Silica nanoparticles functionalised with cation coordination sites and fluorophores for the differential sensing of anions in a quencher displacement assay (QDA)

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In conjunction with quenching metal ions, silica nanoparticles carrying coordination sites and fluorophores were employed for the differential fluorometric recognition of anions.

The development of chromo- and fluorogenic chemosensors for anions has received growing interest during the past years due to the fundamental roles that anions play in chemical and biological processes. The majority of these chemosensors incorporate molecular receptors of supramolecular chemistry design. Traditionally, supramolecular anion sensing utilizes either colour and/or fluorescence changes in probes with a "binding site--signalling subunit" architecture or undergoing chemodosimetrical reactions and indicator release in displacement assays (IDAs) for signal transduction. However, recently a novel trend in chemosensor development has emerged which uses organic-inorganic hybrid materials, in many cases nanoparticles. These hybrid systems often show synergistic effects which are hardly achievable with molecular systems or inorganic solids alone.

Among different inorganic supports, silica nanoparticles have been extensively used for hybrid sensing materials due to their straightforward preparation and surface functionalization and high stability in aqueous suspensions. As an alternative to chromo- and fluorogenic probes, self-assembled structures on surfaces in particular have attracted attention recently. Especially the independent functionalization of inorganic surfaces with receptor and fluorophore units is an appealing approach that overcomes the synthetic problems usually immanent to the classical “binding site--signalling subunit” architecture. Moreover, signal amplification and cooperative binding effects associated with the independent anchoring of receptors and fluorophores in close proximity to each other and to the surface of the support are noticeable features of this strategy, allowing for instance intercommunication between both subunits without the need for a direct covalent link between them. Employing silica nanoparticles functionalised with fluorophores and suitable coordination sites, this approach has recently been used for the selective recognition of metal cations. In contrast, hybrid nanoparticles for the optical recognition of anions are much less explored.

Our present hybrid system for the optical differential recognition of anions is based on the competitive coordination to a metal centre. Since one of our interests in such assays is the facile regeneration of the sensing material, we modified the conventional IDA concept. Instead of displacing an indicator dye with the analyte from a binding site, which requires a two-step regeneration procedure of analyte removal and subsequent indicator reloading (and sometimes also the spatial separation of indicator from the host–guest complex for detection), we attached a fluorophore (F) and a receptor (R), which does not bind to the analyte but to a mediator, separately to the support. A quenching metal ion (Q), which is at the same time a good binder for the target anion, then served as the mediator, i.e., as the displacable species, arriving at a quencher displacement assay (QDA, Scheme 1). In the first step of the protocol, the R units of the highly fluorescent receptor- and fluorophore-functionalised nanoparticles (RFNs) are loaded with Q, switching off the hybrid’s fluorescence. Second, addition of the target analyte, which shows a higher affinity for Q than R does, displaces Q from R, leading to revival of the fluorescence of the RFNs. In a third step, regeneration of the material is simply accomplished by reloading Q onto the RFNs. For the title material, terpyridine units as R for the quencher metal ions Q and sulforhodamine B as signalling units F were independently anchored onto the surface of the silica nanoparticles.

The synthesis of the reactive precursors 1 and 2 (Scheme 1)
and the functionalised nanoparticles is described in the ESI. The contents of terpyridine and rhodamine amount to 30.84 and 3.9 mmol (mol[SiO₂])⁻¹ in the terpyridine–sulfhorodamine-functionalized nanoparticles (TSNP) and that of rhodamine to 3.9 mmol (mol[SiO₂])⁻¹ in the nanoparticles functionalized only with 2 (SNPs). Both, TSNP and SNP show a homogenous particle size of 20 ± 2 nm (Fig. 1). As a rough estimate, each coated TSNP carries up to 2100 attached molecules. The average distance between two subunits thus amounts to ca. 8 Å.

The critical role of the terpyridine receptor was demonstrated in parallel studies using the model SNPs that are only functionalised with F but do not contain R. The addition of metal ions to acetonitrile suspensions of SNPs induced negligible changes in the emission of the dye, indicating that the presence of the R units in SNP is critical for the quenching mechanism. Only when coordinated to terpyridine are the metal ions in a spatial proximity with the fluorophore that is close enough to induce quenching.

Having established the signalling mechanism the corresponding TSNP–Q hybrid materials (Q = Cu²⁺, Fe³⁺, Hg²⁺, Ni²⁺ and Pb²⁺) were tested as sensing systems for anions. The underlying idea of our approach relies on the ability of the anion to successfully compete with the terpyridine units for the metal ion and displace the latter form the R units, resulting in an enhancement of the fluorescence. This enhancement should be a direct consequence of a delicate balance between the binding strength of the metal ion with terpyridine and the affinity of the cation for the added anion.

