Journal Name

ARTICLE

Supporting Information

Cite this: DOI: 10.1039/x0xx00000x

Received ooth January 2012, Accepted ooth January 2012

DOI: 10.1039/x0xx00000x

www.rsc.org/

Biotin-decorated silica coated PbS nanocrystals emitting in the second biological near infrared window for bioimaging

M. Corricelli,^{‡,a,b} N. Depalo,^{‡,b} E. Di Carlo,^a E. Fanizza,^{a,b} V. Laquintana,^c N. Denora,^c A. Agostiano,^{a,b} M. Striccoli,^b M. L. Curri,^{*,b}



Figure S1: HRTEM micrograph of PbS NCs (scale bar = 2nm).

HRTEM analysis has been carried out by a Jeol 2200FS microscope with spherical aberration-corrected objective lens, equipped with a Field Emission Gun (FEG) and working at an acceleration voltage of 200 kV. The HRTEM micrograph has been recorded by a Gatan Ultrascan 1000 CCD camera.



Figure S2: Size distribution histograms of PbS@SiO₂ samples prepared with 700 μ L of Igepal, 400 μ L of ammonia, 30 μ L of TEOS achieving a PbS NC concentration of $6.1 \cdot 10^{-6}$ M (a) and $9.5 \cdot 10^{-6}$ M (b). Size distribution histogram of PbS@SiO₂ sample prepared with 700 μ L of Igepal, 400 μ L of ammonia, 80 μ L of TEOS, achieving a PbS NC concentration of $6.1 \cdot 10^{-6}$ M (c).



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Figure S3: TLC plate with deposited drops of and amine-functionalized silica PbS NPs after spaying the ninhydrin/2,6-lutidine solution in acetone.

The presence of the amine groups onto silica coated PbS NP surface after functionalization by means of APS has been detected by performing a thin layer chromatography (TLC) analysis and using a suitable ninhydrin solution. In particular, a ninhydrin test has been here suitably employed for the characterization of amine-functionalized silica coated NPs¹. The ninhydrin solution has been preliminary prepared by dissolving 110 mg of ninhydrin in 16 mL of acetone (0.68% w/v) and adding 4 mL of 2,6-lutidine. For the qualitative TLC analysis, a drop of the amine functionalized PbS@SiO₂ NP solution has been loaded on the TLC plate and sprayed with the ninhydrin solution. The presence of amino groups has been confirmed by the formation of a blue coloured spot ascribed to the Ruhemanns by-product (Figure S2).

	Size	PDI	ζ-potential
	(nm)		(mV)
PbS@SIO ₂ NPs	36±2	0.27±0.02	-29.5±0.8
PbS@SIO ₂ /biotin NPs	41±3	0.34±0.02	-22.7±0.3
PbS@SIO ₂ /biotin/streptavidin	48±1	0.22±0.01	-28.4±0.4
NPs			

Notes and references

^{*a*} Dipartimento di Chimica, Università degli Studi di Bari, Via Orabona 4, I-70126, Bari, Italy.

^b CNR-IPCF c/o Dipartimento di Chimica, Università degli Studi di Bari,
Via Orabona 4, I-70125, Bari, Italy. * Tel.: +39 (0)80 5442027. E-mail:
lucia.curri@ba.ipcf.cnr.it.

^c Dipartimento di Farmacia - Scienze del farmaco, Università degli Studi di Bari, Via Orabona 4, I-70126, Bari, Italy.

These authors contributed equally.

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