Herein we report the synthesis of a library of thioalkyl- and thioaryl-pyridazinediones for thiol-based self-immolative release of cargo. A bisthioaryl-pyridazinedione is shown to be stable to serum protein albumin but unstable in intracellular conditions. A derivatised analogue underwent self-immolative degradation in cellular thiol conditions as evidenced by LC-MS/release of a turn-on fluorescence fluorophore; versatility of the thiol-pyridazinedione is demonstrated through synthesis of SMDC precursors that contain three different functional groups on the same central molecule.

Traditional chemotherapy relies on the premise that administration of highly potent cytotoxic drugs will preferentially neutralise rapidly dividing cells over normal ones. However, this form of therapy is invariably associated with a lack of specificity for tumours, often resulting in undesirable side effects. Furthermore, traditional low molecular weight cytotoxic agents utilised in such treatments display non-preferential or minimal accumulation at the tumour site in vivo, which has been explained by factors such as a tumour's irregular vasculature, the high interstitial pressure at the site, and the tumour's overexpression of multidrug-resistant proteins. In an effort to overcome the unfavourable characteristics associated with using conventional chemotherapy, important advances have been made in the past decades on tumour-targeted drug delivery systems (TTDDs), which are designed to selectively deliver toxic cargo in intracellular conditions through the action of a cleavable linker. From the outset, we wanted to exploit the fact that thio-substituted PDs can undergo substitution (addition–elimination) in the presence of glutathione (GSH) via a novel thiol-induced self-immolative linker that could be used for the site-selective modification of proteins in various contexts. In this work, we developed the pyridazinedione platform for application in an SMDC setting (Fig. 1), including the creation of a novel thiol-induced self-immolative linker that could be used for the attachment and subsequent controlled release of cargo.

Two key considerations for an SMDC are that it is stable to blood thiois (primarily serum albumin) and is capable of releasing cargo in intracellular conditions through the action of a cleavable linker. From the outset, we wanted to exploit the fact that thio-substituted PDs can undergo substitution (via addition–elimination) in the presence of glutathione (GSH); as GSH exists in high concentration intracellularly but in a low concentration extracellularly, it is often used as the trigger in the design of a cleavable linker. Thus, if the thiol groups on a thio-substituted PD could be manipulated into being self-immolative linkers, there exists an opportunity for an intracellular release mechanism of cargo if the thio-substituted PD is stable to reaction with serum albumin; there is also literature precedent on thiolate released self-immolative linkers.

Electronic supplementary information (ESI) available. See DOI: 10.1039/c9cc08744c

\[ \text{Targeting entity} \quad \text{Spacer} \quad \text{Linker} \quad \text{Cytotoxic drug} \]

Fig. 1 Schematic representation of an SMDC, typically consisting of a targeting ligand, a linker and/or spacer, and a cytotoxic drug payload.
In view of the above, the balance between stability of the thio-substituted PDs to blood thiols whilst maintaining reactivity to intracellular GSH conditions was of great importance (Scheme 1). As such, our study began with the synthesis of mono-BrPD and di-BrPD (see ESL† see Fig. S2 and S3), as conduits to generate and appraise a variety of ary1, alkyl and mixed ary1-alkyl thio-substituted PDs (Fig. 2). This selection was made in order to gauge which groups were more stable in serum and whether the mono- or di-substitution would significantly affect reactivity. The five thio-substituted PDs (2–6) were synthesised from the BrPDs in moderate to excellent yields without any optimisation (see ESL† Fig. S4–S9).

After the formation of PDs 2–6 (Fig. 2), we appraised their stability in serum-mimicking conditions. All five synthesised PDs 2–6 were incubated with human serum albumin (HSA), the blood’s most abundant thiol, at pH 7.4 for 4 h, to determine if there was any reactivity with the protein’s accessible cysteine residue. Gratifyingly, no reaction was observed between HSA and all of the trialled PDs, corroborating previously reported results that alkyl thio-PDs are stable in blood-mimicking conditions and unveiling any thio-PDs to also be stable in the time-frame of the experiment. As a positive control, maleimide was incubated with HSA alongside PDs 2–6 under identical conditions. Maleimides are substantially more reactive towards thiols than PDs and are known to react with HSA, therefore conjugation of HSA by maleimide was expected, and indeed observed. As some SMDCs, such as those based on folic acid for instance, have a blood circulation half-life of only 26 min,25 the substantially longer incubation time of 4 h demonstrates excellent stability of all the PDs in human blood-serum-like conditions in this context.

