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# Magnetoliposomes as carriers for promising antitumor thieno[3,2-*b*]pyridin-7-arylamines: photophysical and biological studies†

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Magnetoliposomes containing superparamagnetic manganese ferrite nanoparticles were tested as nanocarriers for two new promising antitumor drugs, a *N*-(3-methoxyphenyl)thieno[3,2-*b*]pyridin-7-amine (**1**) and a *N*-(2-methoxy-phenyl)thieno[3,2-*b*]pyridin-7-amine (**2**). The fluorescence emission of both compounds was studied in different polar and non-polar media, evidencing a strong intramolecular charge transfer character of the excited state of both compounds. These *in vitro* potent antitumor thienopyridine derivatives were successfully incorporated in both aqueous and solid magnetoliposomes, with encapsulation efficiencies higher than 75%. The magnetic properties of magnetoliposomes containing manganese ferrite nanoparticles were measured for the first time, proving a superparamagnetic behaviour. Growth inhibition assays on several human tumor cell lines showed very low GI<sub>50</sub> values for drug-loaded aqueous magnetoliposomes, comparing in most cell lines with the ones previously obtained using the neat compounds. These results are important for future drug delivery applications using magnetoliposomes in oncology, through a dual therapeutic approach (simultaneous chemotherapy and magnetic hyperthermia).

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## Introduction

The high potential of magnetic nanoparticles for biomedical applications has been widely recognized due to their unique size and physicochemical properties. Nanoparticles with superparamagnetic behavior are preferred in biomedicine, as they exhibit a strong magnetization only when an external magnetic field is applied.<sup>1,2</sup>

Liposomes entrapping magnetic nanoparticles (magnetoliposomes) are of large importance in drug delivery, as they can be guided and localized to the therapeutic site of interest by external magnetic field gradients and used in cancer treatment by hyperthermia.<sup>3,4</sup> Magnetoliposomes have been proposed as T2 contrast agents (negative contrast enhancement) in MRI,<sup>5</sup> while in therapy they have been used as a chemotherapy

alternative through magnetic-controlled drug delivery and thermotherapy.<sup>6–8</sup> Considering the wide applications of magnetoliposomes, much attention has been paid to the synthesis of different kinds of magnetic nanoparticles<sup>9–11</sup> and liposomes.<sup>12–15</sup>

Magnetoliposomes are specially promising as nanocarriers for potential antitumor drugs. Thienopyridine derivatives have been reported as possessing antiangiogenic,<sup>16–21</sup> antitumor,<sup>21–29</sup> or both activities.<sup>30</sup> Among the potential antitumor di(hetero)arylamines in the thieno[3,2-*b*]pyridine series, prepared earlier by some of us, the ones with an *o*-methoxy or *m*-methoxy groups relative to the NH (compounds **1** and **2**, Fig. 1) presented the lowest growth inhibitory concentration (GI<sub>50</sub>) values in several human tumor cell lines, between 0.09 and 0.31 μM for compound **1** and between 1.40 and 5.91 μM for compound **2**.<sup>29</sup> Compound **1** revealed to be more active

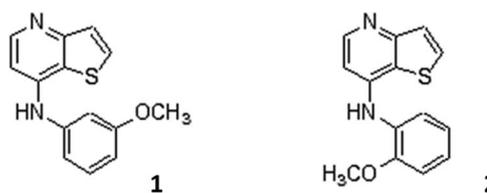


Fig. 1 Structure of compound **1**, *N*-(3-methoxyphenyl)thieno[3,2-*b*]pyridin-7-amine, and compound **2**, *N*-(2-methoxyphenyl)thieno[3,2-*b*]pyridin-7-amine.

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than the well-known anticancer agent ellipticine against the cell lines MCF-7 (breast adenocarcinoma), NCI-H460 (non-small cell lung carcinoma), HepG2 (hepatocellular carcinoma), HCT15 (colon adenocarcinoma), and HeLa (cervical carcinoma), while compound 2 showed the lowest  $GI_{50}$  against HeLa cells. However, these compounds also presented low  $GI_{50}$  values in non-tumor PLP2 cells (1.94  $\mu\text{M}$  for compound 1 and 6.56  $\mu\text{M}$  for compound 2).<sup>29</sup>

Considering future applications of these new compounds as antitumor drugs, they were incorporated in magnetoliposomes, either aqueous or solid, containing manganese ferrite nanoparticles, suitable for biomedical applications.<sup>31</sup>

Magnetoliposomes may constitute ideal nanocarrier systems for encapsulation and transport of these drugs, as they can be guided and localized in tumor cells by a magnetic field gradient, and act simultaneously as chemotherapy/hyperthermia agents in a synergistic approach, reducing drug dosage due to the dual therapeutic effect.

## Experimental

All the solutions were prepared using spectroscopic grade solvents and ultrapure water (Milli-Q grade).

### Preparation of magnetoliposomes

Manganese ferrite nanoparticles (NPs) were synthesized by the coprecipitation method, as previously described.<sup>31</sup> For magnetoliposomes preparation, the lipids egg yolk phosphatidylcholine (Egg-PC), dipalmitoylphosphatidylcholine (DPPC) and cholesterol (Ch), from Sigma-Aldrich, were used. For aqueous magnetoliposomes preparation, a 10 mM lipid solution in ethanol was injected, under vigorous vortexing, to an aqueous solution of magnetic nanoparticles, above the melting transition temperature of the lipids (ethanolic injection method).<sup>32</sup> After encapsulation, the ferrofluid was washed with water and purified by ultracentrifugation to remove all the non-encapsulated NPs.

