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Cite this: *Inorg. Chem. Front.*, 2022, 9, 1490The Mössbauer effect using ^{57}Fe -ferrabisdicarbollide ($[\text{o-}^{57}\text{FESAN}]^-$): a glance into the potential of a low-dose approach for glioblastoma radiotherapy†Ana B. Buades,^a Laura C. J. Pereira,^b Bruno J. C. Vieira,^b Ana C. Cerdeira,^b João C. Waerenborgh,^b Teresa Pinheiro,^c António P. A. Matos,^d Catarina G. Pinto,^b Joana F. Guerreiro,^b Filipa Mendes,^b Srecko Valic,^e Francesc Teixidor,^b Clara Viñas^{b,*a} and Fernanda Marques^{b,*b}

Although a variety of cancers are initially susceptible to chemotherapy, they eventually develop multi-drug resistance. To overcome this situation, more effective and selective treatments are necessary using anti-tumour agents that act in two or more ways and offer greater therapeutic benefits over single-mechanism entities. In this study, we report on treating cancer with $\text{Na}[\text{3,3'}\text{-}^{57}\text{Fe}(\text{1,2-C}_2\text{B}_9\text{H}_{11})_2]$, which offers the possibility of dual action (radiation–drug combinations) to improve the clinical benefits and reduce healthy tissue toxicity. An approach to evaluating the potential of $[\text{o-}^{57}\text{FESAN}]^-$ to treat glioblastoma using the Mössbauer effect is presented. As the therapeutic outcomes rely on the amount and distribution of $[\text{o-}^{57}\text{FESAN}]^-$ inside the cells, several studies, using magnetization, Mössbauer spectroscopy and nuclear microscopy techniques, were performed to ascertain the uptake of $[\text{o-}^{57}\text{FESAN}]^-$ in U87 glioblastoma cells. $[\text{o-}^{57}\text{FESAN}]^-$ was found to be within the cells; 29% of its uptake was in the nuclear fraction, which is a particularly desirable target, because the nucleus is the cell's control centre where DNA and the transcription machinery reside. Irradiation studies with 2D and 3D cellular models of U87 cells showed that the growth inhibition effect observed was more pronounced when $[\text{o-}^{57}\text{FESAN}]^-$ was used in combination with the Mössbauer effect in low total dose regimens, suggesting that this procedure either alone or as adjuvant may be useful for glioblastoma treatment.

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Introduction

Cancer is a multifactorial disorder, resulting from the combined effect of genetic and environmental factors. Predictions indicate an increase of up to 21.6 million cases of cancer every

year until 2030, making cancer the leading cause of death worldwide.

The hallmarks of cancer include independence from growth signals, non-response to signals that stop cell division, uncontrolled replication and the evasion of apoptosis. Mössbauer spectroscopy, based on the recoilless emission and absorption of gamma rays in solids, is a versatile technique that provides important information regarding the chemical, structural and magnetic properties of a material in many areas of science, such as Physics, Chemistry, and Biology.¹ The applications of Mössbauer spectroscopy in biomedical research have become established through at least fifty years of experience in this field.

For iron-containing coordination complexes, Mössbauer spectroscopy is effective in determining the iron oxidation state (Fe^{2+} and Fe^{3+}), as well as the electronic spin-state, coordination number and environment, effectively discriminating different iron sites.

Iron is an ideal element for Mössbauer spectroscopy.² A source of ^{57}Co ($t_{1/2} = 270$ d) decays by electron capture (EC from the K-shell) to the 136 keV nuclear level of ^{57}Fe . This

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excited state decays to the ground state of ^{57}Fe (15%) to the 14.4 keV level (85%). The half-life of this level ($t_{1/2}$ ca. 100 ns) and the emission of the 14.4 keV photon are ideally suited for ^{57}Fe Mössbauer spectroscopy. A fraction of the emitted 14.4 keV photons is recoil free and can be absorbed by a ^{57}Fe nucleus in a sample. Since the Mössbauer effect is only observed in the ^{57}Fe nuclei, which account for 2% of natural Fe, the study of samples with very low Fe content requires enrichment of the ^{57}Fe isotope.

In fact, in biological systems, iron is by far one of the most structurally and relevant metal ions and ^{57}Fe Mössbauer spectroscopy has been very useful in the elucidation of the electronic structures of iron-containing biomolecules, the role of iron in living processes and pathological changes in living systems, organs and tissues.^{3–6}

Radiation therapy (RT) is the most frequently used therapeutic option for the treatment of cancer but high doses of ionizing radiation are delivered to tumours since only a small fraction of the absorbed dose can damage cancer cells. Moreover, the efficacy of RT is limited by the resistance of tumours as well as by the toxicity to normal cells and tissues.⁷ The use of radiosensitizers has been suggested as a promising option to overcome the adverse side effects of RT and make this treatment more effective without increasing the dose of radiation, which may be detrimental to normal tissues. The radiosensitizing effect of platinum(II) cisplatin, whose primary target in cells is DNA,⁸ has encouraged the design of many other transition metal coordination complexes or organometallic complexes and their assessment as anticancer candidates.^{9–11}

Mössbauer absorption represents a method of increasing the radiosensitivity of tumours in terms of orders of magnitude *via* selective energy deposition in cancer cells. The secondary radiation accompanying resonance absorption of Mössbauer radiation in ^{57}Fe nuclei may be able to produce a powerful local effect on biological tissues. After gamma ray absorption, the excited nucleus decay is accompanied by a cascade of Auger or low-energy (of the order of 1 keV) conversion electrons that have much higher biological efficiency than the primary quanta. Therefore, a targeted radiation cancer treatment is foreseen using a method based on the use of secondary radiation excited in a ^{57}Fe target by a Mössbauer source. Moreover, irradiated tumour cells preloaded with iron (^{57}Fe) compounds with resonant Mössbauer γ rays could result in DNA damage through the production of Auger electrons and X-rays. The DNA double-strand breaks are more efficiently produced by Auger electrons (1–10 eV) generated in proximity to DNA. The energy of these electrons is dissipated within a cell diameter and produces multiple double-strand DNA breaks. A single Auger event can be lethal to a cell and is comparable with more than 10^5 photon absorption events in conventional radiotherapy. $^{57}\text{Fe(III)}$ -Bleomycin, administered to malignant cells *in vitro* and *in vivo* and irradiated with resonant Mössbauer gamma rays (14.4 keV), causes ablation of the malignant cells, presumably by Auger cascade, with extremely small radiation doses, about 10^{-5} Gy. As a basis for com-

parison, a dose of about 5 Gy is necessary to achieve a similar effect with conventional radiotherapy.¹²

