

# Non-viral Suicide Gene Therapy: Cytosine Deaminase Gene Directed by VEGF Promoter and 5-fluorocytosine as a Gene Directed Enzyme/prodrug System in Breast Cancer Model

## Authors

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## ABSTRACT

The present study investigated the potential of vascular endothelial growth factor (VEGF) promoter to derive cytosine deaminase (CD) transfected by polyamidoamine (G4-PAMAM) dendrimers to 4T1 murine breast cancer cell line as gene-directed enzyme/prodrug therapy. The VEGF promoter and cytosine deaminase gene were cloned into the pEGFP-N1 vector from the genomic DNA of 4T1 and *E. coli*, respectively. The frequency of transfection for VEGF-CD-pEGFP-N1 and pEGFP-N1-CD treated groups was  $35 \pm 3$  and  $36 \pm 4$ , respectively. MTT assay was performed to evaluate the cytotoxic effects of converted 5-fluorocytosine on 4T1 cells. Also, the optimal concentration of 5-FC in 4T1 cells transfected by VEGF-CD-pEGFP-N1 plasmid was evaluated. The GFP expression of transfected 4T1 cells by VEGF-CD-pEGFP-N1 were observed by fluorescent microscopy and flow cytometry. Results demonstrated that the suicide CD gene was successfully expressed in 4T1 cells determined by RT-PCR and GFP expression. A concentration of 200 µg/ml 5-FC was identified as optimal dose of prodrug. Furthermore, the CD/5-FC enzyme/prodrug system not only demonstrated toxicity on transformed 4T1 cells but also exerted a 'bystander effect' determined by MTT assay. The results showed that by 35% transfection with VEGF-CD-pEGFP-N1 and CD-pEGFP-N1 plasmids, 80% and 90% inhibition of the cells growth occurred, respectively.

## Introduction

Targeted therapy is a type of cancer treatment for precisely identify and attack the cancer cells. It can be used by itself or in combination with other treatments, such as standard chemotherapy, surgery or radiation therapy [1–3]. In recent years, several studies including

molecular therapy, antiangiogenesis therapy [4], immunotherapy [5], apoptosis regulation [6], signal-transduction therapy [7], differentiation therapy [8], targeted radionuclide therapy [9] and gene therapy [10] are considered for targeted cancer therapy. Current gene therapy strategies for the treatment of tumors are restricted

by the lack of targeting the malignant cells. One of the main problems for their *in vivo* applications is low transduction (or transfection) efficiency of the available gene transferring systems [11]. The cytotoxic systems convert a prodrug to an active drug intracellularly. Suicide gene therapy trials can be more promising than the other available gene therapy strategies [12]. The safety and effectiveness are the promising advantages of this approach for cancer treatment [13]. In the suicide gene therapy known as Gene-Directed Enzyme/Prodrug Therapy (GDEPT), the therapeutic transgenes have the ability to convert a non-toxic prodrug into cytotoxic drug or to express the toxic gene expression product [14–16]. Ideally, the gene should be expressed exclusively in the tumor cells not into the normal cells. The enzyme must reach a sufficient concentration for catalytic activity to activate the prodrug under physiological conditions for its clinical use. Because *in vivo* expression of the foreign enzymes will not occur in all cells of a targeted tumor, a bystander effect is required, whereby the prodrug is cleaved to an active drug that kills not only the tumor cells in which it is formed but also neighboring tumor cells that do not express the foreign enzyme [17].

Among many suicide genes, *E. coli* Cytosine Deaminase (CD) and Herpes Simplex Virus thymidine kinase (HSV-tk) are well documented for their strong therapeutic efficacy in cancer treatment [18]. In CD/5-fluorocytosine (5-FC) system, a suicide gene, CD, catalyzes the conversion of the anti-fungal substrate 5-FC into 5-fluorouracil (5-FU), which is a lethal drug [19]. Suicide gene therapy, utilizing the CD/5-FC system, is an efficient approach for targeted therapy in cancer study [20–22]. Most suicide genes under investigation mediate sensitivity by encoding bacterial or fungal enzymes that convert inactive forms of a drug, into toxic metabolites capable of inhibiting nucleic acid synthesis [23, 24]. The cellular enzymes process the formulation to three cytotoxic antimetabolites: 1) 5-FdUTP, 2) 5-FUTP and 3) 5-FdUMP. The cytotoxic effects of CD suicide therapy are based on three properties: a) formation of (5-FU) RNA, b) 5-FU DNA complexes and c) thymidylate synthase. The mitochondrial pathways are down-regulated through the control of Bcl-2 [25]. Also, heat shock protein 90-beta is activated, contributing to tumor regression [26].