We next explored the stability of PDs 2–6 in the presence of GSH 1, due to it being predominant in the intracellular milieu and its ability to cleave SMDCs to liberate free drug in other systems.10,31,32 It was hoped that PDs 2–6 would undergo an addition–elimination mechanism with GSH 1 to expel the thioaryl and thioalkyl moieties. As the concentration of intracellular GSH 1 falls in the range of 1 mM to 10 mM,31,34–36 an intermediate intracellular GSH concentration of 5 mM was considered appropriate for the study. Moreover, since the HSA studies employed an excess of PDs (10 eq.) to demonstrate their stability in blood, their expected instability when exposed to intracellular GSH called for GSH 1 to be used in excess (10 eq.), to mimic the large excess of this thiol inside a cell relative to the intracellular drug concentration. Thus, it was chosen for PDs 2–6 to be at a concentration of 0.5 mM relative to GSH 1. As many SMDCs are internalised by cells via receptor-mediated endocytosis (RME), the drug conjugate must necessarily pass through the early endosome before being degraded in the late endosome/lysosome. The pH extremes of the early endosome and late endosome/lysosome are around 6.5 and 4.9–5.0 respectively and thus it was decided to incubate PDs 2–6 with GSH 1 at pHs 6.5 and 5.0 with LC-MS readings taken at time points up to 24 h.

The studies at pH 6.5 were conducted first (see ESL† Fig. S32–S56), as it was reasoned that if substitution were to take place, as was desired, it would have a greater chance of occurring at the higher of the endosome pH extremes. Bis-thioaryl PD 4 proved to be the superior candidate of all the five tested (Scheme 2). Reaction of PD 4 could give rise to two possible PD substitution products: mono-GSH PD and bis-GSH PD 7. After 4 h, only bis-GSH PD 7 was observed in the LC-MS trace, suggesting that the reaction had reached completion.

When all the studies were conducted at the lower pH extreme of 5.0, the consumption of 2, 3, 5 and 6 was negligible relative to that of PD 4 (see ESL† Fig. S77–S81).

Both pH studies thus confirmed bis-thioaryl PD 4 as the best candidate of the five to be brought forward in derivatising the thioaryl moieties of its PD backbone for the purpose of creating a GSH-triggered cleavable, self-immolative linker. To do this, we considered the work of Van de Bittner et al., who developed a self-immolative phenolic-based construct that releases luciferin.41 Gois et al. modified this phenolic-based linker to release umbelliferone upon self-immolative degradation.42 Taking inspiration from both of these systems, we decided to apply a related, but novel, strategy to the thiophenol system of PD 4, which required the synthesis of umbelliferone–PD 13. To this end, we synthesised PD 8, which could be brominated using N-bromosuccinimide (NBS) and azobisisobutyronitrile (AIBN) to make di-bromo intermediate 11. Displacement of the bromine groups with the phenolic functionality on umbelliferone 12, a commonly used “turn-on” fluorescent probe (not fluorescent when phenol is substituted with anything other than H), was then carried out to obtain umbelliferone–PD 13 (Scheme 3).

As the GSH studies had confirmed that these intracellular thiol conditions did displace the aryl moieties of related PD 4, it was anticipated that subjecting umbelliferone–PD 13 to 5 mM GSH at pH 6.5 would displace its thioaryl groups in a similar manner. If so, thiophenol-based construct 14 should be expelled, which we hoped would undergo self-immolative cleavage to yield fluorescent umbelliferone (Scheme 4). Gratifyingly, when umbelliferone–PD 13 was incubated with GSH 1 at pH 6.5 for 24 h, expulsion of umbelliferone 12 was observed by LC-MS, confirming the self-immolative mechanism based on thiophenol. It was also pleasing that construct 14 was...
Disappointingly however, a reaction suspension was observed upon incubation of PD 13 with GSH 1, suggesting that umbelliferone–PD 13 was not fully soluble in a THF/ammonium acetate buffer mixture; this may also have gone some way to explain the lack of full conversion of PD 13 after 24 h under the reaction conditions. As such, it was pertinent to tailor the structure of umbelliferone–PD 13 in an effort to make it more water-soluble.