In the field of tumour pathology, as an alternative to or in support of conventional radiotherapy, cancer cell inactivation by Mössbauer effect could also have future applications. Patients with chronic myelogenous leukemia (CML) could benefit from this therapeutic modality. Experiments done with bone marrow cells collected from normal donors and K562 leukaemic cells demonstrated a growth inhibition effect after treatment with hematin and low-energy γ rays from a ^{57}Co Mössbauer source.¹³ Thus, Mössbauer cancer therapy promises the advantages of selective radiation without systemic exposure of normal tissue, as well as higher killing effect per event secondary to the production of Auger electrons.

Radiotherapy is a common step during the treatment of glioblastoma multiforme (GBM), the most malignant brain cancer among adults and the second most common pediatric malignant tumour, with a very poor prognosis.¹⁴ Rapid proliferation, high invasiveness and radiation resistance are the main reasons behind the poor therapeutic outcomes. To increase the effectiveness of radiotherapy, the use of radiosensitizers can enhance the radiosensitivity of tumour cells and reduce the radiation dose and the adverse effects on normal brain tissue. The radiosensitization effect of high-Z metals has been evaluated. It was found that the radiation dose was enhanced due to the generation of secondary X-rays, photoelectrons and Auger electrons with subsequently higher energy deposition in the cancer cells.¹⁵ At present, the conventional radiotherapy sensitizers used in the clinic include 5-fluorouracil, temozolomide and platinum drugs, which can enhance the radiotherapy sensitivity of tumour cells through different mechanisms of action. However, these conventional radiotherapy sensitizers can cause many adverse side effects and acquired resistance.¹⁶

Recent studies have demonstrated that 3D hybrid organic-inorganic compounds, known as carboranes, can have a role in the development of novel small molecules for the treatment of various types of cancer.^{17–21} The synthetic flexibility of carboranes, along with their rigid structure and chemical stability, makes them appealing moieties for their use in cancer drug development.^{22,23} Although carborane-based drugs are yet to be approved for therapeutic purposes, many compounds have shown promising properties for biomedical applications, in particular for boron neutron capture therapy (BNCT).^{24–30} In other biomedical fields, cobaltabis(dicarbollide) complexes known as $[o\text{-COSAN}]^-$ species have shown important selective HIV-1 protease inhibition activity and, if combined with specific biomolecules, could be promising for targeted therapy.³¹

A study by Mössbauer spectroscopy of intramolecular rotations and electronic states of Fe in the salt $\text{Cs}[o\text{-FESAN}]$ and the zwitterion boron compound $[\text{Fe}(\text{8}-(\text{C}_2\text{H}_4\text{O})_2\text{-1,2-C}_2\text{B}_9\text{H}_{10})(1',2'\text{-C}_2\text{B}_9\text{H}_{11})]$ was recently published.³² In the present work, we aim to evaluate the potential of ^{57}Fe -ferrabis(dicarbollide), herein $[o\text{-}^{57}\text{FESAN}]^-$, as a radiosensitizer *via* the Mössbauer effect to treat glioblastoma. As the therapeutic



efficiency relies on the cellular amount and distribution of $[o\text{-}^{57}\text{FESAN}]^-$, we have evaluated quantitatively the net iron uptake in both GBM and normal astrocyte cell lines (U87 and rat astrocyte RA, respectively). Furthermore, the spatial subcellular distribution of iron has been measured with nuclear microscopy techniques, which rely on energetic ion beams focused to micrometre dimensions.³³ These techniques offer the distinct possibility of imaging the morphology of single cells with relatively high resolution and quantitating elemental distribution with high-sensitivity ($\mu\text{g g}^{-1}$ range) in conditions close to those found *in vivo*, without the need for stains or contrast agents.³⁴ These possibilities are of the utmost importance for the preservation of the original physiological conditions to ensure realistic two-dimensional microanalysis in single cells, enabling Fe distribution to be visualized and its uptake quantified.

Experimental

Synthesis of $\text{Na}[o\text{-}^{57}\text{FESAN}]$

The first step was the synthesis of $[\text{NMe}_4][o\text{-}^{57}\text{FESAN}]$ (see ESI†), which was achieved by the complexation reaction of the dicarbollide ligand ($[\text{7,8-C}_2\text{B}_9\text{H}_{11}]^{2-}$) with $^{57}\text{FeCl}_2$ using a modified method³⁵ for the original synthesis by Hawthorne *et al.*³⁶ The sodium salt of the anionic $[o\text{-}^{57}\text{FESAN}]^-$ was obtained by means of the cationic exchange resin of its caesium salt as displayed in Scheme 1. The Na^+ salt was characterized by ^1H and ^{11}B NMR, EPR as well as by MALDI-TOF, as reported in the ESI.†

Paramagnetic properties of $\text{Na}[o\text{-FESAN}]$

As the anionic $[o\text{-FESAN}]^-$ complex is an $S = 1/2$ species, to assess the magnetic properties of the paramagnetic $\text{Na}[o\text{-FESAN}]$, EPR measurements on $\text{Na}[o\text{-FESAN}]$ were also run in the solid state and in acetone solution. The EPR spectra of the recently reported low-spin $[\text{NMe}_4][o\text{-FESAN}]$ salt³⁷ and ferrocene were also run for comparison.

