Pegylation of phenothiazine – A synthetic route towards potent anticancer drugs

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Highlights
- Antitumor activity of two PEGylated phenothiazines was investigated
- The compounds showed cytotoxic activity against six tumor lines
- They inhibited the tumor growth in experimental mice
- The PEGylation improved the phenothiazine biocompatibility
- A synergistic effect of PEG and phenothiazine toward properties improvement was proved

Abstract

Introduction: Cancer is a big challenge of the 21st century, whose defeat requires efficient antitumor drugs.

Objectives: The paper aims to investigate the synergistic effect of two structural building blocks, phenothiazine and poly(ethylene glycol), toward efficient antitumor drugs.

Methods: Two PEGylated phenothiazine derivatives were synthesized by attaching poly(ethylene glycol) of 550 Da to the nitrogen atom of phenothiazine by ether or ester linkage. Their antitumor activity has been investigated on five human tumor lines and a mouse tumor line as well, by determination of IC50. The in vivo toxicity was determined by measuring the LD50 in BALB/c mice by the sequential method and the in vivo antitumor potential was measured by the tumors growth test. The antitumor mechanism was investigated by complexation studies of zinc and magnesium ions characteristic to the farnesyltransferase enzyme, by studies of self-aggregation in the cells proximity and by investigation of the antitumor properties of the acid species resulted by enzymatic cleavage of the PEGylated derivatives.

Results: The two compounds showed antitumor activity, with IC50 against mouse colon carcinoma cell line comparable with that of the traditional antitumor drugs 5-Fluorouracil and doxorubicin. The phenothiazine PEGylation resulted in a significant toxicity diminishing, the LD50 in BALB/c mice increasing from 952.38 up to 1450 mg/kg, in phenothiazine equivalents. Both compounds inflicted a 92% inhibition of tumor growth in mice. The antitumor mechanism was investigated by complexation studies of zinc and magnesium ions characteristic to the farnesyltransferase enzyme.
of the tumour growth for doses much smaller than LD50. The investigation of the possible tumour inhibition mechanism suggested the nanoaggregate formation and the cleavage of ester bonds as key factors for the inhibition of cancer cell proliferation and biocompatibility improvement.

**Conclusion:** Phenothiazine and PEG building blocks have a synergetic effect working for both tumour growth inhibition and biocompatibility improvement. All these findings recommend the PEGylated phenothiazine derivatives as a valuable workbench for a next generation of antitumor drugs.

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**Introduction**

Cancer, an affliction known to humanity for thousands of years, became the second leading cause of death with a continuous growing of incidence and mortality rate, predicted to grow by as much 70% in the next 20 years [1]. Occurring in a high variety of different types, it has been recognized that its increasing incidence is close inter-connected with the technological development, becoming thus the big challenge of the 21 century [2]. To defeat cancer, many possible therapies were considered, from traditional medicine [3] to DNA transfection [4]. Nevertheless, chemotherapy demonstrated the most efficient results, despite the high frequency and severity of adverse effects and low success rate caused by the reduced selectivity of the clinical chemodrugs. In this view, the discoveries of new more effective antitumor drugs with lesser side effects remained a desideratum of the contemporary society.

Looking to the chemical structure of the antitumor drugs, many moieties promoted antitumor activity. Among them, phenothiazine heterocycle is a building block with versatile biologic activity, which confirmed cytotoxicity against some cancer lines [5]. It was postulated that its cytotoxicity can be selectively improved by structural modifications, mainly via substitution at the nitrogen atom [6,7]. Considering the biologic versatility of phenothiazine (such as analgesic, antipsychotic, immunosuppressive, anti-inflammatory, bactericide, fungicide, antimarial, antifilarial, trypanocidal, anticonvulsant), it can be envisaged that it may represents a valuable building block for the development of multifunctional antitumor drugs [8]. On the other hand, polyethylene glycol, a synthetic polymer approved by FDA for indwelling bioapplications, proved significant improvement of the pharmaceutical value of antitumor drugs [9]. This was possible due to its ability to enhance the retention time by protection against various degradation mechanisms which are active inside the tissues or cells. Besides, PEG has no interaction with the blood components, because very low plasma protein affinity [10].

