



**An Overview on Nanoparticles Commonly Used in
Fluorescent Bioimaging**

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<p>Cite this: <i>Chem. Soc. Rev.</i>, 2015,</p> <p>Received ... Sep. 2014 DOI: 10.1039/xxxx</p> <p>www.rsc.org/csr</p>	<h2 style="margin: 0;">An Overview on Nanoparticles Commonly Used in Fluorescent Bioimaging</h2> <p style="margin: 0;">Otto S. Wolfbeis</p> <p style="margin: 0;">The article gives an overview on the various kinds of nanoparticles (NPs) that are widely used for purposes of fluorescent imaging, mainly of cells and tissue. Following an introduction and a discussion of merits of fluorescent NPs compared to molecular fluorophores, labels and probes, the article assesses the kinds and specific features of nanomaterials often used in bioimaging. These include fluorescently doped silicas and sol-gels, hydrophilic polymers (hydrogels), hydrophobic organic polymers, semiconducting polymer dots, quantum dots, carbon dots, other carbonaceous nanomaterials, upconversion NPs, noble metal NPs (mainly gold and silver), various others nanomaterials, and dendrimers. Another section covers coatings and methods for surface modification of NPs. Specific examples on the use of nanoparticles in (a) plain fluorescence imaging of cells, (b) targeted imaging, (c) imaging of chemical species, and (d) imaging of temperature are given next. A final section covers aspects of multimodal imaging (such as fluorescence/nmr), imaging combined with drug and gene delivery, or imaging combined with therapy or diagnosis. A <i>Supporting Information</i> gives specific examples for materials and methods used in imaging, sensing, multimodal imaging and theranostics such as imaging combined with drug delivery or photodynamic therapy. The article contains 270 references in the main part, and 157 references in the <i>Supporting Information</i>.</p>
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1. Introduction

The term imaging can be understood in many ways. Imaging is kind of photographing in most people's perception. Scientific imaging goes far beyond this. Images can additionally be created by such diverse methods as (near) infrared and Raman spectroscopy, nuclear magnetic resonance (often referred to as magnetic resonance imaging; MRI), radioimaging using respective nuclides, CT imaging, positron emission tomography, electrochemical imaging using rastering electrodes, by mechanical methods such as AFM, and by even more sophisticated scanning methods such as laser ablation ICP-MS MALDI-MS and the like. It has become accepted that virtually any method yielding a 2-dimensional picture (that, ideally, is presented in pseudo-colors) can be referred to as "imaging". Many of these methods are destructive or require extensive sample preparation, but others are not and therefore well applicable to living systems or intact tissue. The use of nanoparticles as contrast agents for in-vivo bioimaging using MRI probably is the largest single field of applications but this topic is not covered in this review. The purpose of this tutorial is to give the reader an overview on the wealth of nanomaterials that do exist for use in fluorescent imaging, and to assist in making decision as to which material may be selected to solve a specific problem.

Fluorescence (and phosphorescence^{*}) based imaging have found particular interest because these spectroscopies are sensitive, selective, rich in contrast, and versatile. The past 20 years also have experienced

^{*} The common term *fluorescence* is used here for the sake of simplicity even if the term *phosphorescence* may apply.

an enormous increase in resolution that has arrived at the single nanometer scale. One may first differentiate between two kinds of fluorescent imaging. The first involves imaging based on *intrinsically* fluorescent (bio)chemical species (such as NADH in tissue, crude oil in - and on - seawater, or chlorophyll in all kinds of plants and the open sea). The second covers methods for imaging of samples or cells that have been made fluorescent by adding synthetic fluorescent probes, labels, nanoparticles or nanosensors. The use of such probes is, in fact, indispensable in order to detect species that are not amenable to direct fluorometric imaging (such as of pH) but this also implies the risk of local perturbation by the probe or material added.

The acquisition of images of biological matter by using fluorescent probes or fluorescent labels and nanomaterials is generally referred to as bioimaging and forms a large field of its own. Letting aside conventional (light) microscopy and MRI, fluorescence imaging probably is the most widespread method in biosciences. Respective pictures are attractive, easily comprehensible, and can be found in a good fraction of research papers and magazines. Reviews cover topics such as fluorescent nanostructures for bioimaging,¹ quantum dots in bio-imaging^{2,3} and single molecule imaging ("one quantum dot at a time"),⁴ gold nanoclusters with tunable fluorescence as bioimaging probes,⁵ aggregation-induced emission-based fluorescent nanoparticles,⁶ nanocomposite particles for bioapplications including imaging,⁷ on nanoparticles in drug delivery, therapeutics, diagnostics and imaging,⁸ on quantum dots and polymer hybrid composites as fluorescent switches and turn-on probes for sensing anions,⁹ on aspects of deep tissue microscopy and optical imaging,¹⁰ on controlled synthesis, spectra and bioapplications of lanthanide-doped luminescent nanoprobles,¹¹ on advances in (NP-based) fingerprint imaging,¹² on the intersection of CMOS microsystems and upconversion NPs for bioimaging and bioassays,¹³ to mention only a few. A review by Bünzli¹⁴ on lanthanide luminescence for biomedical analyses and imaging contains a section 5.6. on improving sensitivity by using certain nanoparticles. Others are cited later in the respective sections and in the Electronic Supporting Material.

The term fluorescence does not imply a single spectroscopic method but rather includes a variety of techniques in that images can be acquired by measurement of intensity, decay time (lifetime) and polarization, but also by studying effects caused by resonance energy transfer, (dynamic) quenching, or photo-induced electron transfer. Optical imaging was limited to resolutions of a few 100 nm until about 1995, but substantial thrust in terms of resolution resulted from the availability of fluorescent methods of imaging on a nanoscale by methods such as STED, PALM, or STORM, all based on laser technology. Chemo- and bioluminescence also can generate images¹⁵ but chemiluminescence requires the addition of reagents and usually is unidirectional in that an *increase* in the concentration of an analyte can only be monitored. Both methods are time-dependent.

When focusing now on bioimaging based on synthetic fluorescent probes and nanoparticles, one may differentiate between three techniques. (1) In the most simple one, a strong fluorophore or fluorescent nanoparticles are internalized into cells so that they can be imaged. The only purpose of such fluorophores and nanomaterials is to render cells or tissue fluorescent. They do not possess (and are not expected to possess) affinity for a specific site, nor do they respond (like indicator probes) to the presence of chemical species such as certain ions or organic molecules. (2) The second technique is referred to as "targeted bioimaging". It enables specific domains or species to be detected, very much like immunostaining or fluorescence in-situ hybridization. In order to accomplish this, fluorophores or nanoparticles are applied whose surface has been properly functionalized, for example with receptors, ligands, antibodies or oligomers so to recognize the specific counterpart. Examples include targeting of tumor markers, genes, mitochondria, membranes, or the amyloidic plaques in Alzheimer-associated tissue. (3) The third technique is making use of probes and nanomaterials with sensing capability. This enables (bio)chemical species to be imaged that are to not intrinsically fluorescent. Examples include imaging of the distribution of chemical species such as pH values, glucose, calcium(II) or oxygen in the living and metabolizing cell, if not in tumor cells or in cells exposed to candidate drugs. This group also involves nanosensors for temperature. Representative examples for each of these techniques will be presented in Section 5.

2. Fluorescent nanomaterials and nanoparticles versus molecular fluorophores, labels and probes

The availability of nanomaterials for purposes of imaging has generated a variety of methods for imaging, with features including improved brightness (defined as absorbance times quantum yield), inertness to their

microenvironment and a more even distribution (unless targeted imaging of certain domains is desired, of course). Nanoparticles (NPs), in contrast to molecular probes, often are not cytotoxic and do not suffer from nonspecific binding by cellular biomacromolecules or unwanted sequestration. Binding of molecular probes by cellular proteins (or sites) can affect both the optical properties of the probe and even the function of the protein or binding site. Dyed NPs, or intrinsically fluorescent NPs, in contrast, are virtually inert and do hardly interact with cellular proteins nor are their optical properties affected by the proteins outside. Not surprisingly, all known NPs have photostabilities that are distinctly better than those of molecular probes.

Many NPs can be easily internalized into cells and tissue (depending on charge and surface chemistry; positive charged facilitate internalization) and can be even targeted to specific sites. Compared to fluorescent proteins one notes the more simple handling of NPs and more predictable results. Many kinds of NPs are commercially available. The simplicity of loading or labeling with fluorophores or NPs is a particular issue if hundreds of cell lines are being handled simultaneously, for example in high-throughput screening. Second-harmonic generation (which results in low background noise) is more easily accomplished with NPs as demonstrated in the review by Dempsey et al.¹⁶ on respective nanocrystals for in vivo imaging, in particular on nanodiamonds, nonlinear crystals, quantum dots and SERS NPs. Finally, it is fair to say that nanosensors have calibrations plots that are quite similar if not identical if acquired in vitro and in vivo. Molecular probes, in contrast, are quite different in that respect. It is a matter of fact that one must never use a calibration plot that was established in plain buffer solution to quantify a parameter with data acquired in a cellular system.

3. Kinds of nanomaterials often used in bioimaging, and their specific features

A complete coverage of all the nanomaterials used so far in bioimaging would by far exceed the frame of this review but rather fill a book. The following is a discussion of the nanomaterials most often used for purposes of bioimaging. These include NPs made from silica and organically modified silica, hydrophobic and hydrophilic organic polymers, semiconducting organic polymers, quantum dots, carbonaceous nanomaterials including carbon (quantum) dots, carbon nanoclusters and nanotubes, nanodiamonds, upconversion materials, metal particles, metal oxides and others. The discussion on the potential cytotoxicity of NPs is going on, and numerous studies have been performed as to the potentially harmful or perturbing effect of NPs on physiological systems.¹⁷

3.1. Fluorescently doped silicas and sol-gels

Silica nanoparticles (SiNPs) were among the first one to be used in bioimaging. An excessive literature on their uses does exist.¹⁸ The group of Wiesner have reviewed¹⁹ the state of the art in SiNPs for use in sensing/imaging and have described numerous kinds of SiNPs, among them the one-pot synthesis of PEGylated mesoporous and fluorescent SiNPs possessing a single pore, tunable sizes of around 9 nm, and narrow size distributions.²⁰ In fact, mesoporous silica (in addition to materials related to "*controlled porous glass*") are more often used now because they can be heavily loaded with fluorescent dyes, photosensitizers or diagnostic reagents as will also be shown in Section 6. Fluorescent mesoporous silica can be obtained via hydrothermal reaction and functionalized, if desired, with reactive siloxanes.²¹ Such particles possess low cytotoxicity and excellent cell imaging capability.

SiNPs can be easily doped with various kinds of organic, metal-organic and metallic fluorophores, and emission wavelengths range from 300 to 1,000 nm, with a trend towards NPs possessing longwave (> 600 nm) emissions²² because the fluorescence of NPs at wavelengths of >500 nm is often interfered by the autofluorescence of cells. Color, decay times and size are widely tunable, and dopants can be hydrophobic, hydrophilic, or ionic. SiNPs also may be coated with fluorophores but attention has to be paid to possible aggregation effects that usually are accompanied by self-quenching. Two-photon excitation has been demonstrated but this depends on the kind of fluorescent label. SiNPs and their aggregates of >300 nm in size cause strong light scattering. SiNPs are fairly well biocompatible (i.e., not harmful to cells and tissue). Cell permeability depends on zeta potential. If negatively charged, they hardly pass cell membranes. They are virtually nontoxic, easily excreted (unless particle sizes exceed 50 nm), but also quickly coated by intracellular proteins and attacked by the immunosystem. Their surface can be easily modified with a variety of coatings and using standard silica (and sol-gel) surface chemistry. Particles do not swell but tend to aggregate in presence of bivalent ions unless inert shells are deposited on the surface. Mesoporous silica structures warrant strongly increased surface areas and enable high loading of cargo for cellular imaging and

targeting. The state of the art in designing and characterizing fluorophore-doped SiNPs for bioapplications has been reviewed.²³

SiNPs and other NPs are often doped with luminescence lanthanides. Their use in bioimaging has been reviewed by Chen et al.,²⁴ and the fabrication of down-converting and up-converting luminescent probes for optical imaging by Zhang et al.²⁵ Lanthanide-based fluorophores have attractive features such as (a) relatively long decay times which facilitate time-resolved fluorometry and, thus, efficient background suppression; (b) fairly narrow emissions; (c) single-photon and two-photon excitation; and (d) upconversion luminescence and downconversion luminescence. Lanthanide doped SiNPs are obtained by incorporating lanthanide ions, or by grafting the SiNPs with complexes such as Eu(III), Tb(III) or Gd(III).²⁶ Such fluorescent NPs have decay times between 0.35 and 1.87 ms (which facilitates gated spectroscopy), do not photobleach, and display the typical narrow emission bands of lanthanide ions. Other examples for lanthanide doped NPs (also with materials other than silica) will be given in the following sections and in the Tables. In addition to silica, NPs made from titania and zirconia and doped with organic and inorganic fluorophores also are known but less often used. They are treated in section 3.11 (on *Other Nanomaterials*).

Sol-gels and organically modified sol-gels (*ormosils*) may also be used for form NPs. A large variety of materials are known. Their porosity can be governed by proper choice of materials and by varying experimental conditions such as acid or base catalysis. Sol-gels are prepared by polycondensation of tetraalkoxysilanes, and ormosils by copolymerization of mixtures of mixtures of tetraalkoxysilanes with alkyl-alkoxysilanes in varying ratios, or from alkyl-alkoxysilanes only. A large variety of materials is known depending on whether mono-, di- or trialkylsilanes of general formula $(R_1)_x(R_2O)_{3-x}Si-R_1$ (where R_1 is alkyl or aryl, and R_2 is alkyl) are being used. The NPs usually are made fluorescent by non-covalent doping with various kinds of fluorophores. Unless pore sizes are small, covalent immobilization is advised, however. Colors, decay times and particle size are widely tunable. As with SiNPs and others, fluorescence at >500 nm is often interfered by autofluorescence of biomatter. Beads and bead aggregates of >300 nm in size cause strong light scattering. Sol-gel NPs are well biocompatible which can be improved by PEGylation (see Section 4). Cell permeability depends on zeta potential in that NPs are hardly cell permeable if negatively charged. Sol-gel NPs are nontoxic and – if small enough – are easily excreted. Their surface can be modified ex-vivo with a variety of coatings, and the surface chemistry of silica and sol-gels is well established. Particles do not swell but tend to aggregate in presence of bivalent ions if their zeta potential is negative. Photobleaching of the dopant dye can be an issue even though this is often weaker than in case of dyes that are molecularly distributed in cells.

3.2. Hydrophilic polymers

Hydrophilic materials for use in NP-based imaging include the various kinds of hydrogels, but also natural products such as cellulose. Nanogels are soft and usually water soluble. If NPs are to be formed, they have to be crosslinked or mixed with another material in order to form hybrid materials. Hydrogels are well permeable to ions and hydrophilic organic species such as glucose or amino acids, but not to large biomolecules usually. Typical polymers include polyacrylamide (PAA), polyurethanes, poly(hydroxyethyl methacrylamide) (pHEMA), certain poly(ethylene glycols) or specialty polymers such as PluronicTM [a commercial poly(ethylene glycol)-co-poly-ethyleneoxide)] widely used in NP-assisted drug delivery. The design of fluorescent organic nanostructures for bioimaging has been reviewed.²⁷ Depending on the kind of polymer and the degree of crosslinking, such NPs undergo substantial (an ionic strength-dependent) swelling in water, but they do not tend to aggregate. The emission wavelengths of respective NPs can be adjusted to almost any wavelength between 300 and 1,000 nm via the dopant, usually an organic or metal-organic fluorophore whose color, fluorescence, decay time and size is widely tunable.

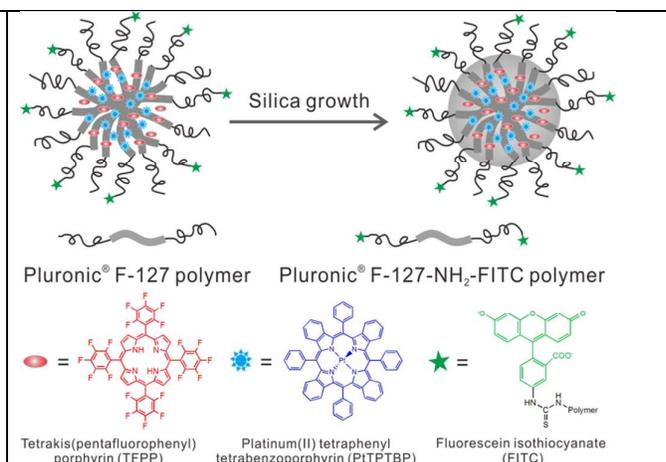
NPs made from organic monomers can be crosslinked to a various extent, and this determines many of their properties including swellability and solubility. The fluorescence of particles at <500 nm often interfered by autofluorescence of most cells. NPs prepared from hydrogels are biocompatible, cell permeable (depending, as always, on the zeta potential), nontoxic, slowly excreted, fairly quickly coated by intracellular proteins and attacked by the immunosystem. Many of them can be degraded by intracellular enzymes. Their surface cannot be easily chemically modified with additional coatings. Functionalities such as amino groups are better introduced by adding a functional monomer to the main monomer and then to initiate radical polymerization. Techniques are known to prepare organic polymer core/shell NPs. Fluorescent (and other) dopants tend to leach into the aqueous environment of the particle unless firmly retained (electrostatically or

covalently). Amino-modified cellulose was applied in a luminescent sensor for high-resolution imaging of pH *in vivo*.²⁸ pH values were imaged by detecting the green fluorescence of the pH probe fluorescein covalently linked to aminocellulose. A ruthenium phenanthroline complex was incorporated in poly-acrylonitrile beads to give a pH-independent red reference signal. The beads were immobilized in a polyurethane hydrogel on a thin transparent support. Both *in-vitro* and *in-vivo* experiments revealed the versatility of the method during physiological and chronic cutaneous wound healing. The method was later extended to simultaneously image extracellular wound pH and oxygenation *in-vivo*.²⁹ The same pH beads were used, and poly(styrene-co-acrylonitrile) particles dyed with Pd(II)-meso-tetraphenyl-tetrabenzoporphyrin were added to give a near-infrared signal that depends on local oxygen partial pressure.

A typical recent example of the use of a PAA hydrogel is provided by nanoparticles containing free amino groups that were prepared by copolymerization of acrylamide and methylene-bisacrylamide with 3-aminopropyl-acrylamide and labeling the terminal amino groups with pH probes such as fluorescein, Oregon Green, Alexa 633, and others. The resulting sensor NPs cover a wide range of pH (4.0 - 8.0) which is needed in certain situations even in cellular imaging.³⁰ In addition to the widely used crosslinked polyacrylamides (PAAs), other acrylamides including polymethacrylamide or N-alkylacrylamides were employed. For example, core-shell microgels containing indicators were fabricated³¹ by two-stage free radical precipitation polymerization of N-isopropylacrylamide. The shell of the microgel exhibits a low critical solution temperature and undergoes a transition from a swollen state to a de-swollen state, associated with a hydrodynamic radius of ~450 nm at 25°C (*in vitro*) and of ~190 nm at 37°C (*in vivo*). The microgel readily enters the cytosol which makes it a potential candidate for the delivery of indicator probes into the cytosol.

The PluronicTM hydrogel (see above) was used to fabricate nanosensors for fluorescent imaging of physiological pH values.³² Features include (a) very small diameters (12 nm); (b) biocompatibility due to the use of a hydrogel kind of material, and (c) lack of toxicity. The nanosensors were incorporated into an agar film to enable continuous monitoring of the pH value of bacterial cultures, and thus of their growth. Dually responding nanosensor particles were reported that were prepared from an organic-inorganic composite (PluronicTM reinforced with silica) and used for simultaneous imaging of oxygen and pH in the cellular cytosol.³³ Fig. 1 shows a schematic of the preparation of the dual nanosensor for oxygen and pH, the architecture of the NPs, and the chemical structures of the probes used. Other multiple (bio)sensors, *i.e.* sensors capable of two or more analytes simultaneously have been reviewed,³⁴ but only a moderate fraction of them is making use of nanomaterials.

