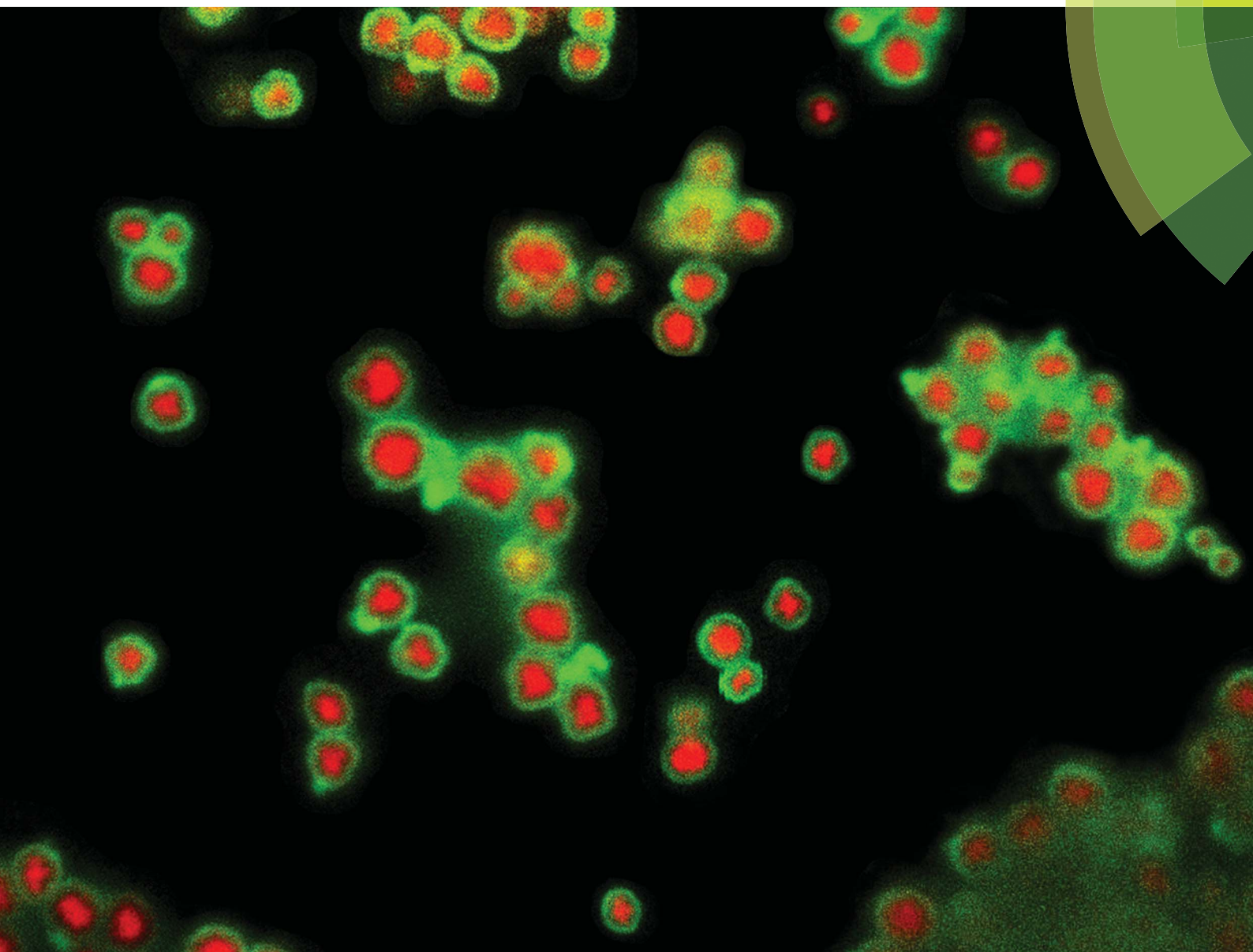


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Cellular uptake of ribonuclease A-functionalised core–shell silica microspheres†

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Analysis of protein function in a cellular context ideally requires physiologically representative levels of that protein. Thus conventional nucleic acid-based transfection methods are far from ideal owing to the over expression that generally results. Likewise, fusions with protein transduction domains can be problematic whilst delivery *via* liposomes/nanoparticles typically results in endosomal localisation. Recently, polymer microspheres have been reported to be highly effective at delivering proteins into cells and thus provide a viable new alternative for protein delivery (protein transduction). Herein we describe the successful delivery of active ribonuclease A into HeLa cells *via* novel polymer core–silica shell microspheres. Specifically, poly(styrene-*co*-vinylbenzylisothiuronium chloride) core particles, generated by dispersion polymerisation, were coated with a poly(styrene-*co*-trimethoxysilylpropyl methacrylate) shell. The resultant core–shell morphology was characterised by transmission electron, scanning electron and confocal fluorescence microscopies, whilst size and surface charge was assessed by dynamic light scattering and zeta-potential measurements, respectively. Subsequently, ribonuclease A was coupled to the microspheres using simple carbodiimide chemistry. Gel electrophoresis confirmed and quantified the activity of the immobilised enzyme against purified HeLa RNA. Finally, the polymer–protein particles were evaluated as protein-transduction vectors *in vitro* to deliver active ribonuclease A to HeLa cells. Cellular uptake of the microspheres was successful and resulted in reduced levels of both intracellular RNA and cell viability.

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Introduction

The study of a protein's activity within living cells is a fundamental requirement for understanding the role of that protein. Moreover, such studies also offer insight into how to modify this role and even how to prevent the protein from fulfilling its normal function altogether. To this end, a number of techniques for introducing proteins into cells have been developed. The most common of which are transfection-based methods,¹ protein fusion-based methods,² and liposomal/nanoparticle-based methods.³

Unfortunately, for the delivery of functional proteins each of these method types is problematic. Transfection of DNA followed by intracellular protein expression, for example, whilst effective at introducing functionally active proteins into a cell, relies on normal transcription and translation processes and as such both precludes careful quantitative analysis and typically results in intracellular protein levels being far higher than normal. As an alternative approach, proteins can be produced as fusions with peptide sequences that possess known cell penetrating capabilities. An example of such a sequence is YGRKKRRQRRR which is the protein transduction domain of the HIV-TAT protein.⁴ Unfortunately, when protein-fusions produced in this way are incubated with cells, they typically enter the cells by one of a number of different endocytosis-based mechanisms *e.g.* receptor-mediated endocytosis, macropinocytosis *etc.* Regardless of which precise endocytosis mechanism is used to effect cellular entry, the process normally results in fusion with lysosomes, cellular organelles whose specific role is to degrade materials either no longer needed by the cell or else which have been taken up by the cell.⁵ A similar fate can befall entities delivered in liposomal preparations or indeed in/on nanoparticle preparations where, again, endocytosis is the typical mechanism of uptake for these delivery vectors. Consequently, whilst relatively robust small organic