Starting from these premises, our group designed PEGylated phenothiazine derivatives as new building blocks for anticancer drugs. PEG has been bonded to the phenothiazine heterocycle at the nitrogen atom and the cytotoxicity of the resulted compounds was investigated on normal cells and six tumour lines. It was demonstrated that the simple linking of PEG to phenothiazine lead to the enhancement of the cytotoxicity against the tumour cells compared to normal cells. Moreover, the in vivo investigation on experimental tumours in mice demonstrated that the studied PEGylated phenothiazines inhibited the tumour growth, promising to be a valuable workbench towards a new generation of more friendly multifunctional antitumor drugs.

**Experimental**

**Materials**

Phenothiazine 98%, ethyl bromoacetate 98%, sodium hydride 95%, methoxy poly(ethylene glycol) (550 Da, polymerization degree: 11–13), p-toluenesulfonyl chloride 98%, pyridine 99.8%, triethylamine (TEA) 99%, N-hydroxysuccinimide (NHS) 98%, N-(3-dimethylaminopropyl)-N′-ethylcarbodiimide hydrochloride (EDC, HCl) 98%, sodium hydroxide 97%, magnesium acetate tetrahydrate 99% and zinc acetate dehydrate 98% were purchased from Aldrich (Verviers, Belgium), fetal bovine serum (non-USA origins) from Sigma Aldrich (Schnelldorf, Germany), TrypLE™ Express Enzyme and StemProAccutase from Gibco (Langley, Virginia, USA), LIVE/DEAD Viability/Cytotoxicity Kit and phosphate buffered saline (PBS) from Invitrogen (Eugene, Oregon, USA), CellTiter 96® Aqueous One Solution Cell Proliferation Assay (MTS) from Promega (Madison, Wisconsin, USA), CytoOne® well plates from StarLab (Hamburg, Germany).

The compounds under study were synthesized using protocols from our previous study [11]. Briefly, the synthesis of 10-(methoxy poly(ethylene glycol))-10H-phenothiazine (PP) was realized by N-alkylation of phenothiazine with 2-methoxy poly(ethylene glycol) 4-methylbenzenesulfonate in DMF, using sodium hydride for phenothiazine deprotonation [11]. The product was purified by column chromatography (DCM:methanol 10:1, v/v), giving a deep red viscous liquid. FT-IR (KBr, cm⁻¹): 3060 (vCH aromatic), 2870 (vCH aliphatic), 1593, 1570 (vC = C), 1460 (δCH₂), 1292 (vC-N), 1110 (vO-C-O), 755 (δH-H). **¹H NMR (400 MHz, DMSO-d₆, ppm)** δ = 7.22–7.13 (t,d, 4H, H₁, H₂, H₈, H₉), 7.06–7.04 (d, 2H, H₄, H₆), 6.97–6.93 (t, 2H, H₃, H₇), 4.07–4.04 (t, 2H, H₁₁), 3.76–3.73 (t, 2H, H₁₂), 3.50–3.42 (m, 48H, H₁₃, H₁₄), 3.24 (t, 3H, H₁₅).

The synthesis of methoxy poly(ethylene glycol) 2-(10H-phenothiazin-10-yl) acetate (PPO) was realized by an esterification reaction of the 2-(10H-phenothiazin-10-yl) acetic acid with methoxy poly(ethylene glycol) in DCM, in the presence of DMAP and DCC [11]. The product was purified by column chromatography (DCM:methanol, 10:1, v/v), when an orange viscous liquid was obtained. FT-IR (KBr, cm⁻¹): 3097–3063 (vCH aromatic), 2904 (vCH₃), 2875 (vCH₂), 1742 (vC = O), 1592–1560 (vC = Cαr), 1467 (δCH₂) 1193 (vO-C-O), 1109 (vO-C-O), **¹H NMR (400 MHz, CDCl₃, ppm)** δ = 7.10–7.07 (m, 4H, H₁, H₂, H₈, H₉), 6.92–6.89 (t, 2H, H₃, H₇), 6.62–6.60 (d, 2H, H₄, H₆), 4.55 (s, 2H, H₁₁), 4.42–4.40 (t, 2H, H₁₃), 3.74–3.71 (t, 2H, H₁₄), 3.65–3.60 (m, 30H, H₁₅), 3.37 (s, 3H, H₁₆).
Equipment and methods

NMR spectra were obtained on a Bruker Avance DRX 400 MHz spectrometer equipped with a 5 mm QNP direct detection probe and z-gradients, at room temperature, with an accumulation of 64 scans. The chemical shifts were reported as δ values (ppm) relative to the residual peak of the solvent.