Magnetization and Mössbauer spectroscopy studies

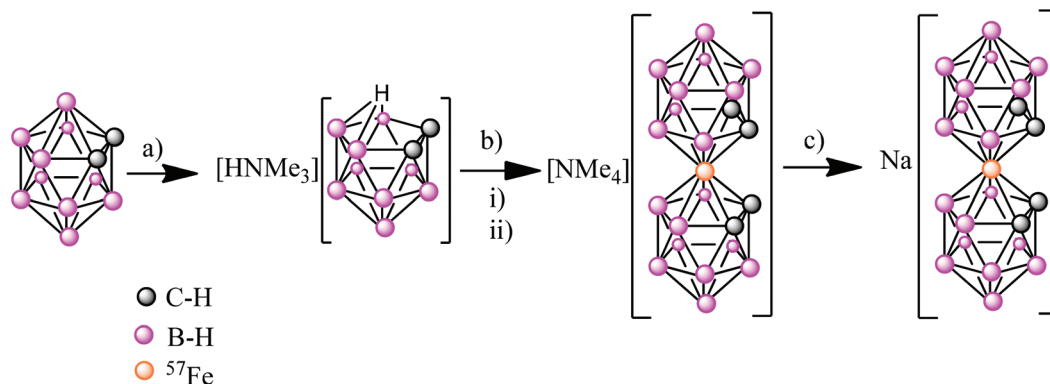
Magnetic measurements of the parent ferrabis(dicarbollide), $[o\text{-}^{57}\text{FESANE}]^-$ anion, as well as of the U87 glioblastoma and RA cells exposed to $[o\text{-}^{57}\text{FESAN}]$, were performed using a SQUID magnetometer S700X (Cryogenic Ltd) in the 4–300 K temperature range under fields of 0.1 T (iron compound) and 1 T (cell samples). Mössbauer spectra of parent $\text{Na}[o\text{-}^{57}\text{FESAN}]$ and U87 and RA cells loaded with $\text{Na}[o\text{-}^{57}\text{FESAN}]$ (300 μM , 6 h incubation) were collected at 80 K using a conventional constant acceleration spectrometer and a ^{57}Co (Rh) source. The velocity scale was calibrated with an $\alpha\text{-Fe}$ foil. The spectra were fitted to Lorentzian lines using a non-linear least-squares method.³⁸ The measurements run at 80 K were obtained with the samples immersed in He gas in a bath cryostat.

Irradiation studies

For cell irradiation with the ^{57}Co source, the velocity range of the spectrometer transducer was restricted to the Doppler frequencies where resonant γ -ray absorption was observed in the Mössbauer spectra of the cells. This procedure implies that the sample was permanently subjected to the γ -ray absorption *via* the Mössbauer effect during the irradiation time of ~ 2 h. The dose received by the sample due to all the γ - and X-rays emitted by the ^{57}Co source was ~ 0.4 mGy.

Biological studies

Cell viability assays. The glioblastoma cell line U87 (ATCC) and the rat astrocyte (RA) cell line (Sigma-Aldrich) were cultured in MEM (U87) containing GlutaMax-I supplemented with 10% FBS and in astrocyte medium (RA) (Gibco, ThermoFisher) at 37 °C in a humidified atmosphere of 5% CO_2 (Heraeus, Germany). Cells were seeded in 96-well plates at a density of 2×10^4 cells (U87) and 5×10^4 cells (RA) per well in 200 μL of culture medium and left to incubate overnight for optimal adherence. After careful removal of the medium, 200 μL of a dilution series of the $\text{Na}[o\text{-}^{57}\text{FESAN}]$ compound (from a stock solution of 1 mM in medium) were added and



Scheme 1 Synthesis of $\text{Na}[o\text{-}^{57}\text{FESAN}]$. (a) Partial deboronation of the neutral *closo* 1,2- $\text{C}_2\text{B}_{10}\text{H}_{12}$. (b) Complexation reaction of the monoanionic *nido* $[\text{7,8-C}_2\text{B}_9\text{H}_{11}]^{2-}$ ligand in two steps: (i) deprotonation reaction with $\text{K}[\text{t-BuO}]$ in dry THF and (ii) complexation of the dianionic *nido* $[\text{7,8-C}_2\text{B}_9\text{H}_{11}]^{2-}$ with $^{57}\text{FeCl}_2$ to obtain $[\text{3,3'}\text{-}^{57}\text{Fe}(\text{1,2-C}_2\text{B}_9\text{H}_{11})_2]^-$ (abbreviated as $[o\text{-}^{57}\text{FESAN}]^-$) that was precipitated as a salt of $[\text{NMe}_4]^+$. (c) The Na^+ salt was obtained from the $[\text{NMe}_4]^+$ using cationic exchange resin.



Imaging Fe distribution in U87 cells. U87 cells ($5-6 \times 10^5$ cells) were prepared for nuclear microscopy experiments by seeding the cells on silicon frames with 5×3 windows of 0.56 mm^2 covered with a 100 nm thick silicon nitride membrane (Silson Ltd, UK) in 6-well plates and incubated overnight. The silicon nitride windows were previously sterilized with UV light. Cells were incubated with $\text{Na}[o\text{-}^{57}\text{FESAN}]$ for 24 h at $50 \text{ }\mu\text{M}$ (a non-cytotoxic dose) at $37 \text{ }^\circ\text{C}$ in a $5\% \text{ CO}_2$ incubator. After incubation, the medium was removed and the windows were washed with cold PBS. Samples were frozen at $-80 \text{ }^\circ\text{C}$ and then dried in a cryostat at $-25 \text{ }^\circ\text{C}$ for subsequent analysis as described elsewhere.³⁴ Nuclear microscopy analysis of single cells was carried out in a vacuum with a 2.0 MeV proton beam focused to $2 \text{ }\mu\text{m}$ dimensions at the nuclear microprobe facility (Oxford Microbeams Ltd, UK) of IST.³³ By scanning the beam across the target cell on the silicon nitride window, several techniques can be used simultaneously, enabling visualization of the distribution of several elements and their localization and quantitation in the cell. Two-dimensional images of Fe and physiological elements in the cell can be obtained with PIXE, together with cell morphological details using scanning transmission ion microscopy (STIM) to obtain density maps and matrix composition estimation by analyzing the elastic backscattering spectrometry (EBS) spectra, which is necessary for PIXE mass normalization and to obtain quantitative data ($\mu\text{g per g dry material}$).³³ Images of whole U87 cells were taken with scan sizes ranging between 53×53 and $25 \times 25 \text{ }\mu\text{m}^2$. Data acquisition, image processing and concentration calculation were performed using the OMDAQ-2007 system.⁴³