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† Electronic supplementary information (ESI) available: A visual guide to the fluorescently-labelled microspheres used in this study, a scheme of the full synthesis of RNase-functionalised microspheres **6**, characterisation of surface functionality, FTIR spectroscopy data, laser diffractometry data, TEM and energy dispersive X-ray analysis data, fluorescent labelling of RNase and micrographs that confirm internalisation of microspheres into cells. See DOI: 10.1039/c4tb01130a



molecules, such as pharmaceutical molecules, can be delivered effectively *in vivo* using transduction domains, liposomes, nanoparticles *etc.*, functionally-active proteins are far more prone to lysosomal degradation and cannot be delivered reliably in this manner.

It has been shown recently, by chemical analysis, microscopic analysis and gene expression, that polymer-based microspheres are taken up readily by all sorts of different cell lines *via* a non-endocytosis-reliant mechanism.⁶ Although it is difficult to discern the precise mechanism of uptake, it is thought to be *via* passive diffusion that involves a rearrangement of the cell membrane to allow the microsphere to pass through. This finding provides the opportunity for a good solution to the problem of functional protein delivery to cells. Originally observed and reported by the Bradley group,⁷ we and others have now also been able to show that polystyrene-based microspheres enter non-phagocytic cell lines easily and with very high efficiencies, typically more than 90% of the cells take up microspheres following a simple incubation process. This is a remarkable phenomenon given the relative size of the microspheres involved (typically 1–2 microns in diameter) to the size of cells (typically eukaryotic cells are 10–20 microns). Moreover, the microspheres can carry molecular cargoes, such as siRNAs⁸ and even proteins,^{9,10} with them into the cells. Pivotal for the delivery of functional proteins is the fact that this microsphere-based delivery method, termed beadfection, is non-endocytosis-based and thus enables the delivered proteins to avoid the harsh lysosomal environment.⁶

To date the only types of microsphere that have been employed in beadfection have been polystyrene-based. Silica-containing materials have not yet been evaluated. This would be a logical extension since silica particles have been used for drug delivery and are noted for their biocompatibility.¹¹ Indeed, highly cross-linked silica-based nano- and microspheres, generated using sol-gel chemistry, have been used for both chemical sensing^{12–16} and biomedical applications.^{17–29} Silica can also be used to coat polymer particles to form hybrid materials suitable for use in the body.¹¹ Moreover, the functionalised core-shell architecture that this approach creates is particularly attractive for beadfection since core and shell may bear different entities. For example, it would enable a reporter element, such as a fluorescent probe, to be located in the core whilst a protein may be attached to the shell.

Our previous work on beadfection had centred on delivering green fluorescent protein (GFP) into cells, primarily so we could use the inherent fluorescence of the protein to study the beadfection process.^{9,10} Whilst an excellent reporter element in its own right, GFP does not chemically modify other molecules within the cell.³⁰ Accordingly, a better test of the beadfection process for importing functional proteins would be to import a pharmaceutically relevant protein. Ribonuclease A (RNase) is just such a protein and degrades RNA by cleavage of the phosphodiester bond in the backbone following pyrimidine bases. Moreover, RNase has been identified by several groups as a potential anti-cancer agent.^{31,32}

Herein we describe our new and refined approach to beadfection for the delivery of functional proteins into cells.

Specifically, the synthesis of polymeric-silica core-shell microspheres, prepared *via* seeded-emulsion and sol-gel polymerisation techniques is described. The multi-functionality of this system is exemplified by concomitant conjugation of two fluorescent probes and the functional protein, RNase. Finally, the delivery of the functional protein *in vitro* has been investigated with HeLa cells and demonstrates that the microspheres we have developed are an excellent platform with which to study functional protein transduction *in vitro*.

Results and discussion

Microsphere synthesis

Dual-functionalised microspheres were synthesised by dispersion polymerisation of styrene-based core microspheres, followed by seeded-emulsion and sol-gel polymerisation of an outer shell (Scheme 1). Amine-functionalised microspheres 3 produced in this way were characterised by scanning electron microscopy (SEM) and confocal fluorescence microscopy (Fig. 1). The former showed the microspheres were roughly spherical and approximately 1 micron in diameter, whilst the latter confirmed the core-shell structure of the particles. Transmission electron microscopy (TEM) also supported the core-shell structure (see ESI, Fig. S5†). Energy dispersive X-ray analysis (EDXA) of a single bead provided confirmation that the beads contained a high degree of both silica and oxygen (see ESI, Fig. S6†).

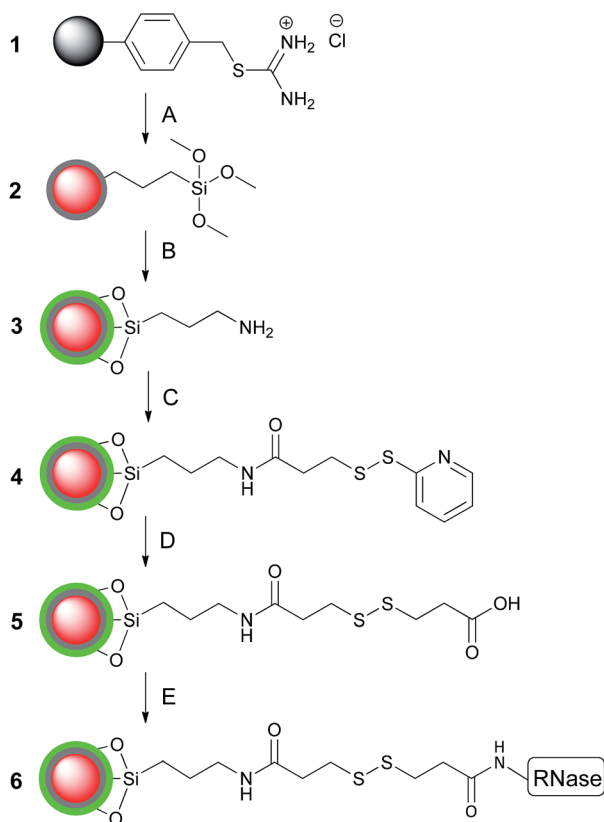
The core-shell synthetic strategy permits orthogonal functionality where thiuronium-functionalised core and amine-functionalised shell can be exploited independently. This was exemplified by further derivatisation of amine-functionalised microspheres 3 to produce RNase-functionalised microspheres 6 with two fluorescent markers (Texas Red in the core with a FITC-containing shell).