Infrared spectra were recorded on a FTIR Bruker Vertex 70 Spectrometer, at room temperature, using KBr pellets.

The absorbance for CellTiter 96® Aqueous One Solution Cell Proliferation Assay (MTS) was measured using FLUOstar Omega Filter-based multi-mode microplate reader from BMG LABTECH (Offenburg, Germany).

Images for live/dead staining were acquired with a Leica DMI 3000B inverted microscope (Wetzlar, Germany). Live cells images, colored in green, were obtained using a GFP filter and dead cells images, coloured in red, using Texas Red (TX2) filter.

Cell culture: Cells were cultivated in complete MEM containing 1% Penicillin-Streptomycin-Amphotericin B mixture and 10% fetal bovine serum under 5% CO2 humidified atmosphere at 37 °C. 

TrypLE® Express Enzyme was used for passaging NHDF, HOS, NHDF cell line in 100 cm2, approximately 103 cells/well for MeWo, HOS, HeLa and HepG2 cell line, 100 mg/kg in PBS to obtain 0.4 mL solution containing 2 mg active substance, meaning 100 mg active substance/kg bodyweight (Table 1). For LD50 testing, PP and PPO derivatives were solubilized in distilled water to obtain 100 mg/mL stock solutions.

Experimental design for the in vivo antitumor effect testing: BALB/c mice were used to assess the antitumor effect and to determine median lethal dose LDL 50. The animals were purchased from the Cantacuzino Institute in Bucharest, 10-week-old nulliparous females, with an average weight of 20 ± 0.45 g. The acclimatization of the mice was done under identical conditions of temperature (22 ± 0.7 °C) and humidity (50 ± 10%), and the light/dark cycle provided was 12 h. Each experimental group was housed in autoclavable polycarbonate cages, 1500 cm2, approximately 300 cm2/mouse. The animals had permanent access to water (ad libitum) (autoclavable bottles with drip system) and standardized food (provided by Cantacuzino Institute) with the following composition: 23% protein, 10% fat, 50% carbohydrates, 8% crude fiber and 9% vitamin-mineral premix, calcium carbonate and phosphate, amino acids. The animals were kept 7 days in the laboratory for accommodation and they were monitored daily to record any disease conditions and abnormal behaviour. Those that did not meet the required health criteria were removed from the experiment.

For the determination of Lethal Dose 50 (LD50) the sequential method has been used (UP and DOWN PROCEDURE-OEC 425). It was preferred because the number of animals is minimal and the estimated toxicity range is smaller. The method consists in the administration of a dose (d) to a single mouse. The dose was calculated in mg/kg and diluted in PBS to an intraperitoneal injection volume of 0.4 mL. If the mouse survived for 24–48 h (the monitoring interval was determined by the onset, duration and severity of toxic signs), another dose 1.3 times higher (d × 1.3) was administered to another mouse. If it also survived, it was continued with d × 1.3 × 1.3 ... , until the experimental animal died. If the first mouse dies, the test continued by decreasing the dose by 1.3 times (d/1.3), until reaching the dose at which the mouse lived. Testing stopped when 3 out of 5 mice survived consecutively at the upper limit. Any changes in behaviour and clinical manifestations were recorded. It was considered less likely to be an effect of treatment if: there was no obvious dose response, the measurement of the

| Table 1  | Protocol for administration of PTZ, PP and PPO compounds to BALB/C mice. |
|----------|-------------------------------------------------------------------|
| Group    | Range of administration | No. of administrations | Dose: mg active substance i.e. PTZ / kg bodyweight |
| PBS      | Once daily              | 10                      | 100 mg/kg; 2 mg/mouse                           |
| PTZ      |                       |                         | 100 mg/kg; 2 mg/mouse                           |
| PP       |                       |                         | 100 mg/kg; 2 mg/mouse                           |
| PPO      |                       |                         | 100 mg/kg; 2 mg/mouse                           |
endpoint being evaluated was inherently inaccurate, it was in a normal biological variation (e.g. within reference values), there was a lack of biological plausibility.