Fig. 1. Structure of a nanosensor for dual sensing of oxygen and pH. Its core consists of Pluronic F-127, a nonionic, surfactant triblock copolymer composed of a central hydrophobic chain of poly(propylene oxide) flanked by two hydrophilic chains of poly(ethylene glycol) (PEG) and reinforced with silica. The NPs are capped with PEG. The oxygen probe (PtTBTBP) and the reference fluorophore TFPP are located in the core, and the pH probe (FITC) is conjugated to the terminal ends of the PEG capping. Reproduced from ref.³³ (X. Wang et al., *J. Am. Chem. Soc.* (2012), 134, 17011-17014) with permission (2014) of the Am. Chem. Soc. *****



Nanogels (like NPs) are of interest in being extremely soft materials that take up water in fraction between 10 and 90%. The gels are well permeable to hydrophilic species and can be made fluorescent by simple labeling with inert labels and made responsive by attaching a fluorescent probe. Nanogel particles are well suited to image pH values inside cells.³⁵ In a method termed CLARITY, nanoporous hydrogel-hybridized forms of intact mouse brain were prepared and crosslinked to a three-dimensional network of hydrophilic polymers.³⁶ They are optically transparent and permeable macromolecules. Tissue imaging is said to reveal local circuit wiring, cellular relationships, subcellular structures, protein complexes, nucleic acids and neurotransmitters. CLARITY also enables intact-tissue *in-situ* hybridization,

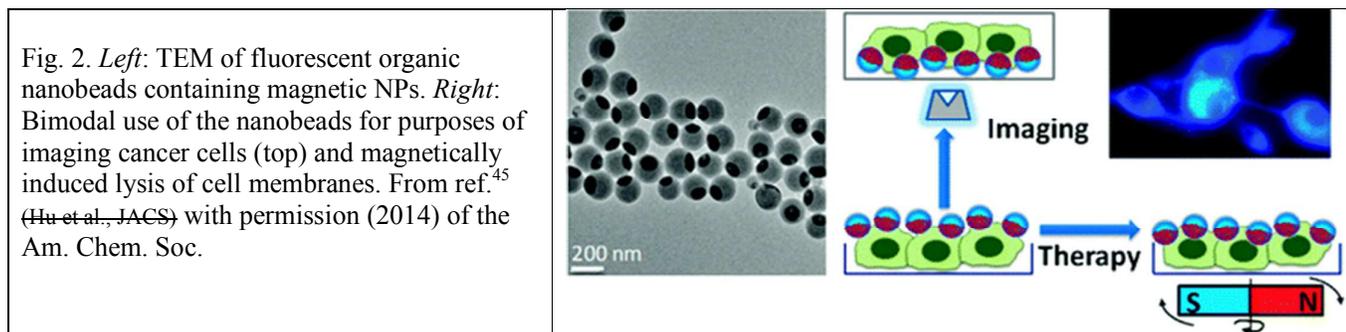
immunohistochemistry in non-sectioned tissue, and antibody labelling. Fluorophore-labeled polymeric nanogels for sensing temperature (T) have attracted much interest because they pave the way to sense T inside cells. The topic has been extensively reviewed.³⁷

3.3. Hydrophobic organic polymers

Polystyrene nanoparticles (PS-NPs) are highly hydrophobic and can be doped with apolar fluorophores with emission peak wavelengths that range from the near UV to beyond 1000 nm. Their color, decay times and size also are widely tunable. Doping with lipophilic materials is preferred because ionic probes are poorly soluble in hydrophobic NPs from which the probes tend to leach out. As in all kinds of NPs for use in imaging, any fluorescence occurring at below 500 nm is interfered by the autofluorescence of biomatter. An (organic) dopant fluorophore can photobleach if exposed to strong laser light. PS-NPs are fairly biocompatible (i.e., not harmful to cells and tissue), fairly well cell permeable, nontoxic, and their excretion is slow. If placed inside cells, they are only slowly coated by intracellular proteins and hardly attacked by the immunosystem. The modification of their surface is limited to certain functional groups. Functionalities (such as amino groups) are best introduced by addition of co-reagents containing such groups to the monomer before starting emulsion polymerization. Post-modification and additional coating is rather difficult. PS-NPs do not measurably swell in water and do not readily aggregate, but this depends on their charge and zeta potential. One of the first nanomaterials for sensing purposes consisted of (pH-insensitive) fluorescent PS beads coated with polyaniline whose absorbance of pH dependent over a large range of pH values. Depending on the actual pH value, the coating screens off the emission of the beads.³⁸ PS-NPs are well permeable to gases but impermeable to charged species including proteins. Particles with an average diameter of 85 nm were loaded with an oxygen-quenchable luminescent ruthenium complex and then used to image oxygen inside cells following 2-photon excitation.³⁹

Polyacrylonitrile (PAN) NPs can be doped with fluorophores with emission wavelengths ranging from 300 to 1,000 nm, preferably with hydrophilic dyes. Color, decay times and size widely tunable. Beads and aggregates of >300 nm in size cause light scattering. PAN is fairly biocompatible (not harmful to cells and tissue) and fairly well cell permeable, nontoxic, easily excreted but slowly coated by intracellular proteins. It is hardly attacked by the immunosystem. Its surface is rather inert and cannot be readily modified once the particles have been formed, usually by precipitation by adding water to a solution of PAN in dimethylformamide. PAN particles do not swell but there is a tendency to aggregation. Fluorophore-doped PAN-NPs were applied, for example, to referenced imaging of pH and temperature with sub- μm spatial resolution.⁴⁰

Biocompatible fluorescent organic NPs with tunable photoluminescence were obtained via one-pot oxidation of polydopamine and subsequently utilized for cell imaging,⁴¹ and water dispersible red fluorescent organic NPs for use in cell imaging were reported by Luo.⁴² Quantum-dots conjugated to dopamine function as redox coupled assemblies and can be applied to in-vitro and intracellular pH sensing.⁴³ Other polymers include poly(vinyl butyral)⁴⁴ that was labeled with a perylene dyes that is easily taken up without coating and does not display in vitro cytotoxicity on human cancer cells. Hu et al.⁴⁵ have introduced a class of organic nanocomposites with functionalities for both fluorescence imaging and magnetic therapy. Magnetic NPs (mNPs; 5 nm in diameter) were incorporated into the amphiphilic block copolymer poly(styrene-*b*-allyl alcohol) that was labeled with pyrene. The fluorescence of the resulting NPs (200 nm i.d.) was exploited when imaging cancer cells, while magnetically controlled mechanical damage of cell membranes represents a way for cancer cell treatment (referred to as magnetolytic therapy). Magnetic field induced heating may pave, in future, the way to hyperthermal cancer therapy. This is schematically shown in Fig. 2. For numerous other examples, see section 6.



3.4. Semiconducting (organic) polymer dots (P-dots)

These come in addition to more conventional (dye) NPs such as those made from polystyrene, polyacrylamide etc. The polymer usually is prepared from aromatic precursors possessing polymerizable double or triple bonds. Particles (as needed for imaging) are prepared by either emulsion polymerization or nanoprecipitations.⁴⁶ Doping with fluorophores is not needed. The backbone of conjugated polymers behaves like an array of light-harvesting units that exhibit a larger optical cross section compared to small organic molecule dyes. Photobleaching was not reported so far. Fine-tuning of the conjugated polymer structure and polymeric encapsulation matrix leads to fluorescent probes with specific spectral properties and targeting capability. P-dots display strong fluorescence that often extends far into the NIR, are highly inert and do not swell in water. Little is known about biocompatibility, internalization, and excretion from tissue. Their uses in imaging and therapy have been reviewed.⁴⁷

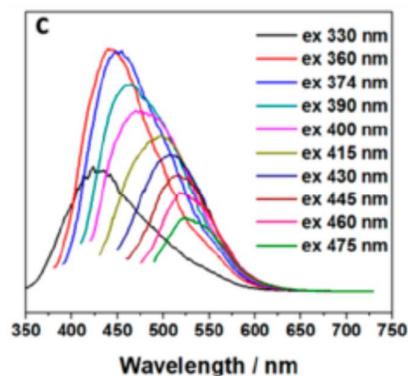
Fluorescent nanodots consisting of semiconducting polymer blends can be attached to peptides (such as chlorotoxin)⁴⁸ and then can be used for targeted imaging (of malignant brain tumors, for example) in clinical diagnosis. By coupling the pH-indicator fluorescein to P-dots, a material is obtained that display two fluorescence peaks, one being pH sensitive, the other not so that it can act as an internal reference. Fully reversible pH sensing was demonstrated⁴⁹ for the pH 5.0 to 8.0 range. Intracellular pH values were determined by imaging of HeLa cells following the uptake of the P-dots by endocytosis. Tetraphenylethene-based fluorescent organic NPs undergo aggregation-induced emission inside cells and this was monitored via cell imaging.⁵⁰ The fluorescence of conjugated polymers, particularly if anionic, can be quenched by ions such as Cu(II).⁵¹ Semiconducting P-dots (20 to 50 nm) also can serve as photoacoustic probes for real-time imaging of reactive oxygen species in living mice tissue where they accumulate quite readily.⁵²

3.5. Carbon dots

Carbon dots (C-dots)⁵³, first reported in 2006, are said to be clusters of carbon atoms (for definitions see ref.⁵⁴) with diameters of typically 2 to 8 nm, but also contain substantial fractions of oxygen and hydrogen if not nitrogen. They do not measurably swell in aqueous solution but aggregation was occasionally observed. C-dots can be made strongly fluorescent and need not be doped or labeled. Their emission color can be tuned to some extent by varying the experimental conditions of synthesis. Both the excitation and emission spectra are very wide and usually extend from the UV to the red (650 nm), a fact that virtually excludes their use in multiplexing. A fine review on the synthesis and photophysical properties (and uses in bioimaging) of C-dots is available.⁵⁵ It includes the very true statement that "C-dots have a much more comprehensive definition compared to graphene quantum dots." The QYs of C-dots range from 5 to 30%,

Fig. 3 shows the emission spectra of carbon dots at different excitation wavelengths from 330 to 475 nm. Their strongest fluorescence is blue, but longwave excitation (at >460 nm) induces green to yellow emission. A review on the synthesis, properties and applications of C-dots contains an interesting section on the origins of their excitation wavelength dependent emission, and particularly the controversial upconverted luminescence.⁵⁶ The emissions also are likely to be pH-dependent. Single-particle fluorescence intensity fluctuation ("blinking") has been reported recently.⁵⁷ Decay times are in the nanosecond time regime and do not vary much. The fluorescence of C-dots can be of the upconversion and the down-conversion type.

Fig. 3. Wavelength dependence of aqueous solutions of carbon dots in water at excitation wavelengths between 330 and 475 nm. Reprinted from ref.⁵⁸ with permission (2014) of the Am. Chem. Soc.



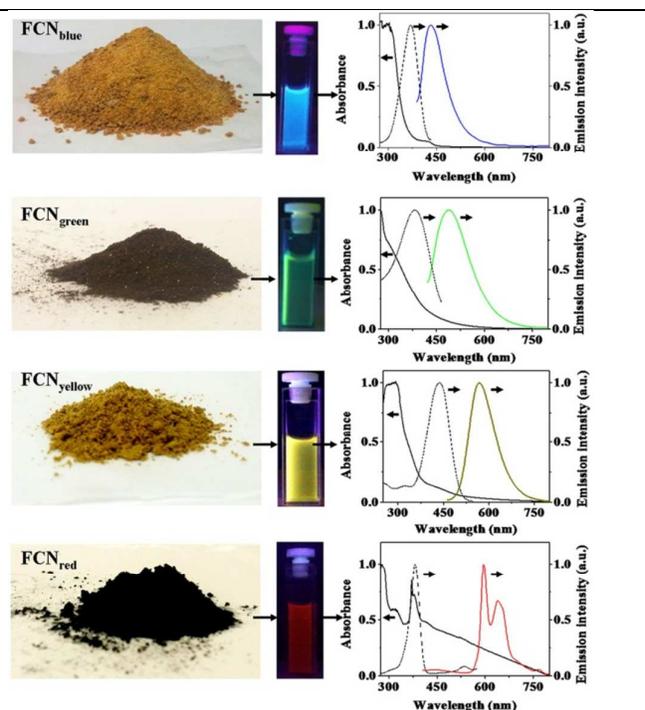
C-dots, and carbon nanoparticles in general, can be single-photon excited and multi-photon excited. C-dot-based inorganic–organic nanosystems were applied, for example, to two-photon imaging and biosensing of pH variation in living cells and tissues.⁵⁹ C-dots are fairly well biocompatible (i.e., not harmful to cells and tissue within a few days), fairly well cell permeable, not known to be toxic, easily excreted, weakly interacting with proteins, and hardly attacked by the immunosystem. Functional surface engineering for purposes of bioconjugation and imaging is more difficult than in case of Q-dots but possible.⁶⁰ They neither swell nor photobleach. Their fluorescence is pH dependent and quenched by iodide⁶¹ (and probably by other notorious quenchers too). A recent review covers the subject.⁶²

Both top-down and bottom-up synthetic approaches are known. The resulting C-dots, in fact, always contain substantial fractions of oxygen (up to 50%) and also nitrogen if a nitrogen-containing substance such as an amino acid is added during synthesis. And yet, they are often confused or termed – mainly by Chinese authors – with graphene quantum dots even though graphene by definition consists of C and H only and is nonfluorescent. Examples where the application of "graphenes" is claimed but materials other than sp²-graphenes have been used include, for example, intracellular fluorescence imaging with a "graphene"-based fluorescent probe,⁶³ the use of highly biocompatible "graphene" nanosheets for cellular imaging.⁶⁴ A particularly confusing example is represented⁶⁵ by an article entitled *The in-vivo and in-vitro toxicity of graphene quantum dots* that has nothing to do with graphene (which is free of oxygen by definition and non-fluorescent). The authors have prepared the highly fluorescent(!) graphene material by oxidation (!) and also claim it to possess a particularly high oxygen content(!).

C-dots have been prepared from numerous organic materials and natural products containing carbon in various form, one example being⁵⁸ (Wang *et al.*, *Anal. Chem.*, 2014) the preparation of 3-nm blue fluorescent C-dots from cow milk by heating it to 180 °C for 2 h. The particles can be used to image U87 cells. If C-dots are doped with nitrogen,⁶⁶ they are even more strongly fluorescent. Both hydrophilic and hydrophobic C-dots are known. Hydrophilic materials are preferred in imaging.⁶⁷ Hydrophobic materials are less often used but also available by microwave synthesis.⁶⁸ Raw C-dots are mainly prepared by microwave induced thermal carbonization of molecular precursors such as glucose (and other carbohydrates), citrate, (poly)glycols, often in the presence of a nitrogen source (such as tryptophan or EDTA). Surface passivated (und, therefore, bright) C-dots can be directly synthesized by microwave induced pyrolysis of glycerol in the presence of 4,7,10-trioxa-1,13-tridecanediamine.⁶⁹ C-dots also can be isolated from soot, or prepared from glucose with P₂O₅ at room temperature.⁷⁰ C-dots may be rendered to more strongly fluorescent by alkali or acid-assisted ultrasonic treatment.⁷¹

Photoluminescent C-dots also have been produced by laser ablation of graphite followed by oxidation with nitric acid and functionalization with diamine-terminated poly(ethylene glycol).⁷² They show multicolor fluorescence.⁷³ C-dots can be produced inexpensively and on a large scale. Fluorescence is conferred or improved by chemical treatment (or passivation) of the surface, for example by oxidation, doping with inorganics, or capping. Water dispersible C-dots with tunable photoluminescence can also be synthesized⁷⁴ via hydrothermal oxidation of nanodiamonds and were subsequently utilized for cell imaging. Carbon nanoparticles (10 nm i.d.) for use in fluorescent bioimaging can be obtained now⁷⁵ on the milligram to gram scale by carbohydrate carbonization (even though in our experience this method is difficult to reproduce not the least because of an inadequate experimental part). Table 1 gives examples of CNPs (prepared on a gram scale) along with colors of emission (from blue to red). Red-fluorescent CNPs are preferred because autofluorescence of biomatter is weaker in this spectral range.

Fig. 4. Images of gram scale solid samples of fluorescent carbon nanoparticles (FCNPs), of their solutions under appropriate excitations, and their absorption (—), excitation (---) and emission (color lines) spectra. Emission spectra have been measured by exciting at 370 nm for FCNP_{blue}, by exciting at 400 nm for FCNP_{green}, by exciting at 425 nm for FCNP_{yellow} and by exciting at 385 nm for FCNP_{red}. From ref.⁷⁵ (Bhunia et al.) with permission (2014) by Nature Publ. Group.



Both C-dots with invariable and with continuously tunable emission are known.⁷⁶ They enable ratiometric sensing of pH values via the ratio of the intensities of the excitation-independent and pH-independent blue emission and the excitation-dependent and pH dependent full-color emissions. Ratiometric (blue and green) fluorescent nanosensors have been described⁷⁷ that are based on water soluble carbon nanodots with multiple sensing capacities. This, however, is an euphemism for poor selectivity in that the dots respond to temperature, pH, Fe(III) ions, all of which mutually interfere.

Table 1. Properties of fluorescent carbon nanoparticles (FCNPs) with blue, green, yellow and red emission; containing between 10 and 33% oxygen and 1 to 8 % of nitrogen. From ref. ref.⁷⁵ (Bhunia et al.)

FCNP	Molecular mass or particle size	Composition (C:H:N:O; in %)	Emission peak and(excitation) wavelength	Molar absorbance at excitation wavelength	Quantum yield (%)
FCNP _{blue}	400 – 2200 Da	65:6:8:21	440 nm (370 nm)	$\sim 2 \times 10^3$	6 – 30
FCNP _{green}	2500 – 14000 Da, 2 – 4 nm	75:10:5:10	500 nm (400 nm)	$\sim 5 \times 10^4$	14
FCNP _{yellow}	1 – 4 nm	50:15:2:33	560 nm (425 nm)	$\sim 4 \times 10^3$	12
FCNP _{red}	~ 4 – 10 nm	70:5:1:24	600 nm (385 nm)	$\sim 7 \times 10^5$	7

3.6. Other Carbonaceous materials

Other nanosized carbon allotropes include fullerenes (C₇₀ mainly; these being much smaller than C-dots), and the larger species including carbon nanotubes (CNTs; single walled and multiwalled), nanodiamonds, graphene (which is nonfluorescent), and the oxidized species graphene oxide, reduced graphene oxide, graphite oxide, graphene quantum dots (often synonymously used for C-dots; see below) and the like. Other authors refer to their materials as graphene (or graphene dots) even if the material was prepared by reduction of graphite oxide (by Hummers method) and still contains large fractions of oxygen. It may be better termed "reduced graphite oxide". There is much confusion.

The fluorescence of graphite oxide⁷⁸ (like that of C-dots⁷⁹) is sensitive to pH. Like C-dots, other carbon nanomaterials need not be doped with fluorophores and are extremely photostable. The colors of emission of all known variants depend on the wavelength of excitation. Excitation in the UV (350 – 380 nm) often results in good brightness and blue fluorescence, but excitation wavelengths can be as long as 650 nm and fluorescence then occurs in the near IR as is shown below. The decay times of fluorescence are in the order

of nanoseconds. Nano-sized fluorescent graphite oxides (nano-GOs) with different size distribution were prepared via one-pot hydrothermal route using ultra-small graphite powder as starting material and subsequently separated using dialysis tubes with different molecular weight cutoff.⁸⁰ Such nano-GOs were found to be readily internalized by A549 cells and then located in the cytoplasm. They display size-dependent photoluminescence (green, yellow, red) and excellent biocompatibility.

Graphene⁸¹ and (color-tunable) fullerenes⁸² have not often been used for purposes of imaging because of their lack (or rather weak) fluorescence. Fullerene C₇₀ (unlike C₆₀ which is nonfluorescent) displays normal and delayed fluorescence which is strongly quenched by oxygen⁸³ and highly dependent on temperature.⁸⁴ Gong et al.⁸⁵ have isolated red, green and blue fluorescent hollow carbon NPs from chromatographic fractions and demonstrated them to be excellent (multi-color) probes for cellular imaging.