Imaging cellular morphological alterations (TEM). U87 cells seeded into 6-well plates at approx. 70% confluence were treated with Na[*o*-⁵⁷Fe-FESAN] at 50 μM for 24 h. After incubation, cells were irradiated with the ⁵⁷Co source (0.4 mGy, 2 h) and processed following a standard procedure described previously.³⁴

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Gol'danskii Karyagin effect as discussed in detail in $[\text{NMe}_4][o\text{-FESAN}]$.⁴⁷

The spectrum obtained for the cells incubated with $\text{Na}[o\text{-}^{57}\text{FESAN}]$ consists of a symmetric quadrupole doublet (Fig. 3). The small relative transmission for the cell measurements is due to the very small amount of Fe in the cells. The recoilless factor and consequently the relative transmission do not change significantly between 80 and 4 K. The IS is the same as in $\text{Na}[o\text{-}^{57}\text{FESAN}]$, showing that the electronic state of Fe in the cells is the same as in the starting $\text{Na}[o\text{-}^{57}\text{FESAN}]$.

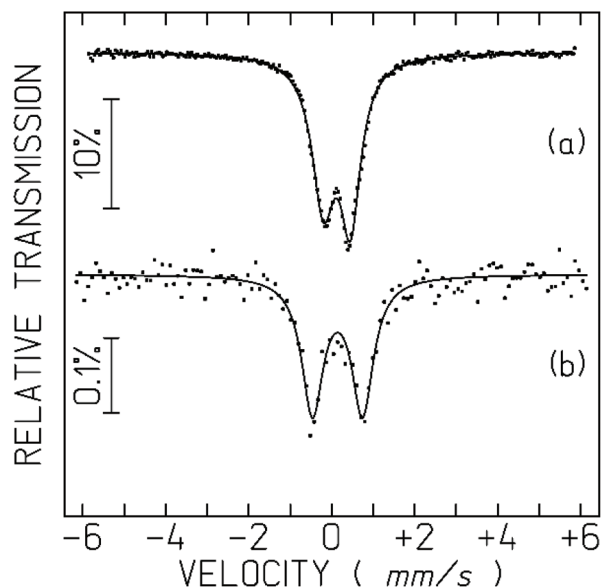


Fig. 3 Mössbauer spectra of $\text{Na}[o\text{-}^{57}\text{FESAN}]$ (a), and the U87 cells incubated with $\text{Na}[o\text{-}^{57}\text{FESAN}]$ (b) taken at 80 K in the solid state. The lines over the experimental points are fitted quadrupole doublets. The estimated parameters are collected in Table 1.

The strong increase in the quadrupole splitting, QS, confirms that the Fe^{3+} environment within the cells is different from $[o\text{-}^{57}\text{FESAN}]^-$ in the starting $\text{Na}[o\text{-}^{57}\text{FESAN}]$. This may be related to the fact that $[o\text{-FESAN}]^-$ produces intercalative interaction with DNA.⁴⁹

Cytotoxic activity in 2 D cell model

The concentrations of $[o\text{-}^{57}\text{FESAN}]^-$ for the irradiation studies were previously optimized by calculating the IC_{50} values at 24 h of incubation. The effect of $[o\text{-}^{57}\text{FESAN}]^-$ on the cellular viability of U87 cells and of RA cells, used as non-tumour, normal glial cells, was evaluated using the MTT assay, and the results are depicted in Fig. 4. As can be seen, the IC_{50} values obtained for both cell lines indicated that, after 24 h of treatment, $\text{Na}[o\text{-}^{57}\text{FESAN}]$ concentrations lower than 100 μM did not induce severe cytotoxic effects and, accordingly, were selected for the subsequent studies with the 2D cell model.

Cellular uptake by PIXE

The radiosensitizing potential of $\text{Na}[o\text{-}^{57}\text{FESAN}]$ could strongly depend on its uptake by the cells. Therefore, studies were carried out to quantify the levels of Fe inside the cells using PIXE and to evaluate any difference in the uptake of $[o\text{-}^{57}\text{FeSAN}]^-$, between the U87 and RA cells. PIXE is a multi-elemental technique that enables quantification of other elements inside the cells, such as K, Ca, Cu and Zn, and thus gives insights into any imbalance in these physiological elements. The results depicted in Fig. 5 show that the cells loaded with $[o\text{-}^{57}\text{FESAN}]^-$ present higher Fe levels than the control (non-treated cells) for both cell lines. However, the uptake in the U87 cells is approximately four times higher than that in the RA cells, in either the whole cells or the nuclear fractions.

These results support the entry of $[o\text{-}^{57}\text{FESAN}]^-$ into the cytoplasm of the cell, reaching the nucleus, which is a particu-

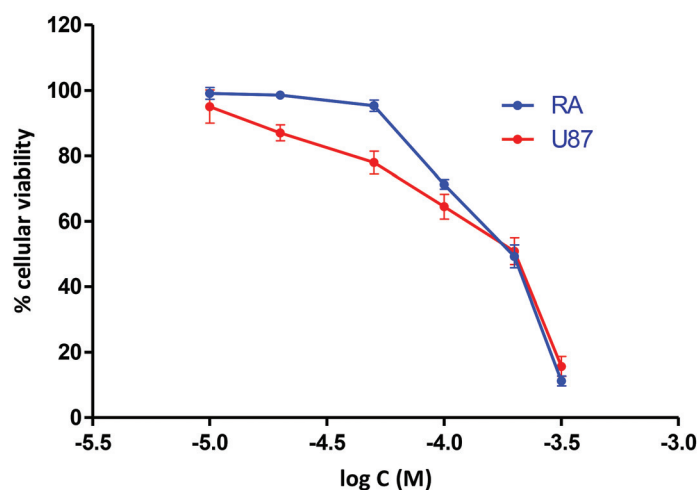


Fig. 4 Cell viability of U87 and RA cells after 24 h exposure to $\text{Na}[o\text{-}^{57}\text{FESAN}]$. The IC_{50} values were calculated from dose–response curves analysed with the GraphPad Prism software (vs. 5.0).