Solid magnetoliposomes (SMLs) were prepared by a method previously developed.<sup>33</sup> First, 10  $\mu\text{L}$  of the synthesized  $\text{MnFe}_2\text{O}_4$  NPs were dispersed in 3 mL of water and centrifuged. Then, the deposited particles were dispersed in 10  $\mu\text{L}$  of water in an ultrasonicator, for one minute at 189 W, and 3 mL of chloroform were added to the aqueous dispersion of NPs. After vigorous agitation, 165  $\mu\text{L}$  of a 20 mM solution of dipalmitoylphosphatidylcholine (DPPC) were added under vortexing, to form the first lipid layer of the SMLs. The particles were washed twice by magnetic decantation with pure water, in order to remove the lipid that was not attached to the NPs. The second lipid layer was then formed by the injection of 165  $\mu\text{L}$  of DPPC (20 mM), under vortexing, in a 3 mL aqueous dispersion of the particles with the first layer. The resulting SMLs were then washed and purified with ultrapure water by centrifugation.

Compounds 1 and 2 were incorporated into aqueous magnetoliposomes by the co-injection method, as already described.<sup>34</sup> In solid magnetoliposomes, the diarylamines 1 and 2 were incorporated by injection of an ethanolic solution together with the formation of the second lipid layer.

### Spectroscopic measurements

**General methods.** Absorption spectra were recorded in a Shimadzu UV-3101PC UV-Vis-NIR spectrophotometer. Fluorescence measurements were performed using a Fluorolog 3 spectrofluorimeter, equipped with double monochromators in both excitation and emission and a temperature controlled cuvette holder. Fluorescence spectra were corrected for the instrumental response of the system.

The fluorescence quantum yields,  $\Phi_s$ , were determined by the standard method (eqn (1)),<sup>35,36</sup>

$$\Phi_s = [(A_r F_s n_s^2)/(A_s F_r n_r^2)] \Phi_r \quad (1)$$

where  $A$  is the absorbance at the excitation wavelength,  $F$  the integrated emission area and  $n$  is the refraction index of the solvents. Subscripts refer to the reference (r) or sample (s) compound. The absorbance value at excitation wavelength was always less than 0.1, in order to avoid inner filter effects.

**Fluorescence anisotropy measurements.** The steady-state fluorescence anisotropy,  $r$ , is calculated by

$$r = \frac{I_{VV} - GI_{VH}}{I_{VV} + 2GI_{VH}} \quad (2)$$

where  $I_{VV}$  and  $I_{VH}$  are the intensities of the emission spectra obtained with vertical and horizontal polarization, respectively (for vertically polarized excitation light), and  $G = I_{HV}/I_{HH}$  is the instrument correction factor, where  $I_{HV}$  and  $I_{HH}$  are the emission intensities obtained with vertical and horizontal polarization (for horizontally polarized excitation light).

**FRET measurements.** The formation of a DPPC bilayer around manganese ferrite nanoparticles was investigated by Förster Resonance Energy Transfer (FRET), using the labeled lipids NBD- $\text{C}_6$ -HPC (1-palmitoyl-2-[6-[(7-nitro-2-1,3-benzoxadiazol-4-yl)amino]hexanoyl]-*sn*-glycero-3-phosphocholine) and rhodamine B-DOPE (*N*-(lissamine rhodamine B sulfonyl)-1,2-dioleoyl-*sn*-3-phosphatidylethanolamine (ammonium salt)), both from Avanti Polar Lipids (Fig. 2).

FRET efficiency,  $\Phi_{\text{RET}}$ , defined as the proportion of donor molecules that have transferred their excess energy to acceptor molecules, can be obtained by taking the ratio of the donor integrated fluorescence intensities in the presence of acceptor ( $F_{\text{DA}}$ ) and in the absence of acceptor ( $F_{\text{D}}$ ) (eqn (3)),<sup>37</sup>

$$\Phi_{\text{RET}} = 1 - \frac{F_{\text{DA}}}{F_{\text{D}}} \quad (3)$$

The distance between donor and acceptor molecules can be determined through the FRET efficiency (eqn (4)),

$$r_{\text{AD}} = R_0 \left[ \frac{1 - \Phi_{\text{RET}}}{\Phi_{\text{RET}}} \right]^{1/6} \quad (4)$$

where  $R_0$  is the Förster radius (critical distance), that can be obtained by the spectral overlap,  $J(\lambda)$ , between the donor emission and the acceptor absorption, according to eqn (5) and (6) (with  $R_0$  in  $\text{\AA}$ ,  $\lambda$  in nm,  $\epsilon_A(\lambda)$  in  $\text{M}^{-1} \text{cm}^{-1}$ ),<sup>37</sup>

$$R_0 = 0.2108 [k^2 \Phi_{\text{D}}^0 n^{-4} J(\lambda)]^{1/6} \quad (5)$$



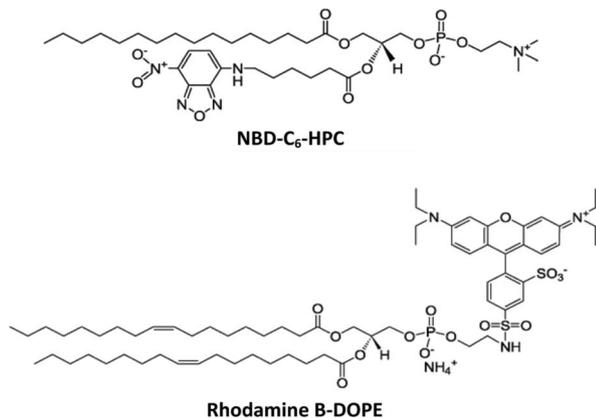


Fig. 2 Structure of the labeled lipids NBD-C<sub>6</sub>-HPC and rhodamine B-DOPE.