For antitumor tests, the animals were divided into 4 groups (5 mice/group), as follows: Group PBS: negative control group injected with phosphate buffer solution (PBS); Group PTZ: positive control group injected with phenothiazine (PTZ); Group PP: injected with PP derivative solution; Group PPO: injected with PPO derivative solution. All mice were acclimatized at least one week before the start of the experiment. CT26 colorectal cancer cells were transplanted on a growth medium [i.e., Dulbecco’s Modified Eagle’s Medium (DMEM)] containing 10% Fetal Bovine Serum (FBS), 1 mM sodium pyruvate, 1% glutamine, and 1% streptomycin for multiplication. Tumour cells were cultured at 37 °C and 5% CO₂. The inoculum was administered subcutaneously in the thoracic-dorsal (intrascapular) region of the mice anesthetized with 3% isoflurane, in a volume of 0.25 mL containing a number of 500 000 tumour cells. Prior to inoculation, the administration site was shaved and disinfected with povidone iodine. Then, the mice were monitored daily, to assess the degree of tumour development, their general condition, behaviour, respiration, recording the clinical score [15,16], the toxic effects of long-term administration such as cyanosis, anorexia, and jaundice or any other sign of pain. All mice developed subcutaneous tumours. The tumour became palpable on day 5 after injection of CT26 cells. The in vivo antitumor test started in the day 7 after inoculation, when the tumour volume was around 120 mm³.

The administration protocol was given in Table 1. All mice were injected intraperitoneally [IP] using a 27G insulin syringe, one administration daily, for 10 days. The dose was calculated considering the phenothiazine unit as active substance, 100 mg/kg body-weight, meaning approximately 2 mg /mouse, respectively, e.g. 2 mg PTZ/mouse; 7.33 mg PP/mouse; 7.83 mg PPO/mouse. The compounds were diluted in PBS so that the final volume of the solution injected intraperitoneally was 0.4 mL/mouse [17,18]. Tumour measurements were performed daily before administration to establish a basis for comparison. The tumour volume was determined by measuring 3 diameters and subsequently calculated using the formula for a hemielipsoid: Volume = 0.52 × W (width-diameter perpendicular to length) × H (height) [19].

In order to appreciate the efficiency of studied compounds compared to other findings reported in literature, the percent of tumour growth inhibition (Tik%) was calculated with the equation: Tik% = (1 - D(d) / D(c)) × 100, where D(d) – tumour dimension developed in the treated mice and D(c) - tumour dimension developed in the control mice.

Statistical analyses were performed using GraphPad Prism software version 7.00 for Windows (GraphPad Software, San Diego, California). The obtained results represent the mean ± standard deviation (S.D.) of three different experiments, and the differences between groups were done with 2 way ANOVA Tabular results and multiple comparisons. P-values < 0.05 were considered as significant (*p < 0.05, **p < 0.001, ***p < 0.0001), p > 0.05 the differences between data are not significant.

Ethical implications

The study was conducted in accordance with national and international regulations on animal welfare, identification, control and elimination of factors causing physiological and behavioural disorders: Directive EC86 / 609 EU; Government Ordinance no. 37/2002, approved by Law no. 471/2002; Law 205/2004 on animal protection, amended and supplemented by Law no. 9/2008; Joint Order of ANSVSA and of the Ministry of Interior and Administrative Reform no. 523/2008 for the approval of the Methodological Norms for the application of Law 205/2004 on animal protection.