IC_{50} (μM), 24h	
U87	RA
150 ± 23	182 ± 57



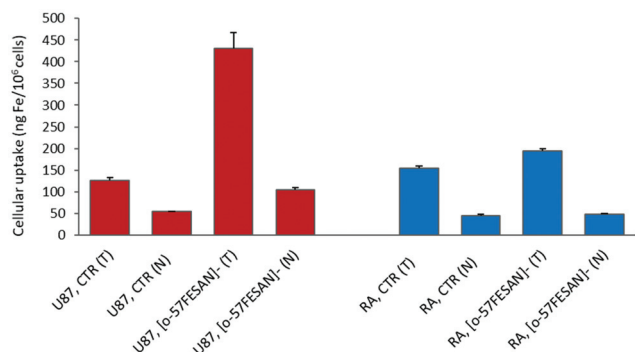


Fig. 5 Concentrations of Fe in U87 and RA whole cell (T) and nuclear (N) fractions after treatment with 50 μM Na[o-⁵⁷FESAN]. The Fe concentrations in the whole-cell and nuclear for the control cells are also plotted. Values are shown as the mean $\pm$ SD.

larly desirable target because the nucleus is the cell's control centre, in which DNA and the transcription machinery reside. Actually, *in silico* studies (Monte Carlo simulations) have shown that the cell-killing effect is more efficient if the energy deposition of the radiation reaction occurs in the nucleus of cells compared with a uniform distribution within the whole cell.⁵⁰ To date, only studies related to DNA interaction with Na[o-FESAN] that demonstrate a strong intercalative interaction between Na[o-FESAN] and CT-dsDNA have been reported.⁴⁹ The high cellular uptake of small anionic [o-⁵⁷FESAN][−] molecules by U87 cells and their accumulation in the nucleus (29%) support earlier theoretical results of the effect of the charge and hydrophobicity properties on the translocation of model particles through the nuclear pore complex.⁵¹ The generic hydrophobicity and negative charge of model particles are necessary for their easy translocation through nuclear membranes.⁵¹

Concerning the relevant physiological elements (see Table S1 at the ESI†), the following data were observed: (i) in both U87 and RA whole cells, the Zn levels increased twice after treatment with Na[o-⁵⁷FESAN] (in ng per 10⁶ cells, U87 controls *vs.* treated, 99 $\pm$ 7 and 180 $\pm$ 17; RA controls *vs.* treated, 28 $\pm$ 2 and 48 $\pm$ 3); while (ii) in the respective nuclear fractions an opposite effect was observed for Zn and Ca (in ng per 10⁶ cells, U87 control *vs.* treated, Zn = 60 $\pm$ 2 and 20 $\pm$ 2; Ca = 460 $\pm$ 10 and 180 $\pm$ 4; RA controls *vs.* treated, Zn = 57 $\pm$ 4 and 26 $\pm$ 2, Ca = 415 $\pm$ 8 and 237 $\pm$ 4). In treated RA whole cells, K decreased by 40% (controls 2000 $\pm$ 100, treated 1125 $\pm$ 37). However, in both cell lines, Cu remained essentially unchanged. This observation indicates a similar response of U87 and RA cells towards [o-⁵⁷FESAN][−]. The changes in K, Ca and Zn contents in glial cells may reflect disturbances in the K⁺ channels involved in the membrane potential, intracellular Ca²⁺ signalling and impaired nuclear activity, after treatment with Na[o-⁵⁷FESAN]. These changes may have consequences for the level of growth control, cell volume and activity regulation in cancer cells.^{52–54} Nevertheless, the interpretation of these results requires further experiments.

Imaging and elemental distribution by nuclear microscopy

The U87 cells that showed a higher uptake towards [o-⁵⁷FESAN][−] were inspected by nuclear microscopy techniques to visualize the micro distribution of Fe and other physiological elements, such as P, K, Ca and Zn. Fig. 6 depicts representative images of mass density and elemental distributions in U87 cells under the control and 50 μM Na[o-⁵⁷FESAN] conditions.

The images of mass density showed that U87 control cells are larger with a denser central region, probably corresponding to the nucleus. In treated U87 cells, the density images showed a reduction in the cell size that can translate to a volume decrease.

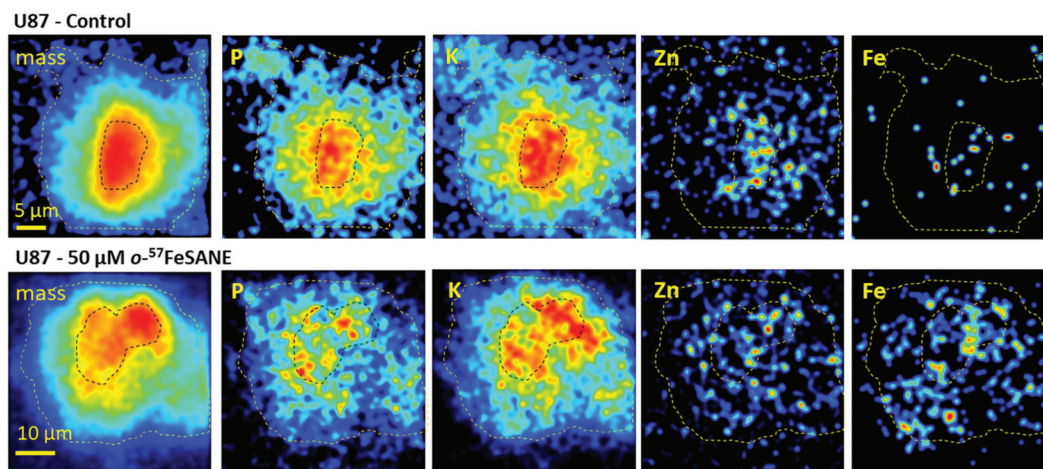


Fig. 6 Mass density (ass) and 2D elemental maps of the distribution of P, K, Zn and Fe in whole U87 cells: control and 50 μM Na[o-⁵⁷FESAN] treated conditions. The dotted lines indicate the cell contour and the central nuclear region. The mass density and elemental distributions are represented by a colour gradient with a dynamic scale: high level, red, to the low level, dark blue.