$$J(\lambda) = \int_0^{\infty} I_D(\lambda) \varepsilon_A(\lambda) \lambda^4 d\lambda \quad (6)$$

where  $k^2 = 2/3$  is the orientational factor assuming random orientation of the dyes,  $\Phi_D^0$  is the fluorescence quantum yield of the donor in the absence of energy transfer,  $n$  is the refraction index of the medium,  $I_D(\lambda)$  is the fluorescence spectrum of the donor normalized so that  $\int_0^{\infty} I_D(\lambda) d\lambda = 1$ , and  $\varepsilon_A(\lambda)$  is the molar absorption coefficient of the acceptor.

For determination of fluorescence quantum yield of NBD-C<sub>6</sub>-HPC (energy donor) in magnetoliposomes, this fluorescent labeled lipid incorporated in lipid membranes was used as reference,  $\Phi_r = 0.32$  at 25 °C, as reported by Invitrogen.<sup>38</sup>

### Compound encapsulation efficiency

The encapsulation efficiency, EE (%), of antitumor compounds was determined through fluorescence emission measurements. After preparation, drug-loaded magnetoliposomes (MLs) were subjected to centrifugation at 11 000 rpm for 60 minutes. The supernatant was pipetted out and its fluorescence was measured, allowing to determine the compound concentration using a calibration curve previously obtained. Three independent measurements were performed for each system and standard deviations (SD) were calculated. The encapsulation efficiency of compounds 1 and 2 was determined using the following equation:

$$EE (\%) = \frac{(\text{total amount} - \text{amount of non encapsulated compound})}{\text{total amount}} \times 100 \quad (7)$$

### Scanning electron microscopy (SEM)

Scanning electron microscopy (SEM) images of solid magnetoliposomes were recorded using a Scanning Electron Microscope FEI – Nova 200 NanoSEM. For the negative staining procedure, a 2% aqueous solution of ammonium molybdate tetrahydrate was prepared. Then, 20  $\mu\text{L}$  of sample and 20  $\mu\text{L}$  of staining

solution were mixed and a drop of this mixture was placed onto a Formvar grid, held by tweezers. After 20 seconds, almost all the solution was removed with filter paper and left dry.

### Magnetic properties of magnetoliposomes

Magnetic measurements were performed at room temperature in a Superconducting Quantum Interference Device (SQUID) magnetometer (Quantum Design MPMS5XL), using applied magnetic fields up to 5.5 T.

The magnetization hysteresis loop measurements were performed by fixing the temperature and measuring the magnetization at a series of different applied magnetic fields. This type of study gives information about the saturation magnetization, the degree to which the sample remains magnetized when the applied field is removed, and how easily the sample magnetization can be reversed, the so-called coercive field. The magnetization was corrected for the diamagnetic contribution from the lipids and water and normalized by the mass of the samples, which was determined after drying them.

### Growth inhibitory activity of drug-loaded AMLs on human tumor cell lines and on porcine liver primary cells

Two types of aqueous magnetoliposomes with different compositions (100% Egg-PC and 95 : 5 Egg-PC : DSPE-PEG2000-Folate, from Avanti Polar Lipids) were loaded with compound 1 or compound 2, with different compound concentrations (0.03  $\mu\text{M}$  to 7.5  $\mu\text{M}$ ). *In vitro* cytotoxicity evaluation was assessed for four human tumor cell lines, namely MCF-7 (breast adenocarcinoma), NCI-H460 (non-small cell lung cancer), HeLa (cervical carcinoma) and T3M4 (pancreatic cancer). The cell line PLP2 (non-tumor cells) was used to evaluate the toxicity to healthy tissues. The cell lines were obtained from Leibniz Institute DSMZ – German Collection of Microorganisms and Cell Cultures.

The cells were routinely maintained as adherent cell cultures in RPMI-1640 medium containing 10% heat-inactivated FBS, at 37 °C, in a humidified air incubator containing 5% CO<sub>2</sub>. Each cell line was plated at an appropriate density ( $1.0 \times 10^4$  cells per well) in 96-well plates and allowed to attach for 24 h. The cells were then treated for 48 h with the different solutions. Following this incubation period, the adherent cells were fixed by adding cold 10% TCA (100  $\mu\text{L}$ ) and incubated for 60 min at 4 °C. Plates were then washed with deionized water and dried. A sulforhodamine B (SRB) solution (0.1% in 1% acetic acid, 100  $\mu\text{L}$ ) was then added to each plate-well and incubated for 30 min at room temperature. Unbound SRB was removed by washing with 1% acetic acid. The plates were air-dried and the bound SRB was solubilized with 10 mM Tris-HCl buffer (200  $\mu\text{L}$ , pH = 7.4). The absorbance was then measured at 540 nm in a microplate reader.<sup>39,40</sup> The results were expressed in GI<sub>50</sub> values (concentration that inhibited 50% of net cell growth).

## Results and discussion

### Photophysical properties in homogeneous solution

The promising antitumor properties of the di(hetero)arylamines 1 and 2 (Fig. 1)<sup>29</sup> inspired us to study the photophysical



behaviour of these compounds in homogeneous solution, as they exhibit intrinsic fluorescence. Thus, the absorption and fluorescence properties of the thieno[3,2-*b*]pyridine derivatives **1** and **2** were studied in several solvents of different polarity. The maximum absorption ( $\lambda_{\text{abs}}$ ) and emission wavelengths ( $\lambda_{\text{em}}$ ), molar absorption coefficients and fluorescence quantum yields are shown in Table 1. The normalized fluorescence spectra are displayed in Fig. 3 and 4 (examples of absorption spectra are also shown).