In a separate experiment, conjugation of RNase to the surface of the microspheres was assessed by fluorescence microscopy. This was achieved by the synthesis of core-shell microspheres without fluorescein isothiocyanate (FITC) in the silica shell 3a. After which, RNase partially labelled with FITC (for protocol see ESI†), was conjugated to these microspheres as depicted in Scheme 2. Confocal fluorescence microscopy confirmed the conjugation of RNase to the surface of the particles (Fig. 2). As a control experiment, the same procedure was conducted without the addition of the EDAC coupling reagent. Subsequent analysis of these control microspheres showed no fluorescence from the labelled RNase, verifying that non-specific adsorption of RNase did not occur (see ESI,† Fig. S7).

Characterisation of surface functionality

Amine functionality. The sol-gel process presents a simple way of altering the extent of amine-functionalisation of the silica shell. Simply increasing the amount of APTMS, up to a 1 : 1 ratio with TEOS, results in an increased loading of amine groups. This was quantified by a two-step assay³⁵ with amine-functionalised microspheres 3b. Firstly, Fmoc-Lys-(Fmoc)-OH





Scheme 1 Preparation of RNase-functionalised microspheres **6** with a TR (red)-labelled core and FITC (green)-labelled shell. (A) (i) Texas Red maleimide (TR), DMF, 16 h, (ii) TMSPM, styrene, DVB, AIBN, SDS, H₂O, N₂, 70 °C, 5 h to yield trimethoxysilane-functionalised microspheres **2**. (B) APTMS, TEOS, FITC, NH₄⁺OH⁻, H₂O, 50 °C, 2 h to yield amine-functionalised microspheres **3**. (C) SPDP, DMF, PBS, 1 h to yield pyridyl-functionalised microspheres **4**. (D) 3-mercaptopropionic acid, DMF, 1 h to yield carboxylic acid-functionalised microspheres **5**. (E) (i) EDAC, MES pH 5.5, H₂O, 1 h, (ii) RNase, MES pH 5.5, H₂O, 2 h to yield RNase-functionalised microspheres **6**.

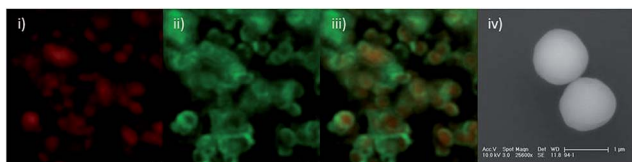
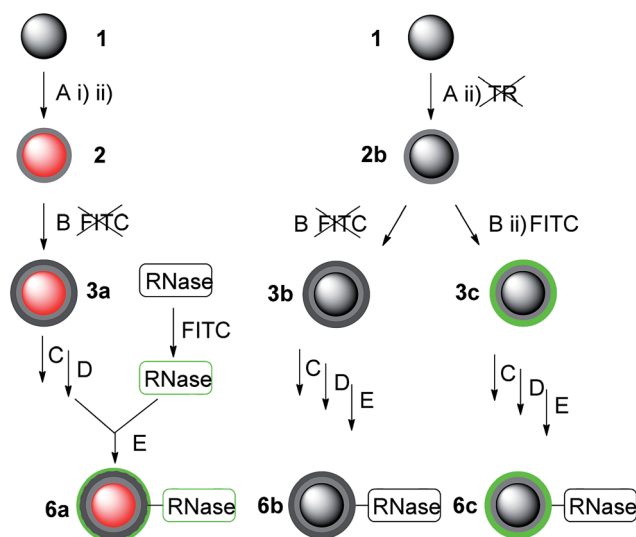


Fig. 1 Confocal fluorescence microscopy and SEM micrographs of amine-functionalised core-shell microspheres **3** with TR (red)-labelled cores and FITC (green)-labelled shells. Confocal fluorescence images were obtained using a 488 nm excitation laser at 15% power, and a 100× oil emersion objective with 12× zoom; (i) fluorescence emission collected in Channel 1 between 590 nm and 630 nm, (ii) fluorescence emission collected in Channel 2 between 500 nm and 560 nm, (iii) overlay of Channel 1 and Channel 2, (iv) SEM micrograph, scale bar represents 1 μm.

was conjugated to the amine groups. Secondly, both Fmoc moieties were cleaved with DBU and amino acid loading quantified by the characteristic absorbance at 294 and 305 nm of the Fmoc cleavage product.³⁵ These absorbance values were



Scheme 2 Preparation of variants of RNase-functionalised microspheres used in different experiments. (**6a**) TR (red)-labelled core, unlabelled shell with FITC (green)-labelled RNase, (**6b**) no labels and (**6c**) unlabelled core and FITC (green)-labelled shell. Thionium-functionalised microspheres **1** synthesised by dispersion polymerisation with 4-VBTU, styrene, DVB, AIBN, PVP, EtOH, N₂, 70 °C, 16 h. Followed by (A) (i) TR, DMF, 16 h, (ii) TMSPM, styrene, DVB, AIBN, SDS, H₂O, N₂, 70 °C, 5 h to yield trimethoxysilane-functionalised microspheres **2**. (B) APTMS [(B) (ii) with FITC], TEOS, NH₄⁺OH⁻, H₂O, 50 °C, 2 h to yield amine-functionalised microspheres **3a**, **3b** and **3c**. (C) SPDP, DMF, PBS, 1 h to yield pyridyl-functionalised microspheres **4a**, **4b** and **4c**. (D) 3-mercaptopropionic acid, DMF, 1 h to yield carboxylic acid-functionalised microspheres **5a**, **5b** and **5c**. (E) (i) EDAC, MES pH 5.5, H₂O, 1 h, (ii) RNase [6a with FITC (green)-labelled RNase], MES pH 5.5, H₂O, 2 h to yield RNase-functionalised microspheres **6a**, **6b** and **6c**.