Stabilized graphene oxides were applied to cellular imaging,⁶³ (Peng) some after having been made highly biocompatible⁸⁶ or stabilized and biofunctionalized.⁸⁷ Blue and green fluorescent carbon NPs derived with vitamin B1 also are described to be very bright and applicable to cell imaging.⁸⁸ Luo et al. have reviewed applications of carbon-based quantum dots for fluorescence imaging of cells and tissues.⁸⁹ Fluorescent graphene quantum dots (GQDs) have been synthesized⁹⁰ that display both upconverted and excitation-independent downconverted photoluminescence. Doping of GQDs with almost any kind of heteroatom including nitrogen,⁹¹ boron,⁹² sulfur,⁹³ fluorine⁹⁴ and chlorine⁹⁵ enhance their brightness. By coupling them to photocatalytically active rutile (TiO₂/GQD) and anatase (TiO₂/GQD) systems, the complete visible spectrum of sunlight can be harnessed. Strong two-photon-induced fluorescence was reported⁹⁶ for photostable, biocompatible nitrogen-doped graphene quantum dots (N-GQDs; possibly C-dots?) for cellular and deep-tissue imaging. Their two-photon absorption cross-section reaches 48,000 Göppert-Mayer (GM) units, which far surpasses that of many organic dyes. It is comparable to that of the high performance semiconductor Q-dots and represents the highest value ever reported for carbon-based nanomaterials. The penetration depth in phantom tissue revealed an imaging depth as deep as 1.8 mm. Zhu et al.⁹⁷ have presented a study on surface chemistry based routes to modulate the photoluminescence of GQDs, how to govern fluorescence mechanism to induce up-conversion fluorescence, and on bioimaging applications.

Single-walled and multi-walled CNTs are fluorescent in the NIR but have low quantum yield.⁹⁸ The one-dimensional electronic structure of nanotubes results in sharp interband transitions in the absorption spectra of SWNTs, and in photoluminescence in the NIR region (800 – 1600 nm). These wavelengths include the tissue-transparent region of the electromagnetic spectrum. Their other properties are comparable to those of C-dots. Like C-dots, they do not photobleach. While traceable in living cells, their cytotoxicity is still not refuted,^{99,100} but carbon nanotubes encapsulated by a DNA oligonucleotide remain functional in live cells for up to three months.¹⁰¹ CNTs have been applied to Raman imaging, near-infrared (NIR) fluorescence imaging and photoacoustic imaging but still are less often used than some other fluorescent nanomaterials. The group of Strano¹⁰² have reviewed advances in molecular recognition based on single-walled CNTs and respective nanoengineered platforms. These were used to fluorescently sense species such as ATP, NO, H₂O₂, and glucose in cells.

Three-dimensional tracking of single-walled CNTs with an orbital tracking microscope was demonstrated.¹⁰³ The technique was applied to determine the viscosity regimes within live HeLa cells, and this was used to spatially map corral volumes (0.27–1.32 μm³), to determine active transport velocity (455 nm/s), and to calculate local viscosities (54 – 179 cP) within the cell. The NIR emission of CNTs (with their second window at 950 – 1400 nm) is attractive for in-vivo fluorescence imaging due to its deep penetration depth in tissues and low tissue autofluorescence. Genetically engineered multifunctional M13 phage were shown to assemble single-walled CNTs and ligands for targeted fluorescence imaging of tumors.¹⁰⁴

Nanodiamonds (NDs) like C-dots are being produced, as powders, by detonation synthesis on a commercial scale. They have low or no cytotoxicity.^{105,106} NDs doped with nitrogen are particularly bright. Synthetic NDs form fluorescent centers by thermal annealing and then have rather longwave peaks of excitation (~560 nm) and red emission (~700 nm). The intrinsic red fluorescence is strong enough to detect a single 35-nm ND in a cell.¹⁰⁷ Others types of NDs possess green fluorescence (peaking at 531 nm) and represent a promising alternative to semiconductor quantum dots (see above) because they are photostable, hardly toxic, easily excreted, and can be fairly easily bioconjugated. Hydrophobic derivatives of NDs possessing blue emission have been obtained by modifying the surface with long-chain alkyl groups.¹⁰⁸ NDs do not swell, and their size can be hardly tuned by chemical means.

3.7. Metal chalcogenide quantum dots (classical Q-dots)

These are typically made from combinations of zinc(II), cadmium(II), selenide and sulfide. Numerous additional components and dopants are known, and sophisticated methods have been developed to modify surfaces and to create additional shells. Q-dots have experienced an unprecedented success in imaging because of their unique properties. They need not be doped with fluorophores because their fluorescence results from a photonic quantum effect. The color of emission and their size are well tunable and interdependent. Decay times (in the low ns time domain), in contrast, do not vary much. Q-dots display Gaussian emission spectra (with FWHMs of typically 30 nm) and therefore have multiplexing capacity (like upconverting NPs but unlike C-dots). All Q-dots require photoexcitation at <500 nm where biomatter often strongly absorbs (this causing an inner filter effect), and fluorescence intensity strongly varies over time ("blinks"; however, non-blinking Q-dots have been described recently). The quantum yield (QY) of Cd/Zn based Q-dots are rather high (0.3 – 1.0) which is distinctly better than the QYs of upconversion NPs (see section 3.8.), for example. Dots and aggregates of >300 nm in size cause strong light scattering. Single photon, 2-photon and, recently,¹⁰⁹ 3-photon excitation and imaging have been demonstrated, and cross sections can be as large as 60,000 GM units.

Q-dots are cell toxic unless coated with inert shells, but passivation and reduced toxicity of CdS dots also was accomplished by coating them with DNA.¹¹⁰ They are fairly well cell permeable but clearance from tissue is difficult, partially because of their interaction with thiol groups of cysteines in proteins.¹¹¹ Their surface chemistry is well established, and several kinds of surface-modified Q-dots are commercially available. They do not swell or photobleach but fluorescence depends on temperature. They are uniquely suited for high resolution and multiplexed imaging of cells. Good reviews cover aspects such as on applications to fluorescence spectroscopy and imaging,¹¹² rendering them biocompatible,¹¹³ on the cytotoxicity of cadmium-based Q-dots,¹¹⁴ or on nucleic acid-passivated Q-dots acting as biomolecular templates of varying form and function.¹¹⁵ Recently work includes the application of Q-dot nanosensors to fluorescence lifetime imaging microscopy of intracellular pH,¹¹⁶ or of Q-dots loaded with fluorescent liposomes in order to perform fluorescence resonance energy transfer studies and NIR in-vivo imaging of mouse tissue.¹¹⁷ The fluorescence decay time of Q-dots becomes pH-dependent on coating them with NIR fluorescent dyes.¹¹⁸ The fluorescence of certain Q-dots is quenched Zn(II) and Cd(II) ions,¹¹⁹ and this paves the way for imaging such ions intracellularly. Q-dots may additionally be doped with other metal ions to give, for example, brightly fluorescent Mn-doped ZnS, Mn-doped ZnSe, or Cu-doped InZnS particles (10–80 nm in diameter) which represent a new class of fluorescent probes with low toxicity.¹²⁰

Aside from Q-dots composed of Zn(II), Cd(II), sulfide and selenide there are numerous other kinds of such particles. It is difficult to keep track with the variety of materials that have been presented in recent years. NPs with quantum effect also can consist of group III-V elements. Some are brightly fluorescent, one example being InP Q-dots functionalized with a Ln(III) chelate and coated with a cell-penetrating peptide for use as bimodal imaging agents (MRI and confocal microscopy).¹²¹ The reader is referred to some of the many reviews that exist on the use of Q-dots in bioimaging. Also see Tables 2 – 5.

3.8. Upconversion nanoparticles (UCNPs)

UCNPs (also referred to as upconversion nanocrystals) consist mostly of hexagonal NaYF₄ nanocrystals doped with trivalent lanthanide ions such as Er(III), Yb(III) or Tm(III). The dopant is the emitter and additional doping with fluorophores is not needed. UCNPs display several emission colors (with at least two strong bands in the visible) whose peak wavelengths depend on the kind of lanthanide dopant. However, single color emitting UCNPs consisting of lanthanide-doped KMnF₃ nanocrystals also have been reported.¹²² The size of UCNPs is widely tunable (typical sizes ranging from 10 to 100 nm) and affects quantum yields. The control of size and of the emission and excitation spectra is still a challenge.¹²³ The fact that UCNPs display multi-photon (bicolor) emissions also paves the way to method to increasing the resolution in microscopy.¹²⁴ Their dual (or multiple) emissions often enable referenced (2-wavelength) sensing and imaging but unfortunately the ratio of the two emissions (often green and red) varies with the coating and undergoes a change if hydrophobic particles (of the oleate type) are converted into hydrophilic particles (as preferably used in bioimaging). The use of upconversion NPs in bioimaging, therapy, drug delivery and bioassays has been reviewed.¹²⁵ Very recently it was shown¹²⁶ that coating UCNPs with a layer of silver causes metal-enhanced (plasmon) upconversion and a 30-fold increase in brightness compared to NPs without silver core. The NPs with a used for imaging of HeLa cells have 3-layer core-shell-shell architecture

of the type $\text{Ag}@\text{SiO}_2@\text{Lu}_2\text{O}_3:\text{Gd},\text{Yb},\text{Er}$. The silver induced plasmon enhancement mechanism in $\text{NaYF}_4:\text{Yb},\text{Er}$ NPs (Maxwell versus Förster) was studied in some detail.¹²⁷

Unlike in case of carbon dots, the color of the emission of UCNPs is independent of the excitation wavelength which is rather longwave (750 – 1000 nm). UCNPs with oleate capping (as obtained by the most popular method of synthesis) possess moderate brightness only, but those modified with hydrophilic coatings are much less bright (with QYs that hardly exceed 0.5% in water solution). QYs of 1 – 3 % have also been reported but only for bulk materials or for dried and aggregated particles. The seemingly poor QYs of UCNPs are still acceptable because their anti-Stokes emission allows fluorescence images to be acquired against a black background.

Remarkably, and unlike in case of Q-dots, the QYs of UCNPs also depend on the power density (Watt cm^{-2}) of the (cw) laser and on particles size, with smaller particles of the same kind displaying smaller QYs.¹²⁸ It is reminded that the QY of UCNPs is poorly defined.¹²⁹ If data are given, it shall also be stated whether these refer to a single emission band or the total emission (in either Stokes and anti-Stokes mode). In addition, the size of the particles investigated must be given along with power density, and how the inner filter effect of water solutions (under 980-nm excitation) has been taken into account. UCNPs, if coated with a shell of undoped NaYF_4 (and even silica), are much brighter in water than uncoated UCNPs. The group of Resch-Genger¹³⁰ has determined, in a solid study, the QY of oleic acid-coated UCNPs of type $\text{NaYF}_4:\text{Yb},\text{Er}$ to be 0.35% under well defined experimental conditions. Gargas et al.¹³¹ report on seemingly highly attractive nanocrystals (5 - 8 nm i.d.) for single-molecule imaging. The brightness under single-particle imaging conditions is said to be much higher than that of other compositions. However, the power density applied in the experiments is as high as $>10^6 \text{ W/cm}^2$ which hardly tolerated by any living organism. If such a power density is applied to watery samples, strong local heating will occur. In fact, if excited with conventional power density, the luminescence of these UCNPs is so weak that no spectra can be acquired.

UCNPs may also be applied to optically encode and to multiplexed imaging of cells and microspheres.¹³² The method may be extended to lifetime-based encoding by exploiting their tunable luminescence lifetimes which are in the microsecond time regime (in case of $\text{NaYF}_4:\text{Yb},\text{Er}$).¹³³ By exciting a single color band, one can generate more than ten excited state populations with lifetimes ranging from 25.6 μs to 662 μs and decode their well-separated lifetime identities which are independent of both color and intensity.

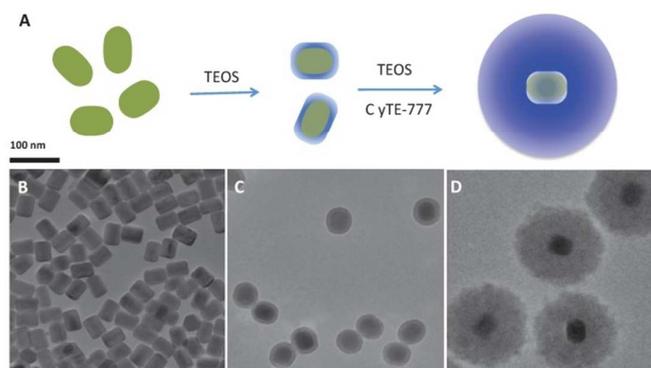
Host crystals other than NaYF_4 have been studied recently. Light management in UCNPs was demonstrated for ultrasmall NaGdF_4 nanoparticles core-shell architectures to tune their emission color.¹³⁴ In another example,¹³⁵ the brightness of core-shell nanocrystals ($\text{NaLuF}_4:\text{Gd},\text{Yb},\text{Er}$ coated with a shell of $\text{NaLuF}_4:\text{Yb}$) was found to be remarkable higher than that of inert-shell coated nanocrystals. The particles can be used to image HeLa cells. Even more complex UCNPs of the type $\text{NaLuF}_4:\text{Gd},\text{Yb},\text{Er}$ were synthesized recently and applied to bioimaging.¹³⁶

UCNPs are perfectly suited for bioimaging because fluorescence is not at all interfered by the autofluorescence of cells.¹³⁷ Moreover, they are fairly biocompatible and – if small enough – cell permeable.¹³⁸ They are not known to be toxic and easily excreted. They usually are weakly interacting with proteins but can be well conjugated to them.^{139,140} UCNPs are hardly attacked by the immunosystem, and their surface can be easily modified. They do not swell but tend to aggregate in presence of bivalent ions. Photobleaching cannot occur. Their luminescence is highly dependent on temperature¹⁴¹ and (rather unselectively) quenched by several heavy metal ions.¹⁴² It is reminded that photoexcitation of UCNPs with lasers of wavelengths above 800 nm can lead to local heating which can represent a substantial source of error in quantitative fluorometry.

If excited with NIR lasers, UCNPs heat up. This can be desired or not. It is not desired in case of bioimaging, and an excitation wavelength of 915 nm was recommended to reduce laser induced heating of UCNPs of the type $\text{NaYbF}_4:\text{Tm},\text{Er},\text{Ho}$ for deeper in-vivo imaging.¹⁴³ Heating may, however, also be desired because local heating can be exploited in cancer therapy. In a typical application, oleate-capped UCNPs were coated first with a shell of plain silica (also in order to make them more stable in water solution) and then with a layer of silica doped with a blue carbocyanine dye. The optical emission of the upconverting NP was absorbed by the dye to cause a local heating by up to 21 °C and this causes cells to disrupt. Fig. 5 shows a schematic of the preparation of such UCNPs and respective TEMs. Even shorter excitation wavelengths can be used as shown¹⁴⁴ with core-shell UCNPs doped with Nd(III) ions as sensitizers. The upconversion effect already occurs at excitation wavelengths of around 800 – 820 nm, and this can strongly reduce sample heating and overtone IR absorption by water. Others have shifted the excitation wavelength for upconversion

to 1490 nm by using $\text{LiYF}_4:\text{Er}$ nanocrystals.¹⁴⁵

Fig. 5. (A): Preparation of UCNPs for use in imaging and hyperthermal cancer treatment. (A): Coating of the green UCNPs first with silica and then with silica doped with the blue and NIR emitting dye Cite-777. (B – D): TEM images of the UNNPs, $\text{UCNP}@SiO_2$ core/shell particles, and $\text{UCNP}@SiO_2$ particles coated with the dye-silica nanocomposite. From ref.¹⁴³ 266 (Zhan et al., 2011) with permission.



Most syntheses yield water-insoluble NPs which first have to be surface-modified so to enable phase transfer to aqueous solutions. Reviews are available on the design, nanochemistry and applications of UCNPs in theranostics,¹⁴⁶ on the surface chemistry of UCNPs, and how to make them hydrophilic¹⁴⁷. Any surface modification has, however, an effect on their luminescence and colloidal properties.¹⁴⁸ The upconversion fluorescence of NPs of the type $\text{Ag}@SiO_2@Y_2O_3:\text{Er}$ can be fine-tuned by the size of the silver core.¹⁴⁹ The emission color of lanthanide doped NaYF_4 UCNPs also changes during the transformation of crystalline phases from α , a transition state of α mixed with β , and finally to the β phase (from red to yellow and finally to green). Coated with polyethylenimine, such particles were used to image cells.¹⁵⁰

Examples for applications include the kinetic determination of the activity of the enzyme phospholipase.¹⁵¹ A mixture of a PEGylated phospholipid and a rhodamine-labeled phospholipid was deposited on an UCNP composed of hexagonal NaYF_4 doped with 20% Yb(III) and 2% Er(III). The 540-nm emission of the UCNP is used to photoexcite rhodamine B close to the NP. If, however, the phosphodiester is hydrolyzed by the enzyme phospholipase inside a cell, the rhodamine is released from the surface of the UCNP, and this leads to the suppression of the fluorescence of the rhodamine. The red emission peaking at 655 nm is not affected by this process and can serve as an internal standard. Earlier work in the use of UCNPs in chemical sensing and biosensing has been reviewed.¹⁵² A recent review covers their application to bioassays and in bioimaging.¹⁵³

3.9. Noble metal nanoparticles

Specific features of such NPs include excellent photostability, water-solubility, size-dependent colors, lack of swelling, sharp contrast, ease of characterization by other means such as TEM or SEM, and an established surface chemistry (often thiol-based) which is useful if targeted imaging or biosensing/imaging is desired. The uptake of engineered gold nanoparticles by mammalian cells has been reviewed by Dykman & Khlebtsov.¹⁵⁴ Gold and silver NPs also can be coupled to plasmonic detection quite readily or used to generate fluorescence patterns through differential release of fluorescent polymers.¹⁵⁵

Single gold or silver NPs display rather weak fluorescence but were used to image HeLa cells.¹⁵⁶ While one-photon luminescence is weak, two-photon luminescence of gold NPs is strong under excitation at 514 and 633 nm.¹⁵⁷ Two-photon luminescence imaging of cancer cells down to 75 μm depth and using molecularly targeted gold nanorods¹⁵⁸ and of silver NPs¹⁵⁹ has been reported. On the other hand, gold NPs can quench fluorescence by phase induced radiative rate suppression.¹⁶⁰ It was relatively late when it was discovered that metal clusters made from metallic gold, silver, copper, for example, display strong intrinsic fluorescence.¹⁶¹ Noble metals are preferred for their inertness. The surface of these clusters (gold and silver in particular) can be protected with alkanethiolate monolayers. If properly modified, they enable plain imaging and targeted imaging. While luminescence is often attributed to particle size effects that cause size-dependent fluorescence,¹⁶² structural parameters such as surface ligands, valence states of metal atoms and crystallinity of NPs also affect spectra and decay times. Gold NPs and clusters can be composed of a few to millions of atoms.¹⁶³ Such "quantum" clusters may also be protected or made targetable by coating them with respective proteins.¹⁶⁴ For example, gold NPs were functionalized with luminescent ruthenium(II) polypyridyl to endow DNA binding capability and applicability to cellular imaging.¹⁶⁵ These structures bind

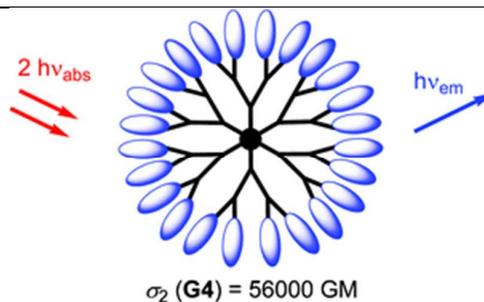
to DNA and undergo rapid cellular uptake, being localized within the cell cytoplasm and nucleus within 4 h. Various kinds of fluorescent silver nanoclusters (with green, red and yellow luminescences) have reported by Díez et al.¹⁶⁶ In a smart sensing approach towards probing phosphate ion,¹⁶⁷ gold nanoclusters (NCs) were capped with 11-mercaptoundecanoic acid and loaded with Eu(III) ions. The red fluorescence of the gold NCs is quenched by the Eu(III) ions, but fluorescence is restored on addition of phosphate.