Results and discussions

The paper focuses on the investigation of the antitumor activity of two PEGylated phenothiazine derivatives, for which PEG has been bonded to the phenothiazine by ether or ester unit (Scheme 1). It should be stressed that, following well established synthetic protocols, the compounds were obtained with high purity (higher than 99%). They presented excellent solubility in water and common organic solvents (Figure S1), and were hydrolytically stable on an investigation period of 7 days. However, a deeper investigation of their water solutions by DLS and UV–vis showed that they self-assembled into nanoaggregates, at the critical aggregation concentration of 8.7×10⁻⁴ and 9.23×10⁻⁴ mM for PP and PPO respectively (Figure S2, Figure S3) [11]. The hydrodynamic diameter of the nanoaggregates was dependent on the solution concentration, decreasing from 289 to 106 nm for PP, and from 322 to 63 nm for PPO when the concentration decreased from 10 to 0.1 mM, rationally explained by the variation of the molecules density which influenced the aggregate size, i.e. a higher molecules density favoured a higher aggregate size (Figure S2) [20].

In vitro investigation of the antitumor activity

The investigation of the antitumor activity of the PP and PPO derivatives was performed by determining the cytotoxicity on five human cancer cell lines: cervical carcinoma (HeLa); malignant melanoma (MeWo); osteosarcoma (HOS), breast cancer (MCF7) and liver cancer (HepG2) versus a normal cell line (NHDF). Having in mind the in vivo investigation of the antitumor activity on mice, mouse colon carcinoma cell line (CT26) was also included in the study. The cytotoxicity tests were done for solutions with concentration from 1 μM up to 1 mM, and the IC₅₀ was determined for each cell line. The dose-responsive curves were represented in Fig. 1a-l, as graphs of the relative cell viability against the concentration of PP and PPO.

At first glance, it can be observed that among the human tumor cell lines, for the concentration of 64 μM at which the NHDF viability was around 80%, the PP showed the higher cytotoxic effect against MeWo (cell viability 47%), and the PPO against HeLa (cell viability 40%). Compared to the human cell lines, the viability of CT29 tumor cells decreased to<40% for a similar concentration of both PP and PPO compounds.

These observations were reflected in a more accurate way in the values of IC₅₀ parameter, calculated for both compounds on every cell line (Fig. 1m, Table S1). It can be seen that PP and PPO exhibited high cytotoxic activity against CT26 cells, and IC₅₀ values were 6 and 12 fold lower compared to that against NHDF cells indicating good selectivity. Even thought that CT26 cells are specific for mice
tumours and not human ones, by comparing the results with other data reported for therapeutic antitumor drugs or other compounds investigated as potential anticancer drugs, the antitumor potency of PP and PPO compounds appears remarkable (Table 2). As an example, it can be seen that IC50 of PP and PPO is comparable with that of traditional antitumor drugs 5-Fluorouracil and doxorubicin, therefore encouraging further in vivo tests. Moreover, it should be remarked that even though some papers reported significant higher IC50 values, they did not report IC50 for normal cells, making difficult to appreciate their selectivity.
Regarding the IC₅₀ values obtained on the human cell lines, PP have increased cytotoxic effect against two tumour lines, HeLa (cervical cancer) and MeWo (skin cancer), for which the IC₅₀ gave values of 229.1 μM and 251.9 μM, respectively, compared to the 305.8 μM value obtained for the normal cell line. By comparison, PPO showed higher cytotoxic effect against all the investigated tumour cell lines compared to the normal cells, but significant lower IC₅₀ values were recorded for HepG2 (human liver cancer) and MCF7 (breast cancer): 161.3 μM and 131.7 μM, respectively. These values were one third lower and one-half lower, respectively, compared to that for normal cells (286.2 μM). However, the selectivity index SI(IC₅₀) calculated using NHDF as control showed low values indicating low selectivity (Table S2) [34,35]. Compared to traditional therapeutic drugs, IC₅₀ values of PP and PPO were mostly higher and sometimes lower, encouraging further investigations (Table S3).

Fluorescent imaging confirmed the MTS assay results, as can be seen in Fig. 2, where live cells were stained an intense, uniform green, while dead cells had a predominantly nuclear red fluorescence [36]. Characteristics of apoptotic cells like cell shrinkage, rounding, partial detachment and lobulated appearance can be easily observed for HeLa, MCF7 and HepG2 cells treated with PP, and NHDF, MCF7 and HepG2 cells treated with PPO [37].