It was envisaged that adding a carboxylic acid group to the 3-position of each umbelliferone moiety in umbelliferone–PD 13 would aid water solubility. To this end, 3-tert-butyl carboxylic acid umbelliferone was synthesised prior to base-promoted reaction with di-bromo species 11 to afford a tert-butyl protected umbelliferone–PD, which was then stirred in 25% TFA/CH₂Cl₂ at 21 °C for 4 h to yield the desired carboxylic acid–umbelliferone–PD 16 (see ESI, † Fig. S12–S14).

With carboxylic acid–umbelliferone–PD 16 in hand, it was then appropriate to repeat the incubation with GSH 1 at pH 6.5, to test whether the addition of the carboxylic acid groups to the structure aided water solubility. Gratifyingly, nearly full consumption of PD 16 had occurred after 4 h, with no solubility issues observed. The analogue of intermediate 14 was observed at t = 0.05 h in low levels, but was consumed by the 4 h time point. Moreover, the main peaks observed by LC-MS after the 4 h time point were bis-GSH PD 7 and the released fluorophore 17 (see ESI, † Fig. S84–S86). This provided confirmation that PD 16 has a similar reactivity profile to model PD 4, and that greater solubility was key to this correlation. As well as the aforementioned structures being observed by LC-MS, the fluorescence emission spectrum of the reaction was also monitored, with the fluorescence increasing sharply between the t = 0.05 h and 4 h time points, with a less significant increase between the t = 4 h and 24 h time points (Fig. 3). Taken together, the LC-MS and fluorescence emission data confirm the turn-on fluorescence nature of this reaction, which could have important applications in cell imaging.

With the self-immolative breakdown of a PD’s thiolate moieties demonstrated, attention turned to substituting the N-alkyl portions of a BrPD into useful functional moieties. To this end an acid group and a simple PEG were chosen (Scheme 5).

To synthesise di-bromo PEG acid PD 20, PEG ester hydrazine 18 was synthesised using sequential alkylations of hydrazines followed by refluxing with dibromomaleic acid 19 to yield PD 20 in 74% yield. The reason for adding the PEG functionality was to aid solubility in order to counteract the attachment of any hydrophobic group that may be used in an SMDC (e.g. drug, fluorophore, porphyrin, etc.). The acid group can be readily functionalised with a plethora of commercially or readily available
amine-bearing cargos that are relevant to SMDCs. This functionalisation was demonstrated by synthesising di-bromo PEG azide PD 21 and di-bromo PEG alkyne PD 22, as azide and alkyne groups are very useful for “click” modification. With these amine-functionalised PDS in hand, the final step in synthesising SMDC precursors involved substituting the bromine atoms for a derivatisable thiol, namely 4-methylbenzenethiol 23 (Scheme 6).

PDs 24 and 23 were obtained in 82% and 85% yield respectively. They constitute SMDC precursors where bromination of the methyl thiol (possibly after “click” modification) could be used for fluorophore/drug installation, and where the azide/alkyne groups serve as “click” handles for incorporating tumour-targeting ligands.

In conclusion, we have demonstrated the development of a novel, sulfur-based self-immolative linker that was utilised to incorporate a turn-on fluorescence fluorophore that is only released from the PD construct by GSH under conditions mimicking intracellular conditions, which in some cases is similar to certain tumour microenvironments. The first PD–fluorophore construct (13) suffered from suboptimal buffer solubility, and this issue was solved by installing a thiol (possibly after “click” modification) could be used for fluorophore/drug installation, and where the azide/alkyne groups serve as “click” handles for incorporating tumour-targeting ligands.

Conflicts of interest
There are no conflicts to declare.

Notes and references