3.10. Dendrimers, lipid drops and micelles

Dendrimers (dendrites) are a kind of NPs but much smaller than those treated so far. They can be both hydrophilic and hydrophobic and are easily internalized by cells. Their fluorescence usually originates from a label that has to be attached to the hyperbranched dendrimer skeleton such that no self-quenching does occur. The brightness of dendrimeric NPs can be controlled by size, and color by the terminal fluorophore. Their size exceeds 2 nm in very few cases. Dendrimer chemistry is well established. They have been applied to plain imaging, to targeted imaging, and to sensing/imaging. Features of fluorescent dendrimers include very large molar absorptances that can be in the order to $7,000,000 \text{ M}^{-1} \text{ cm}^{-1}$. The quantum yields of conventional fluorescence range from 0.3 to 0.5 provided the fluorophores are arranged such that they do not undergo self-quenching.

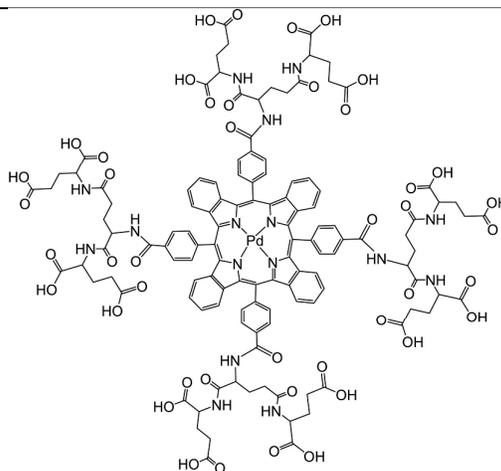
Nanoscope fluorescent dendrimers were designed¹⁶⁸ that carry up to 96 two-photon chromophores and show very large two-photon absorption cross-sections (up to 56,000 GM units). Such organic nanodots are said to represent "brilliant" alternatives to semiconductor quantum dots. A schematic of such a dendrimer particle is shown in Fig. 6. In a typical example of its use in imaging, a dendrimer-based fluorescent pH probe was used to visualize pH values in living HeLa cells.¹⁶⁹ In another example,¹⁷⁰ an amino-terminated generation 5 dendrimer labeled with a fluorescent marker was integrated into a nanogel and used to track it inside cells by fluorescence microscopy.

Fig. 6. Structure of a hyperbranched dendrimer with terminal fluorophores for use in single-photon and two-photon fluoroimaging. From ref.¹⁶⁸ (O. Mongin et al., 2006). © Royal Soc. Chem.)



Vinogradov's group¹⁷¹ has used smart dendrimers labeled with quenchable probes for oxygen. These are most viable (and commercially available) nanoprobe for imaging of oxygen in blood vessels and (cerebral) tissue. A dendrimer termed Oxyphor G2 [which is a two-layer glutamate dendrimer containing the Pd(II) complex of tetra-(4-carboxyphenyl) benzoporphyrin] is now widely used for phosphorescence based assay and imaging of oxygen. Its chemical structure is shown in Fig. 5. Other (and much larger) *Oxyphors* are known, some being more lipophilic, others more hydrophilic and even being charged.

Fig. 5. Chemical structure of the dendrimeric nanoparticles used (under the tradename *Oxyphore G*) for imaging of oxygen. It consists of a Pd(II)benzoporphyrin core and a glutamate-based periphery whose carboxy group are dissociated which imparts water solubility. From ref.¹⁷¹ (Devor... *NeuroMethods* 2014) with permission (2014) by Springer Verlag (Berlin).



In addition to dye labeled dendrimers, there are reports¹⁷² on autofluorescent hyperbranched poly(amido amine) NPs and their application to cell imaging. They display blue emission, are said to be nontoxic, and can be recognized by sialo-glycoprotein receptors on the surface of cancer cells. Similarly, a hyperbranched conjugated polyelectrolyte was reported for use in bioimaging.¹⁷³ It has a core-shell structure, an emission maximum at 565 nm, a quantum yield of 12% and a Stokes shift of 143 nm in water. Its poly(ethylene glycol) shell minimizes nonspecific interaction. Conjugated to anti-HER2 affibody, it was utilized for targeted cellular imaging of HER2-overexpressed cancer cells. A Q-dot-based ratiometric pH nanosensor with a dendrimer coating was constructed for measurement of physiological pH ranges. It is based on CdSe/CdZnS nanocrystals where the pH probe SNARF was conjugated to the poly(amido amine) dendrimer coating.¹⁷⁴ Dendrimer probes for enhanced photostability and localization in fluorescence imaging have been reviewed.¹⁷⁵ The group of Tian¹⁷⁶ described multifunctional nanomicelles for recognition and precisely targeted NIR-light induced cancer therapy. The nanomicelle encapsulates a pH-activatable fluorescent probe and a robust NIR photosensitizer and is functionalized with an aptamer for targeting viable cancer cells. The fluorescent probe can fluorescently detect the lysosomes for real-time imaging. NIR irradiation causes the generation of reactive oxygen species to trigger lysosomal cell death. The Lovell group reported¹⁷⁷ on the development of porphysomes (i.e., nanovesicles formed from self-assembled porphyrin bilayers) with very large extinction coefficients, structure-dependent fluorescence self-quenching, and photothermal properties. Their NIR fluorescence, regenerated on dissociation, results in low-background fluorescence imaging. Porphysomes are biodegradable and induce minimal acute toxicity in mice. Similar to liposomes, the aqueous core of porphysomes can be loaded with drugs or agents. Injected porphysomes accumulate in tumors of xenograft-bearing mice, and laser irradiation induces photothermal tumor ablation.

3.11. Others nanomaterials

Numerous other materials have been described for use as fluorescent NPs, most with the potential of acting as probes for (bio)imaging. These include metal oxides, sulfides, tellurides, fluorides, complex ionic species (including more than one cation and anion). They cannot be discussed here in depth but a selection is given. They usually are covered with additional layers (such as reactive groups and/or PEG) or coated with shells of (inert) solid materials, often silica. The list of conceivable candidates for preparing NPs of various size and optical properties is virtually endless. One typical example is given by Nd(III)-doped LaF₃ nanoparticles that can be applied to imaging in the second optical window of biomatter.¹⁷⁸ Fluoridated hydroxyapatites (HAPs) doped with Eu(III) or Tb(III) that were shown to be viable NPs for cell imaging.¹⁷⁹ They were made water-dispersible by hydrophobic/hydrophilic transformation with the surfactant PluronicTM F127. Similar NPs were obtained by PEGylation of fluoridated HAP nanorods doped with Ln(III) and used for cell imaging.¹⁸⁰ A comparative study was performed on the structure of Tb(III) doped fluorescent HAP nanocrystals¹⁸¹ and showed them to possess good cytocompatibility and cell imaging capability.

NPs consisting of a YVO₄ matrix and doped with trivalent lanthanides and phosphorus enable multicolor tuning at a single wavelength of excitation.¹⁸² Emission wavelengths and intensity ratios can be controlled using the host-activator system and the concentrations of dopants. By coupling luminescent Tb(III) complexes to magnetite (Fe₃O₄), NPs are obtained¹⁸³ that display paramagnetism, low cytotoxicity, and high cell uptake. If coated with folic acid, they can be used for in vitro fluorescence and magnetic resonance targeted imaging of cells that overexpress the folate receptor, for example the HeLa cells.

Titanium dioxide ("titania") is easily prepared and doped, for example with europium(III), and respective TiO₂ hollow nanoshells are viable two-photon nanoprobess. If coated with poly(ethylene imine), they adhere to HeLa cervical cancer cells.¹⁸⁴ Mesoporous titania was deposited on silver/silica core-shell NPs to result in a nanoarchitecture of type Ag@SiO₂@mTiO₂. The metal core acts as an enhancer of fluorescence. The particles were loaded with fluorescent flavin mononucleotide and the fluorescent cancer drug doxorubicin and used for simultaneous bimodal (fluorescence and SERS) imaging of drug delivery.¹⁸⁵ Amino-functionalized zirconia NPs (5 nm i.d.) doped with Ln(III) ions were prepared and used for time-resolved fluorescence resonance energy transfer detection of avidin with a detection limit of 3.0 nM. The ZrO₂ NPs can specifically recognize cancer cells overexpressed with urokinase plasminogen activator receptor.¹⁸⁶

Fluorescent Ag₂S nanoclusters possess tunable photoluminescence that extends from the red to the NIR. They can be made biocompatible and applied to bioimaging by introducing glutathione as the capping

reagent.¹⁸⁷ Two-photon excitable CaF_2 NPs doped with Tm(III) and Yb(III) represent multifunctional nanoprobe for deep-tissue fluorescence bioimaging.¹⁸⁸ On the basis of the strong Tm(III) ion emission (at around 800 nm), tissue penetration depths as large as 2 mm have been demonstrated, which is more than 4 times that of the visible emissions in comparable lanthanide-doped CaF_2 NPs. The outstanding penetration depth, together with the fluorescence thermal sensitivity these NPs well suited as multifunctional nanoprobe for high-contrast and deep-penetrating in-vivo imaging. Other sulfides also display quantum dot like emission, for example 5-nm NPs prepared from lead(II) sulfide which were protected with oleic acid and oleyl amine and made hydrophilic by hydroxylation at the terminal end of the alkyl chains.¹⁸⁹

Nanoscale metal-organic frameworks (often handicapped by hydrolytic decomposition) were shown to enable bioimaging¹⁹⁰ by incorporation of luminescent or high Z element building blocks. They can serve as viable contrast agents. The pores and channels of such frameworks may also be loaded with active agents and drugs. Colloidal and stable silicon (not silica!) NPs with their red fluorescence and being grafted with hyperbranched polyglycerol are water soluble and display good colloidal stability.¹⁹¹ The transformation of the hydrophobic surface of plain silicon NPs to a hydrophilic surface makes them water-soluble and suitable for specific targeting overexpressed cervical cancer cells and glioblastoma cells. Such particles are said to be "bright" but it is difficult to assess brightness (defined as molar absorbance multiplied with quantum yield) if neither of the two is known. Inorganic semiconductor fluorescent NPs consisting of ultrathin silica carbide (SiC) also display pH-dependent luminescence and can be used¹⁹² to probe intracellular pH values in the range from 5.6 to 7.4.

Lipid NPs – like hydrogels NPs – are soft but highly apolar. They resemble hydrophobic dendrimers and can be easily prepared but often lack stability if placed in complex systems. Gravier et al.¹⁹³ have described lipid NPs loaded with fluorophores (NIR dyes including Indocyanine Green) for use in fluorescence imaging (so called "lipidots"). Multichannel in-vivo imaging of lymph nodes in mice was demonstrated for doses as low as 2 pmol of NPs which have very high "molar" absorbance. Their cytotoxicity is very low. An interesting example of a micelle-based NP probe was reported by Zhou et al.¹⁹⁴ who have prepared such NPs from micelle-forming block copolymers with tertiary amine and poly(ethylene oxide) segments. If the local pH value is below the pK_a of the ammonium groups, the micelles dissociate into unimers which is accompanied by a strong increase in fluorescence due to suppression of homo-FRET. These NPs are highly sensitive to pH and were applied to target specific endocytic organelles in living cells and to image pH values. The method of pH-induced micellization was extended to a panel of multicolored nanoparticles with wide emission range (500–820 nm) and different pH transitions.¹⁹⁵ Wang et al.¹⁹⁶ have shown that ultra-small fluorescent nanosensors for oxygen can be obtained by a rather simple method. The materials have a hydrophobic core capable of firmly hosting hydrophobic luminescent oxygen probes and a shell composed of a long-chain poly(ethylene glycol) which renders them cell-membrane impermeable but yet highly sensitive to oxygen. These NPs are highly stable in aqueous solutions, in cell culture media, and in extracellular fluids such as blood, interstitial and brain fluid. Four kinds of nanosensors were presented, whose excitation spectra cover a wide spectral range (395 – 630 nm), thus matching many common laser lines, and with emission maxima ranging from 565 to 800 nm, thereby minimizing interference from background luminescence of biomatter.

4. Coatings and Surface Modifications

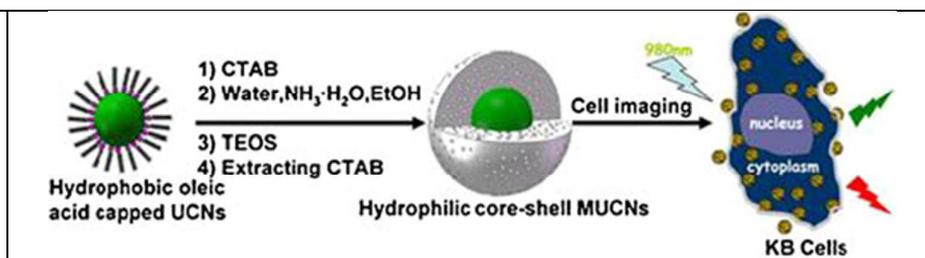
Coatings and surface modifications of NPs can serve two purposes. The first is to render NPs fluorescent, the second is to improve the properties of the NPs, for example by making them brighter, better biocompatible, or cell-permeable. NPs are rendered fluorescent by attaching fluorophores to their surface or by depositing additional coatings which may carry a fluorophore or are rendered fluorescent by immobilizing a fluorophore on their surface. The layer-by-layer technique (LbL) has turned out to be quite viable¹⁹⁷ but the stability may be an issue if the NPs are placed in complex samples such as serum. The LbL technique consists in the deposition of single layers of (polymer) molecules of alternating polarity or charge and works best with alternatively charged ionic materials.

Methods for the functionalization of inorganic NPs for bioimaging applications are rather diverse because conjugation of NPs to specific biomolecules enables targeted imaging, can reduce overall toxicity, and improve brightness. In general, such NPs are modified in two steps, the first being surface activation, the second being conjugation.¹⁹⁸ The latter knows numerous methods depending on the kind of species to be immobilized. These range from polymers such as poly(ethylene glycol) to proteins, fluorophores, oligomers

or other particles. Both physical adsorption and covalent binding have their merits and are widely used. Quality criteria of the final product include colloidal stability in water, biocompatibility, cell permeability, and a brightness that enables even minute quantities of NPs to be discerned from an often strongly fluorescent background.¹⁹⁹ Methods for the functionalization of the surface of metal complex-derived NPs for use in molecular imaging have been reviewed.²⁰⁰ One of the most versatile methods consists in the biomimetic modification of the surface of NPs by phospholipids as described in a milestone paper by the Selvin group.²⁰¹ The approach involves coating the surface of NPs with a monolayer of phospholipids containing different functional terminal groups that can govern polarity and can render the particles water soluble, for example, or conjugatable to biomolecules. These particles have a range of applications in fields such as imaging and quantitative bioanalysis. Specifically, dispersible and functionalized NPs were described for selective imaging of live cancer cells. Also see a related recent article where zwitterionic phospholipid coatings are employed.²⁰²

Many fluorescent NPs are simply coated with a shell of silica. This introduces a negative charge on the surface and – more importantly – protects the core from external quenchers. In addition, the silica coating can also be made fluorescent with a second dye and be further (bio)functionalized. Fig. 6 shows a schematic of a typical process. Methods for coating NPs with silica are well established and well reproducible. Biomedical applications of organically modified bioconjugated silica NPs have been reviewed.²⁰³

Figure 6. Schematic of the common method for coating oleate-capped upconversion nanoparticles (UCNPs) with a shell of silica, and their subsequent internalization into cells. From ref.²⁰⁴ with permission.



When deployed in-vivo, NPs are typically protected from the immune system by coating them with poly(ethylene glycol) (PEG). A wide variety of strategies are known²⁰⁵ to coat and characterize NPs with PEG. The core materials differ strongly in terms of size, shape, density, loading level, molecular weight, charge and method for purification. However, other hydrogels have been used that work as well. Jin et al.²⁰⁶ have coated upconversion nanocrystals of the type NaYF₄:Yb,Er with poly(vinyl pyrrolidone), and then a ligand exchange reaction was performed with polyethylenimine (PEI) and poly(acrylic acid). The coated UCNPs can be dispersed in aqueous medium. The positively charged PEI-coated particles are readily taken up by cells (in comparison to their neutral or negatively charged counterparts). The long-term in-vivo biodistribution of polyacrylic acid-coated upconversion nanophosphors was imaged and their toxicity is low.²⁰⁷ Others have coated UCNPs with the hydrogel poly[(N-isopropylacrylamide)-co-(methacrylic acid)].²⁰⁸ Gelatine, another kind of hydrogel, was used to coat CdTe/CdSe Q-dots to make them hydrophilic, more biocompatible and, thus, suitable for bioimaging.²⁰⁹ The cell permeability of particles is strongly improved if their surface is coated with poly-L-lysine.²¹⁰

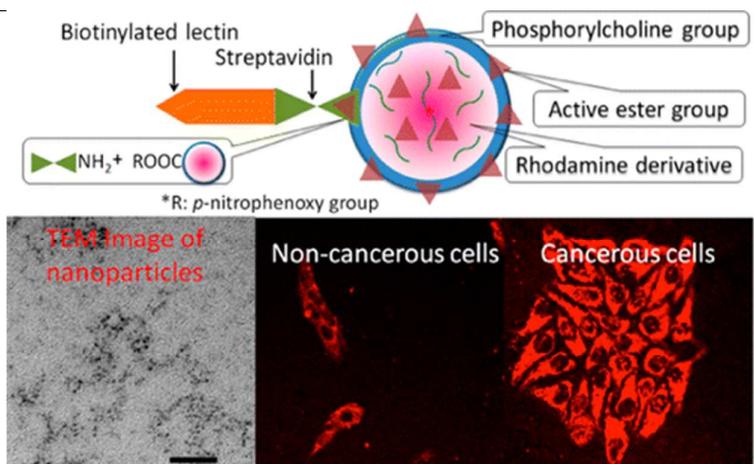
UCNPs are hydrophobic if prepared by the most common method. In order to make them water-soluble, Liras et al.²¹¹ have capped the surface of UCNPs with a polymer by replacing the original oleate ligand by multidentate thiolate grafting. The side chains of the coating extend into the solution and render the UCNPs water-dispersible. The resulting nanohybrids exhibit an emission brighter by a factor of 10 in organic solvent and even in water (by a factor of 2), and their fluorescence is highly thermoresponsive. A review is available on the functionalization of inorganic NPs including metal (Au, Ag), metal oxide (Fe₃O₄), and semiconductor nanocrystals (e.g. quantum dots and magnetic quantum dots) for bioimaging applications.²¹² Muhr et al.¹⁴⁷ have summarized methods for converting hydrophobic UCNPs (as obtained by standard synthetic methods for making particles with low dispersity) to water soluble hydrophilic UCNPs which is almost mandatory if such particles are to be used in bioimaging. Upconversion nanophosphors for small-animal imaging have been reviewed,²¹³ as have been methods for modification and characterization of UCNPs for use in bioanalytical methods.²¹⁴

The delivery of luminescent europium coated NPs into platelets can be controlled via the local pH values and can occur within a few min.²¹⁵ Gold NPs were coated with a luminescent Eu(III) complex (EuL) and the pHLIP peptide to give particles of the type pHLIP/EuL@AuNPs. The 13-nm diameter gold NPs act as a

scaffold for the attachment of both the luminescent probe and the peptide to target delivery. The NPs enter the platelets at low pH conditions only, typically at pH 6.5, but not at pH 7.4, and this is mediated by pHLIP translocation across the membrane. Luminescence microscopy images clearly show the red luminescence of the europium probe.

If NPs are to be used for purposes of targeting (see section 5.2), it is mandatory to modify the surface of the NP such that it can recognize its target. It is known, for example, that the coating of a surface with triphenylphosphonium groups (using reagents such as MitoPTM) will cause the particle to accumulate in the proximity of mitochondria.²¹⁶ This was exploited to visualize oxygen tension at various sites on a microscale.²¹⁷ Depending on the kind of surface (silica, polylysine, or triphenylphosphonium groups), the polystyrene nanosensors are located in the extracellular or intracellular space, or near mitochondria. A good example for the architecture of a NP that can target cancer cells is provided by the work of Cho et al.²¹⁸ who have made rhodamine-dyed NPs (~ 35 nm i.d.) highly biocompatible by coating the surface with phosphocholine and lectin as shown in Fig. 7. Such particles have a high affinity for sialic acid as overexpressed on the surface of tumor cells.

Fig. 7. Architecture of a nanoparticle capable of targeting cancer cells. The surface is rendered biocompatible and negatively charged by coating it with phosphocholine groups. Immobilization of lectin via biotin-streptavidin interaction results in particles that have extremely high affinity for sialic acid that is overexpressed on the surface of cancer cells. The images show, from left to right, a TEM and fluorescence images of non-cancerous and of cancerous cells. From ref.²¹⁸ (Cho et al.; *Biomacromolecules*, 2014) with permission (2014) of the Am. Chem. Soc.