**In vivo toxicity**

In order to establish the doses for safe in vivo use of the PP and PPO, the in vivo toxicity of the compounds was investigated.Usu-
ally, the estimation of the in vivo toxicity of a new substance is preliminary done by determining the acute toxicity - LD50, a dose that causes death in 50% of the studied experimental animals. After testing several dose levels for each compound, as single intraperitoneal administration, according to the protocol described in the Experimental part, it was concluded that the LD50 in BALB/c mice was 952.38 mg/kg for phenothiazine (PTZ) and increased to 1450 mg/kg for PP and 1300 mg/kg for PPO (Table S4). It should be highlighted that LD50 for PP and PPO was expressed as equivalent in phenothiazine (see Experimental part). It was obvious that PEGylation reduced the toxicity of the phenothiazine with >50%.

Contrary to the in vitro findings, the in vivo toxicity of PPO was slightly higher compared to that of PP. The mice in both PP and PPO groups exhibited serious adverse reactions, CNS depression, generalized muscle contractions with
body torsion, and reduced cardiac, respiratory, and central activity, immediately after administration (1–2 min) at all tested levels. These reactions last for 10 min, and then the mice recovered and adopted a normal behaviour. About 20–30 min after this return, a state of suffering occurred especially in the PPO group that leads to the partial immobilization of the animal. To a lesser extent, the phenomena were also observed in animals from the PP group, but they were of lower intensity. In the case of the phenothiazine group, a state of suffering accompanied by depression was recorded only at the highest dosage levels. The adverse effects described above are characteristic to the phenothiazine derivatives used as neuroleptic drugs or investigated as anticancer drugs [7,38], and were ascribed to the ability of the positively charged nitrogen to induce a higher permeability of the blood–brain-barrier (BBB) [39]. On the other hand, the amphiphilic character of the PEGylated phenothiazine derivatives allows the diffusion into the endothelial cells and from there to the brain [40]. This can be the indicative of a better absorption of the PEGylated derivatives, especially of the PPO in which the PEG chain has been bonded to the phenothiazine core by an ester linking group.

### In vivo investigation of the antitumor activity

The next step in the investigation of the antitumor activity was to monitor the inhibitory effect of the phenothiazine derivatives on the tumour growth rate in mice as animal models. Starting from the idea

| Compound name | Structure | Ti (%)* | Reference |
|---------------|-----------|---------|-----------|
| PTZ | ![PTZ structure](image) | 47 | This study |
| PPO | ![PPO structure](image) | 92 | This study |
| PP | ![PP structure](image) | 92 | This study |
| Thioridazine | ![Thioridazine structure](image) | 51 | [42] |
| PTX-MB@PLGA | ![PTX-MB@PLGA structure](image) | 60 | [28] |
| Doxorubicine | ![Doxorubicine structure](image) | 27.27 | [43] |
| 5-Flourouracil | ![5-Flourouracil structure](image) | 75, 82.85, 85.71 | [44] |

*Ti = percent of tumor inhibition during 10 days of treatment calculated with the eq. $T_i = \frac{(1-D(d)/D(c)) \times 100}{100}$, where $D(d)$ - tumour volume developed in the treated mice and $D(c)$ - tumour volume developed in the control mice
that systemic cytotoxic chemotherapy is still the therapeutic basis for many types of cancer, the compounds solutions were intraperitoneal administered one dose per day during 10 days, at doses much lower that LD50 [34,41]. The evolution of the tumour growth in model animals treated with phenothiazine was considered as a positive control and those treated with PBS as negative control. The obtained data are presented in Fig. 3 and Table S5.

In the first day of administration, the average volume of the subcutaneous tumours was around 120 mm$^3$ in all experimental groups. At the end of experiment, the tumour volume recorded in the negative reference (PBS group) increased about 17 times to 2253.72 mm$^3$. In the group treated with phenothiazine (PTZ group) as positive reference, the tumour increased about 9 fold (1190.12 mm$^3$) while in the groups treated with PP and PPO respectively, the tumour increased<1.5 times (173.49 mm$^3$ in the PP group and 170.58 mm$^3$ in the PPO group). This means that the tumour inhibitory (Ti) effect of the PTZ compared to the negative control PBS of 47%, drastically increased to 92% by PEGylation in the case of the mice treated with PP and PPO. Interesting, even if the in vitro tests indicated PPO as more effective than PP in killing the CT26 cells, the in vivo tests showed statistical insignificant differences between them (p > 0.05), possible due to different concentration of the compounds used for measurements.