The thieno[3,2-*b*]pyridines **1** and **2** present moderate to high absorption coefficient values ( $\epsilon \geq 7 \times 10^3 \text{ M}^{-1} \text{ cm}^{-1}$ ) in all solvents, having also very reasonable fluorescence quantum yields (Table 1), a similar behaviour to that observed in other thienopyridine derivatives previously studied.<sup>26,34,42,43</sup>

Additional common features with other thienopyridines are the negligible fluorescence in protic media (due to hydrogen bonding with this type of solvents), the red shifts in emission and band enlargement with increasing solvent polarity,<sup>26,34,42,43</sup> usually attributed to an intramolecular charge transfer (ICT) character of the excited state.<sup>37</sup> The effect is more pronounced for compound **2**, exhibiting larger bathochromic shifts in polar solvents (a red shift of 120 nm between cyclohexane and dimethylsulfoxide, Table 1).

The significant sensitivity of the emission of diarylamines **1** and **2** to their environment can be useful to understand their behaviour in magnetoliposomes of different compositions.

### Magnetoliposomes formulations

Both aqueous (AMLs) and solid (SMLs) magnetoliposomes containing manganese ferrite nanoparticles were prepared. The influence of lipid membrane fluidity in AMLs was investigated using egg-phosphatidylcholine (Egg-PC), a fluid phosphatidylcholine mixture at room temperature,<sup>44</sup> dipalmitoylphosphatidylcholine (DPPC), with a melting transition temperature of 41 °C,<sup>45</sup> and Egg-PC/cholesterol 7 : 3, the latter mixture usually being used as model of biological membranes.<sup>46,47</sup> Neat Egg-PC and Egg-PC : cholesterol aqueous magnetoliposomes containing lauric acid coated MnFe<sub>2</sub>O<sub>4</sub> nanoparticles were also previously prepared by Pradhan and coworkers,<sup>48</sup> the Egg-PC : cholesterol 2 : 1 being

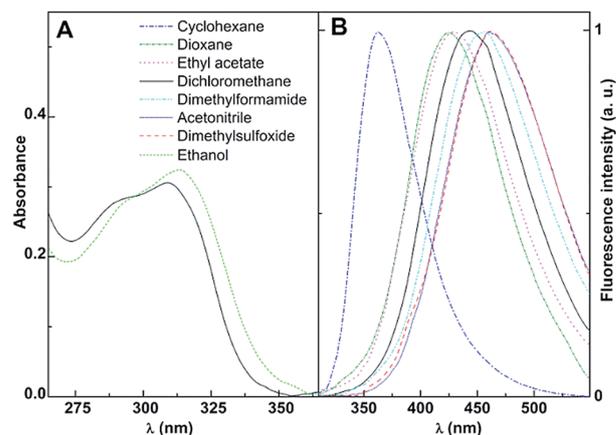


Fig. 3 (A) Absorption spectra of  $2 \times 10^{-5}$  M solutions of compound **1** in dichloromethane and ethanol, as examples. (B) Normalized fluorescence spectra ( $\lambda_{\text{exc}} = 310$  nm) of  $3 \times 10^{-6}$  M solutions of compound **1** in several solvents.

the most promising formulation for hyperthermia treatment of cancer.

Here, DPPC was used for the first time in the preparation of solid magnetoliposomes containing manganese ferrite NPs. The interest of this phospholipid is its melting transition temperature slightly above the physiological temperature, thus being promising for the release of drugs enhanced by the phase transition in combined chemotherapy/hyperthermia strategies. In fact, several studies have demonstrated that the thermal enhancement of the cytotoxic activity of many anticancer agents is maximized at mild hyperthermia temperatures (40.5–43 °C).<sup>49</sup>

The formation of the double DPPC bilayer around a cluster of magnetic nanoparticles was investigated by FRET assays. Thus, the NDB labeled lipid NBD-C<sub>6</sub>-HPC was included in the second lipid layer of SMLs (NBD acting as the energy donor), while the labeled lipid rhodamine B-DOPE (energy acceptor) was incorporated in the first lipid layer. The emission of SMLs containing both donor and acceptor labeled lipids was measured exciting only NBD, and was compared with the emission of SMLs labeled with only the energy donor (Fig. 5). Comparing the fluorescence spectra of the two systems, a decrease in the NBD emission

Table 1 Maximum absorption ( $\lambda_{\text{abs}}$ ) and emission ( $\lambda_{\text{em}}$ ) wavelengths, molar absorption coefficients ( $\epsilon$ ) and fluorescence quantum yields ( $\Phi_{\text{F}}$ ) for the thieno[3,2-*b*]pyridine derivatives **1** and **2** in several solvents