CD/5-FC system has some advantages over other GDEPT systems, including significant distant bystander effect independent of gap junctions [27, 28]. One of the advantages of CD/5-FC system is radiosensitizing ability of 5-FU to enhance tumor cytotoxicity efficiency along with radiotherapy [25, 29]. Some studies have obtained the mutant forms of bacterial CD in order to improve their kinetics including high affinity and lower IC<sub>50</sub> for 5-FC [30].

The bystander effect can improve the therapeutic consequences which extended beyond the transfected tumor cells. There are currently five mechanisms investigated for the regression of untransfected cells due to bystander effects: a) release of soluble formulations, b) passage through gap junctions, c) passive transportation, d) stimulation of local microenvironment, and e) endocytosis of apoptotic vesicles. The CD/5-FU system demonstrates stronger local bystander effect than the other suicide genes. The 5-FU diffuses efficiently within the tumor cells, and does not require cell to cell contact [25].

There are currently two main vehicles used to deliver the suicide genes: viral vectors and non-viral vectors [31]. Non-viral vectors are widely used with three different approaches: 1) naked DNA, 2) phys-

ical methods and 3) chemical methods. Non-viral vectors such as cationic liposomes and cationic polymers are less efficient than viral vectors; they have the advantages of safety, simplicity of preparation and high gene encapsulation capability. These two delivery systems have a positive charge, which interacts with the negative charge of the DNA [32]. PolyAmidoAmine (PAMAM) dendrimers, is used for drug and gene delivery. The positive charge of PAMAM dendrimer surfaces decreases their cytotoxicity and has interesting implications for DNA transfection applications. Depending on its characteristics, such as a high efficiency in delivery, the capabilities of specific targeting, biodegradability, non-toxicity, non-immunogenicity, and capability of limiting DNA degradation, PAMAM dendrimer provides a promising approach for gene delivery [11, 12].

Gene therapy for cancer requires vectors that are selectively expressed in tumor cells. It is achieved by putting the suicide gene under the control of a tumor specific promoter [33]. A wide number of cancer tissue specific promoters' studies have used tissue-specific promoter elements, such as: human telomerase reverse transcriptase (hTERT) promoter, carcino-embryonic antigen (CEA) promoter, and osteocalcin (OC) promoter [34]. In the recent years, various new promising promoters like auxin response factors (ARF) [35], glucose-regulated protein (GRP78) [36], chemokine (C-X-C motif) receptor-4 (CXCR4) [37], and osteopontin (OPN) [38], have been developed. The human cytomegalovirus (HCMV) promoter/enhancer is a flexible promoter that is expressed in many cell types and provides high levels of trans-gene expression [39]. Based on the overexpression of the vascular endothelial growth factor (VEGF) in breast cancer cells [34, 40], a recombinant pEGFPN1 harboring the VEGF promoter and the CD suicide gene (pEGFPN1-VEGFp-CD), was constructed in the present study, for expression in 4T1 cell line. Plasmid constructs modified with the VEGF promoter element may provide improvements in gene expression for cancer cells. In the present study, we designed a tumor-specific promoter system by combining VEGF promoter, which can target to cancer cells with sufficient transcriptional activity. A gene-directed enzyme prodrug therapy (GDEPT) system consisting of the bacterial cytosine deaminase gene (bCD) driven by the VEGF promoter and the prodrug 5-fluorocytosine (5FC) was developed and its cytotoxicity effect was evaluated.

## Materials and Methods

### Cell culture condition, reagents and instruments

4T1 cells, *E. coli* BL21 and *E. coli* TOP10, were obtained from Pasteur institute of Iran and were cultured in LB agar media. 4T1 cells were cultured in RPMI 1640 medium, supplemented with 10% fetal bovine serum and 1% penicillin/streptomycin, and incubated at 37 °C in a 5% CO<sub>2</sub> incubator.

The restriction enzymes were purchased from Takara Bio, Inc., (Otsu, Japan). T4 DNA ligase, Probest DNA polymerase and were purchased from New England Biolabs, Inc., (Ipswich, MA, USA). Agarose gel DNA extraction kit, 5-fluorocytosine (5-FC), MTT and dimethyl sulfoxide (DMSO) were purchased from Sigma-Aldrich (Merck KGaA, Darmstadt, Germany).

The instruments that were used in the present study were: DLS (zetasizer, USA), ELISA Infinite Tecan (USA), Bio-Rad GenePCR

(PerkinElmer, Inc., Waltham, MA, USA), OLYMPUS Ix53 optical microscope (Olympus Corporation, Tokyo, Japan), an OLYMPUS BX61 inverted fluorescence microscope and camera system (Olympus Corporation), BD Calibur Flow Cytometer (BD FACS Calibur, San Jose, CA, USA) and step -one-plus Real time PCR (ABI, USA).