To estimate the efficiency of the phenothizine PEGylation on the tumour growth inhibition, the percent of tumour inhibition (Ti) was compared with those of the other phenothiazine derivatives reported for CT26 cell line (Table 3), or other tumour lines (Table S6) for a similar treatment period. For comparison reasons, the results reported for some therapeutic anticancer drugs were included in Table 3 too. It can be seen that the tumour inhibition of 92% recorded for the studied PEGylated phenothiazines is significantly higher compared to other Ti values, a maximum of almost 60% being reported for quite complicated phenothiazine structures [28]. Moreover, the CT26 tumour inhibition growth was significantly higher compared to 5-fluorouracil and doxorubicine drugs. Considering the simple structure of the studied PEGylated phenothiazines, the lack of toxicity of the PEG building block, and the possibilities to further functionalize the phenothiazine core with potent anticancer units, we can consider that phenothiazine PEGylation is a valuable pathway for designing anticancer drugs.

Cumulating the data of the in vivo tests, it can be concluded that the PEGylated phenothiazine derivatives, especially PP one, have potential of anticancer drugs, and their design can serve as a work bench for further chemical engineering in order to improve the antitumor activity and to minimize the side effects, towards a new class of anticancer drugs.

**Antitumor mechanism**

To further develop the design of the PEGylated phenothiazines, it is important to understand the structure particularities triggering the improvement of antitumor activity and on the other hand those promoting toxicity. As the principal building blocks of the studied compounds are the phenothiazine and PEG, their influence was considered. Moreover, the presence of the ester bond in PPO, which can be easily hydrolysed under the influence of esterases to give the acid precursor (Scheme 2) was examined as well [45,46]. It should be mentioned that the esterases are overexpressed in the tumour cells increasing the probability of ester bond cleavage in their proximity [47].

**Investigation of farnesyltransferase inhibition.** The investigation of the antitumor activity of various phenothiazine based derivatives demonstrated that the mechanism of their tumour growth inhibition involves the inhibition of farnesyltransferase (FT) enzyme, which plays a key role in the tumour cell proliferation [48,49]. More precisely, the inhibition mechanism consists in the binding of thiol units or coordination of Zn$^{2+}$ or Mg$^{2+}$ sites of FT [50,51]. As no specific sites for binding thiol units were present in the structure of the studied PEGylated phenothiazines, their ability to inhibit FT was verified by experiments of complexation with Zn$^{2+}$ or Mg$^{2+}$ metals (see Supporting Information Figures S5-S11 and explanations therein). The phenothiazine product (PAcOH) which can result by enzymatic degradation of PPO (Scheme 2) was investigated too. Neither PP nor PPO were able to bind Zn$^{2+}$ or Mg$^{2+}$ ions, indicating their inability to inhibit FT. Nevertheless, PACOH was able to bind both ions. In the case of magnesium, the bonding consisted in the formation of an ionic compound, were Mg$^{2+}$ coordinated water molecules and the electron-donating phenothiazine played the role of counter ion neutralizing the magnesium charge (Fig. 4) [52]. In the case of zinc, it appeared that Zn$^{2+}$ ions were coordinated by the acid groups of the PACOH. This suggests that in biologic fluids which favour the PPO hydrolysis, the resulted PAcOH can inhibit the farnesyltransferase by bonding the magnesium and/or zinc sites. This hypothesis is in agreement with the higher in vitro antitumor activity of the PPO compared to PP.