Solvent	$\lambda_{\text{abs}}$ (nm) ( $\epsilon/10^4 \text{ M}^{-1} \text{ cm}^{-1}$ )		$\lambda_{\text{em}}$ (nm)		$\Phi_{\text{F}}^a$	
	1	2	1	2	1	2
Cyclohexane	292 (1.29); 312 <i>sh</i>	283 <i>sh</i> ; 314 (0.77)	363	366	0.37	0.42
Dioxane	300 (1.67)	282 <i>sh</i> ; 312 (1.17)	428	436	0.35	0.30
Ethyl acetate <sup>b</sup>	293 <i>sh</i> ; 302 (1.01)	284 (1.30); 315 (1.18)	431	450	0.60	0.59
Dichloromethane	293 <i>sh</i> ; 310 (1.54)	283 <i>sh</i> ; 312 (1.15)	444	461	0.53	0.39
<i>N,N</i> -Dimethylformamide	312 (2.09)	313 (0.78)	457	476	0.43	0.40
Acetonitrile	294 <i>sh</i> ; 312 (2.03)	283 <i>sh</i> ; 318 (1.01)	459	485	0.30	0.18
Dimethylsulfoxide <sup>b</sup>	311 (1.96)	313 (1.39)	461	486	0.48	0.36
Ethanol	313 (1.77)	287 <i>sh</i> ; 313 (1.06)	—	—	—	—

<sup>a</sup> Relative to anthracene in ethanol ( $\Phi_{\text{F}} = 0.27$ ).<sup>41</sup> Error about 10%. <sup>b</sup> Solvents cut-off: ethyl acetate: 265 nm; dimethylsulfoxide: 270 nm; *N,N*-dimethylformamide: 275 nm. *sh*: shoulder.



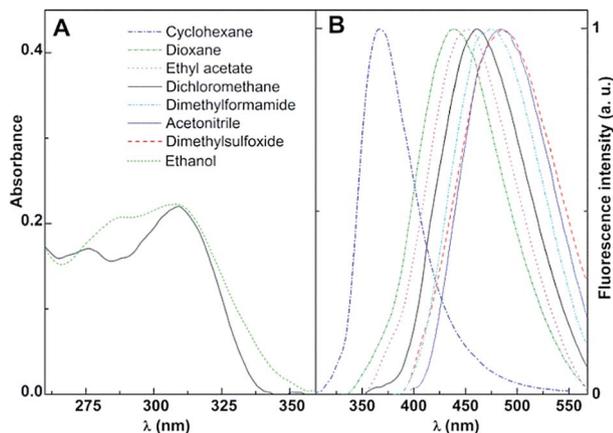


Fig. 4 (A) Absorption spectra of  $2 \times 10^{-5}$  M solutions of compound 2 in dichloromethane and ethanol, as examples. (B) Normalized fluorescence spectra ( $\lambda_{\text{exc}} = 310$  nm) of  $3 \times 10^{-6}$  M solutions of compound 2 in several solvents.

band and the presence of a pronounced rhodamine emission is observed in SMLs containing both fluorophores, evidencing the energy transfer from the excited NBD to rhodamine B.

A FRET efficiency of 85% was calculated from eqn (3) to (6), with a corresponding donor–acceptor distance ( $r$ ) of 5.3 nm. Considering that a cell membrane presents a thickness of 7 to 9 nm,<sup>50</sup> it can be concluded that the labeled lipids rhodamine-DOPE and NBD- $C_6$ -HPC are located in the first and second lipid layer, respectively, around the nanoparticle clusters, confirming the synthesis of the solid magnetoliposomes.

Analogous results were obtained for SMLs of DPPC : cholesterol 2 : 1. These results are also similar to the ones already reported for SMLs of the phospholipid DOPG containing manganese ferrite nanoparticles,<sup>31</sup> indicating that the method previously developed by us for the preparation of DOPG solid magnetoliposomes<sup>33</sup> can also be employed for different lipids. Size distributions of  $124 \pm 23$  nm (by number) and of  $141 \pm 38$  nm (by intensity) were obtained through Dynamic Light Scattering measurements for SMLs

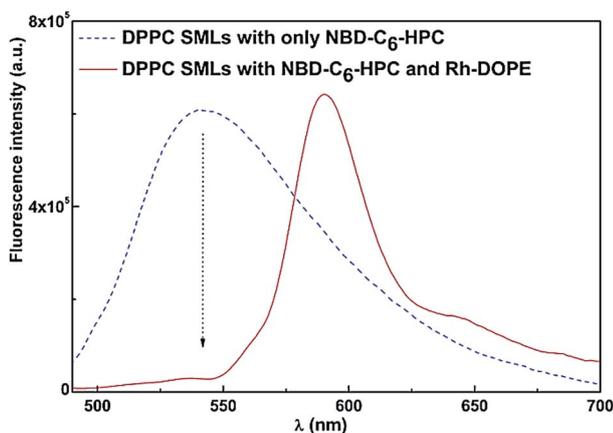


Fig. 5 Fluorescence spectra ( $\lambda_{\text{exc}} = 470$  nm, no rhodamine excitation) of SMLs with DPPC bilayer labeled with only NBD- $C_6$ -HPC and SMLs labeled with both NBD- $C_6$ -HPC and rhodamine B-DOPE.

of phospholipid DPPC (Fig. S1 in ESI<sup>†</sup>), showing an appropriate size for biomedical applications. SEM images of these SMLs also evidence structures with diameters lower than 150 nm (Fig. S2 in ESI<sup>†</sup>), in accordance with DLS data.

### Magnetic properties of magnetoliposomes

Previously synthesized manganese ferrite nanoparticles have shown to be superparamagnetic at room temperature.<sup>31</sup> Here, the magnetic hysteresis cycles of aqueous magnetoliposomes and solid magnetoliposomes were measured at 316 K and are displayed in Fig. 6. The hysteresis loop of neat  $\text{MnFe}_2\text{O}_4$  NPs is also shown, for comparison.