The most remarkable difference in the elemental distribution of treated U87 cells was the Fe uniform distribution and its increased content across the whole cells, contrasting with the scattered distribution in the control, as can be observed in Fig. 6. The average Fe concentration in whole control cells was low, below 150 μg per g dry weight, whereas in treated cells a 6-fold increase to 849 ± 296 μg per g dry weight was observed. In treated cell peripheral regions and in the denser central nuclear region Fe concentrations did not significantly differ (780 ± 320 vs. 950 ± 230 μg per g dry weight). These concentration values are similar to those obtained in the bulk analysis of whole cells and nuclear fractions (concentrations ranging from 600 $\mu\text{g g}^{-1}$ in total cells to 850 $\mu\text{g g}^{-1}$ in the nuclear fraction, on a dry weight basis). Therefore, even for non-toxic concentrations of $\text{Na}[o\text{-}^{57}\text{FESAN}]$, the Fe content in treated cells dramatically increased, which is in line with uptake results. More importantly, Fe does not seem to be compartmentalized in the cell, as its distribution is relatively homogeneous throughout the cell.

The micro distribution of P, K, Ca and Zn in U87 treated cells cannot give a clear definition of cell compartments, contrasting with control cells, where the nuclear region is clearly associated with high contents of P, K and Zn (Fig. 6). In particular, the changes in the microdistribution of Zn reflect those observed in whole cells and nuclear fraction as described above. The rationale behind these elemental fingerprints is the essentiality of the phosphate backbone in the DNA structure,⁵⁵ of the K^+ cation in chromatin stability (although K^+ is mainly a cytoplasm ion)⁵⁶ and of Zn as a constituent of protein domains (zinc fingers) and a cofactor of enzymes intrinsically linked to DNA replication machinery.⁵⁴ Therefore, the alteration of P, K and Zn compartmentalization in U87 treated cells suggests cellular remodeling after exposure to $[o\text{-}^{57}\text{FESAN}]^-$ and the subsequent intracellular increase in the Fe concentration.

Evaluation of the Mössbauer irradiation effect

The radiosensitizing potential of $[o\text{-}^{57}\text{FESAN}]^-$ was evaluated in both cell lines, using the MTT assay before and after Mössbauer irradiation with and without $\text{Na}[o\text{-}^{57}\text{FESAN}]$. Cells were previously treated with $\text{Na}[o\text{-}^{57}\text{FESAN}]^-$ at 50 and 100 μM followed by 24 h incubation before irradiation. Concerning the non-irradiated cells, no significant cytotoxicity was observed after 24 h of incubation at 50 μM $\text{Na}[o\text{-}^{57}\text{FESAN}]$ for U87 cells, as shown in Fig. 7. The exposure to Mössbauer radiation on the U87 cells incubated with 50 μM $\text{Na}[o\text{-}^{57}\text{FESAN}]$ induced a decrease in cellular viability of about 30% compared with the control (irr) and of about 30% compared with $\text{Na}[o\text{-}^{57}\text{FESAN}]$ (non-irr). At higher doses (100 μM $\text{Na}[o\text{-}^{57}\text{FESAN}]$), an enhanced radiosensitizing effect was observed for $\text{Na}[o\text{-}^{57}\text{FESAN}]$ ($19.7 \pm 3.8\%$), 80% higher compared with the control (irr) and about 60% higher compared with $\text{Na}[o\text{-}^{57}\text{FESAN}]$ (non-irr). For the RA cells, only a slight enhancement of the radiation effect was observed at the same higher dose (100 μM $\text{Na}[o\text{-}^{57}\text{FESAN}]$), indicating that this effect was rather specific for cancer cells.

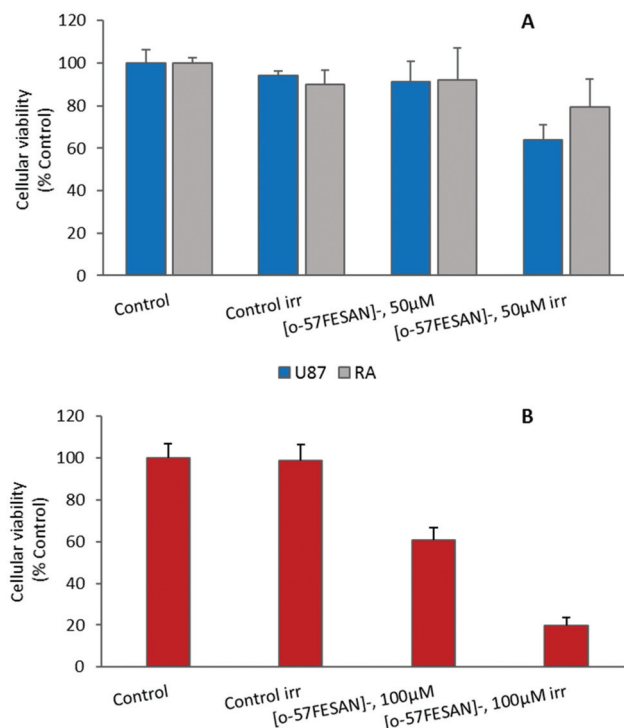


Fig. 7 Cell viability of U87 and RA cells after exposure to Mössbauer irradiation (0.4 mGy, 2 h). (A) U87 (blue colour) and RA (grey colour) cells incubated with $\text{Na}[o\text{-}^{57}\text{FESAN}]$ at 50 μM ; (B) U87 cells incubated with $\text{Na}[o\text{-}^{57}\text{FESAN}]$ at 100 μM . Results are shown as the mean $\pm$ SD of two independent experiments done with at least six replicates per condition.

Further irradiation studies were also performed as proof-of-concept experiments to confirm that the Mössbauer effect was responsible for the cell death observed. ^{57}Co ($t_{1/2} = 272$ d) decays by electron capture to the 136.4 keV level in ^{57}Fe , which is de-excited both by 136.4 keV (12.3%) γ rays and by emission of 122 keV (87.5%) and 14.4 keV γ rays (see the decay scheme below). The 122 keV γ rays have a relative probability of 87.5%. Therefore, X-rays at equivalent energy and dose (0.4 mGy) were used to irradiate cells. According to these results (Fig. 8), X-ray irradiation did not induce significant killing of the cells.