## Molecular cloning of shuttle vector constructions

### Amplification of CD gene

*E. coli* strain BL21 was inoculated into 5 mL liquid LB medium with shaking, for overnight at 37 °C. The genomic DNA of *E. coli* BL21 was extracted by lysis method. It used as the template DNA for the amplification of CD gene by PCR. Specific primers of CD gene were designed based on published sequence (Genebank, NC-000913). The CD gene was amplified from the obtained genomic DNA of *E. coli* BL21 by PCR. The following primers were used: forward primer, 5'- TATAAGCTTGTGTCGAATAACGCTTTAC -3' (Italic part indicates the recognizing site of *Hind*III) and the reverse primer, 5'- ATTAAGCTTCAACGTTTGTATC-GATGG -3' (Italic part indicates the recognizing site of *Hind*III). The PCR product of CD (1302 bp) and the expression vector, VEGF-pEGFP-N1 (5349 bp) made in our previous stud [41], were digested by *Hind*III and then ligated to obtain the final plasmid, VEGF-CD-pEGFP-N1 (6650bp). Also, the pEGFP-N1 plasmid was linearized by *Eco*RI and *Xho*I and the CD gene was inserted in its multiple cloning sites (*Hind*III). Finally, the recombinant plasmids sequences were verified by DNA sequencing, and the plasmid was further amplified and purified by a QIAfilter™ plasmid maxi kit (Qiagen China Co., Ltd., Shanghai, China), according to the manufacturer's protocol.

## Optimization of dendriplex transfection

### Preparation of PAMAM/DNA Dendriplex

G4-PAMAM-D, Sigma-Aldrich, (USA), was kindly provided by Dr. Behzad Baradarn (Tabriz University of Medical Sciences, Tabriz, Iran). Dendriplexes of DNA/PAMAM complexes were prepared at different N/P ratios. The N/P ratios were calculated based on PAMAM nitrogen per nucleic acid phosphate. 2 µg of DNA were mixed in HEPES buffer saline (20 mM) and appropriate amounts of diluted PAMAM by HEPES buffer saline were added to DNA solutions. Solutions were mixed. The resulting dendriplexes were incubated for 30 min at room temperature before use.

## Gel retardation assay

Gel retardation assay was performed to assess the formation of electrostatic complexes between PAMAM and plasmid DNA. Briefly, a 0.1 % agarose gel was prepared in TAE buffer (1X) with 0.5 µg/ml safe stain. Dendriplexes were loaded into the agarose gel wells at N/P ratios of 5, 10, 20, 40, 80, and 160. Samples of dendriplexes were run against free plasmid. Electrophoresis was conducted at 80 v for 45 min. Gel were photographed under UV light.

## Size and zeta potential measurements

20 µg DNA was incorporated with the different amount of PAMAM in 10 mL HEPES buffer saline (20 mM) giving N/P = 10,20,40,80 and 160 polyplex. Dynamic light scattering was used to measure the mean diameters and surface *Zeta* potentials of the polyplexes using Zetasizer-Nano ZS (Malvern Instruments, Worcestershire, UK). The mean diameters and *Zeta* potentials of the dendriplexes were measured three times in triplicate.

## Cell culture and cytotoxicity assay of dendriplexes & prodrug on 4T1 cells

4T1 (murine mammary gland tumor), were cultured in RPMI 1640 medium with 10 % fetal bovine serum (FBS), and were maintained at 37 °C in a 5 % CO<sub>2</sub> incubator. The potential PAMAM-G4 dendriplex cytotoxicity was evaluated on 4T1 cells. This study was carried out by MTT assay of cell, according to the method described by Mosmann [42]. Briefly 5 × 10<sup>4</sup> cells were seeded in a 24 well plate and after 24h the cells were treated with different N/P ratios of dendriplex at 500 µl RPMI1640 media. After incubation at 37 °C for 4–6 h, the growth medium was replaced with fresh FBS + /antibiotics + media and cultured for one day. The culture medium was then removed and a total of 20 µl MTT (5 mg/ml) was added to each well and incubated for 4 h at 37 °C. The culture medium was then removed, followed by the addition of 100 µl DMSO to each well. The absorbance was measured by a plate reader at 570 nm. The cell viability percentages of different N/P ratios were calculated by considering the absorbance of control as 100 % viability. Presented results correspond to n = 3 performed in triplicate and expressed as mean ± SD.