5. Specific examples on the use of nanoparticles

5.1. Plain fluorescence imaging

In this case, the particles are simply added to the cell culture, tissue, or vascular system to be imaged, usually by, but not limited to microscopy. The only purpose of the particles is to render cells fluorescent, and therefore they are expected to display a large brightness, to be detectable in even deep regions of tissue or in blood, not to be toxic, and not to respond to their microenvironment (such as its local pH). NPs that work in the optical window of biomatter, i.e. in the 600 to 900 nm range, are strongly preferred, while probes with UV excitation suffer from inner filter effects on both the excitation and emission, and from strong UV, blue and green background fluorescence of most biomatter. Water-dispersible and fluorescent organic NPs that are made more biocompatible by PEGylation and exploiting the phenomenon of aggregation induced emission enhancement have found particular interest in the past years with respect to cell imaging applications.²¹⁹ Quite a number of articles have been published in recent years on the use of NPs for plain imaging, and a selection is presented in Table S1 in the Supporting information.

5.2. Targeted bioimaging

If specific chemical groups or domains are to be detected, a molecular probe (indicator), a labeling reagent (so to visualize functional groups such as thiols), or domain-recognizing probe (that can target mitochondria or the glycolipids in Alzheimer-associated tissue, for example), is being added. If the species to be targeted is an intracellular protein, Q-dots have been almost exclusively used in the past,²²⁰ but UCNPs and C-dots are on the rise. Such methods are compromised, however, by nonspecific binding, difficulty of intracellular delivery, or endosomal trapping of the NPs. Targeted bioimaging includes classical (not particle-based) methods of intracellular immunostaining and in-situ hybridization. Biju²²¹ has reviewed methods for

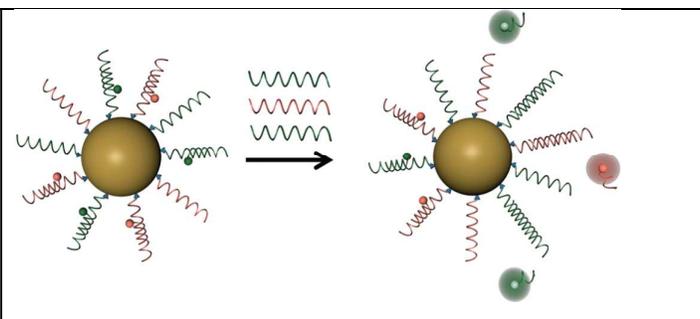
chemical modifications and bioconjugate reactions of nanomaterials such as silica NPs, gold NPs, gold quantum clusters, semiconductor Q-dots, carbon nanotubes, fullerene and graphene for use in (targeted) sensing and imaging.

Internalization is facilitated by modifying the surface with poly(ethylene glycol) (PEG) and/or coating with a cell penetrating peptide (which also avoids endosomal sequestration) and, finally, with a receptor. The art of delivering bioconjugated quantum dots to their targets and their uses for nonspecific extracellular and intracellular imaging has been reviewed.²²² Biocompatibility and internalization are critical issues, and particles usually are coated with PEG to impart or improve biocompatibility during functional live-cell imaging. It was shown more recently that sulfobetaine zwitterionic organic species on quantum dots serve the same purpose.²²³ Specifically, bidentate zwitterionic dihydrolipoic acid–sulfobetaine ligands represent a favorable alternative to PEG-coated NPs since they combine small size, low nonspecific adsorption, and stability over time and a wide range of pH values and salinity. In addition to using antibody-modified NPs, nanomaterials functionalized with DNA aptamers have been applied to targeted imaging of specific sites inside cells.²²⁴

Tumor cells are best recognized by targeting their folate receptor. It was targeted with an aggregation-enhanced fluorescent silica nanoprobe and used for both one-photon and two-photon excited bioimaging.²²⁵ The two-photon technique can overcome the limitations caused by fluorescence quenching due to high chromophore loading and provided 3D cellular-level resolution imaging of up to 350 μm deep in a HeLa tumor. Hyaluronated fullerene with its strong NIR fluorescence also allows for high-resolution fluorescent imaging of tumor sites in-vivo.²²⁶ Dually targeting upconversion nanocrystals were obtained²²⁷ by attaching 3-aminophenylboronic acid and hyaluronated fullerene to them. The two ligands warrant specific targeting of cancer cells. In vivo single-cell pharmacokinetic imaging of PARP-1 inhibitors and model drug behavior was demonstrated²²⁸ under varying conditions. Cancer cells were visualized through expression of H2B-Apple (580 nm; red) and TAMs were visualized by blue and fluorescent NPs internalized into endosomes. The method is said to allow for a better insight into drug action in vivo.

Work on the use of DNA aptamer modified nanomaterials for analysis of intracellular components and metabolites (also including aspects of imaging) has been reviewed.²²⁹ A most useful method for targeted imaging mRNA has been presented by the Mirkin group.²³⁰ Their term "multiplexed nanoflare" stands for a NP agent capable of simultaneously detecting two distinct mRNA targets inside a living cell. These probes are spherical and consist of polynucleotides conjugated to gold NPs as shown in Fig. 8. The oligomers are densely packed and highly oriented, many of which are hybridized to a reporter with a distinct fluorophore label and each complementary to its corresponding mRNA target. If multiplexed nanoflares are exposed to their targets, they provide a sequence specific signal in both extra- and intracellular environments. One of the targets can be used as an internal control, improving detection by accounting for cell-to-cell variations in NP uptake and background. Compared to single-component nanoflares, these structures allow for a precise determination of relative levels of mRNA in individual cells, but also have applications in cell sorting. In yet another new scheme for targeting, bioorthogonal (non-natural) chemistry was used to amplify NP binding and to enhance the sensitivity of cell detection. Antibodies against biomarkers were modified and used as scaffolds to couple NPs to live cells.²³¹ The group of Han²³² have reviewed the use of upconversion NPs as tools for multiscale targeted bioimaging. Other examples are given in Table S2 in the Supporting Information.

Fig. 8. Schematic representation of target detection of mRNA by multiplexed nanoflares. The multiplexed nanoflares bind different target nucleic acids (shown in red and green), displacing the corresponding flare. Once the flare is released, the fluorophore is no longer quenched by the gold surface, and an increase in fluorescence can be measured. Using two different fluorophores, the ratio of each target can be determined in cells. From ref.²³⁰ [A. E. Prigodich et al.; *Anal. Chem.*, 2012] with permission (2014) by the Am. Chem. Soc.



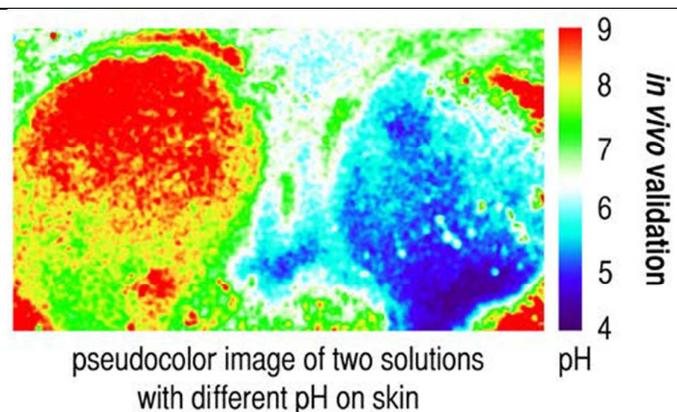
5.3. Imaging of chemical species

There is a substantial interest in visualizing chemical parameters such as pH values or chemical species such as glucose, calcium(II) or oxygen in the living and metabolizing cell, if not in tumor cells or in cells exposed to candidate drugs. Nanosensors have the unique feature of not being affected by intracellular species (that often bind to molecular probes to affect their binding constants or act as quenchers). The use of fluorescent NPs for intracellular sensing has been reviewed by the Hall group.²³³

5.3.1. Imaging of pH values

Sensing of pH is by far the most attractive option of fluorescence imaging because no other method enables pH to be imaged and monitored over time on a nanoscale. Fig. 9 shows, exemplarily, an image of the distribution of local pH values of a wound as imaged via the RGB technique (see above) using sensor particles in a sprayable film. Two kinds of polystyrene particles (one being pH-responsive because of a fluorescent pH probe immobilized on its surface, the other containing an inert reference fluorophore) were applied.²³⁴ A smart approach was presented where photon upconversion of respective NPs was applied to sensitize fluorescent nanoprobe for sensing and imaging of pH.²³⁵ In another approach,²³⁶ a block copolymer labeled with dyes not(!) sensitive to pH and with emission wavelengths from 500 to 820 nm was used to establish a panel of multicolored NPs with different pH transitions. The primary driving force of fluorescence activation between the unimer state and the micellar state is pH-induced micellization. Each kind of NP displays a sharp pH response (on/off within < 0.25 pH units), with pH transition points at pH 5.2, 6.4, 6.9, and 7.2. Incubation of a mixture of multicolored NPs with human H2009 lung cancer cells demonstrated their sequential activation inside endocytic compartments. Table S3 in the Supp. Information gives an overview on the variety of nanomaterials that have been employed to image pH values inside various kinds of living cells, in tissue and on surfaces.

Fig. 9. Pseudocolor images of pH values on an inflamed wound. Bluish colors indicate fairly intact skin (with pH values between 5 and 6), whilst green and red colors represent areas of sustained inflammation and partial granulation. From ref.²³⁴ (Schreml et al., 2012)

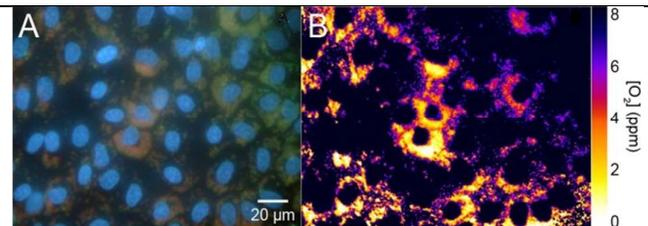


5.3.2. Imaging of oxygen

Aside from pH, there are species that are essential for understanding the function of cells, in (tumor) diagnosis, and in cellular testing including high-throughput screening. Among these, oxygen is of particular significance, and optical methods based on the use of NPs are widely used.²³⁷ Wang et al.²³⁸ have applied the red-green-blue techniques available with digital cameras to measure the distribution of oxygen in normal rat kidney cells. Fig. 10 shows a respective image. Nanosensors were made from polystyrene and containing two luminophores matching the red and the green channels of digital color cameras. The red emission of the oxygen probe is quenched by oxygen, while the blue-green emitting reference fluorophore yields a constant signal. Selected other nanomaterials for use in fluorescent imaging of oxygen are summarized in Table S4 in the Supp. Information.

Fig. 10. RGB image (A) and ratiometric image (B) of the distribution of oxygen in NRK cells using nanosensors for oxygen. The brightnesses of the red luminescence (resulting from the probe for oxygen) and the green luminescence (of the reference dye) is stored in 2 different memories and can be used to calculate the ratio for each single pixel. It can be plotted in pseudo-colours (right) to

reflect the local oxygen partial pressure. From ref.²³⁸
 [Wang, et al., Chem. Sci., 2011]. © Royal Soc. Chem.



5.3.3. Imaging of other (bio)chemical species

Other important classes of species include metal ions (from alkali to heavy metal) and organic species such as glucose. Temperature is another parameter that can be well sensed by using NPs. It shall be noted here, however, that several of the probes reported in the literature (sometimes also referred to as "sensors") do not respond in a reversible way so that they only can be used for acquiring a single moment picture of a cell, but not to monitor the ups and – in particular – the "downs" in the concentration of a species over time. Such probes should not be termed "sensor" and are not included here. Table S5 in the Supp. Information gives examples of nanomaterials for use in imaging of cations such as Mg(II) and Ca(II), of anions such as chloride, phosphate and cyanide, of thiols, ROS and NO, and of metabolites such as glucose.

5.4. Imaging of temperature

Conventional methods for measurement of temperature (T) cannot be applied to cells and other nanostructures. In fact, the use of molecular or NP-based optical probes is the only way to intracellularly sense T . A review on this topic has appeared.³⁷ (Wang) One may differentiate between two kinds of nanomaterials that can be used to sense T over time. The first is making use of one or two fluorophores of which at least one is displaying a highly T -dependent luminescence. If a single probe is used, the determination of lifetime is the (referenced) method of choice. If two fluorophores are used, one acts as the probe while the other serves as a reference or as a partner in a FRET system. An Eu(III) chelate probe in a silica matrix was claimed to possess luminescence that strongly depends on T in the physiological range.²³⁹ The second type is based on the use of fluorescent NPs displaying intrinsic fluorescence, examples being Q-dots,²⁴⁰ C-dots,²⁴¹ or upconverting nanocrystals which have the additional advantage of displaying two emissions bands of different sensitivity to T .²⁴² The group of Uchiyama²⁴³ were the first to report on nanomaterial-based intracellular sensing of temperature by using a fluorescent polymeric thermometer and fluorescence lifetime imaging microscopy. The spatial and temperature resolutions were at the diffraction limited level (200 nm) and 0.18–0.58 °C. The intracellular temperature distribution indicated that the nucleus and centrosome of a COS7 cell showed a significantly higher temperature than the cytoplasm and that the temperature gap between the nucleus and the cytoplasm differed depending on the cell cycle.

It appears that NPs prepared from a poly(methyl methacrylate)-co-1,2-bis(trimethoxysilyl)decane composite and containing a red-luminescent europium(III) complex display the best sensitivity to T at present.²⁴⁴ The NPs also contain a green-emitting and virtually T -independent reference dye. The ratio of the green and a red fluorescence under single-wavelength excitation is highly dependent on T in the 25 - 45 °C range, with a sensitivity of –4.0 % per °C. Given their small size (20 – 30 nm) and biocompatibility (due to the presence of an outer layer of silica), such NPs are likely to be useful nanoprobe for imaging of T inside cells. Silica NPs doped with the T-probe Ru(bpy) were also used²⁴⁵ to image intracellular T but have a smaller sensitivity (expressed as the signal change ΔS per °C). An interesting method, albeit not applied to imaging, is exploiting the T -sensitive fluorescence of NaYF₄:Er,Yb₃ upconversion NPs, where the intensity ratio of the two green bands of the Er(III) dopant changes with temperature.²⁴⁶ It was applied to measure T inside HeLa cells. Representative other nanomaterials and their features in terms of optical sensing of T are compiled in Table S6 in the Supp. Information.

6. Multimodal imaging, and imaging combined with drug and gene delivery, or with (photodynamic) therapy

One very exciting trend a present involves a combination of fluorescent imaging with other methods of

imaging, mainly magnetic resonance imaging (MRI), plasmonic imaging, and positron emission tomography (PET) imaging. This is referred to as bimodal or multimodal imaging. Another one involves the combination of fluorescent imaging with drug or gene delivery but these rely on the release of a diagnostically active species. A third (and closely related one) involves fluorescent imaging combined with photodynamic therapy (PDT) or photothermal therapy (PTD) where agents may be, or may be not, released from the NP. These methods shall be discussed in the following sections, again mainly from a nanomaterials point of view.

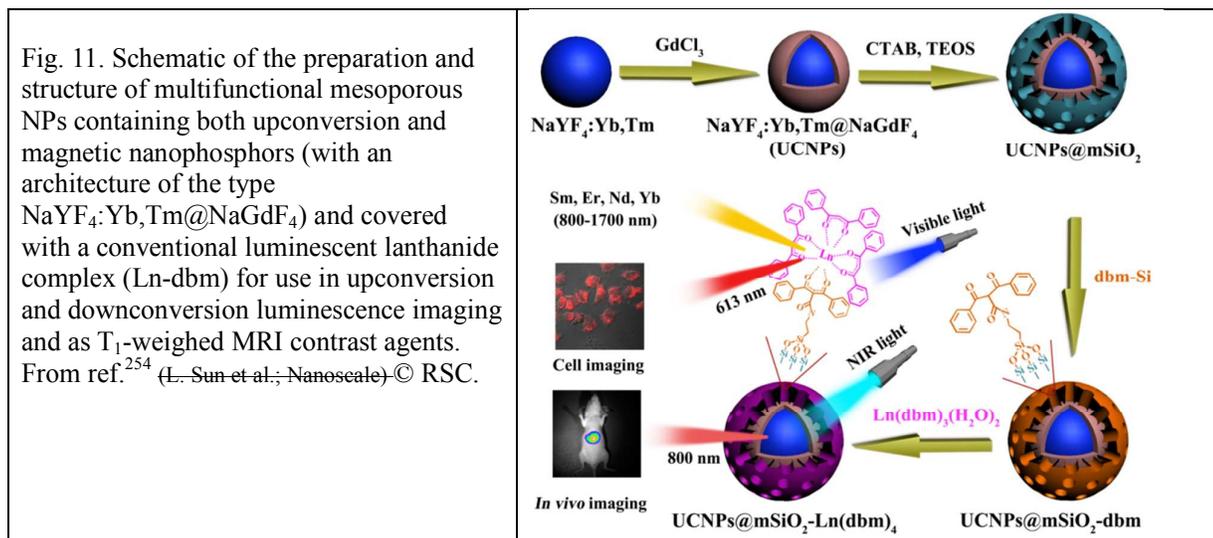
6.1. Multimodal imaging

Fluorescent NPs, magnetic NPs, and plasmonic NPs are the three most common nanoparticles when looking at the particle functionality. Their use in multimodal imaging has been briefly reviewed.²⁴⁷ Depending on the kind of multimodal imaging, a second species is required to be present in the NP in addition to a fluorescent probe that will give a signal in MRI or PET. This section gives examples in addition to those already described earlier in this review (search for bimodal or multimodal). A typical example²⁴⁸ is provided by the use of a nanocomposite consisting of magnetic iron oxide NPs and single-walled CNTs. These heterostructured complexes were utilized as bioimaging agents after encapsulation with oligonucleotides with the sequence d(GT)₁₅ and enrichment using a 0.5 Tesla magnetic array. The resulting nanotube complexes show distinct NIR fluorescence, Raman scattering, and visible/NIR absorbance. Macrophage cells that engulf the DNA-wrapped complexes were imaged using MRI and NIR mapping.

Dendrimer based bimodal imaging with longwave emitting fluorophores (these are preferred for the reasons indicated before) was shown for a fluorinated dendron conjugated to a cyanine dye for bimodal MRI and NIR fluorescence imaging.²⁴⁹ Bimodal upconversion fluorescence and X-ray imaging was also accomplished²⁵⁰ by using hexagonal phase NaLuF₄:Gd/Yb/Er nanorods, and blood vessels of lung were visualized with the aim to improve the diagnosis of lung and pulmonary vascular diseases. It is advised to read the critical comment by van Veggel et al.²⁵¹ on the use of lanthanide-doped NPs for fluorescence and MRI. Dual mode nanoprobe (without obvious cytotoxicity) for targeted bimodal fluorescent imaging and MRI of MCF-7 breast cancer cells were obtained²⁵² by co-encapsulation of an NIR-emissive conjugated polymer and lipid-coated iron oxides (IOs) in a hydrophilic poly(lactic-co-glycolic-acid)-poly(ethylene glycol) composite and coating the 180-nm particles with folic acid.

An important recent finding is the effect of size and phase on the multimodality of co-encapsulated magnetic photon-upconverting polymeric NPs for use in bimodal (fluorescent/magnetic resonance) imaging.²⁵³ In another sophisticated approach, the large potential of multimodal functionality was impressively demonstrated²⁵⁴ by showing that NPs of the layer architecture UCNP@mSiO₂-Ln(dbm)₄ (where mSiO₂ stands for mesoporous silica, Ln for any trivalent ion out of Eu, Sm, Er, Nd, Yb; and dbm stands for the organic ligand complexing the Ln ion). Both downconversion and upconversion luminescence is found, and this results in multicolor emission (covering the spectral region from 450 nm to 1700 nm) under visible-light excitation and 980-nm excitation, respectively. The mesoporous materials were applied to in-vitro imaging based on Eu(III) luminescence (under 405-nm excitation), and to small animal imaging based on Tm(III) luminescence (under 980-nm excitation). In addition, the Gd(III) dopant causes T1 signal enhancement and thus makes them potential contrast agents for MRI. This is schematically shown in Fig. 11. Other multifunctional nanocomposites include those for dual luminescence imaging, especially those showing both upconversion and downconversion luminescence.²⁵⁵

Bimodal surface enhanced Raman spectroscopy along with 3D fluorescence imaging based on the use of NPs is rather new. In the first approach of that kind,²⁵⁶ a rhodamine dye was conjugated to gold NPs which then were applied to live endothelial cells and revealed inhomogeneous distribution in the cytoplasm. Table S7 in the Supp. Information summarizes other representative examples of NPs for use in multimodal imaging.