Effects of $\text{Na}[o\text{-}^{57}\text{FESAN}]$ in 3D culture models

Cytotoxic activity of $\text{Na}[o\text{-}^{57}\text{FESAN}]$ in 3D models. The cytotoxic activity of $\text{Na}[o\text{-}^{57}\text{FESAN}]$ was further evaluated in more advanced culture models, where tumour cells were grown in a 3D environment, forming spheroids. The concentrations of $\text{Na}[o\text{-}^{57}\text{FESAN}]$ selected to evaluate its cytotoxic activity in spheroid cultures of U87 GBM cells were based on IC_{50} values determined for monolayer cultures, using the MTT assay (around 150 μM for both cell lines), with concentrations one-third below and above these values also being taken into consideration. The viability of the spheroids was studied following a 24 h incubation period, as shown in Fig. 9a, using the CellTiter-Glo® 3D Cell Viability Assay, suitable for advanced culture models. In parallel, the spheroids' growth was also



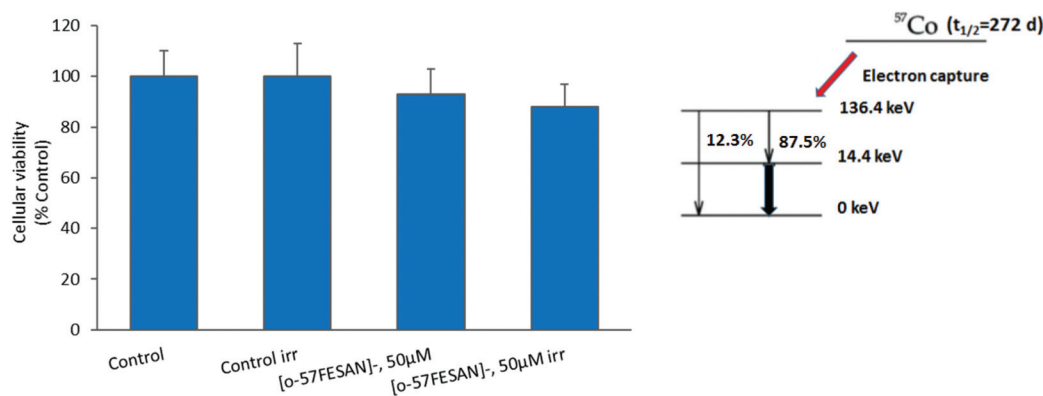


Fig. 8 Cellular viability of U87 GBM cells after irradiation with X-rays in the presence of 50 μM Na[o-⁵⁷FESAN]. Irradiation was done using a Philips MCN 165 X-ray tube and a YXLON 9421 high-voltage generator at a 0.4 mGy radiation dose. ISO beam quality N120 was used ($E_{\text{max}} = 120 \text{ keV}$, $E_{\text{ave}} = 100 \text{ keV}$, filtered with 4 mm Al + 5 mm Cu + 1 mm Sn).

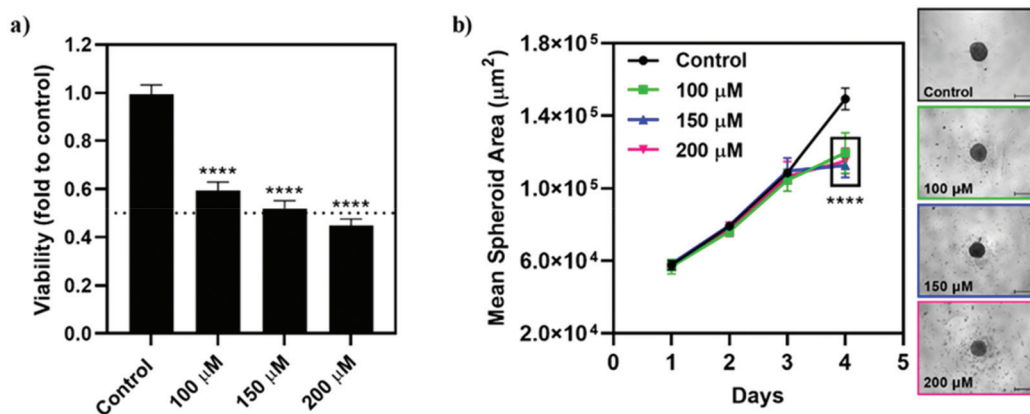


Fig. 9 Effects of the incubation of Na[o-⁵⁷FESAN] on U87 spheroids. (a) Cellular viability of U87 spheroids at 24 h, assessed using the CellTiter-Glo® 3D Cell Viability Assay, and (b) spheroid growth after incubation with Na[o-⁵⁷FESAN] at 100, 150 and 200 μM, for 24 h, on day 3, represented by the mean spheroid area (in μm²) as a function of the number of days in culture. Controls consist of spheroids incubated only with the medium. Data are presented as the average ± SEM of 5/6 replicates. Statistical significance was calculated using one-way ANOVA, followed by Dunnett's test comparing treated spheroids with control spheroids (**** $p \leq 0.0001$).

evaluated throughout the culture period, as depicted in Fig. 9b.

Qualitatively, the incubation of the spheroids with Na[o-⁵⁷FESAN] did not seem to affect the shape and integrity of the spheroids. The circularity of the spheroids was also not severely affected after incubation with Na[o-⁵⁷FESAN], with the spheroids having maintained regular shapes as reflected by circularity values very close to 1 (data not shown). Quantitatively, the growth and size of the spheroids from the U87 cell line were affected after 24 h of incubation with Na[o-⁵⁷FESAN] (Fig. 9b), considering that treated spheroids stopped growing after incubation with the compound, which was not observed for spheroids from the control condition.

The viability results obtained (Fig. 9a) reflect an obvious cytotoxic effect of [o-⁵⁷FESAN]⁻ on the spheroids from U87,

suggesting that [o-⁵⁷FESAN]⁻ can penetrate in more complex cellular systems, affecting tumour resistance mechanisms.

Based on the importance of this type of metallacarborane compound, we thus foresee that Na[o-⁵⁷FESAN] could be explored within the scope of a low-dose approach for glioblastoma radiotherapy. In our previous report on metallabis (dicarbollide), Na[o-COSAN], we demonstrated that this type of compound showed no cytotoxicity in normal cells and no acute toxicity in a mouse model.³⁹ Although the results are preliminary and further studies are needed for the validation of this approach, the present work has lifted the veil on a new type of low-dose radiation glioblastoma therapy with such promising small anionic compounds. Therefore, we consider that the ferrabis(dicarbollide) Na[o-⁵⁷FESAN] described in this study could constitute a promising approach for radiation therapies, also benefiting from its solubility in water, as



solubility is a main challenge in the development of chemotherapeutic drugs.