MTT colorimetric assay was used to evaluate the lethal effect and optimal concentration of 5FC on 4T1 and transfected 4T1 cells. Cells were cultured in 96-well plates (1 × 10<sup>4</sup> cells) in 200 µl RPMI 1640. After 24 h of culture, the concentration of 5-FC was tested at 0, 50, 100, 200, 400, 600, 800 or 1000 µM. The blank control, experimental and control groups were set up in five replicate groups. In total, 24, 48 and 72 h after 5-FC treatment, the cell viability was determined using the MTT assay. The culture medium was then removed and a 20 µl MTT (5 mg/ml) was added to each well and incubated for 4 h at 37 °C. The culture medium was then removed, followed by the addition of 100 µl DMSO to each well. The absorbance (optical density; OD) of each well was measured at a wavelength of 570 nm. The cell survival rate (%) was calculated according to the following formula: (Experimental group OD value/blank control group OD value) × 100. The IC<sub>50</sub> value was determined with a Prism 3.03 program for each group.

## Cell transfection

One day before transfection, cells were seeded in 6-well plates and cultured in growth medium without antibiotics. The cell confluency was approximately 70 % at the time of transfection. 2 µg plasmid per well was used for transfection in 6-well plate. PAMAM and plasmid DNA (pDNA) were freshly diluted to equal volumes with HEPES buffer saline prior to use. The dendriplexes were prepared by adding PAMAM into pDNA using different N/P ratios. The dendriplexes were incubated for 30 minutes at RT. Based on standard PAMAM mediated transfection, the growth medium was replaced by fresh FBS free/antibiotics free medium (2 mL for 6-well) and then the dendriplexes diluted by RPMI1640 medium and added to the culture. After incubation at 37 °C for 4–6 h, the growth medium was replaced with fresh FBS + /antibiotics + media and cultured for one to two days for detection.

## CD and GFP gene expression analysis

### Fluorescent microscopy

To qualitatively evaluation of the transfection efficiency, cells transfected with pEGFP-N1 were observed and imaged using an invert-

ed fluorescent microscope (Olympus BX61) for GFP expression at days 1 and 2 after transfection. After 72 h of cell culture (37 °C), cells were washed with PBS three times, trypsinized and collected for observing by fluorescence microscope. Cells were then centrifuged at 250 × g for 2 min, washed with 4 °C precooled PBS twice, fixed in 4 °C precooled 3 % glutaraldehyde for 4 h and rinsed with PBS for 10 min. Subsequently, they were fixed on slides at room temperature and observed under the fluorescent microscope.

### Flow cytometry analysis

The transfection efficiency was quantitatively analyzed at cellular level by detecting and comparing the GFP excitation in the transfected and non-transfected cells. 2 days after transfection, the cell colonies were dissociated with trypsin and subjected to flowcytometry analysis (BD Biosciences, San Jose, CA, USA). The data were processed using FlowJo vX10 software. The transfection efficiency was determined as the percentage of viable GFP-positive cells among total cells.

### Real Time PCR

After 48 h of transfection, total RNA was extracted from transfected 4T1 CD<sup>+</sup> (CD-transfected 4T1), with TRIzol reagent (Invitrogen; Thermo Fisher Scientific, Inc.) and then one-step reverse transcription (RT)-PCR was performed using a One-step RT-PCR kit, according to the manufacturer's protocol. The forward (5' CAGCGGCTACCGTGAT-TCA 3') and reverse (5' TTGCACATGGCGTTGG 3') primers were designed to amplify a 61 bp fragment of the CD gene. PCR reaction conditions were as follows: 95 °C for 5 min, followed by 30 cycles of 94 °C for 30 sec, 58 °C for 30 sec, 72 °C for 1 min, then 72 °C for 10 min. Differences in gene expression were normalized to the  $\beta$ -actin gene expression with the following primers: forward 5'-aacgagcgggtccgatgccttgag-3'; reverse 5'-tgtcgcttcaccgttcagtt3.

### Bystander effect

After determining the percentage of transfection, the transfected cells were mixed with a certain amount of non-transfused cell and the normalized cells were seeded in 96-well plates 2 days prior prodrug addition. In brief, cells were washed with RPMI1640 and treated by 5-FC. The number of viable cells after 5-FC was measured according to the cell cytotoxicity assay mentioned in MTT assay section [43, 44].

### Evaluation of the effect of cell soup obtained from transfected cells on non-transfected cells

According to the transfection instructions, cells in the 6-cell plate were transfected at N/P ratio 160. 48h after transfection, the supernatant of the transfected cells was collected. 66 % and 33 % of the cell soup were prepared and the cells were treated with 5-FC for 48 hours. Non-treated cells transfected with cell soup were considered as positive control and non-transfected cells untreated with cell soup were considered as negative control.