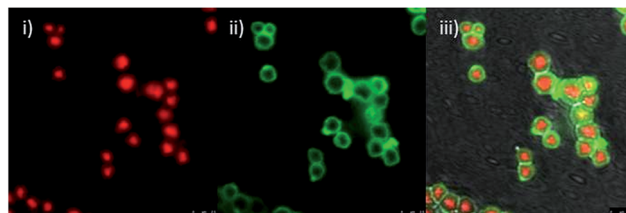


Fig. 2 Confocal microscopy images of core-shell microspheres **6a** having TR (red)-labelled cores, unlabelled shells and FITC (green)-labelled RNase conjugated to the outer shell. Images were obtained using a 488 nm excitation laser at 15% power, and a 100× oil emersion objective with 12× zoom; (i) fluorescence emission collected in Channel 1 between 590 nm and 630 nm, (ii) fluorescence emission collected in Channel 2 between 500 nm and 560 nm, (iii) overlay of Channel 1, Channel 2 and the phase channel.

used to calculate the number of moles of cleaved Fmoc, and by extension the molar loading of our amine-functionalised microspheres **3b**. This strategy also exemplifies this microsphere system as a versatile template for covalent attachment of other desired compounds. The assay confirms that the amine-loading can be tailored readily within a defined range of between 0.20 and 0.47 mmol g⁻¹ (see ESI,† Fig. S2).



Carboxylic acid functionality. The carboxylic acid-functionalised microspheres **5c** were conjugated to an alternative compound in order to verify the presence of carboxylic acid moieties, and, in turn, the success of the prior modification steps. Trimethoxysilyl-functionalised **2b**, amine-functionalised **3c**, pyridyl-functionalised **4c** and carboxylic acid-functionalised **5c** microspheres were all subjected to EDAC coupling to BODIPY, to highlight the difference in surface chemistry of the microspheres. BODIPY is an amine-functionalised fluorophore with characteristic emission at 616 nm. As expected, inspection by confocal fluorescence microscopy showed successful conjugation only with the carboxylic acid-functionalised microspheres **5c** (see ESI,† Fig. S1).

Surface charge. Surface functionalisation of the microspheres at each synthetic step was characterised by zeta potential measurements (Table 1). As expected, thiuronium **1**, pyridyl **4** and RNase-functionalised microspheres **6** were positively charged, whilst trimethoxysilane **2**, amine **3** and carboxylic acid-functionalised microspheres **5** were negatively charged. Amine-functionalised microspheres **3** gave an overall negatively charged zeta potential presumably due to the high number of silanol groups also present on the exterior of the microspheres. The positive charge conferred by the amine groups was identifiable by a zeta potential less negative (−19.99 mV) than that of trimethoxysilane-functionalised microspheres **2** (−44.45 mV).

Functional group analysis. To compliment the characterisation of amine functionality, carboxylic acid functionality and zeta potential, infrared (IR) spectroscopy studies were carried out on microspheres **2b** to **6b**. Surface functional groups were low in number relative to the effective molar mass of the particles making their detection difficult. However, the differences between each spectrum were such that it was possible to make tentative assignments for each of the microspheres **2b** to **6b** based on characteristic peaks. IR spectra are provided in the ESI,† Fig. S3.

Activity of conjugated RNase

One unit of RNase degrades 100 ng of RNA per second and 1 mg of the RNase used in this study has approximately 35 000 units of activity. A preliminary experiment demonstrated that our microspheres have no inherent degradative activity, whilst RNase retains activity when immobilised. Specifically, total RNA from HeLa cells was incubated with both carboxylic acid-functionalised microspheres **5b** and RNase-functionalised

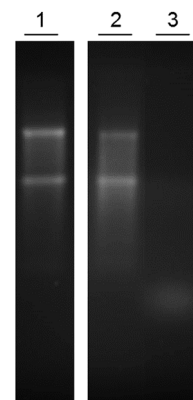


Fig. 3 Gel electrophoresis to compare the activity of carboxylic acid-functionalised microspheres **5b** against RNase-functionalised microspheres **6b**. Each reaction was carried out with 1.5 μg of RNA. Lane (1) no treatment, (2) 5 μg of carboxylic acid-functionalised microspheres **5b** (no RNase) for 5 minutes, (3) 5 μg of RNase-functionalised microspheres **6b** for 5 minutes.

microspheres **6b** and the products compared with intact HeLa RNA using gel electrophoresis (Fig. 3). The similarity between the RNA products in lanes 1 (control) and 2 shows that RNA treated with carboxylic acid-functionalised microspheres **5b** remains intact. In contrast, the small size of the product in lane 3 confirms there is high degradative activity displayed by RNase-functionalised microspheres **6b**, as expected.

To estimate the quantity of accessible, active RNase immobilised on microspheres **6b**, RNA degradation catalysed by decreasing quantities of native RNase was compared with RNA degradation catalysed by decreasing quantities of microspheres **6b**. Specifically, aliquots containing 0.100, 0.075, 0.050, 0.025 and 0.001 units of native RNase (2.86, 2.15, 1.43, 0.72 and 0.03 ng, respectively) were compared with 1000, 100, 10, and 1 ng of RNase-functionalised microspheres **6b** under the conditions outlined in the Experimental section and the products examined by gel electrophoresis (Fig. 4). The smaller the RNA product, the further it migrates in electrophoresis (towards the bottom of the gel). Whilst the two discreet 28S and 18S bands are clearly visible in lanes 1 (untreated RNA) and 10 (1 ng of microspheres **6b**), these bands are clearly eliminated in all other RNase-treated lanes. Moreover, there is clear correlation between the decreasing quantity of RNase (left to right) and increasing size of the RNA products. Visual inspection of Fig. 4 and intensity profile analysis (see ESI, Fig. S9†) suggests similar

Table 1 Zeta potential measurements of microspheres at each stage of synthesis

Microspheres	Zeta potential/mV	FWHM/mV
Thiuronium-functionalised microspheres 1	+7.26	2.17
Trimethoxysilane-functionalised microspheres 2	−44.45	2.82
Amine-functionalised microspheres 3	−19.99	2.27
Pyridyl-functionalised microspheres 4	+12.74	2.91
Carboxylic acid-functionalised microspheres 5	−18.74	2.77
RNase-functionalised microspheres 6	+7.41	2.33