In terms of hysteresis, both nanosystems present a similar behaviour to the one of neat NPs, with small coercive fields of 4.22 Oe and 8.43 Oe for SMLs and AMLs, respectively (Fig. 6). Table 2 summarizes the magnetic data obtained from the loops. The presence of superparamagnetic behaviour can be evidenced by determining the magnetic squareness value, which is the ratio between the remnant magnetization,  $M_r$ , and the saturation magnetization,  $M_s$ . Magnetic squareness values indicative of superparamagnetism are of the order or below 0.1, meaning the loss of more than 90% of the magnetization upon removal of the applied magnetic field.<sup>51,52</sup> The magnetic squareness values for AMLs and SMLs clearly indicate the presence of a superparamagnetic behaviour (Table 2).

As previously reported for magnetoliposomes based on iron oxide nanoparticles,<sup>53,54</sup> the saturation magnetization of the manganese ferrite nanoparticles decreases after encapsulation into liposomes, this decrease being much more pronounced in the case of AMLs. For SMLs, the saturation magnetization is close to the one of neat  $\text{MnFe}_2\text{O}_4$  nanoparticles, while for AMLs a strong decrease is observed (Table 2). This is due to the lower molar ratio between magnetic NPs and lipids for AMLs, when compared to SMLs and bare NPs samples, which contributes to an overall lower saturation magnetization of the combined lipid + nanoparticles system. On the other hand, on AMLs and SMLs,

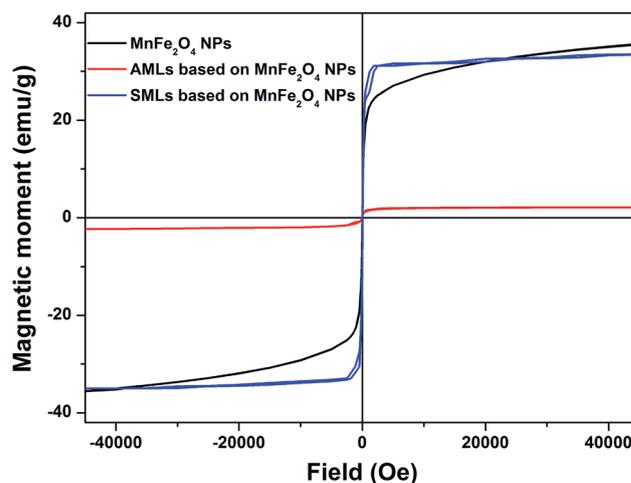


Fig. 6 Magnetization hysteresis cycles, measured at 316 K, of neat  $\text{MnFe}_2\text{O}_4$  nanoparticles, DPPC SMLs and Egg-PC AMLs containing  $\text{MnFe}_2\text{O}_4$  NPs.



**Table 2** Coercive field ( $H_c$ ), saturation magnetization ( $M_s$ ), remnant magnetization ( $M_r$ ) and ratio  $M_r/M_s$  for manganese ferrite NPs and magnetoliposomes

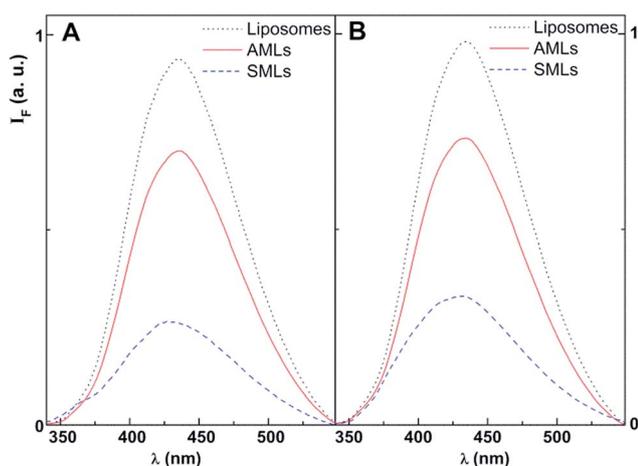
System	$H_c$ (Oe)	$M_s$ (emu g <sup>-1</sup> )	$M_r$ (emu g <sup>-1</sup> )	$M_r/M_s$
MnFe <sub>2</sub> O <sub>4</sub> NPs <sup>29</sup>	6.30	36.00	0.58	0.016
SMLs (DPPC)	4.22	34.16	0.90	0.03
AMLs (Egg-PC)	8.43	1.17	0.08	0.07

a more clear saturation of the hysteresis loops is observed as compared to the bare NPs (Fig. 6). This indicates an increased degree of clustering of the nanoparticles encapsulated in liposomes, which enhances the interparticle magnetic interaction and favors the alignment of their magnetic moments, as is more clearly observed at higher applied magnetic fields.

### Fluorescence studies in (magneto)liposomes

The thienopyridine derivatives **1** and **2** were encapsulated in magnetoliposomes, both aqueous (AMLs) and solid (SMLs). For comparison, studies in liposomes (without magnetic nanoparticles) of the same lipid compositions were also performed.

Examples of fluorescence emission spectra of compounds **1** and **2** in magnetoliposomes are shown in Fig. 7 (maximum emission wavelengths are presented in Table 3). Emission in liposomes (without magnetic nanoparticles and with the same concentration of compound) is also shown for comparison. A quenching effect of compounds emission by the magnetic nanoparticles is observed, proving the incorporation of these potential drugs in the magnetic nanocarriers. As expected, the fluorescence quenching is significantly more pronounced in solid magnetoliposomes, where the magnetic nanoparticles are closer to the antitumor compounds. This behaviour is similar to the previously observed with another antitumor thienopyridine derivative in AMLs and SMLs containing MnFe<sub>2</sub>O<sub>4</sub> NPs.<sup>31</sup>



**Fig. 7** Normalized fluorescence spectra of compounds **1** and **2** ( $3 \times 10^{-6}$  M) in liposomes and magnetoliposomes (Egg-PC AMLs and DPPC SMLs), at room temperature ( $\lambda_{\text{exc}} = 310$  nm). (A) Compound **1**; (B) compound **2**.