### Statistical analysis

Statistical analyses were performed using GraphPad Prism 8.02 (©1992–2019 GraphPad Software, Inc.). Data were expressed as mean  $\pm$  standard deviation of the mean and analyzed by one-way ANOVA. The means were compared using Duncan's multiple range tests.

## Results

### Construction and confirmation of the plasmids

We used *in Silico* Snap Gene design to obtain the schematic structure of plasmids and MCS sites (► Fig. 1).

#### VEGF- CD -pEGFP-N1 plasmid

The CD gene was amplified from *E. coli* BL21 genomic DNA by PCR (► Fig. 2a). The VEGF-pEGFP-N1 vector and CD gene (1302 bp) were digested by *Hind*III restriction enzyme, before ligation and transformation into competent cell (► Fig. 2b). Colony PCR was performed for transformation screening [41] (► Fig. 3a). The recombinant plasmid was extracted and verified by enzymatic digestion (► Fig. 3b).

#### CD -pEGFP-N1 plasmid

The CD gene was amplified from *E. coli* BL21 genomic as mentioned before (► Fig. 2a). The pEGFP-N1 and a 1302 bp fragment of CD sequence were digested, before ligation and transformation into competent cell (► Fig. 4a). Colony PCR was performed for transformation screening (► Fig. 4b). The recombinant plasmid was extracted and verified by enzymatic digestion (► Fig. 4c).

### Gel Retardation assay

Gel retardation assay was performed to determine the most appropriate N/P ratio of dendrimer and plasmid DNA. PAMAM /plasmid complexes were prepared at various N/P ratios ranging from 5, 10, 20, 40, 80 and 160. After 30 min of incubation and dendriplex formation, the samples were electrophoresed (► Fig. 5).

### Transfection optimization

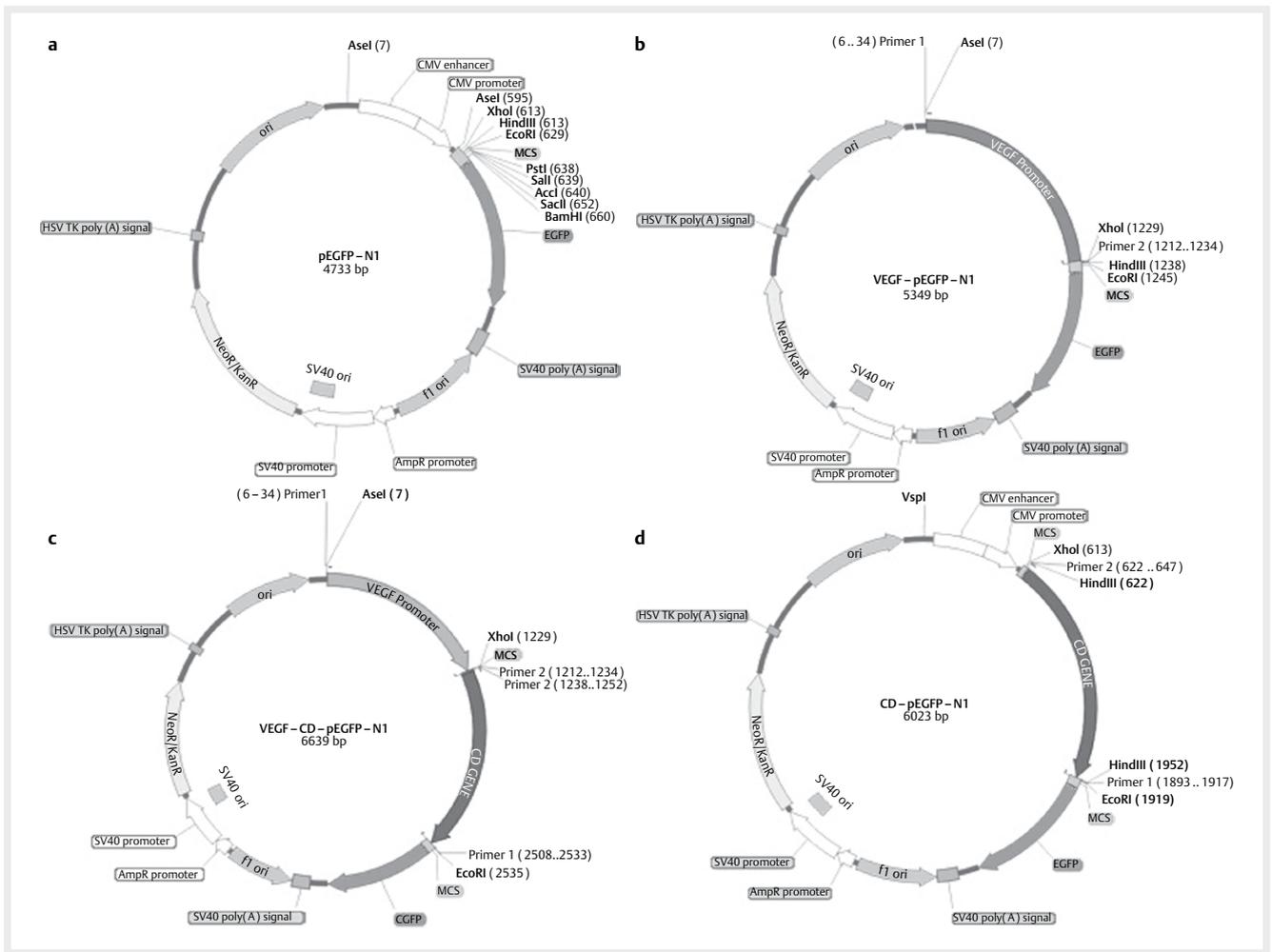
In order to increasing the transfection efficiency, four factors including size, zeta potential, cell cytotoxicity and N/P ratio of dendriplex were investigated. As the N/P ratio increases, the particle size and the zeta potential of dendriplex and also the transfection percentage increase but the cell viability decreases (► Fig. 6).