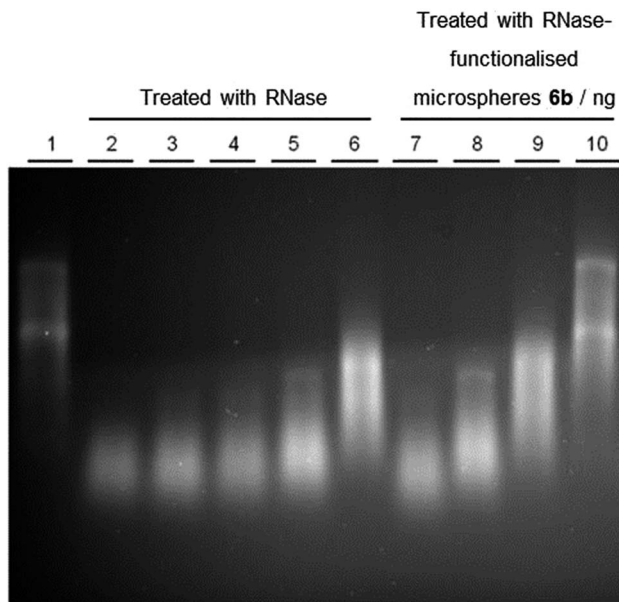


Fig. 4 Gel electrophoresis to compare the activity of native RNase with RNase-functionalised microspheres **6b**. Each reaction was carried out with 1.5 μg of RNA for 3 minutes. Lane (1) control, no treatment, (2) 0.100 units of RNase, (3) 0.075 units of RNase, (4) 0.050 units of RNase, (5) 0.025 units of RNase, (6) 0.001 units of RNase, (7) 1000 ng of RNase-functionalised microspheres **6b** (ca. 0.050 units of RNase), (8) 100 ng of RNase-functionalised microspheres **6b** (ca. 0.025 units of RNase), (9) 10 ng of RNase-functionalised microspheres **6b** (ca. 0.001 units of RNase), (10) 1 ng of RNase-functionalised microspheres **6b**.

RNA degradation between 0.05 units of native RNase (lane 4) and 1000 ng of microspheres (lane 7); 0.025 units of RNase (lane 5) and 100 ng of microspheres (lane 8) and 0.001 units of RNase (lane 6) and 10 ng of microspheres (lane 9).

Cellular delivery

In order to verify the internalisation of microspheres, adherent HeLa cells were incubated with RNase-functionalised microspheres without TR (red)-labelled cores **6c**. After 36 hours, the cells were stained with PKH26 red membrane label according to the manufacturer's instructions, and detached by trypsinisation. Suspended cells were inspected by confocal fluorescence microscopy, and, as expected, RNase-functionalised microspheres were fully internalised by the HeLa cells (for micrographs see ESI, Fig. S10[†]). Previous work by the Bradley group has shown that particles of this size are taken up by the cells using a mechanism that is not endocytosis-based.⁶ Avoidance of endocytosis is desirable for the delivery of functional proteins, since this process typically results in the proteins being targeted to the lysosomes where they are broken down by the harsh degradative environment therein.

Cellular delivery was investigated by incubating HeLa cells with increasing amounts of RNase-functionalised microspheres **6b** (without TR or FITC), specifically 50 μg , 100 μg and 200 μg doses per well of 3 ml of media. Control cells were not incubated with microspheres. After staining with acridine orange, images

were acquired by confocal fluorescence microscopy. Acridine orange is a molecular probe that emits green fluorescence when bound to DNA and emits red fluorescence when bound to RNA.³⁶ Control cells appear healthy (maintaining their characteristic elongated diamond-like shape, with a smooth, defined outline³⁷) in Fig. 5(i)–(iii), with both red and green fluorescence distributed throughout the entirety of each cell.

This suggests that the control cells at the time of imaging were proceeding healthily through stages of mitosis, accounted for by the dividing cells that can be seen with two distinct nuclei. Treated cells appear unhealthy (having an uneven/textured outline as the cell membrane begins to become unstable, they become rounded in shape with a reduced cellular volume³⁷) with a distinct decrease in both green (DNA, nuclear) and red (RNA, cytosolic) fluorescence after treatment with RNase-functionalised microspheres **6b**. The decrease in red fluorescence corresponds to a decrease in cellular RNA, which suggests that the delivered RNase is indeed active. Furthermore,

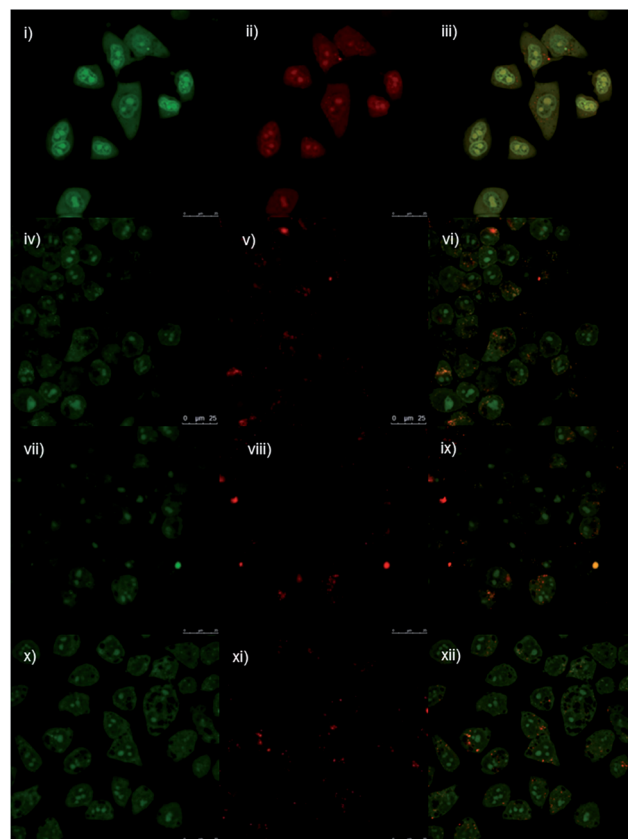


Fig. 5 Confocal microscopy images to show the fluorescence emissions of cells stained with acridine orange. The 488 nm excitation laser was used at 15% power, with 10 \times objective. Cells incubated for 36 hours with (i), (ii) and (iii) no microspheres; (iv), (v) and (vi) 50 μg of RNase-functionalised microspheres **6c**; (vii), (viii) and (ix) 100 μg of RNase-functionalised microspheres **6c**; and (x), (xi) and (xii) 200 μg of RNase-functionalised microspheres **6c**. Left-hand images shows fluorescence emission collected in Channel 1 between 500 nm and 560 nm (DNA); central images show fluorescence emission collected in Channel 2 between 590 nm and 630 nm (RNA); right hand images shows an overlay of the two Channels.



