameters and area at risk for all groups in this series were comparable between groups (Table 1). The sGC activator BAY 60-2770 afforded protection across the concentration range 5 nM–1 μM during early reperfusion from 33.0±2.6% in control hearts to 22.0±2.8% (P < 0.01) for hearts treated with the highest concentration. However, no concentration effect was observed (Figure 4A).

To confirm the haem independence of BAY 60-2770’s action further hearts were co-perfused with 2 μM ODQ. Concomitant perfusion of ODQ with the lowest concentration of BAY 60-2770 (5 nM) limited...
infarct size to 17.6 ± 2.0% (P < 0.01 vs. control) (Figure 4B). Perfusion with ODQ 2 μM alone did not afford protection (32.9 ± 2.0%) (Figure 4B), an effect comparable with that seen in Series 1. Reperfusion with the NO scavenger C-PTIO produced infarct sizes similar to controls (32.0 ± 2.8%) (Figure 4B). Furthermore, C-PTIO did not abrogate the protection afforded by BAY 60-2770 (22.2 ± 2.2% vs. 33.0 ± 2.6%, P < 0.01) (Figure 4B). To investigate the protective effect of targeting both the reduced and oxidized/haem-free forms of sGC, we concomitantly perfused BAY 60-2770 5 nM and BAY 41-2272 1 μM. This combination resulted in only a modest 21% reduction in infarct size (24.8 ± 2.7%, P < 0.05 vs. control) (Figure 4B).

### 3.3.3 Myocardial cGMP concentration

Tissue levels of cGMP were measured in hearts that had been perfused with or without BAY 60-2770 and subjected to 35 min regional ischaemia. Measurements were also made in hearts perfused with BAY 60-2770 concomitantly with ODQ or a sub-maximal concentration of BAY 41-2272. In LV samples, cGMP levels were not elevated in tissue that had been perfused with BAY 60-2770 compared with samples that had not (12.60 ± 1.65 vs. 9.20 ± 0.70 fmol/mg tissue). Tissue samples perfused with concomitant BAY 60-2770 and ODQ had cGMP levels 60% higher than those perfused with BAY 60-2770 alone (20.16 ± 2.25 vs. 12.60 ± 1.65 fmol/mg tissue, P < 0.01). An increase of 36% was also seen in LV samples perfused with both the sGC stimulator and activator compared with the activator alone (17.1 ± 1.90 vs. 12.60 ± 1.65 fmol/mg tissue, P < 0.05), an increase of 86% compared with untreated hearts (17.11 ± 1.90 vs. 9.20 ± 0.70 fmol/mg tissue, P < 0.05) (Figure 4C).

### 4. Discussion

The principal findings of these studies can be summarized as follows.

1. Targeting the reduced form of sGC during early reperfusion with the stimulator BAY 41-2272 afforded concentration-dependent infarct limitation and this protection was independent of endogenous NO, demonstrated with concomitant perfusion of BAY 41-2272 and the NOS inhibitor L-NAME or NO scavenger C-PTIO. Tissue cGMP concentrations during early reperfusion...
were elevated by the sGC stimulator. The observed elevations were partly independent of NO.

(2) Perfusion of the NO donor NOC-9 during early reperfusion demonstrated concentration-dependent infarct limitation. Targeting the oxidized state of sGC with BAY 60-2770 also afforded infarct limitation. However, the protection observed was not increased beyond that seen with the sGC stimulator. Concomitant perfusion of both the sGC stimulator and activator did not afford greater protection than either treatment alone.

(3) Concomitant perfusion of BAY 60-2770 and ODQ elevated total cGMP concentration beyond BAY 60-2770 only perfused myocardium.

(4) Infarct size and left ventricular cGMP content measured at 10 min reperfusion analysed for correlation did not meet significance, suggesting that elevation of total LV cGMP levels are not imperative to afford protection (Figure 5).