### Dendriplexes size and zeta potential

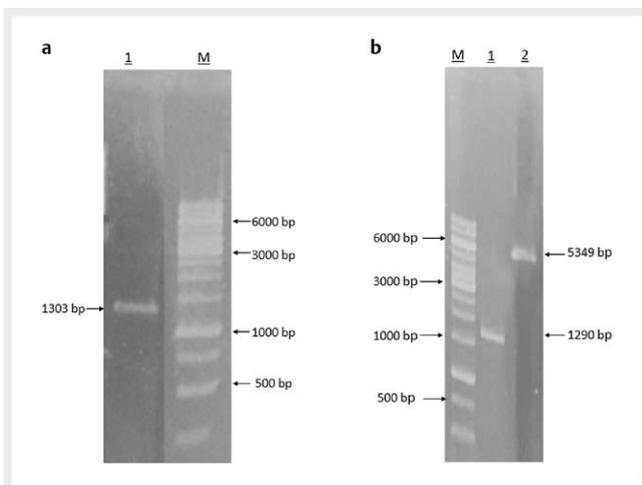
Since PAMAM has a positive charge and DNA has a negative charge in aqueous solution, PAMAM and DNA can form a complex with electrostatic effects in physiological conditions. PAMAM/plasmid complexes (dendriplex) at a most appropriate charge ratio were incubated for transfection of DNA. The particle size, size distribution and particle charge of PAMAM G4 dendrimer was measured by DLS (Zetasizer Nano ZS, Malvern Instruments, UK) (► Fig. 6). Data are expressed as mean  $\pm$  SD. In the following, size and charge of polyplex (DNA + PAMAM) with various N/P ratios were measured by DLS.

### Cellular cytotoxicity and transfection efficiency of dendriplex

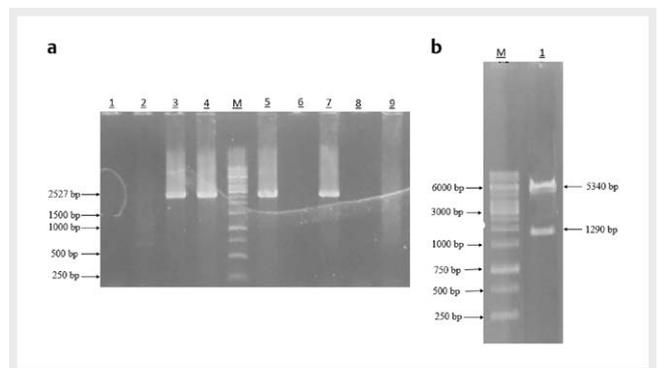
Increasing the dendrimer content of the dendriplex increases its cytotoxicity, but more importantly, increasing the size of the dendriplex may make endocytosis by cells almost impossible. As shown in ► Fig. 6a, with increasing the dendrimer content of the dendriplex, its size also increases, so that in the dendriplex with a N/P ratio of 160, dendriplex size has reached to 400 nm.