Table 2 Fluorescence ratios of cells stained with acridine orange. Treated cells showed a decrease in red fluorescence (stained RNA) relative to green fluorescence (stained DNA)

Treatment	Channel 1 (green) mean value	Channel 2 (red) mean value	Ratio green : red
Control ^a	11.43	9.99	1 : 0.88
50 ^b µg	11.81	5.21	1 : 0.44
100 ^b µg	6.64	3.12	1 : 0.47
200 ^b µg	14.54	4.2	1 : 0.29

^a No microspheres added to control cells. ^b Mass of RNase-functionalised microspheres **6b**.

the ratio of green (DNA) to red (RNA) fluorescence increases. To quantify this, the relative amounts of fluorescence for each channel were calculated with LAS-AF software (Table 2). It should be noted that lower fluorescence values collected in Channels 1 and 2 for 100 µg of RNase-functionalised microspheres are likely due to the inevitable variability of microscopy sample preparation. However, as the conditions are identical for both channels the ratio remains valid. It is also important to note that the ratios are derived from a qualitative method and are therefore semi-quantitative in nature. Treated cells are rounded and deformed and nuclear changes are apparent, consistent with the onset of apoptosis. Green fluorescence

appears to be more evenly distributed throughout the treated cells, with the nucleoli identifiable in each cell.

Whilst by no means optimised, it was intriguing to assess whether delivered RNase had any effect on the viability of the cells to which it had been delivered. Accordingly, a CellTiter-Blue® Cell Viability Assay was performed (see Experimental section). The results of this 'proof of concept' study are shown in Fig. 6 and indicate that beadfection of HeLa cells with RNase-functionalised microspheres **6b** caused a decrease in cell viability, and significantly so, at the highest concentration of microspheres tested. This is a notable finding since a control experiment with the same concentration of microspheres, but critically, with no RNase attached did not appear to have any detrimental effect on cell viability.

Conclusions

Polymer core-silica shell microspheres were prepared *via* seeded-emulsion and sol-gel polymerisation techniques. The particles were successfully labelled with two concomitant fluorescent markers and a functional RNase enzyme to exemplify the multifunctionality of the system. RNase immobilised on microspheres retained high levels of degradative activity. Delivery of the microspheres with the functional protein to HeLa cells decreased the levels of intracellular RNA and moreover caused a significant decrease in cell viability at the highest concentration tested. Thus the core-shell microspheres that we have developed have been shown to be ideal for *in vitro* functional protein transduction.

Experimental section

Materials and instruments

3-Aminopropyl trimethoxysilane (APTMS); ammonium hydroxide in water 28% w/w; azobisisobutyronitrile (AIBN); 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU); *N,N*-dimethylformamide (DMF); divinylbenzene (DVB); Dulbecco's modified Eagle's medium; *N*-ethyl-*N'*-(3-dimethylaminopropyl)carbodiimide hydrochloride (EDAC); fluorescein *isothiocyanate* (FITC); Fmoc protected Lysine (Fmoc-Lys(Fmoc)-OH); hexadecane; hydrochloric acid (HCl); 3-mercaptopropionic acid; 2-(*N*-morpholino)ethanesulphonic acid (MES); phosphate buffered saline (PBS); poly(*N*-vinyl pyrrolidone) (~58 000 g mol⁻¹) (PVP); sodium dodecyl sulphate (SDS); styrene; succinimidyl 3-(2-pyridyldithio)propionate (SPDP); tetraethylorthosilicate (TEOS); tetrahydrofuran (THF); thiourea; 3-(trimethoxysilyl)propyl methacrylate (TMSPM); and 4-vinylbenzyl chloride (4-VBC) were purchased from Sigma-Aldrich and used as received except for styrene, which was washed with sodium hydroxide solution (1 M) and water and dried over magnesium sulfate prior to use. Acridine orange, BODIPY® TR cadaverine (BODIPY), Texas Red® malimide (TR) and TRIZOL® were purchased from Invitrogen. Foetal calf serum and non-essential amino acids were purchased from PAA, Yeovil, UK. 4-Vinylbenzylisothiouonium chloride (4-VBTU) was synthesised in-house according to our previous protocol.³³

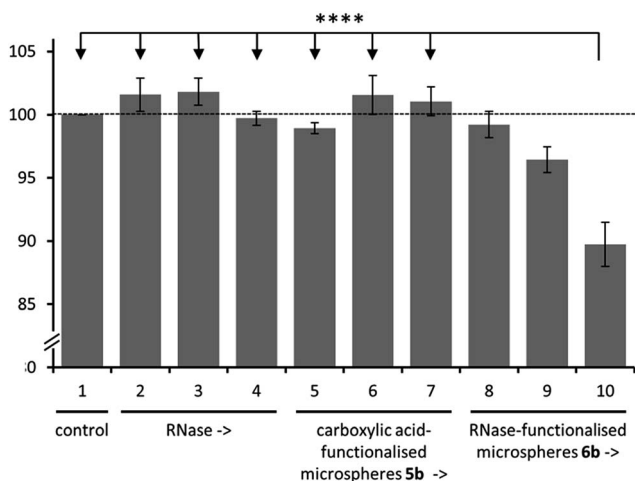


Fig. 6 Graph showing CellTiter-Blue® Cell Viability Assay to compare the effect of native RNase, carboxylic acid-functionalised microspheres **5b** and RNase-functionalised microspheres **6b** upon cell viability. Lane (1) control, no treatment, (2) 84 ng of RNase, (3) 168 ng of RNase, (4) 335 ng of RNase, (5) 1.56 µg of carboxylic acid-functionalised microspheres **5b**, (6) 3.13 µg of carboxylic acid-functionalised microspheres **5b**, (7) 6.25 µg of carboxylic acid-functionalised microspheres **5b**, (8) 1.56 µg of RNase-functionalised microspheres **6b**, (9) 3.13 µg of RNase-functionalised microspheres **6b**, (10) 6.25 µg of RNase-functionalised microspheres **6b**. Data in each case represents the mean percentage results ± S. E. M. of cell fluorescence, calculated using cells from five different passage numbers, performed independently in quadruplet. *P* values, arising from Tukey's multiple comparisons test, of less than 0.0001 are indicated with four (****) asterisks.