NO/sGC/cGMP signalling during early reperfusion has been shown to be a tractable target to limit infarct size. However, the literature is inconsistent in reporting the protective effects of NO donors. Elevating cGMP levels during the first minutes of reperfusion is recognized as a contributory mechanism in the reperfusion injury salvage kinase signalling paradigm. Previous work by us and others has documented
that administration of natriuretic peptides during early reperfusion limits infarct size in both the rabbit and rat. Furthermore, this protection was mediated through elevated cGMP concentration and PKG-dependent mechanisms. The sGC activator BAY 58-2667 (Cinaciguat) has been shown to limit infarct size at reperfusion, again by elevating cGMP concentrations. This demonstrates that the oxidized and/or haem-free forms of sGC are present in the myocardium during reperfusion and can be targeted to elevate cGMP.

To explore the infarct-limiting properties of pharmacological agents that elevate cGMP, we used both a sGC stimulator and a sGC activator. Our initial results demonstrated that perfusion of the sGC stimulator during early reperfusion limited infarct size in a concentration-dependent manner. It was then demonstrated that the protection afforded was at least in part as result of elevated cGMP levels during early reperfusion. To characterize the mechanistic action of BAY 41-2272 further, ODQ was concomitantly perfused which abrogated the protection afforded by the stimulator alone. This is in agreement with the cell-based studies reported by Stasch et al. who reported that sGC stimulation by BAY 41-2272 was haem-dependent requiring the haem moiety to be in the Fe^{2+} state for enzymatic activity.

The relaxant effects of BAY 41-2272 to rat tracheal rings was dampened in preparations pre-incubated with L-NAME or ODQ. To investigate the NO component of BAY 41-2272 action, the sGC stimulator was perfused concomitantly with either L-NAME or C-PTIO. In both instances the infarct-limiting effects of BAY 41-2272 could not be completely abrogated, confirming that in terms of infarct limitation, the stimulator affords protection independently of NO. The results do however suggest that there is an NO component to the protection observed.

Stasch et al. reported that PTIO at high concentrations could not block sGC stimulation. Since reports that exogenous NO can afford infarct limitation when perfused during early reperfusion have been inconsistent, we carried out experiments using NOC-9, a rapid release NO donor. We were able to demonstrate a concentration-dependent action of the NO donor, limiting infarct size by half at the highest concentration. The current data are supported by previous experiments that demonstrate the need to recruit eNOS in reperfusion salvage.

Using the sGC activator BAY 60-2770, we investigated the infarct-limiting effects of targeting what may be regarded as a pathological state of the enzyme. Cohen et al. and Krieg et al. demonstrated that perfusion of the structurally similar BAY 58-2667 could limit infarct size in both the rat and the rabbit in a global model of ischaemia-reperfusion. Our results support this work and confirm that there indeed must be a component of the so-called pathological sGC present during early reperfusion. Infarct size limitation reported by Cohen et al. and Krieg et al. was much more marked than in the present study, matched by a larger percentage increase in cGMP. In contrast to our own studies; however, the basal concentrations of cGMP measured were much lower. Additionally, we perfused hearts with the sGC activator concomitantly with ODQ. These results confirm the haem-independent action of BAY 60-2770 and suggest that the concomitant therapy limited infarct size beyond the activator alone. More convincing were the cGMP measurements made in comparable experiments which showed that oxidizing the haem group with ODQ rendered the enzyme more sensitive to the activator, demonstrated by the larger elevation of total LV cGMP following concomitant perfusion.