6. 2. Imaging combined with drug delivery, gene delivery or chemotherapy

Theranostics with the cancer drug cis-platin has attracted much interest. Nanoparticles consisting of an upconversion fluorescent core and a porous silica shell were applied, for example, to deliver cis-platin to tumors. Such NPs may even be used in synergetic chemo- and radiotherapy by radiosensitization of the cis-platin in the shell, and in magnetic/luminescent bimodal imaging.²⁵⁷ The design of multifunctional upconversion NPs coated with a polymer nanocomposites for use in the delivery of cis-platin and in biomedical imaging was also described.²⁵⁸ In a series of articles, the groups of Qu and Ren have impressively shown how fluorescence nanoparticles prepared from various materials can be applied to simultaneous imaging and delivery of genes and drugs. In one example, polycation-functionalized and water-soluble gold nanoclusters were constructed that act as a platform for simultaneous gene delivery and cell imaging.²⁵⁹ In another, lanthanide-modified hollow mesoporous nanoparticles were also shown to serve this purpose.²⁶⁰ Dextranes, in crosslinked form, also can be used to prepare NPs for use in in-vitro drug release by loading dextran–chitosan NPs with the drug curcumin.²⁶¹ C-dots were shown to be useful nanocarriers for gene delivery and bioimaging. They were prepared by pyrolysis of glycerol in the presence of polyethylenimine which imparts better brightness.²⁶² Advances in lanthanide ion-based upconversion nanomaterials for drug delivery have been reviewed recently.²⁶³ Further examples are given in Table S8 in the Supp. Information.

In impressive work in multimodally functional NPs, chemo-, radio- and photodynamic therapy and simultaneous MRI/UCL imaging has been demonstrated for a single kind of NPs.²⁶⁴ This represents an exciting leap forward. By integrating upconversion NPs and mesoporous silica into a single platform, diagnostic/therapeutic functions were combined to provide a more advanced way for the efficient theranostics of cancer. Specifically, multifunctional Gd(III)-UCNPs with a mesoporous silica shell were constructed for the co-delivery of a radio-/photo-sensitizer hematoporphyrin (HP) and the radiosensitizer and chemodrug docetaxel. Upon NIR excitation and X-ray irradiation, a tumor was eliminated by the synergetic chemo-/radio-/photodynamic tri-modal therapy under the assistance of simultaneous magnetic and luminescent bimodal imaging. In other exciting work,²⁶⁵ upconversion NPs of the type BaGdF_5 ; 10 nm in size were shown (a) to act as carriers for the drug doxorubicin; (b) to act as multimodal probes for simultaneous imaging (optical/magnetic/X-ray/CT) and drug release which can be triggered by low pH. This exciting approach was applied to kill HeLa cells.

6.3. Imaging combined with photodynamic or photothermal therapy

The heating effect exerted by cw-laser light as used for photoexcitation of upconversion NPs is the result of the absorbance of light by such NPs and by their aqueous microenvironment because water has a weak overtone absorption at above 850 nm. This was exploited in a bimodal scheme for UCNP-based imaging and photothermal therapy (PDT).²⁶⁶ Similarly, in-vivo optical imaging and PDT of tumors was accomplished with upconversion NPs with a $\text{NaYF}_4:\text{Yb,Er}$ core and a NaGdF_4 shell combined with the photosensitizer chlorin e6. Tumors were observed both in the luminescence images and via MRI. In-vivo PDT was

simultaneously triggered by irradiation with 980-nm light.²⁶⁷ Related studies were performed where PDT is based on the multicolor emission capability of upconversion NPs at a single excitation wavelength that can simultaneously activate two photosensitizers. Mice were photodynamically treated by direct injection of the NPs into melanoma tumors or by intravenous injection of NPs conjugated to a tumor-targeting agent.²⁶⁸

Dextran based NPs dyed with Indocyanine Green can be used for NIR imaging and PDT in-vitro.²⁶⁹ The dye is retained by electrostatic interactions, and the NPs are well biocompatible and readily internalized because the dextran is block copolymerized with PEG. Even triple-functional core-shell structured NPs were reported more recently.²⁷⁰ The NPs were covalently grafted with a photosensitizer for luminescent, magnetic resonance imaging and PDT in vivo. Representative examples of NPs that have been used for these purposes are summarized in Table S9 in the Supp. Information.

7. Conclusions and outlook

The state of the art in fluorescent imaging is impressive. This is due to the progress made in materials science and in spectroscopy and microscopy. The number of nanomaterials, mainly nanoparticles, for use in fluorescent imaging is entangling, however, and a newcomer may find it difficult to make the proper selection. Depending on the kind of application, different kinds of criteria may apply when selecting particles. Nanoparticles always are preferred in intracellular studies, and often preferred (over microparticles) in extracellular studies. Table 2 gives an overview on figures of merit for the classes of NPs that are more widely used. The following criteria apply when selecting nanoparticles for use in fluorescence imaging:

- (a) Plain imaging requires bright particles that are non-toxic and – if this is needed – well cell permeable. Candidate materials include upconversion nanocrystals, C-dots, Q-dots (if properly coated), dyed particles consisting of (mesoporous) silica or polystyrene, and semiconductor P-dots.
- (b) Targeted (cellular) imaging requires bright particles that are not (cell-)toxic and, if needed, cell permeable. As their surface has to be coated with a ligand binding to the target (the receptor), their surface has to be modified which is easy in case of silica particles and silica-coated upconversion materials, also with C-dots and particles made from PAA, but less so with P-dots and with most hydrophobic organic particles.
- (c) NPs for use in chemical sensing are supposed, first of all, to be selective and, ideally, hardly affected by temperature. Biocompatibility is needed in case of imaging biosystems, of course. Sensors working in the optical (spectral) window of biomatter (i.e. between 600 and 900 nm) are preferred. Such sensors also should be calibratable in-vivo.
- (d) NPs for use in imaging of temperature (T) are, first of all, expected to be completely inert except for the T -dependence of their luminescence. Depending on the application, biocompatibility and cell permeability may be required. They are supposed to respond to T with a large signal change which results in a good resolution. In medical and bioapplications, the resolution is supposed to be better than ± 0.2 °C in the 20 – 55 °C range. Other nanosensors are supposed to work over a wide range of T s, and this will not require such a good resolution.
- (e) NPs for use in multimodal imaging requires nanomaterials of high sophistication. Such NPs are expected to display bright fluorescence and, in parallel, to be applicable to a second kind of imaging, MRI for example. Hence, additional functional materials have to be incorporated in the core or shell of the NPs however without compromising other properties including size and cell viability.
- (f) NPs for use in optical imaging along with (cancer) therapy also require nanomaterials of high sophistication. Particles are preferably covered with porous shells that can be loaded with a drug, a gene or a photodynamic or photothermal agent. Drugs and genes can be released once the NPs have arrived at the site of action. Controlled release can be induced by light or by changes of the local pH value, for example.

Table 2. Properties and simplified assessment of commonly used nanoparticles (NPs) or shell materials with respect to their properties and applicability to bioimaging, in multimodal imaging, and in theranostics

Kind of NP	Smallest size (nm)	Particles best characterized by ^{e)}	Exc/em wavelengths (nm)	Brightness	Photo-stability	Surface chemistry	Toxicity	Imaging & drug delivery	Imaging and PDT
Silica NPs (dye doped)	15	TEM, DLS,	from UV to NIR ^{f)}	high	fair	facile	low	not suited	not suited
Porous silicas (dye doped)	20	TEM, DLS,	from UV to NIR	high	fair	facile	low	well suited	well suited
Organic polymer dots (dye doped)	15	DLS	from UV to NIR	high	fair	fair	low	hardly demonstrated but likely	hardly demonstrated but likely
P-dots ^{a)}	25	DLS, AFM	(320 – 450)/ (400 – 600)	moderate	high	difficult	- ^{h)}	not yet demonstrated	not yet demonstrated
C-dots ^{b)}	2 – 3	AFM, XPS, HRTEM	(350 – 500)/ (450 – 700)	good	high	complex	said to be low	possibly suited	possibly suited
Q-dots ^{c)}	8	SEM, TEM, DLS,	(330 – 450)/ (420 – 700)	high	high	moderately facile	low if properly coated	not recommended	not recommended
UCNPs ^{d)}	8	TEM, DLS, XRD, ICP-AES	(780 – 980)/ (350 – NIR) ^{g)}	poor	high	fairly facile	low	well suited	well suited
Gold NP clusters	20	SEM, TEM, UV/Vis	(500 – 570)/ (560 – 680)	moderate	high	facile	low	well suited	well suited

^{a)} Semiconducting (organic) polymer dots; ^{b)} carbon dots; ^{a)} quantum dots based on the use of heavy metal ions; ^{d)} upconversion nanoparticles; ^{e)} TEM: transmission electron microscopy; DLS: dynamic light scattering; AFM: atomic force microscopy; XPS: X-ray photoelectron spectroscopy (for elemental analysis); HRTEM: high-resolution TEM; SEM: scanning electron microscopy; ICP-AES: inductively-coupled plasma atomic emission spectroscopy; ^{f)} depending on the dye used; ^{g)} usually two or more emission bands; ^{h)} no reliable data found.

Given the number of conceivable applications of nanoparticles in the very large medical field (cancer diagnosis and therapy mainly), in bioanalysis, biology, the military, security (from documents to banknotes) and in various kinds technical nanosystems, it is difficult to make general recommendation but it is hoped that readers will find this review to be of help when making decisions as to the material to be chosen.

References

- ¹ M. Chen and M. Yin, *Progr. Polym. Sci.* 2014, *39*, 365–395
- ² J. Li and J.J. Zhu, *Analyst*, 2013, *138*, 2506-2515
- ³ X. Michalet, F.F. Pinaud, L.A. Bentolila, J.M. Tsay, S. Doose, J. J. Li, A. M. Wu, S. S. Gabhir and S. Weiss, *Science* 2005, *307*, 538-544
- ⁴ F. Pinaud, S. Clarke, A. Sittner and M. Dahan. *Nature Meth.* 2010, *7*, 275–285
- ⁵ (a) S. Palmal and N.R. Jana, *Wiley Interdisc. Rev.: Nanomed. Nanobiotechnol.* 2014, *6*, 102-110; (b) C. Zhou, S. Yang, J. Liu, M. Yu, J. Sheng, *Exp. Biol. Med.* 2013, *238*, 1199-1209
- ⁶ X. Zhang, X. Zhang, L. Tao, Z. Chi, J. Xu and Y. Wei, *J. Mater. Chem. B*, 2014, *2*, 4398-4414
- ⁷ T. Trindade, A. L. D. da Silva; eds., *Nanocomposite Particles for Bio-applications*, Pan Stanford Pte. Publ., Singapore, 2011. ISBN 978-981-4267-78-6
- ⁸ S. Parveen, R. Misra and S.K. Sahoo, *Nanomedicine: Nanotechnol., Biol. Med.* 2012, *8*, 147-166
- ⁹ G. Yang, P. Shen, K. Tan and Y. Xia, *Microchim. Acta*, 2014, *181*, 607-613
- ¹⁰ V. Ntziachristos, *Nature Meth.* 2010, *7*, 603–614
- ¹¹ Y. Liu, D. Tu, H. Zhu, X. Chen, *Chem. Soc. Rev.*, 2013, *42*, 6924-6958
- ¹² P. Hazarika and D. A. Russell, *Angew. Chem. Int. Ed.* 2012, *51*, 3524-3531.
- ¹³ L. Wei, S. Doughan, Y. Han, M.V. DaCosta, U.J. Krull and D. Ho, *Sensors*, 2014, *14*, 16829-16855
- ¹⁴ J.C.G. Bünzli, *Chem. Rev.* 2010, *110*, 2729–2755
- ¹⁵ T. Ozawa, H. Yoshimura and S. B. Kim, *Anal. Chem.* 2013, *85*, 590-609
- ¹⁶ W. P. Dempsey, S. E. Fraser, P. Pantazis, *Bioessays* 2014, *34*, 351–360
- ¹⁷ Y. Zhang, Y. Bai, J. Jia, N. Gao, Y. Li, R. Zhang, G. Jiang, B. Yan, *Chem. Soc. Rev.* 2014, *43*, 3762-3809.
- ¹⁸ K. Wang, X. He, X. Yang, H. Shi, *Acc. Chem. Res.* 2013, *46*, 1367-1376.
- ¹⁹ A. Burns, P. Sengupta, T. Zedayko, B. Baird and U. Wiesner, *Small* 2006, *2*, 723–726
- ²⁰ K. Ma, H. Sai and U. Wiesner, *J. Am. Chem. Soc.* 2012, *4*, 13180–13183
- ²¹ H. Zhou, X. Lv, L. Zhang, A. Gong, A. Wu, Z. Liang, G. Peng, H. Lin, *J. Mater. Chem. C*, 2014, Accepted Manuscript. DOI: 10.1039/C4TC02163K
- ²² (a) H. S. Mader, X. Li, S. M. Saleh, M. Link, P. Kele, O. S Wolfbeis; *Annals New York Acad. Sci.* 2008, *1130*, 218-223; (b) S. M. Saleh, R. Ali and O. S. Wolfbeis; *Microchim Acta* **2011**, *174*, 429-434
- ²³ S.W. Bae, W. Tan and J.I. Hong, *Chem. Commun.*, 2012, *48*, 2270-2282
- ²⁴ X Chen, Y. Liu and D. Tu, Lanthanide Doped Luminescent Nanomaterials, in: *Nanomedicine and Nanotoxicology*, Springer Verlag, Berlin, 2014, chapter 7, pp 145-164. DOI: 10.1007/978-3-642-40364-4_7.
- ²⁵ Y. Zhang, W. Wei, G. K. Das and T. T. Y. Tan, *J. Photochem. Photobiol. C: Photochem. Rev.* 2014, *20*, 71–96
- ²⁶ S.L.C. Pinho, H. Faneca, C.F.G.C. Geraldés, M.H. Delville, L. D. Carlos and J. Rocha, *Biomaterials*, 2012, *33*, 925–935
- ²⁷ M. Chen and M Yin, *Progr. Polym. Sci.*, 2014, *39*, 395
- ²⁸ S. Schreml, R. J. Meier, O. S. Wolfbeis, M. Landthaler, R. M. Szeimies and P. Babilas, *Proc. Natl. Acad. Sci. U.S.A.* 2011, *108*, 2432–2437
- ²⁹ S. Schreml, R. J. Meier, M. Kirschbaum, O. S. Wolfbeis, M. Landthaler, P. Babilas, *Theranostics* 2014, *4*, 721-735
- ³⁰ H. H. Sun, K. Almdal and T. L. Andresen, *Chem. Commun.* 2011, *47*, 5268–5270
- ³¹ X. Zhou, F. Su, Y. Tian and D.R. Meldrum, *PLoS One*, 2014, *9*, e88185. DOI: 10.1371/journal.pone.0088185
- ³² X. Wang, R.J. Meier, O.S. Wolfbeis, *Angew. Chem. Intl. Ed.* 2013, *52*, 406-409
- ³³ X. Wang, J. A. Stolwijk, T. Lang, M. Sperber, R. J. Meier, J. Wegener and O. S. Wolfbeis. *J. Am. Chem. Soc.* 2012, *134*, 17011-17014
- ³⁴ M. I. J. Stich, L. H. Fischer and O. S. Wolfbeis; *Chem. Soc. Rev.* 2010, *39*, 3102-3114
- ³⁵ H. Peng, J. A. Stolwijk, L. Sun, J. Wegener and O. S. Wolfbeis; *Angew. Chem. Intl. Ed.* **2010**, *49*, 4246-4249
- ³⁶ K. Chung, J. Wallace, S. Kim, S. Kalyanasundaram, A. S. Andalman, T. J. Davidson, J. J. Mirzabekov, K. A. Zalocusky, J. Mattis, A. K. Denisin, S. Pak, H. Bernstein, C. Ramakrishnan, L. Gosenick, V. Gradinaru and K. Deisseroth, *Nature* 2013, *497*, 332–337
- ³⁷ X. Wang, O. S. Wolfbeis and R. J. Meier, *Chem. Soc. Rev.*, 2013, *42*, 7834-7869
- ³⁸ E. Pringsheim, D. Zimin and O. S. Wolfbeis; *Adv. Mat.* 2001, *13*, 819-822
- ³⁹ X. Wang, D. E. Achatz, C. Hupf, Sperber, J. Wegener, S. Bange, J. M. Lupton, O. S. Wolfbeis. *Sensors Actuat. B*, 2013, *188*, 257-262
- ⁴⁰ X. Wang, R. J. Meier and O. S. Wolfbeis. *Adv. Funct. Mat.* 2012, *22*, 4202-4207
- ⁴¹ X Zhang, S Wang, L Xu, L Feng, Y Ji, L Tao, S Li and Y Wei, *Nanoscale*, 2012, *4*, 5581-5584
- ⁴² M Luo, *Bull. Korean Chem. Soc.* 2014, *35*, 1732-1736
- ⁴³ I. L. Medintz, M. H. Stewart, S. A. Trammell, K. Susumu, J. B. Delehanty, B. C. Mei, J. S. Melinger, J. B. Blanco-Canosa, P. E. Dawson, H. Mattoussi, *Nature Mater.* 2010, *9*, 676–684
- ⁴⁴ D. Posavec, A. Dorsch, U. Bogner, G. Bernhardt and S. Nagl, *Microchim. Acta* 2011, *173*, 391-399