Microspheres and HeLa cells were characterised by confocal fluorescence microscopy on a Leica TCS SPS II Broadband confocal microscope with Leica Application Suite-Advanced Fluorescence (LAS-AF) software. The 488 nm excitation laser was used at 15% power, and fluorescence emission collected in two channels. One collected fluorescence in the red region between 590 nm and 630 nm, and the other collected fluorescence in the green region between 500 nm and 560 nm. SEM with Energy Dispersive X-ray Analysis (EDXA) was carried out with a Link System AN10000 and Cambridge Scanning Electron Microscope Stereoscan S90. The quantity of accessible amine groups present on amine-functionalised microspheres **3b** was characterised by UV-vis absorbance spectroscopy on a Perkin Elmer UV-vis Spectrometer Lambda 12. RNase activity was characterised by gel electrophoresis, and images obtained on a Syn-gene G:BOX HR1.4 with GeneSys software. Zeta potential measurements were recorded on a Brookhaven Zeta Plus.

Synthesis of core-shell microspheres

Dispersion polymerisation – synthesis of thiuronium-functionalised polystyrene core microspheres 1. PVP (0.61 g, 1.05×10^{-2} mmol) was dissolved in ethanol (50 ml) by sonication. AIBN (86 mg, 0.525 mmol) and 4-VBTU (120 mg, 0.525 mmol) were dissolved in a mixture of styrene (6.0 ml, 52.5 mmol) and DVB (75 μ l, 0.525 mmol). These solutions were subsequently combined with stirring at room temperature. The mixture and vessel were purged with N_2 for 20 minutes, and subsequently heated to 70 °C. After 16 hours the reaction was cooled to room temperature. To wash a microsphere latex dispersion, a simple three-step process was repeated. Firstly, the dispersion was centrifuged at $7000 \times g$ for 10 minutes, to make the polymer particles form a pellet. Secondly, the supernatant was decanted off. Finally, fresh solvent was added to the pellet and microspheres were re-dispersed by vortex and/or sonication. However, sonication is incompatible with RNase-functionalised microspheres due to potential denaturation of the protein. The latex product was washed twice with methanol (2 \times 100 ml) to remove any unreacted monomer and twice with DMF (2 \times 100 ml). Finally, thiuronium-functionalised polystyrene microspheres **1** were re-suspended in DMF at a concentration of 500 mg per 50 ml.³³

Labelling of thiuronium-functionalised polystyrene core microspheres 1 with Texas Red maleimide*. TR in DMF (1 mg in 1 ml) was added to thiuronium-functionalised polystyrene microspheres **1** in DMF (500 mg, 50 ml) with shaking at room temperature for 16 hours. The TR (red)-labelled microspheres were washed twice with DMF (2 \times 40 ml), twice with methanol (2 \times 40 ml), twice with water (2 \times 40 ml), and finally re-suspended in SDS solution (20 mg SDS per 25 ml of water) at a concentration of 500 mg of microspheres in 25 ml. *It is important to note that when using microspheres in delivery to HeLa cells for confocal fluorescence microscopy assays TR was NOT included in the microsphere core.

Seeded-emulsion polymerisation – synthesis of poly[3-(trimethoxysilyl)propyl methacrylate] shell 2. AIBN (63 mg, 3.8×10^{-4} mol) was dissolved in styrene (0.880 ml, 7.68×10^{-3} mol),

DVB (54.5 μ l, 3.8×10^{-4} mol), TMSPM (191 μ l, 8.7×10^{-4} mol) and hexadecane (129 μ l, 4.4×10^{-4} mol). This initiator-monomer mixture was added to thiuronium-functionalised polystyrene microspheres **1** in SDS solution (500 mg, 25 ml) and mixed with continuous stirring at room temperature. Both mixture and vessel were purged with N_2 for 20 minutes, and subsequently heated to 70 °C for 5 hours. After cooling to room temperature, the trimethoxysilane-functionalised core-shell microspheres **2** were washed twice with water (2 \times 40 ml), twice with methanol (2 \times 40 ml), twice with water (2 \times 40 ml), then finally re-suspended in water (420 mg, 42 ml).

Sol-gel extension of silica shell with tetraethylorthosilicate and aminopropyl trimethoxysilane – synthesis of amine-functionalised microspheres 3. An aqueous solution of ammonium hydroxide 28% w/w (400 μ l) was added to trimethoxysilane-functionalised core-shell microspheres **2** in water (200 mg, 40 ml) with stirring at room temperature. 200 μ g of FITC* was added to APTMS (200 μ l) and shaken for 1 hour. After this time, TEOS (200 μ l) was mixed with the FITC-APTMS, and the mixture immediately added to the microsphere **2** suspension. Subsequently, the vessel was heated to 50 °C for 2 hours with continual stirring. The reaction mixture was cooled to room temperature, and the resultant amine-functionalised microspheres **3** were then washed three times with water (3 \times 10 ml) and finally re-suspended in water (100 mg, 10 ml). *It is important to note that when using microspheres in delivery to HeLa cells for confocal fluorescence microscopy assays, and microspheres used with FITC (green)-labelled RNase, FITC was NOT included in the silica shell.

Modification of core-shell microspheres and conjugation of RNase

Reaction of amine-functionalised silica shell with succinimidyl 3-(2-pyridyldithio)propionate. SPDP in DMF (2 mg, 200 μ l) was added to amine-functionalised microspheres **3** in PBS (100 mg, 10 ml) suspension with shaking for 1 hour at room temperature. The pyridyl-functionalised microspheres **4** were washed twice with water, once with DMF and finally re-suspended in DMF (100 mg, 10 ml).

Disulfide exchange of pyridyl functionalisation with carboxylic acid. 3-Mercaptopropionic acid (40 μ l) was added to pyridyl-functionalised microspheres **4** in DMF (100 mg, 10 ml) suspension and mixed with shaking for 1 hour at room temperature. The carboxylic acid-functionalised microspheres **5** were washed twice with DMF, once with MES and re-suspended in MES (10%, pH 5.5) (100 mg, 10 ml).