Radiosensitizing effect of $\text{Na}[o\text{-}^{57}\text{FESAN}]$ on spheroids

The radiosensitizing effect of $[o\text{-}^{57}\text{FESAN}]^-$ was further evaluated in spheroid cultures from the tumour cell line, U87, through the CellTiter-Glo® 3D Cell Viability Assay. The concentrations used were chosen based on the IC_{50} values determined for monolayer cultures, using the MTT assay, while concentrations one-third below and above these values were also used, as before. Spheroids were incubated with $\text{Na}[o\text{-}^{57}\text{FESAN}]$ for 24 h, after which they were irradiated and left to recover for an additional 24 h period, followed by the viability assessment (Fig. 10a). In

parallel, the spheroids' growth was evaluated throughout the culture period, as depicted in Fig. 10b. As before, the circularity of the spheroids was not severely affected by incubation with $\text{Na}[o\text{-}^{57}\text{FESAN}]$ and irradiation (data not shown).

Spheroids from the tumour cell line stopped growing after being subjected to incubation with $\text{Na}[o\text{-}^{57}\text{FESAN}]$ and Mössbauer irradiation (Fig. 10a). However, the irradiation effect on its own is not significant when comparing irradiated spheroids with spheroids that were not irradiated, both for the control and for the $\text{Na}[o\text{-}^{57}\text{FESAN}]$ treated cells.

The viability results obtained suggest that there is a cytotoxic effect on U87 spheroids that is slightly increased after irradiation (Fig. 10a).

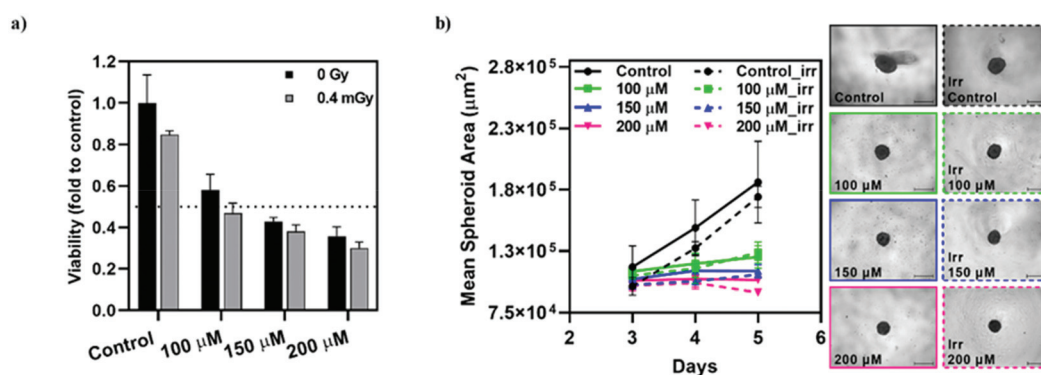


Fig. 10 Radiosensitizing effects of $\text{Na}[o\text{-}^{57}\text{FESAN}]$ on U87 spheroids. (a) Cellular viability of U87 spheroids incubated with $\text{Na}[o\text{-}^{57}\text{FESAN}]$ for 24 h and irradiated, assessed using the CellTiter-Glo® 3D Cell Viability Assay. (b) U87 spheroid growth after incubation with $\text{Na}[o\text{-}^{57}\text{FESAN}]$ at 100, 150 and 200 μM, for 24 h, on day 3, followed by irradiation on day 4 and viability assessment on day 5, represented by the mean spheroid area (in μm^2) as a function of the number of days in culture. Controls consist of spheroids incubated only with the medium. Data are presented as the average $\pm$ SEM of 2 replicates.

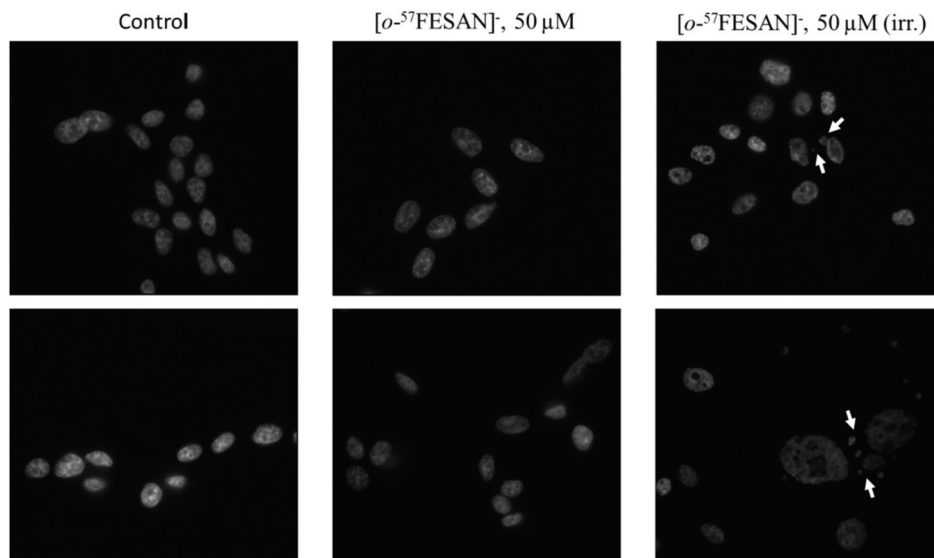


Fig. 11 Fluorescence microscopy for visualizing cellular responses in non-treated cells (control; left panel), 50 μM $\text{Na}[o\text{-}^{57}\text{FESAN}]$ -treated but non-irradiated cell (middle panel), or 50 μM $\text{Na}[o\text{-}^{57}\text{FESAN}]$ -treated cells that were also subjected to Mössbauer irradiation (right panel). Morphological damage to the nucleus was observed as indicated by the presence of nuclear fragments and abnormalities (white arrows).



Author contributions

Conflicts of interest

Acknowledgements

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