- ⁴⁵ S.H. Hu, X.H. Gao, *J. Am. Chem. Soc.* 2010, *132*, 7234–7237
- ⁴⁶ K Li, B Liu, *J. Mater. Chem.* 2012, *22*, 1257–1264
- ⁴⁷ R Tang, X Feng, *Can. Chem. Trans.* 2013, *1*, 78–84.
- ⁴⁸ C. Wu, S. J. Hansen, Q. Hou, J. Yu, M. Zeigler, Y. Jin, D. R. Burnham, J. D. McNeill, J. M. Olson, D. T. Chiu, *Angew. Chem. Int. Ed.* 2011, *50*, 3430–3434
- ⁴⁹ Y.H. Chan, C. Wu, F. Ye, Y. Jin, P.B. Smith, D.T. Chiu, *Anal. Chem.* 2011, *833*, 1448–1455
- ⁵⁰ X. Zhang, M. Liu, B. Yang, X. Zhang, Y. Wei, *Coll. Surf. B: Biointerf.* 2013, *112*, 81–86
- ⁵¹ L. Sun, D. Hao, W. Shen, Z. Qian, C. Zhu, *Microchim. Acta*, 2012, *177*, 357–364
- ⁵² K Pu, AJ Shuhendler, JV Jokerst, J Mei, S. S. Gambhir, Z. Bao, J. Rao, *Nature Nanotechnol.* 2014, *9*, 233–239
- ⁵³ S.N. Baker, G.A. Baker, *Angew. Chem. Int. Ed.* 2010, *49*, 6726–6744
- ⁵⁴ M. Bacon, S.J. Bradley, T. Nann, *Part. Part. Syst. Charact.*, 2014, *31*, 415–428
- ⁵⁵ Y. Song, S. Zhu and B. Yang, *RSC Adv.* 2014, *4*, 27184–27200
- ⁵⁶ H. Li, Z. Kang, Y. Liu, S.T. Lee, *J. Mat. Chem.* 2012, *22*, 24230–24253
- ⁵⁷ S. K. Das, Y. Liu, S. Yeom, D. Y. Kim and C. I. Richards, *Nano Lett.* 2014, *14*, 620–625
- ⁵⁸ L. Wang and H. S. Zhou, *Anal. Chem.*, 2014, *86*, 8902–8905; and references 4, 17 and 21–25 cited therein).
- ⁵⁹ B. Kong, A. W. Zhu, C. Q. Ding, X. M. Zhao, B. Li B and Y. Tian Y., *Adv. Mater.* 2012, *24*, 5844–5848.
- ⁶⁰ C. Ding, A. Zhu and Y. Tian; *Acc. Chem. Res.*, 2014, *47*, 20–30
- ⁶¹ F. Du, F. Zeng, Y. Ming, S. Wu, *Microchim. Acta* 2013, *180*, 453–460
- ⁶² P.G. Luo, S. Sahu, S.T. Yang, S.K. Sonkar, J. Wang, H. Wang, G. E. LeCroy, L. Cao and Y. Sun, *J. Mater. Chem. B*, 2013, *1*, 2116–2127
- ⁶³ C. Peng, W.B. Hu, Y.T. Zhou, C.H. Fan and Q. Huang, *Small*, 2010, *6*, 1686–1692
- ⁶⁴ K.P. Liu, J.J. Zhang, F.F. Cheng, T.T. Zheng, C.M. Wang and J.J. Zhu, *J. Mater. Chem.* 2011, *21*, 12034–12040
- ⁶⁵ Y. Chong, Y. Ma, H. Shen, X. Tu, X. Zhou, J. Xu, J. Dai, S. Fan, Z. Zhan, *Biomaterials* 2014, *35*, 5041–5048
- ⁶⁶ C.F. Hu, Y.L. Liu, Y.H. Yang, J.H. Cui, Z.R. Huang, Y.L. Wang and L.F. Yang, *J. Mater. Chem. B* 2013, *1*, 39
- ⁶⁷ J. C.G. Esteves da Silva and H. M.R. Goncalves, *Trends Anal. Chem.* 2011, *30*, 1327–1336
- ⁶⁸ S. Mitra, S. Chandra, T. Kundu, R. Banerjee, P. Pramanik and A. Goswami; *RSC Advances*, 2012, *2*, 12129–12131
- ⁶⁹ C. Liu P. Zhang, F. Tian, W. Li, F. Li, W. Liu, *J. Mat. Chem.* 2011, *21*, 13163–13167
- ⁷⁰ P. Zuo, D. Xiao, M. Gao, J. Peng, R. Pan, Y. Xia, H. He, *Microchim. Acta* 2014, *181*, 1309–1316
- ⁷¹ H. Li, X. He, Y. Liu, H. Huang, S. Lian, S. Lee and Z. Kang, *Carbon* 2011, *49*, 605–609
- ⁷² Y. Sun, B. Zhou, Y. Lin, W. Wang, K. A. S. Fernando, P. Pathak, M. J. Meziani, B. A. Harruff, X. Wang, H. Wang, P. G. Luo, H. Yang, M. E. Kose, B. Chen, L. M. Veca and S. Xie, *J. Am. Chem. Soc.* 2006, *128*, 7756–7757
- ⁷³ H.P. Liu, T. Ye and C.D. Mao, *Angew. Chem. Int. Ed.* 2007, *46*, 6473–6475
- ⁷⁴ X. Zhang, S. Wang, C. Zhu, M. Liu, Y. Ji, L. Feng, L. Tao and Y. Wei, *J. Coll. Interf. Sci.* 2013, *397*, 39–44
- ⁷⁵ S.K. Bhunia, A. Saha, A.R. Maity, S.C. Ray and N.R. Jana, *Scientific Reports*, 2013, *3*, Article number 1473. DOI: 10.1038/srep01473
- ⁷⁶ H. Nie, M. Li, Q. Li, S. Liang, Y. Tan, L. Sheng, W. Shi and S. X. Zhang, *Chem. Mater.*, 2014, *26*, 3104–3112
- ⁷⁷ S. Qu, H. Chen, X. Zheng, J. Cao and X. Liu, *Nanoscale* 2013, *5*, 5514–5518
- ⁷⁸ S. Kochmann, T. Hirscher and O. S. Wolfbeis; *J. Fluoresc.* 2012, *22*, 849–855
- ⁷⁹ H. Goncalves, A. Duarte and J.C.G. Esteves da Silva, *Biosens. Bioelectron.* 2010, *26*, 1302
- ⁸⁰ X. Zhang, S. Wang, M. Liu, B. Yang, L. Feng, Y. Ji, L. Tao and Y. Wei, *Phys. Chem. Chem. Phys.*, 2013, *15*, 19013–19018
- ⁸¹ G. Gollavelli and Y.C. Ling, *Biomaterials*, 2012, *33*, 2532–2545
- ⁸² J. Jeong, J. Jung, M. Choi, J.W. Kim, S.J. Chung, S. Lim, H. Lee and B. H. Chung, *Adv. Mat.* 2012, *24*, 1999–2003
- ⁸³ C. Baleizão, S. Nagl, M. Schaeferling, M. N. Berberan-Santos and O. S. Wolfbeis; *Anal. Chem.* 2008, *80*, 6449–6457
- ⁸⁴ C. Baleizao, S. Nagl, S. M. Borisov, M. Schaeferling, O. S. Wolfbeis and M. N. Berberan-Santos, *Chem. Eur. J.* 2007, *13*, 3643–3651
- ⁸⁵ X. Gong, Q. Hu, M.C. Paa, Y. Zhang, S. Shuang, C. Dong and M.M.F. Choi, *Nanoscale*, 2014, *6*, 8162–8170
- ⁸⁶ K.P. Liu, J.J. Zhang, F.F. Cheng, T.T. Zheng, C.M. Wang and J.J. Zhu, *J. Mater. Chem.* 2011, *21*, 12034–12040
- ⁸⁷ B.J. Hong, O.C. Compton, Z. An, I. Eryazici and S.T. Nguyen; *ACS Nano* 2012, *6*, 63–73
- ⁸⁸ S.K. Bhunia, N. Pradhan, N.R. Jana, *ACS Appl. Mater. Interf.*, 2014, *6*, 7672
- ⁸⁹ P.G. Luo, F. Yang, S.T. Yang, S.K. Sonkar, L. Yang, J. J. Broglie, Y. Liu and Y. Sun, *RSC Adv.* 2014, *4*, 10791–10807
- ⁹⁰ S. Zhuo, M. Shao and S.T. Lee, *ACS Nano*, 2012, *6*, 1059–1064
- ⁹¹ (a) Y. Li, Y. Zhao, H. H. Cheng, Y. Hu, G. Q. Shi, L. M. Dai, L. T. Qu, *J. Am. Chem. Soc.*, 2012, *134*, 15; (b) C. F. Hu, Y. L. Liu, Y. H. Yang, J. H. Cui, Z. R. Huang, *J. Mater. Chem. B*, 2013, *1*, 39; (c) Q. Liu, B. D. Guo, Z. Y. Rao, B. H. Zhang, J. R. Gong, *Nano Lett.*, 2013, *13*, 2436; (d) J. Ju and W. Chen, *Biosens. Bioelectron.*, 2014, *58*, 219
- ⁹² (a) L. Zhang, Z. Y. Zhang, R. P. Liang, Y. H. Li, J. D. Qiu, *Anal. Chem.*, 2014, *86*, 4423; (b) Z. T. Fan, Y. C. Li, X. H. Li, L. Z. Fan, S. X. Zhou, D. C. Fang, S. H. Yang, *Carbon* 2014, *70*, 149
- ⁹³ D. Qu, M. Zheng, P. Du, Y. Zhou, L. G. Zhang, D. Li, H. Q. Tan, Z. Zhao, Z. G. Xie, Z. C. Sun, *Nanoscale*, 2013, *5*, 12272
- ⁹⁴ Q. Feng, Q. Q. Cao, M. Li, F. C. Liu, N. J. Tang, Y. W. Du, *Appl. Phys. Lett.*, 2013, *102*, 013111
- ⁹⁵ X. M. Li, S. P. Lau, L. B. Tang, R. B. Ji, P. Z. Yang, *J. Mater. Chem. C*, 2013, *1*, 7308
- ⁹⁶ Q. Liu, B. Guo, Z. Rao, B. Zhang and J.R. Gong, *Nano Lett.* 2013, *13*, 2436–2441
- ⁹⁷ S. Zhu, J. Zhang, S. Tang, C. Qiao, L. Wang, H. Wang, X. Liu, B. Li, Y. Li, W. Yu, X. Wang, H. Sun and B. Yang, *Adv. Funct. Mater.* 2012, *22*, 4732–4740
- ⁹⁸ Z. Liu, K. Yang and S.T. Lee, *J. Mater. Chem.*, 2011, *21*, 586–598
- ⁹⁹ P. Cherukuri, C. J. Gannon, T. K. Leeuw, H. K. Schmidt, R. E. Smalley, S. A. Curley and R. B. Weisman, *Proc. Natl. Acad. Sci.*

- U.S.A.* 2006, *103*, 18882–18886
- 100 S. Dang, Q. Liu, X. Zhang, K. He, C. Wang and X. Fang, *J. Nanosci. Nanotechnol.* 2012, *12*, 4478–4484
- 101 DA Heller, S Baik, TE Eurell and MS Strano, *Adv. Mater.* 2005, *17*, 2793–2799
- 102 B. Mu, J. Zhang, T. P. McNicholas, N. F. Reuel, S. Kruss and M. S. Strano, *Acc. Chem. Res.*, 2014, *47*, 979–988
- 103 N.F. Reuel, A Dupont, O Thouvenin, DC Lamb and MS Strano, *ACS Nano*, 2012, *6*, 5420–5428
- 104 H. Yi, D Ghosh, MH Ham, J Qi, PW Barone, MS Strano and A. M. Belcher, *Nano Lett.*, 2012, *12*, 1176–1183
- 105 H. Huang, E. Pierstorff, E. Osawa and D. Ho, *Nano Lett.* 2007, *7*, 3305–3314
- 106 R. Lam, M. Chen, E. Pierstorff, H. Huang, E. J. Osawa and D. Ho, *ACS Nano* 2008, *2*, 2095–2102
- 107 C. C. Fu, H. Y. Lee, K. Chen, T. S. Lim, H. Y. Wu, P. K. Lin, P. K. Wei, P. H. Tsao, H. C. Chang and W. Fann, *Proc. Natl. Acad. Sci. U.S.A.* 2007, *104*, 727–732
- 108 V. N. Mochalin and Y. Gogotsi, *J. Am. Chem. Soc.* 2009, *131*, 4594–4595
- 109 J. H. Yu, S.-H. Kwon, Z. Petráček, O. K. Park, S. W. Jun, K. Shin, M. Choi, Y. I. Park, K. Park, H. B. Na, N. Lee, D. W. Lee, J. H. Kim, P. Schwillie and T. Hyeo, *Nature Mat.* 2013, *12*, 359–366
- 110 N. Ma, J. Yang, K.M. Stewart, S.O. Kelley, *Langmuir* 2007, *23*, 12783–12787
- 111 M. Cao, C. Cao, M. Liu and P. Wang, C. Zhu, *Microchim. Acta* 2009, *165*, 341–346
- 112 E. Petryayeva, W.R. Algar and I.L. Medintz, *Appl. Spectrosc.* 2013, *67*, 215–251
- 113 S.J. Rosenthal, J.C. Chang, O. Kovtun, J.R. McBride and I. D. Tomlinson, *Chem. Biol.* 2011, *18*, 10–24
- 114 N. Chen, Y. He, Y. Su, X. Li, Q. Huang, H. Wang, X. Zhang, R. Tai and C. Fan, *Biomaterials*, 2012, *33*, 1238–1244
- 115 N. Ma, G. Tikhomirov and S.O. Kelley, *Acc. Chem. Res.* 2010, *43*, 173–180
- 116 A. Orte, J. M. Alvarez-Pez and M. J. Ruedas-Rama, *ACS Nano* 2013, *7*, 6387–6395
- 117 C. Ye, Y. Wang, C. Li, J. Yu, Y. Hu, *Microchim. Acta* 2013, *180*, 117–125
- 118 R. Tang, H. Lee and S. Achilefu, *J. Am. Chem. Soc.* 2012, *134*, 4545–4548
- 119 X. Luo, W. Wu, F. Deng, D. Chen, S. Luo, C. Au, *Microchim. Acta* 2014, *181*, 1361–1367
- 120 A.R. Maity, S. Palmal, S.K. Basiruddin, N.S. Karan, S. Sarkar, N. Pradhan and N.R. Jana, *Nanoscale*, 2013, *5*, 5506–5513
- 121 G. J. Stasiuk, S. Tamang, D. Imbert, C. Poillot, M. Giardiello, C. Tisseyre, E.L. Barbier, P.H. Fries, M. de Waard, P. Reiss, M. Mazzanti, *ACS Nano* 2011, *5*, 8193–8201
- 122 J. Wang, F. Wang, C. Wang, Z. Liu and X. G. Liu, *Angew. Chem. Int. Ed.* 2011, *50*, 10369–10372
- 123 L.-D. Sun, Y.-F. Wang and C.-H. Yan, *Acc. Chem. Res.*, 2014, *47*, 1001–1009
- 124 H. Liu, C.T. Xu, S. Andersson-Engels, *Optics Express*, 2014, *22*, 17782–17790
- 125 Y. Yang, *Microchim. Acta* 2014, *181*, 263–294
- 126 D. Yin, C. Wang, J. Ouyang, X.Y. Zhang, Z. Jiao, Y. Feng, K. Song and M. Wu, *ACS Appl. Mater. Interf.*, on the web. DOI: 10.1021/am505633g
- 127 D. Lu, S. K. Cho, S. Ahn, L. Brun, C. J. Summers, W. Park, *ACS Nano*, 2014, *8*, 7780–7792
- 128 M. Pollnau, D. R. Gamelin, S. R. Luthi, H. U. Guedel, M. P. Hehlen, *Phys. Rev. B*, 2000, *61*, 3337
- ¹²⁹ J.-C. Boyer and F. C. J. M. van Veggel, *Nanoscale* 2010, *2*, 1417–1419
- ¹³⁰ C. Würth, J. Pauli, C. Lochmann, M. Spieles and U. Resch-Genger, *Anal. Chem.* 2012, *84*, 1345–1352
- ¹³¹ D. J. Gargas, E. M. Chan, A. D. Ostrowski, S. Aloni, M. V. P. Altoe, E. S. Barnard, B. Sani, J. J. Urban, D. J. Milliron, B. E. Cohen and P. J. Schuck; *Nature Nanotechnol.* 2014, *9*, 300–305
- ¹³² H.-H. Gorris, O. S. Wolfbeis; *Angew. Chem. Intl. Ed.* 2013, *52*, 3584–3600.
- ¹³³ Y. Lu, J. Zhao, R. Zhang, Y. Liu, D. Liu, E. M. Goldys, X. Yang, P. Xi, A. Sunna, J. Lu, Y. Shi, R. C. Leif, Y. Huo, J. Shen, J. A. Piper, J. P. Robinson and D. Jin, *Nature Photonics* 2014, *8*, 32–36
- 134 M. Quintanilla, F. Ren, D. Ma, F. Vetrone, *ACS Photonics*, 2014, *1*, 662–669 .
- 135 Y. Ou, D. Yin, X. Cao, C. Wang, K. Song, B. Liu, L. Zhang, Y. Han, M. Wu, *Dalton Trans.*, 2014, *43*, 14001–14008
- 136 D. Yin, X. Cao, *Dalton Trans.* 2014, *43*, 14001–14008
- 137 H. S. Mader, P. Kele, S. M. Saleh, O. S. Wolfbeis; *Curr. Opin. Chem. Biol.* 2010, *14*, 582–596.
- ¹³⁸ E. Hemmer, F. Vetrone and K. Soga, *Mater. Res. Soc.* 2014, *39*, 960–964
- 139 S. Wilhelm, T. Hirsch, W. M. Patterson, E. Scheucher, T. Mayr, O. S. Wolfbeis; *Theranostics* 2012, *3*, 239–248
- 140 R. B. Liebherr, T. Soukka, O. S. Wolfbeis, H. H. Gorris; *Nanotechnol.* 2012, *23*, 485103 (7pp)
- 141 A. Sedlmeier, D. E. Achatz, L. H. Fischer, H. H. Gorris, O. S. Wolfbeis. *Nanoscale* 2012, *4*, 7090–7096
- 142 S. M. Saleh, R. Ali, O. S. Wolfbeis; *Chem. – Eur. J.* 2011, *17*, 14611–14617
- 143 Q. Zhan, J. Qian, H. Liang, G. Somesfalean, D. Wang, S. He, Z. Zhang, S. Andersson-Engels, *ACS Nano* 2011, *5*, 3744–3757
- 144 (a) Y. Wang, G. Liu, L. Sun, J. Xiao, J. Zhou, C. Yan, *ACS Nano* 2013, *7*, 7200–7206; (b) H. Wen, H. Zhu, X. Chen, T. F. Hung, B. Wang, G. Zhu, S. F. Yu, F. Wang, *Angew. Chem. Int. Ed.* 2013, *52*, 13419–13423
- 145 G. Y. Chen, T. Y. Ohulchanskyy, A. Kachynski, H. Agren and P. N. Prasad, *ACS Nano* 2011, *5*, 4981 – 4986
- 146 G. Chen, H. Qiu, P. N. Prasad, X. Chen, *Chem. Rev.* 2014, *114*, 5161–5214
- 147 V. Muhr, Th. Hirsch, S. Wilhelm, O. S. Wolfbeis; *Acc. Chem. Res.* 2014; on the web. DOI: 10.1021/ar500253g
- 148 S. Wilhelm, M. Kaiser, C. Würth, J. Heiland, C. Carrillo-Carrion, V. Muhr, O. S. Wolfbeis, W. J. Parak, U. Resch-Genger, T. Hirsch, *Nanoscale* 2015, submitted.
- 149 F. Zhang, G. B. Braun, Y. Shi, Y. Zhang, X. Sun, N. O. Reich, D. Zhao, G. Stucky, *J. Am. Chem. Soc.*, 2010, *132*, 2850–2851.
- 150 Y. Hu, Y. Sun, Y. Li, S.K. Sun, J. Huo and X.J. Zhao, *RSC Adv.* 2014, *4*, 43653–43660
- 151 Y. Cen, Y. Wu, X. Kong, S. Wu, R. Yu, X. Chu, *Anal. Chem.*, 2014, *86*, 7119–7127
- 152 D. E. Achatz, R. Ali, O. S. Wolfbeis; *Topics Curr. Chem.* 2011, *300*, 29–50
- 153 M. V. DaCosta, S. Doughan, Y. Han, U. J. Krull, *Anal Chim Acta* 2014, *832*, 1–33
- 154 L. A. Dykman and N. G. Khlebtsov, *Chem. Rev.* 2014, *114*, 1258–1288