**Figure 3** Infarct size data for NOC-9 concentration response (A), expressed as infarct to-risk ratio %. Data are means ± SEM. *P < 0.05, **P < 0.01, ***P < 0.001 vs. control (one-way ANOVA). Total cGMP concentrations in LV (solid) and RV (open) myocardial tissue samples (B). *P < 0.05, **P < 0.01, and ***P < 0.001 vs. 10’R BAY 41 (one-way ANOVA) n = 5 – 12.
To explore the benefits of targeting both redox states of sGC during early reperfusion, we co-perfused submaximal concentrations of both the sGC stimulator and the activator to ascertain whether targeting both redox states could afford greater infarct limitation than either treatment alone. Our results demonstrate that in terms of infarct limitation, concomitant perfusion of both compounds does not potentiate the protection observed by the other compound. Furthermore, infarct size was greater when both compounds were perfused together compared with the sGC activator alone. A possible explanation for these results could lie in physical or chemical interactions of the compounds at the target site on sGC. Although the sites of action of both compounds differ, they may impede each other’s action when in close proximity. However, the subsequent results refute this speculation with elevation of LV cGMP measured in comparable experiments. Concentrations were significantly elevated above those of BAY 60-2270 only treated hearts, comparable with those of hearts perfused with the sGC stimulator BAY41-2272. These data suggest that both compounds were able to elicit a response in the presence of the other. Previous work from our laboratory demonstrated that the cGMP analogue 8-Br-cGMP had a deleterious effect on infarct size at high concentrations, suggesting that excess cGMP may inhibit the protective effect of lower concentrations.²²

Figure 4 Infarct size data for BAY 60-2770 concentration response (A), concomitant perfusion of the haem site oxidiser ODQ, NO scavenger C-PTIO, and BAY 41-2272 (B) expressed as infarct to-risk ratio %. Data are means ± SEM. *P < 0.05, **P < 0.01 vs. control (one-way ANOVA). Total cGMP concentrations in LV (solid) and RV (open) myocardial tissue samples (C). ***P < 0.01 and ****P < 0.001 vs. 10'R BAY 60, ##P < 0.05 and ###P < 0.01 vs. 10'R (one-way ANOVA) n = 5–18.
For the first time, we plotted total LV cGMP levels against infarct size across a range of corresponding treatment groups (Figure 5). Although there is a trend towards infarct limitation with increased total LV cGMP concentration, correlation analysis did not reach significance ($P = 0.07$). This data set suggests that affording protection goes beyond gross elevation of LV cGMP. Most surprising are the results for LV cGMP measurements in hearts perfused concomitantly with BAY 41-2272 and either L-NAME or C-PTIO. Although a modest protective state is achieved, cGMP levels are not elevated above baseline. This suggests that the gross elevation seen in BAY 41-2272 only treated hearts requires the presence of NO. Similarly, hearts protected with BAY 60-2770 did not show a significant elevation in total LV cGMP, whereas those perfused concomitantly with ODQ did.

There is a growing body of evidence to suggest that cGMP is compartmentalized within discrete subcellular domains, regulated by specific PDEs and guanylyl cyclases. We can only speculate that affording infarct limitation by targeting cGMP requires specific targeting of discrete pools and not gross LV elevation. What we may be missing in the current data set is that total LV cGMP may not be elevated, yet protective specific pools maybe, without net elevation. Conversely, the ability to resolve local protective cGMP elevation may be masked in groups where total LV cGMP is elevated. This has led us to make reference to the notion that there may be specific ‘cytoprotective cGMP pools’.

This thesis is yet to be explored in the ischaemia-reperfusion setting as the ability to visualize/quantify changes in discrete subcellular cGMP is in its infancy. In addition, how the redox balance of sGC shifts during early reperfusion and how it affects distribution and production of cGMP would also be worthy of exploration and support this thesis.

In conclusion, these studies demonstrate that targeting sGC during early reperfusion following simulated AMI is a tractable target to limit infarct size. Using cGMP elevating compounds provides a more specific target in comparison with NO donors which are associated with vascular tolerance and non-desirable effects. The current study corroborates in vitro mechanistic studies. These data do not directly support the hypothesis that targeting the so-called pathological state of sGC is a more attractive target during early reperfusion; nevertheless, quantifying the spatial differences of the sGC redox state, the extent of shift during the critical moments and the specific localized changes in cGMP concentration remains desirable.

**Supplementary material**

Supplementary material is available at Cardiovascular Research online.

**Conflict of interest:** J.-P.S. is currently a full-time employee at Bayer Pharma AG. J.-P.S. holds more than 60 patent applications related to sGC stimulators, such as BAY 41-2272, and sGC activators, such as BAY 58-2667.

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