To test the ability of the TSNP–Q ensembles for differential recognition of small monovalent anions, acetonitrile solutions of TSNP–Q chemosensors were prepared by adding 1 equivalent (with respect to the 2.0 × 10⁻⁴ mmol R units contained, see ESI for details) of the corresponding metal ion to quench the rhodamine emission. Then, 0.1–10 equivalents of the anions H₃PO₄⁻, HSO₄⁻, F⁻, Cl⁻, Br⁻, I⁻ and NO₃⁻ were added to the TSNP–Q suspensions. Except in one case (see below), no uniquely selective fluorescence changes were found. However, as a general trend the addition of H₃PO₄ and F⁻ induced a remarkable fluorescence enhancement, addition of Cl⁻, Br⁻ and HSO₄⁻ induced a moderate increase of the emission whereas the addition of I⁻ and NO₃⁻ induced only poor fluorescence changes. In addition, despite this general observation a clear modulation of the relative response in the presence of anions was found, depending on the metal ion employed. This behaviour is reminiscent of other probe–metal ion–anion ternary complex systems and is a well-suited prerequisite for differential recognition. Qualitative analysis of this differential response for anions using PCA (Principal

![Scheme 1](image) Design concept of the quencher displacement assay (QDA) involving terpyridine–sulfhorodamine-functionalised nanoparticles (TSNPs) and metal ion quenchers (Q).

![Fig. 1](image) a) TEM image of TSNPs, average diameter = 20 nm; b) fluorescence spectra of TSNP and SNP suspended and sulfhorodamine B dissolved in acetonitrile.
Component Analysis) algorithms yielded the results summarized in Fig. 2. As can be deduced from the figure, recognition patterns can be identified for all the anions studied even for those that display a rather poor change in fluorescence. Fluorescence lifetime studies of the ternary systems revealed that upon addition of anions, the lifetime features of the TSNP–M$^{2+}$ ensembles are gradually turned into the features of TSNP, i.e., the major 2.2 ns component of the sulforhodamine is recovered, pointing to an actual displacement of the metal ion from the terpyridine units.

![Fig. 2 Principal component analysis (PCA) score plot for the anions as indicted using TSNP–Q (Q = Fe$^{3+}$, Hg$^{2+}$, Cu$^{2+}$, Ni$^{2+}$ and Pb$^{2+}$) ensembles. Data shown from three different trials and 1:5 TSNP–Q to anion ratios. PC axes are calculated to lie along lines of diminishing levels of variance in the data set.](image)

Whereas the TSNP–Q systems, with Q = Cu$^{2+}$, Fe$^{3+}$, Hg$^{2+}$ or Ni$^{2+}$, show a differential but no selective response in the presence of small inorganic anions, TSNP–Pb$^{2+}$ behaves different, i.e., only the addition of H$_2$PO$_4^-$ revives the sulforhodamine fluorescence whereas other anions remain largely passive (Fig. 3). Using this procedure, concentrations as low as 5 ppm of H$_2$PO$_4^-$ can be detected.

A final control experiment revealed that the addition of anions to acetonitrile suspensions of TSNP (in the absence of Q) or SNPs did not result in changes of the dye’s fluorescence, pointing out the importance of the quenching metal ion as the mediator in the sensing paradigm. A further benefit of the system is obvious from Scheme 1. Once an analytical reaction has been performed and Q displaced, regeneration is easily accomplished by addition of an appropriate metal ion salt with a non-coordinating anion like perchlorate.

In summary, we have prepared a hybrid organic-inorganic sensing ensemble by grafting fluorophores and receptors onto the surface of silica nanoparticles and its use in anion recognition. Coordination of metal ions at the terpyridine units induced quenching of the neighbouring fluorophores. Upon addition of anions, coordination to the metal centres and a partial recovery of the fluorescence was observed. The final response of a certain TSNP–Q system to a particular anion is a delicate balance between the binding strength of the metal ion with the terpyridine and the affinity of the cation for the added anion. This allows differential recognition of small inorganic anions in a quencher displacement assay. Furthermore, regeneration of the system is possible in a straightforward manner. The facile and independent functionalisation of silica surfaces with various chemical entities and the possible use of a number of different coordination units and metal ions make this approach highly appealing for the search for new chemosensors for anions.

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### Notes and references

12. Higher valent anions naturally induce similar changes as verified with [(CH$_3$)$_2$N]$_2$(SO$_4$). However, because of the significantly lower solubility of such salts in CH/CH$_2$, the interference is negligible.