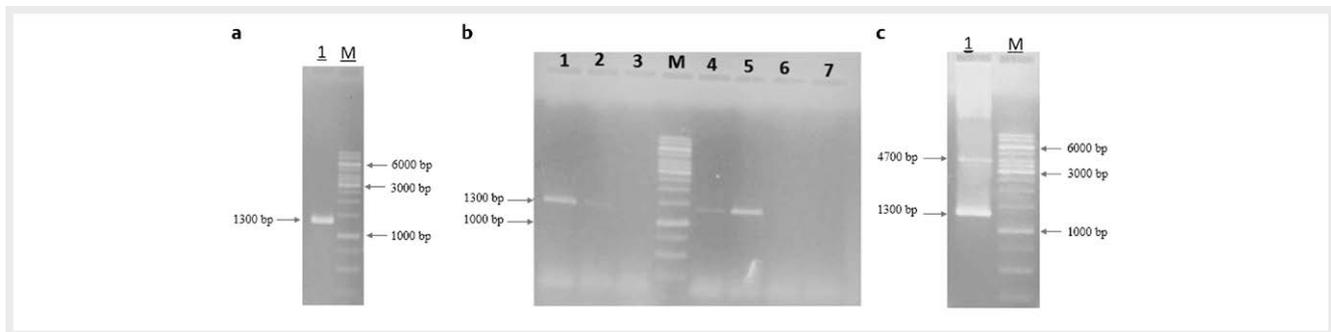
► **Fig. 1** *In Silico* SnapGene constructs: **a** native pEGFP-N1 (with CMV promoter and total size 4733 bp), **b** VEGF-pEGFP-N1- (substitute VEGF<sub>promoter</sub> with CMV, size : 5349 bp), **c** VEGF-CD-pEGFP-N1-, final construct, size : 6639 bp), **d** CD-pEGFP-N1- ( size : 6023 bp).



► **Fig. 2** **a)** Gel electrophoresis of PCR product of CD gene; M: 1 Kb Ladder (Fermentas Co.), **1**: CD PCR product. **b)** Gel electrophoresis of CD & pEGFPN1-VEGF enzymatic digestion by *HindIII*), M: 1 Kb Ladder (Fermentas Co.), lane**1**: cytosine deaminase, lane **2**: VEGF- pEGFP-N1.



► **Fig. 3** **a)**: Gel electrophoresis of colony PCR products after ligation and transformation, lane M, 1 Kb ladder (Fermentas Co.), lanes 1 to 12, colony PCR products of random selected colonies. Clones of 2, 5,6,7,9 and 12 are positive transformed clones. **b)**: Molecular confirmation of correct orientation of inserted CD into VEGF- pEGFP-N1 by enzymatic digestion by *XhoI*, *HindIII* and electrophoresis: lane M: 1 Kb ladder (Fermentas Co.), lane**1**: uncut construct, lane **2**: cut VEGF-pEGFP-N1 (5349 bp) and CD (1300 bp).



► **Fig. 4** **a** Gel electrophoresis of PCR product of CD gene; M: 1 Kb Ladder (Fermentas Co.), 1 CD PCR product. **b** Gel electrophoresis of colony PCR products after ligation and transformation, lane M, 1 Kb ladder (Fermentas Co.), lanes 1 to 7, colony PCR products of random selected colonies. Clones of 1, 4 and 5 are positive transformed clones. **c** Molecular confirmation of inserted CD into VEGF- pEGFP-N1 by enzymatic digestion and electrophoresis: lane M: 1 Kb ladder (Fermentas Co), lane 1: cut CD- pEGFP-N1 (4733bp) and CD (1302 bp).



► **Fig. 5** Dendriplex Gel Retardation assay: Lane 1: control (pEGFP-N1); lanes of 2 to 7: dendriplex (pEGFP-N1 + PAMAM) with different N/P ratios (5, 10, 20, 40, 80, 160).

## Fluorescence microscope and flowcytometry analysis

The constructs expresses a fluorescent reporter (green fluorescent protein) and a suicide cytosine deaminase gene, under control of VEGF promoter. The GFP expression was detected in 4T1 cells at different N/P ratios of dendriplexes (► **Fig. 7**). These show that dendriplexes were not cytotoxic in 4T1 tumor cells and were uptaken by these cells. The increasing N/P ratio may be influencing the total number of cells was GFP positive and the transfection efficiency. This hypothesis has been demonstrated qualitatively and quantitatively by fluorescent microscopy (► **Fig. 7**) and flowcytometry (► **Fig. 8**), respectively.

## Expression of CD driven by the VEGF promoter or CMV promoter inhibits 4T1 cells growth

To investigate the level of cell proliferation in these enzyme/prodrug systems (VEGF-pEGFP-N1, VEGF-CD-pEGFP-N1 and CD-pEGFP-N1), 4T1 cells were transfected with these recombinant plasmids and its sensitivity to the 5-FC was evaluated by MTT assay. Cell viability % versus 5-FC was drowned using GraphPad Prism from the dose-cell viability curves. From the  $IC_{50}$  comparisons shown in ► **Fig. 9**, it is clear that the cell growth inhibition rates in the VEGF-CD-pEGFP-N1 and CD-pEGFP-N1 treated cell groups was observed

while the cytotoxicity in the VEGF-pEGFP-N1 treated group was observed under the same experimental conditions. Cell growth inhibition rates were progressively elevated, may be due to 'bystander effect'. Therefore, the presence of bystander effect was investigated in the next experiment.