- 155 E. Boisselier, D. Astruc, *Chem. Soc. Rev.*, 2009, 38, 1759-1782
- 156 H. He, C. Xie, J. Ren, *Anal. Chem.*, 2008, 80, 5951-5957
- 157 A. Tcherniak, S. Dominguez-Medina, W. Chang, P. Swanglap, L. S. Slaughter, C. F. Landes, S. Link, *J. Phys. Chem. C*, 2011, 115, 15938-15949
- 158 N. J. Durr, T. Larson, D.K. Smith, B.A. Korgel, K. Sokolov, A. Ben-Yakar, *Nano Lett.*, 2007, 7, 941-945
- 159 J. Jiang, H. Gu, H. Shao, E. Devlin, G. C. Papaefthymiou, J. Y. Ying, *Adv. Mat.* 2008, 20, 4403-4407
- 160 E. Dulkeith, M. Ringler, T.A. Klar, J. Feldmann, A. Muñoz Javier, W. J. Parak, *Nano Lett.*, 2005, 5, 585-589
- 161 Y. Lu, W. Chen, *Chem. Soc. Rev.*, 2012, 41, 3594-3623
- 162 L. Shang, S. Dong, G.U. Nienhaus, *Nano Today*, 2011, 6, 401-418
- 163 J. Zheng, C. Zhou, M. Yu, J. Liu, *Nanoscale*, 2012, 4, 4073-4083
- 164 P.L. Xavier, K. Chaudhari, A. Baksi, T. Pradeep, *Nano Rev.*, 2012, 3, 14767_1 - 14767_16. DOI: 10.3402/nano.v3i0.14767
- 165 R. B. P. Elmes, K. N. Orange, S. M. Cloonan, D. C. Williams, T. Gunnlaugsson, *J. Am. Chem. Soc.*, 2011, 133, 15862-15865
- 166 (a) L. Díez, R. H. A. Ras, *Nanoscale* 2011, 3, 1963; (b) I. Díez, M.I. Kanyuk, A.P. Demchenko, A. Walther, H. Jiang, O. Ikkala, R. H. A. Ras, *Nanoscale*, 2012, 4, 4434-4437
- 167 S. N. Ding, C. M. Li, B. H. Gao, O. Kargbo, N. Wan, X. Chen and C. Zhou, *Microchim. Acta*, 2014, on the web. : 10.1007/s00604-014-1299-1
- 168 O. Mongin, T. R. Krishna, M. H. V. Werts, A.-M. Caminade, J.-P. Majoral, M. Blanchard-Desce, *Chem. Comm.*, 2006, 42, 915-917
- 169 L. Albertazzi, B. Storti, L. Marchetti, F. Beltram, *J. Am. Chem. Soc.*, 2010, 132, 18158-18167
- 170 M. Gonçalves, D Maciel, D Capêlo, S Xiao, W. Sun, X. Shi, J. Rodrigues, H. Tomas, Y. Li, *Biomacromolecules*, 2014, 15, 492-499
- 171 A. Devor, S. Sakadžić, M. A. Yaseen, E. Roussakis, P. Tian, H. Slovin, I. Vanzetta, I. Teng, P. A. Saisan, L. E. Sinks, A. M. Dale, S. A. Vinogradov, D. A. Boas, *NeuroMethods* 2014, 85, 225-253.
- 172 W. Yang, C. Pan, X. Liu, J Wang, *Biomacromolecules*, 2011, 12, 1523-1531
- 173 K.-Y. Pu, J. Shi, L. Cai, K. Li, B. Liu, *Biomacromolecules*, 2011, 12, 2966-2974
- 174 R. C. Somers, R. M. Lanning, P. T. Snee, A. B. Greytak, R. K. Jain, M. G. Bawendi, D. G. Nocera, *Chem. Sci.* 2012, 3, 2980-2985
- 175 Y. Kim, S.H. Kim, M. Tanyeri, J.A. Katzenellenbogen, C. M. Schroeder, *Biophys. J.* 2013, 104, 1566-1575
- 176 J. Tian, L. Ding, H. Ju, Y. Yang, X. Li, Z. Shen, Z. Zhu, J. Yu and C. J. Yang, *Angew. Chem. Int. Ed.* 2014, 53, 9544-9548
- 177 J. F. Lovell, C. S. Jin, E. Huynh, H. Jin, C. Kim, J. L. Rubinstein, W. C. W. Chan, W. Cao, L. V. Wang and G. Zheng, *Nat. Mater.* 2011, 10, 324-332
- 178 U. Rocha, K.U. Kumar, C. Jacinto, I. Villa, F. Sanz-Rodríguez, M. de la Cruz, A. Juarranz, E. Carrasco, F. C. J. M. van Veggel, E. Bovero, J. García Solé and D. Jaque, *Small* 2014, 10, 1141-1154
- 179 J. Hui, X. Zhang, Z. Zhang, S. Wang, L. Tao, Y. Wei, X. Wang, *Nanoscale*, 2012, 4, 6967-6970
- 180 X. Zhang, J. Hui, B. Yang, Y. Yang, D. Fan, M. Liu, L. Tao, Y. Wei, *Polym. Chem.*, 2013, 4, 4120-4125
- 181 X. Li, H. Zeng, L. Teng, H. Chen, *Mater. Lett.*, 2014, 125, 78-81
- 182 F. Wang, X. Xue, X. Liu, *Angew. Chem. Int. Ed.* 2008, 47, 906-909
- 183 B. Wang, J. Hai, Q. Wang, T. Li, Z. Yang, *Angew. Chem. Int. Ed.*, 2011, 50, 3063-3066
- 184 S. Sandoval, J. Yang, J.G. Alfaro, A. Liberman, M. Makale, CE Chiang, IK Schuller, AC Kummel, WC Trogler, *Chem. Mater.*, 2012, 24, 4222-4230
- 185 Y. Wang, L. Chen, P. Liu, *Chem. Eur. J.* 2012, 18, 5935-5943
- 186 Y. Liu, S. Zhou, D. Tu, Z. Chen, M. Huang, H. Zhu, E. Ma, X. Chen, *J. Am. Chem. Soc.*, 2012, 134, 15083-15090
- 187 C. Wang, Y. Wang, L. Xu, D. Zhang, M. Liu, X. Li, H. Sun, Q. Lin, B. Yang, *Small*, 2012, 8, 3137-3142
- 188 N. Dong, M. Pedroni, F. Piccinelli, G. Conti, A. Sbarbati, J. Ramírez-Hernández, L. Martínez Maestro, M. C. Iglesias-de la Cruz, F. Sanz-Rodríguez, A. Juarranz, F. Chen, F. Vetrone, J. A. Capobianco, J. G. Solé, M. Bettinelli, D. Jaque, A. Speghini, *ACS Nano*, 2011, 5, 8665-8671
- 189 D. Zherebetsky, M. Scheele, Y. Zhang, N. Bronstein, C. Thompson, D. Britt, M. Salmeron, P. Alivisatos, and L.-W. Wang, *Science* 2014, 344, 1380-1382.
- 190 J. Della Rocca, D. Liu, W. Lin, *Acc. Chem. Res.*, 2011, 44, 957-968
- 191 P. Das, N.R. Jana, *ACS Appl. Mater. Interf.*, 2014, 6, 4301-4305
- 192 J. H. Guo, S. J. Xiong, X. L. Wu, J. C. Shen, P. K. Chu, *Biomaterials* 2013, 34, 9183-9189
- 193 J. Gravier, F.P. Navarro, T. Delmas, F. Mittler, A.-C. Couffin, F. Vinet, I. Texier, *J. Biomed. Opt.* 2011, 16, 096013
- 194 K. J. Zhou, Y. G. Wang, X. N. Huang, K. Luby-Phelps, B. D. Sumer, J. M. Gao; *Angew. Chem. Int. Ed.* 2011, 50, 6109-6114
- 195 K. J. Zhou, H. M. Liu, S. R. Zhang, X. N. Huang, Y. G. Wang, G. Huang, B. D. Sumer and J. M. Gao, *J. Am. Chem. Soc.* 2012, 134, 7803-7811
- 196 X. Wang, J. A. Stolwijk, M. Sperber, R. J. Meier, J. Wegener, O. S. Wolfbeis; *Meth. Appl. Fluoresc. (IOP Publ.; London)* 2013, 1, 035002 (7 pp)
- 197 R. R. Costa, J. F. Mano, *Chem. Soc. Rev.* 2014, 43, 3453-3479.
- 198 G.T. Hermanson, *Bioconjugate Techniques*, Academic Press, New York, 2013
- 199 N. Erathodiyil, J.Y. Ying, *Acc. Chem. Res.*, 2011, 44, 925-935
- 200 C.R. Maldonado, L. Salassa, N. Gomez-Blanco, J. C. Mareque-Rivas, *Coord. Chem. Rev.*, 2013, 257, 2668-2688.
- 201 L. Li, R. Zhang, L. Yin, K. Zheng, W. Qin, P. R. Selvin, Y. Lu; *Angew. Chem. Int. Ed.* 2012, 51, 6121-6125
- 202 C. Yao, P. Wang, L. Zhou, R. Wang, X. Li, D. Zhao, F. Zhang, *Anal. Chem.* 2014, 86, 9749-9757
- 203 K. Paulkumar, R. Arunachalam, G. Annadurai, *Int. J. Nanotechnol.*, 2011, 8, 653-663

- 204 L. Sun, T. Liu, Y. Qiu, J. Liu, F. Li, L. Shi, O. S. Wolfbeis, *Microchim. Acta* 2014, *181*, 775-782
- 205 J.V. Jokerst, T. Lobovkina, R.N. Zare, S.S. Gambhir; *Nanomedicine*, 2011, *6*, 715-728
- 206 J. Jin, Y.J. Gu, C.W.Y. Man, J. Cheng, Z. Xu, Y. Zhang, H. Wang, V. H. Lee, S. H. Cheng, W. Wong, *ACS Nano*, 2011, *5*, 7838-7847
- 207 L. Xiong, T. Yang, Y. Yang, C. Xu, F. Li, *Biomaterials*, 2010, *27*, 7078-7085
- 208 Y. Dai, P. Ma, Z. Cheng, X. Kang, X. Zhang, Z. Hou, C. Li, D. Yang, X. Zhai, J. Lin, *ACS Nano*, 2012, *6*, 3327-3338
- 209 R. G. Aswathy, B. Sivakumar, D. Brahatheeshwaran, T. Ukai, Y. Tomofumi; M. Yasuhiko; T. Maekawa, S. D. Kumar, *Mater. Express*, 2011, *1*, 291-298
- 210 X.H. Wang, H.S. Peng, Z. Chang, L.L. Hou, F.T. You, F. Teng, H.W. Song, B. Dong, *Microchim. Acta* 2012, *178*, 147-152
- 211 M. Liras, M. González-Béjar, E. Peinado, L. Francés-Soriano, J. Pérez-Prieto, I. Quijada-Garrido, O. García, *Chem. Mater.*, 2014, *26*, 4014-4022
- 212 N. Erathodiyil, J.Y. Ying, *Acc. Chem. Res.*, 2011, *44*, 925-935.
- 213 J. Zhou, Z. Liu and F. Li, *Chem. Soc. Rev.*, 2012, *41*, 1323-1349
- 214 A. Sedlmeier and H. H. Gorris, *Chem Soc Rev* 2014, on the web. DOI: 10.1039/C4CS00186A
- 215 A. Davies, D.J. Lewis, S.P. Watson, S. G. Thomas and Z. Pikramenou, *PNAS USA*, 2012, *109*, 1862-1867
- 216 (a) T. Murase, T. Yoshihara, S. Tobita, *Chem. Lett.* 2012, *41*, 262-263; E. T. Chouchani, C. Methner, S. M. Nadtochiy, A. Logan, V. R. Pell, S. Ding, A. M James, H. M. Cochemé, J. Reinhold, K. S. Lilley, L. Partridge, I. M. Fearnley, A. J Robinson, R. C. Hartley, R. A. J. Smith, T. Krieg, P. S. Brookes, M. P. Murphy, *Nature Med.* 2013, *19*, 753-759
- 217 X. Wang, H. Peng, L. Yang, F. You, F. Teng, L. Hou, O. S. Wolfbeis; *Angew. Chem. Int. Ed.* 2014, *53*, 12471-12475.
- 218 J. Cho, K. Kushihiro, Y. Teramura, M. Takai, *Biomacromolecules*, 2014, *15*, 2012-2018
- 219 X Zhang, X Zhang, B Yang, Y Zhang, M Liu, Y. Chen., Y. Wei, *Coll. Surf. B: Biointerf.* 2014, *113*, 435-441
- 220 Y. Choi, K. Kim, S. Hong, H. Kim, Y.J. Kwon, R. Song, *Bioconj. Chem.* 2011, *22*, 1576-1586
- 221 V. Biju, *Chem. Soc. Rev.*, 2014, *43*, 744-764
- 222 V. Biju, T. Itoh, M. Ishikawa, *Chem. Soc. Rev.* 2010, *39*, 3031-3056
- 223 E. Muro, T. Pons, N. Lequeux, A. Fragola, N. Sanson, Z. Lenkei, B. Dubertret, *J. Am. Chem. Soc.*, 2010, *132*, 4556-4557
- 224 H. Xing, N.Y. Wong, Y. Xiang, Y. Lu, *Curr. Opin. Chem. Biol.*, 2012, *16*, 429-435
- 225 X. Wang, A.R. Morales, T. Urakami, L. Zhang, M.V. Bondar, M. Komatsu, K. D. Belfield, *Bioconj. Chem.*, 2011, *22*, 1438-1450.
- 226 D.S. Kwag, K. Park, K.T. Oh, E.S. Lee, *Chem. Commun.*, 2013, *49*, 282-284
- 227 X. Wang, C.X. Yang, J.T. Chen, X.P. Yan, *Anal. Chem.* 2014, *86*, 3263-3267
- 228 G.M. Thurber, K.S. Yang, T. Reiner, R.H. Kohler, P. Sorger, T. Mitchison, R. Weissleder, *Nature Comm.* 2013, *4*, article number 1504
- 229 H. Xing, N.Y. Wong, Y. Xiang, Y. Lu, *Curr. Opin. Chem. Biol.* 2012, *16*, 429-435
- 230 A. E. Prigodich, P. S. Randeria, W. E. Briley, N. J. Kim, W. L. Daniel, D. A. Giljohann, C. A. Mirkin, *Anal. Chem.*, 2012, *84*, 2062-2066
- 231 J. B. Haun, N.K. Deveraj, SA Hilderbrand, H Lee, R Weissleder, *Nature Nanotechnol.* 2010, *5*, 660-665
- 232 X. Wu, G. Chen, J. Shen, Z. Li, Y. Zhang and G. Han; *Bioconj. Chem.*, on the web; DOI: 10.1021/bc5003967
- 233 M.J. Ruedas-Rama, J.D. Walters, A. Orte, E.A.H. Hall, *Anal. Chim. Acta*, 2012, *751*, 1-23
- 234 S. Schreml, R. J. Meier, J. Cattani, D. Flittner, S. Gehmert, O. S. Wolfbeis, M. Landthaler, P. Babilas; *Exptl. Dermatol.* **2012**, *21*, 942-970.
- 235 R. Arppe, T. Näreoja, S. Nylund, L. Mattsson, S. Koho, J. M. Rosenholm, T. Soukka, M. Schaeferling; *Nanoscale*, 2014, *6*, 6837-6843
- 236 K. Zhou, H. Liu, S. Zhang, X. Huang, Y. Wang, G. Huang, B. D. Sumer and J. M. Gao, *J. Am. Chem. Soc.* 2012, *134*, 7803-7811
- 237 X. Wang, O. S. Wolfbeis, *Chem. Soc. Rev.* 2014, *43*, 3666-3761
- 238 X. D. Wang, H. H. Gorris, J. A. Stolwijk, R. J. Meier, D. B. M. Groegel, J. Wegener, O. S. Wolfbeis, *Chem. Sci.* 2011, *2*, 901-906
- 239 H. Peng, M. I. J. Stich, J. Yu, L. Sun, L. H. Fischer and O. S. Wolfbeis; *Adv. Mat.* 2010, *22*, 716-719
- 240 L. M. Maestro, E. M. Rodriguez, F. S. Rodriguez, M. C. Iglesias-de la Cruz, A. Juarranz, R. Naccache, F. Vetrone, D. Jaque, J. A. Capobianco and J. G. Solé, *Nano Lett.* 2010, *10*, 5109-5115
- 241 P. Yu, X. Wen, Y.R. Toh, J. Tang, *J. Phys. Chem. C*, 2012, *116*, 25552-25557
- 242 A. Sedlmeier, D. E. Achatz, L. H. Fischer, H. H. Gorris, O. S. Wolfbeis. *Nanoscale* 2012, *4*, 7090-7096
- 243 K. Okabe, N. Inada, C. Gota, Y. Harada, T. Funatsu and S. Uchiyama, *Nature Comm.* 2012, *3*, article number 705.
- 244 H. Peng, S. Huang, O. S. Wolfbeis. *J. Nanoparticle Res.* 2010, *12*, 2729-2733
- 245 L. Yang, H.S. Peng, H. Ding, F.T. You, L.L. Hou, F. Teng, *Microchim. Acta* 2014, *181*, 743-749
- 246 F. Vetrone, R. Naccache, A. Zamarrón, A. J. de la Fuente, F. Sanz-Rodríguez, L. M. Maestro, E. M. Rodriguez, D. Jaque, J. G. Solé and J. A. Capobianco, *ACS Nano* 2010, *4*, 3254-3258.
- 247 N.C. Bigall, W.J. Parak, D. Dorfs, *Nano Today*, 2012, *7*, 532-563
- 248 J. H. Choi, F. T. Nguyen, P. W. Barone, D.A. Heller, A.E. Moll, D. Patel, S.A. Boppart and M.S. Strano, *Nano Lett.*, 2007, *7*, 861-867
- 249 Z. Wang, X. Yue, Y. Wang, C. Qian, P. Huang, M. Litzak, G. Niu, F. Wang, P. Rong, D.O. Kiesewetter, Y. Ma, X. Chen, *Adv. Healthcare Mat.* 2014, *3*, 1326-1333
- 250 S. Zeng, H. Wang, W. Lu, Z. Yi, L. Rao, H. Liu, and J. Hao, *Biomaterials*, 2014, *35*, 2934-2941.
- 251 F. C. J. M. van Veggel, C. Dong, N. J. J. Johnson, J. Pichaandi, *Nanoscale* 2012, *4*, 7309-7321

-
- 252 K. Li, D. Ding, D. Huo, K.Y. Pu, N.N.P. Thao, Y. Hu, Z. Li and B. Liu, *Adv. Funct. Mat.* 2012, 22, 3107–3115.
- 253 M. Challenor, P. Gong, D. Lorensen, M. J. House, R. C. Woodward, T. S. Pierre, M. Fitzgerald, S. A. Dunlop, D. D. Sampson and K. S. Iyer, *Dalton Trans.*, 2014, 43, 16780-16787
- 254 L. Sun, X. Ge, J. Liu, Y. Qiu, Z. Wei, B. Tian, L. Shi, *Nanoscale*, 2014. On the web. DOI: 10.1039/C4NR04258A
- 255 Y. Ding, X. Zhang, H. Zhu and J.-J. Zhu, *J. Mater. Chem. C*, 2014, 2, 946.
- 256 A. Jaworska, T. Wojcik, K. Malek, U. Kwolek, M. Kepczynski, A. A. Ansary, S. Chlopicki, M. Baranska, *Microchim. Acta*, 2014, on the web. DOI: 10.1007/s00604-014-1307-5
- 257 W. Fan, B. Shen, W. Bu, F. Chen and J. Shi, *J. Am. Chem. Soc.* 2013, 135, 6494-6503
- 258 P. Ma, H. Xiao, X. Li, and J. Lin, *Adv. Mater.* 2013, 25, 4898-4905
- 259 Y. Tao, Z. Li, E. Ju, J. Ren and X. Qu, *Nanoscale* 2013, 5, 6154
- 260 Z. Li, Y. Tao, S. Huang, N. Gao, J. Ren, X. Qu, *Chem. Comm.* 2013, 49, 7129
- 261 A. Anitha, V.G. Deepagan, V.V. Divya Rani, D. Menon, S.V. Nair and R. Jayakumar, *Carbohydr. Polym.* 2011, 1158–1164
- 262 C. Liu, P. Zhang, X. Zhai, F. Tian, W. Li, J. Yang, Y. Liu, H. Wang, W. Wang and W. Liu, *Biomaterials* 2012, 33, 3604-3613
- 263 D Yang, Z Hou, Z Cheng, C Li, J Lin, *Chem. Soc. Rev.* 2014, Advance Article. DOI: 10.1039/C4CS00155A
- 264 W. Fan, B. Shen, W. Bu, F. Chen, Q. He, K. Zhao, S. Zhang, L. Zhou, W. Peng, Q. Xiao, D. Ni, J. Liu, J. Shi, *Biomaterials*, 2014, 32, 8992-9002.
- ²⁶⁵ D. Yang, Y. Dai, J. Liu, Y. Zhou, Y. Chen, C. Li, P. Ma, J. Lin, *Biomaterials*, 2014, 35, 2011–2023
- ²⁶⁶ G. Shan, R. Weissleder, S.A. Hilderbrand, *Theranostics*, 2013, 3, 267–274
- ²⁶⁷ Y. I. Park, H. M. Kim, J. H. Kim, T. Hyeon, *Adv. Mater.* 2012, 24, 5755-5761
- ²⁶⁸ N. M. Idris, M. K. Gnanasammandhan, J. Zhang, P. C. Ho, R. Mahendran and Y. Zhang, *Nat. Med.* 2012, 18, 1580-1585
- ²⁶⁹ P. Liu, C. Yue, B. Shi, G. Gao, M. Li, B. Wang, Y. Ma and L. Cai, *Chem. Comm.* 2013, 49, 6143-6145
- ²⁷⁰ X.F. Qiao, J.C. Zhou, J.W. Xiao, Y.F. Wang, L.D. Sun, C.H. Yan, *Nanoscale*, 2012, 4, 4611-4623