Relevant information about the location of these antitumor compounds in (magneto)liposomes can be obtained through fluorescence anisotropy ( $r$ ) measurements, as the  $r$  value increases with the rotational correlational time of the fluorescent molecule (and, thus, with the viscosity of the environment, eqn (8))<sup>37</sup>

$$\frac{1}{r} = \frac{1}{r_0} \left( 1 + \frac{\tau}{\tau_c} \right) \quad (8)$$

where  $r_0$  is the fundamental anisotropy,  $\tau$  is the excited-state lifetime and  $\tau_c$  is the rotational correlation time.

The fluorescence (steady-state) anisotropies of these diarylamines in the several systems are displayed in Table 3. For comparison, the fluorescence anisotropy values in the highly viscous solvent glycerol ( $\eta = 993.4$  cP at 25 °C)<sup>55</sup> at room temperature were also measured (Table 3). Upon temperature increase, the excited-state lifetime of these molecules is predicted to decrease, due to the increment of the non-radiative deactivation pathways (mainly the rate constant for internal conversion  $S_1 \rightarrow S_0$ ). According to eqn (8), an increase of the fluorescence anisotropy is expected from a diminution of the excited-state lifetime. The anisotropy values for **1** and **2** in DPPC-based systems decrease significantly at 55 °C (Table 3) and this behaviour can be attributed to a decrease in the rotational correlation time of the fluorophore. This originates from the decrease of membrane microviscosity upon changing from the gel to the liquid-crystalline phase of DPPC above 41 °C (melting transition temperature).

The differences in anisotropy values between the compounds incorporated in lipid systems and in glycerol at room temperature can also be attributed to the distinct environment viscosities, as lipid membranes, exhibiting viscosity values around 100–200 cP,<sup>56,57</sup> are much less viscous than glycerol. These results indicate that both compounds are mainly located in the lipid bilayers.

The effect of cholesterol (Ch) in Egg-PC membranes, as reported by fluorescence anisotropy values, is different for compounds **1** and **2** (a slight increase in fluidity reported by compound **2** and a diminution for compound **1**). However, this effect is small and may be related with slightly different locations of these compounds in membranes. From the values of maximum emission wavelengths in AMLs and SMLs, an environment with polarity similar to dioxane or ethyl acetate is anticipated (Tables 1 and 3).

It was reported that membrane viscosity decreases from the outside to inside the membrane<sup>58,59</sup> and this may justify, besides possible differences in compound excited-state lifetimes, the lower anisotropies for both compounds in solid magnetoliposomes, when compared to AMLs. Maximum emission wavelength values point to less polar environments felt by compounds in SMLs (Table 3), corroborating that these potential drugs are deeply located in SMLs membranes.

The encapsulation efficiencies of both compounds in AMLs and SMLs are presented in Table 4. Values of encapsulation efficiency are larger than 75%, the lowest EE% being observed for compound **2** in aqueous magnetoliposomes.



**Table 3** Steady-state fluorescence anisotropy ( $r$ ) values and maximum emission wavelengths ( $\lambda_{em}$ ) for diarylamines **1** and **2** in several liposome and magnetoliposome formulations, at 25 °C and 55 °C

	Lipid formulation	$T$ (°C)	$\lambda_{em}$ (nm)		$r$	
			<b>1</b>	<b>2</b>	<b>1</b>	<b>2</b>
Liposomes (without NPs)	Egg-PC (100%)	25	432	433	0.195	0.222
	Egg-PC : Ch (7 : 3)	25	433	434	0.212	0.202
	DPPC (100%)	25	432	433	0.200	0.222
AMLs		55	430	431	0.141	0.101
		25	432	433	0.191	0.230
		25	433	435	0.220	0.201
		25	432	433	0.202	0.287
SMLs	DPPC (100%)	55	432	432	0.153	0.140
		25	430	427	0.159	0.135
		55	428	424	0.101	0.051
Glycerol	—	25	—	—	0.304	0.321

These high encapsulation efficiencies point to a promising use of these nanocarriers in magnetically-guided transport of compounds **1** and **2**, and as agents for simultaneous chemotherapy and hyperthermia in oncological therapy. These potentialities will allow reducing the blood circulating time of the drugs and lowering side effects, which is especially important as both compounds are hydrophobic and present some cytotoxicity for normal PLP2 cells, with low  $GI_{50}$  values of  $1.94 \pm 0.16 \mu\text{M}$  for compound **1** and  $6.56 \pm 0.2 \mu\text{M}$  for compound **2**.<sup>29</sup>

#### Growth inhibitory activity of drug-loaded AMLs on human tumor cell lines and on porcine liver primary cells

The tumor cell growth inhibitory activity of AMLs containing compounds **1** and **2** was evaluated in four human tumor cell lines, specifically MCF-7 (breast adenocarcinoma), HeLa (cervical carcinoma), T3M4 (pancreatic cancer) and NCI-H460 (non-small cell lung cancer). Moreover, the toxicity of the drug-loaded AMLs towards non-tumor cells was studied using porcine liver primary cells (PLP2). This allowed the determination of the  $GI_{50}$  values ( $\mu\text{M}$ ), corresponding to the compound concentration which inhibited 50% of cell growth (Table 5). Besides the neat Egg-PC AMLs, the influence of 5% PEG-folate in AMLs formulation was also investigated, as folate receptors are generally overexpressed in tumor cells.