EDAC coupling of RNase to carboxylic acid-functionalised microspheres. EDAC in water (1 mg, 1 ml) was added to carboxylic acid-functionalised microspheres **5** in MES solution (10%, pH 5.5) (50 mg, 10 ml) and mixed with shaking for 1 hour at room temperature. The microspheres were washed twice with MES (10%, pH 5.5) and re-suspended in MES (10%, pH 5.5). 2 mg of RNase in 1 ml MES (10%, pH 5.5) was added to the EDAC-activated microspheres and mixed with shaking for 2 hours at room temperature. The RNase-functionalised microspheres **6** were washed twice with ethanol to sterilise the



microspheres, and finally re-suspended in DMEM at a concentration of 1 mg ml⁻¹.

Gel electrophoresis

The enzymatic activity of RNase-functionalised microspheres **6b** was assessed by reaction with purified RNA, and characterised by gel electrophoresis. Specifically, HeLa cells were seeded at 1 million cells per well, in 6-well plates in DMEM. After 24 hours, the RNA was harvested and purified with TRIzol® according to the manufacturer's instructions. For each reaction, a 1 ml centrifuge tube was charged with 1.5 µg of RNA in 15 µl of 200 mM sodium acetate, 10 mM Tris HCl buffer (pH 7.5). 5 µg of carboxylic acid-functionalised microspheres **5b**, or 5, 1, 0.1, 0.01, or 0.001 µg of RNase-functionalised microspheres **6b**, or 2.86, 2.15, 1.43, 0.72 or 0.03 ng of native RNase, in 15 µl of 200 mM sodium acetate, 10 mM Tris HCl buffer (pH 7.5) were added to the RNA, mixed well and incubated for 1, 3 or 5 minutes at room temperature. After this time, 1 µl of sodium hypochlorite solution (11–15% available chlorine) was added (with mixing) to each to terminate the degradation reaction. Subsequently, 4 µl of RNA loading buffer was added to each to give the final reaction mixture for gel electrophoresis. A gel electrophoresis system, suitable for RNA, was made based on a protocol described previously by Jorcyk and co-workers.³⁴ Specifically, a stock solution of TAE buffer was made up with 120 g of Tris base dissolved in 300 ml of deionised water, and to this 28 ml of anhydrous acetic acid and 50 ml of EDTA (500 mM) solution. 20 ml of TAE buffer was added to 980 ml of deionised water and mixed thoroughly. 200 ml of this was transferred to a 500 ml Schott bottle with magnetic stirrer bar inside, and 2 g of agarose added. The mixture was stirred magnetically for 1 minute, and 2 ml of sodium hypochlorite solution (11–15% available chlorine) was added. After this, the Schott bottle was removed from the stirrer plate and placed in the microwave with the lid fitted loosely, for a total of 2 minutes at 600 W, with gentle stirring at 20 second intervals. Subsequently, the gel was allowed to cool down slowly on the bench top and stirred, until around 50 °C (comfortable to touch). After this time 5 µl of ethidium bromide (10 mg ml⁻¹ in H₂O) was added and stirred thoroughly before casting into the gel mould. The gel was covered, allowed to set by cooling to room temperature, then placed in the electrophoresis tray, and covered with the remaining 800 ml of TAE 1% buffer. Each degradation reaction mixture was loaded into a lane in the gel and electrophoresed at a constant voltage of 8 V cm⁻¹. The addition of hypochlorite, or bleach, inactivates all RNases in the system, to prevent any further degradation of RNA during the electrophoresis process.³⁴

Acridine orange assay with HeLa cells

HeLa 229 (human epithelial cells) were cultured in Dulbecco's modified Eagle's medium/C high glucose (4.5 mg ml⁻¹) supplemented with 10% v/v foetal calf serum and 100 IU ml⁻¹ penicillin and 100 mg ml⁻¹ streptomycin supplemented with non-essential amino acids (the complete mixture is referred to as DMEM). Glass coverslips (12 mm diameter) were sterilised and one placed in the bottom of each well in a 6-well plate. HeLa

cells were seeded at 250 000 cells per well, on top of the coverslips, in 3 ml of DMEM. After 12 hours incubation at 37 °C, 5% CO₂, the cells had adhered to the coverslips. RNase-functionalised microspheres **6b** were prepared without TR or FITC fluorophores and suspended in DMEM. 0 µg (control), 50 µg, 100 µg or 200 µg of RNase-functionalised microspheres **6b** were then added to the wells and mixed evenly throughout the media. After 36 hours of incubation at 37 °C, 5% CO₂, excess media was removed from the well, and the cells on the coverslips were washed gently three times with 3 ml of PBS, using a pipette. Excess PBS was removed, and 150 µl of acridine orange in PBS (10 mg ml⁻¹) was pipetted onto the cells on the coverslip and left for 5 minutes. The cells were again washed gently with PBS three times. Subsequently, the coverslips were carefully removed from the well with tweezers and placed cell-side down on microscope slides. The live cells were imaged immediately by confocal fluorescence microscopy.

CellTiter-Blue® Cell Viability Assay

HeLa cells were seeded in quadruplet into 96-well plates at 5000 cells per well in 200 µl of DMEM, and incubated for 24 hours at 37 °C, 5% CO₂. After this time, cells were treated with microspheres; 6.25, 3.13 or 1.56 µg of carboxylic acid-functionalised microspheres **5b** or RNase-functionalised microspheres **6b**, or 335, 168 or 84 ng of native RNase was added to the wells, and control cells were left untreated. After 48 hours of incubation at 37 °C, 5% CO₂, cells were washed (3 × 100 µl) with phenol red-free DMEM, with an auto-pipette. CellTiter-Blue® reagent was diluted with phenol red-free DMEM in a 1 : 6 ratio, and 120 µl of the solution added to each well (20 µl CellTiter-Blue® reagent, 100 µl phenol red-free DMEM per well). Cells were incubated for a further 4 hours at 37 °C, 5% CO₂ and absorbance measured at 570 nm. Results were averaged and used to represent percentage cell viability. Absorption of treated cells was expressed as a percentage of control cells, which were taken as 100% viable. This experiment was repeated for five different passage numbers of cells.

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