**Investigation of the PEGylation effect.** Recent advances in the tumour therapy demonstrated that PEG is a great building block for enhancing the drugs efficiency by protecting them against var-

![Fig. 4. Single crystal X-ray structure of the product obtained by mixing PACOH with Mg(Ac)$_2$.](image)

![Fig. 5. Brightfield microscopy images of PPO in the blank medium (in the inset it was displayed a magnified region of the picture). The scale represents 100 μm.](image)
Nanoaggregates had the tendency to accumulate in the proximity of the cells (Fig. 6). These results suggest that the antitumor effect of PPO is the result of a synergistic effect of the PEG and ester units. On a hand, the presence of PEG assures the phenothiazine protection inside the nanoaggregates improving its biocompatibility and on the other hand the ester units permit the enzymatic cleavage of the phenothiazine acid precursor which can inhibit the FT enzyme of the tumour cells [47]. This hypothesis is supported by the literature data which indicate that the cellular uptake of PEGylated derivatives follows a different mechanism compared to small molecule drugs. Thus, while the small drug molecules penetrate inside the cell by diffusion, the PEGylated compounds penetrate the cell membrane by endocytosis. In this process a vacuole is formed and the polymeric compound is degraded, in the case of PPO probably by the cleavage of the ester function. As a result, the PAcOH acid is formed and interacts with FT [56,57].

To further investigate the feasibility of this hypothesis, the antitumor activity of the PAcOH phenothiazine precursor was investigated on HeLa cells, for which PPO showed the best results. It was found that for a wide concentration range, from $10^{-6}$ to $10^{-3}$ mM, the viability of the normal cells was not affected while that of the tumour cells was around 70% (Fig. 7). This means that in the scenario of a slow release of the PAcOH by enzymatic degradation, the tumour cell killing occurs even for very low PAcOH concentrations. Moreover, the IC50 on HeLa cells was 13.2 μM while for NHDF cells was 498.5 μM, giving a high selectivity index of 37, indicating a high selectivity towards tumour cells.

Conclusions

The investigation of in vitro and in vivo antitumor activity of two PEGylated phenothiazine derivatives revealed a synergism of the phenothiazine and PEG building blocks towards an improved tumour inhibition. Thus, the two compounds showed half maximal inhibitory concentration against mouse colon carcinoma cell line (CT26) comparable with that of the traditional antitumor drugs 5-Fluouracil and doxorubicin. The median lethal dose in BALB/c mice significantly increased from 952.38 mg/kg for phenothiazine to 1450 mg/kg and 1300 mg/kg (in phenothiazine equivalents) for PEGylated derivatives, highlighting the role of PEG in the biocompatibility improvement. Both compounds inflicted a 92% inhibition of the tumour growth compared with a maximum of 60% reported for other phenothiazine compounds with quite complicated chemical structures. Furthermore, the investigation of possible tumour inhibition mechanisms suggested that the binding of PEG to the phenothiazine via an ester linkage was favourable for (i) the nanoaggregate formation with protective role for phenothiazine units and (ii) enzymatic cleavage releasing the acid phenothiazine species, which inhibit farnesyltransferase, leading to the inhibition of the cancer cell proliferation. All these findings recommend the PEGylated phenothiazine derivatives as a valuable workbench for a next generation of antitumor drugs.

Compliance with ethics requirements

The study was conducted in accordance with national and international regulations on animal welfare, identification, control and elimination of factors causing physiological and behavioral disorders: Directive EC86 / 609 EU; Government Ordinance no. 37/2002, approved by Law no. 471/2002; Law 205/2004 on animal protection, amended and supplemented by Law no. 9/2008; Joint Order of ANSVSA and of the Ministry of Interior and Administrative Reform no. 523/2008 for the approval of the Methodological Norms for the application of Law 205/2004 on animal protection.
Credit author statement

L. Marin, M. Mares and S. Cibotaru contributed to the conception, design, analysis, interpretation, writing and critical revision of the manuscript. S. Cibotaru performed the synthesis of the compounds, their structural characterization, and investigation of the antitumor mechanism and drafting of the manuscript. V. Nastasa and A.C. Bostanaru performed the in vivo tests on BALB/c mice and drafted the manuscript. I.A. Sandu performed the in vitro tests of antitumor activity and drafted the manuscript. All authors gave final approval and agreed to be accountable for all aspects of the work.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at https://doi.org/10.1016/j.jare.2021.07.003.

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