Both aqueous and solid magnetoliposomes (without compounds) evidenced no inhibitory activity on the same cell

lines, including the non-tumor PLP2, when tested in the same conditions used with the drug-loaded systems. From the results obtained (Table 5), it can be concluded that the drug-loaded AMLs are efficient in inhibiting tumor cells, presenting  $GI_{50}$  values similar to the ones determined previously using only the compounds,<sup>29</sup> except for compound **2** in these formulations, that presents very much low  $GI_{50}$  values for the NCI-H460 lung cancer cell line. The presence of folate has only a significant effect for compound **1** in HeLa tumor cell line. For compound **2**, the presence of folate in magnetoliposomes does not influence noticeably the results (Table 5). Nevertheless, the  $GI_{50}$  values clearly show that aqueous magnetoliposomes are able to release the encapsulated compounds **1** and **2** and to inhibit the growth of tumor cells. It is also noteworthy the very low  $GI_{50}$  values obtained with compounds **1** and **2** using the pancreatic tumor cell line T3M4, where the single compounds had not been previously tested.

In general, the results in PLP2 cells (Table 5) show that the encapsulated compounds are not toxic for the  $GI_{50}$  values in all tumor cell lines, apart from the compound **2**-loaded systems in MCF-7 cells, that present  $GI_{50}$  values higher than for PLP2.

The low  $GI_{50}$  values obtained for non-tumor PLP2 cells reinforce the importance of using a magnetic field gradient to guide and target these nanosystems for tumors *in vivo*, taking also advantage of hyperthermia capability.

Unfortunately, for the drug-loaded DPPC solid magnetoliposomes (SMLs), no satisfactory results were obtained, with very high  $GI_{50}$  values (above  $7.5 \mu\text{M}$ ) in all tumor cell lines (Table S1 in ESI†). This is possibly due to the fact that the phospholipid DPPC is in the rigid gel phase at room temperature, thus preventing the release of the compounds. The addition of 5% DSPE-PEG-Folate did not improve the results (Table S1†). Other lipid formulations for SMLs will be tested in the near future, as well as assays at mild hyperthermia temperatures will be carried out. Also, confocal microscopy measurements will be performed to assess internalization of magnetoliposomes in cells.

**Table 4** Encapsulation efficiencies (EE%) of compounds **1** and **2** in magnetoliposomes (SD: standard deviation)

System	EE% $\pm$ SD	
	Compound <b>1</b>	Compound <b>2</b>
AMLs (Egg-PC)	$89.0 \pm 4.5$	$75.9 \pm 6.8$
SMLs (DPPC)	$88.1 \pm 4.9$	$82.1 \pm 6.1$



**Table 5** Growth inhibitory activity of drug-loaded aqueous magnetoliposomes on various human tumor cell lines and non-tumor porcine liver primary cells (PLP2)

Cell line	GI <sub>50</sub> values <sup>a</sup> (μM) for compound 1			GI <sub>50</sub> values <sup>a</sup> (μM) for compound 2		
	Only compound 1 (ref. 29)	Egg-PC AMLs with compound 1	Egg-PC/PEG-Fol (95 : 5) AMLs with compound 1	Only compound 2 (ref. 29)	Egg-PC AMLs with compound 2	Egg-PC/PEG-Fol (95 : 5) AMLs with compound 2
HeLa	0.09 ± 0.00	0.15 ± 0.02	0.09 ± 0.01	1.40 ± 0.06	1.32 ± 0.15	1.44 ± 0.07
MCF7	0.25 ± 0.03	0.45 ± 0.01	0.44 ± 0.0003	5.88 ± 0.86	5.67 ± 0.62	5.56 ± 0.11
T3M4	—	0.26 ± 0.02	0.30 ± 0.002	—	1.56 ± 0.05	1.68 ± 0.07
NCI-H460	0.28 ± 0.03	0.37 ± 0.03	0.35 ± 0.07	5.40 ± 0.89	1.66 ± 0.22	1.79 ± 0.12
PLP2	1.94 ± 0.16	1.46 ± 0.03	1.37 ± 0.14	6.56 ± 0.2	3.48 ± 0.25	3.46 ± 0.13

<sup>a</sup> GI<sub>50</sub> values correspond to the concentration which inhibited 50% of cell growth. Results are from three independent experiments (performed in triplicate), and are expressed as mean ± standard deviation (SD).

## Conclusions

In this work, the fluorescence emission of two new promising antitumor compounds, a *N*-(3-methoxyphenyl)thieno[3,2-*b*]pyridin-7-amine **1** and a *N*-(2-methoxyphenyl)thieno[3,2-*b*]pyridin-7-amine **2**, was studied in different environments. These *in vitro* potent antitumor thienopyridin-7-arylamine derivatives were successfully incorporated in both aqueous and solid magnetoliposomes based on superparamagnetic manganese ferrite nanoparticles, with high encapsulation efficiencies. The intrinsic fluorescence of the potential antitumor drugs was used as a tool to obtain information about compounds location in magnetoliposomes.

The magnetic properties of the manganese ferrite-based magnetoliposomes were measured for the first time, evidencing a superparamagnetic behaviour.

Growth inhibitory concentrations on tumor cell lines obtained with aqueous magnetoliposomes with encapsulated compounds **1** and **2** point to promising future drug delivery applications of these anticancer compounds *in vivo*, using magnetoliposomes simultaneously as drug nanocarriers and hyperthermia agents, to increase the efficacy of anticancer therapies.

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