## Bystander effect

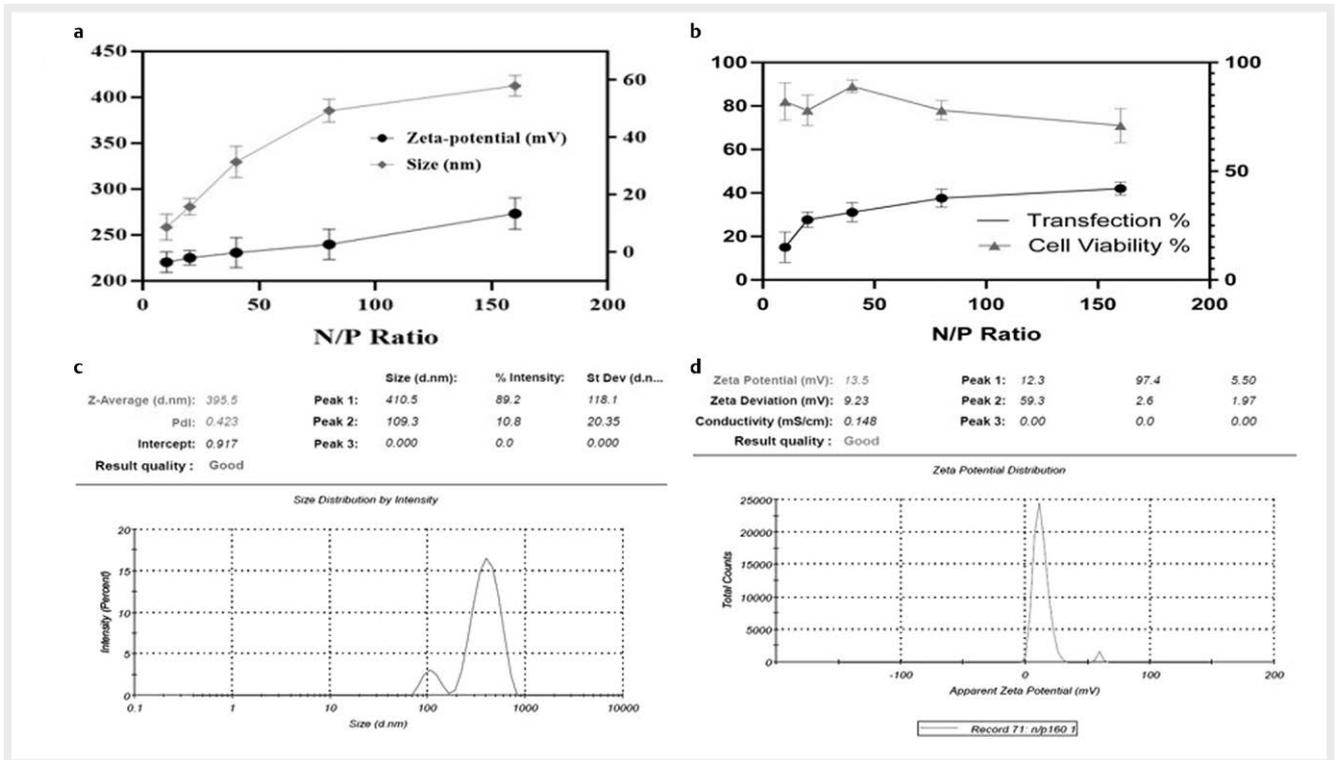
For investigation of the bystander effect of each group in response to 5-FC treatment, co-cultures consisting of transfected and non-transfected 4T1 cells were used. First, transfection efficiency was measured by flowcytometry and normalized cell transfection % was prepared by co-culturing of transfected and non-transfected 4T1 cells for each group of 4T1 cells. All of cell groups was treated by 100 $\mu$ M 5-FC, and cell survival rates were calculated. Frequency of transfection for VEGF-CD-pEGFP-N1 and pEGFP-N1- CD treated group cells was  $35 \pm 3$  and  $36 \pm 4$  (► **Fig. 10**). By increasing in the transfection rates of VEGF-CD- pEGFP-N1 and CD - pEGFP-N1 treated group cells, cell viability % were progressively decreased, may be due to 'bystander effect'. However, the most prominent 'bystander effect' was observed in the CD-pEGFP-N1 treated groups.

## Effect of transfected cell supernatant on non-transfected cells

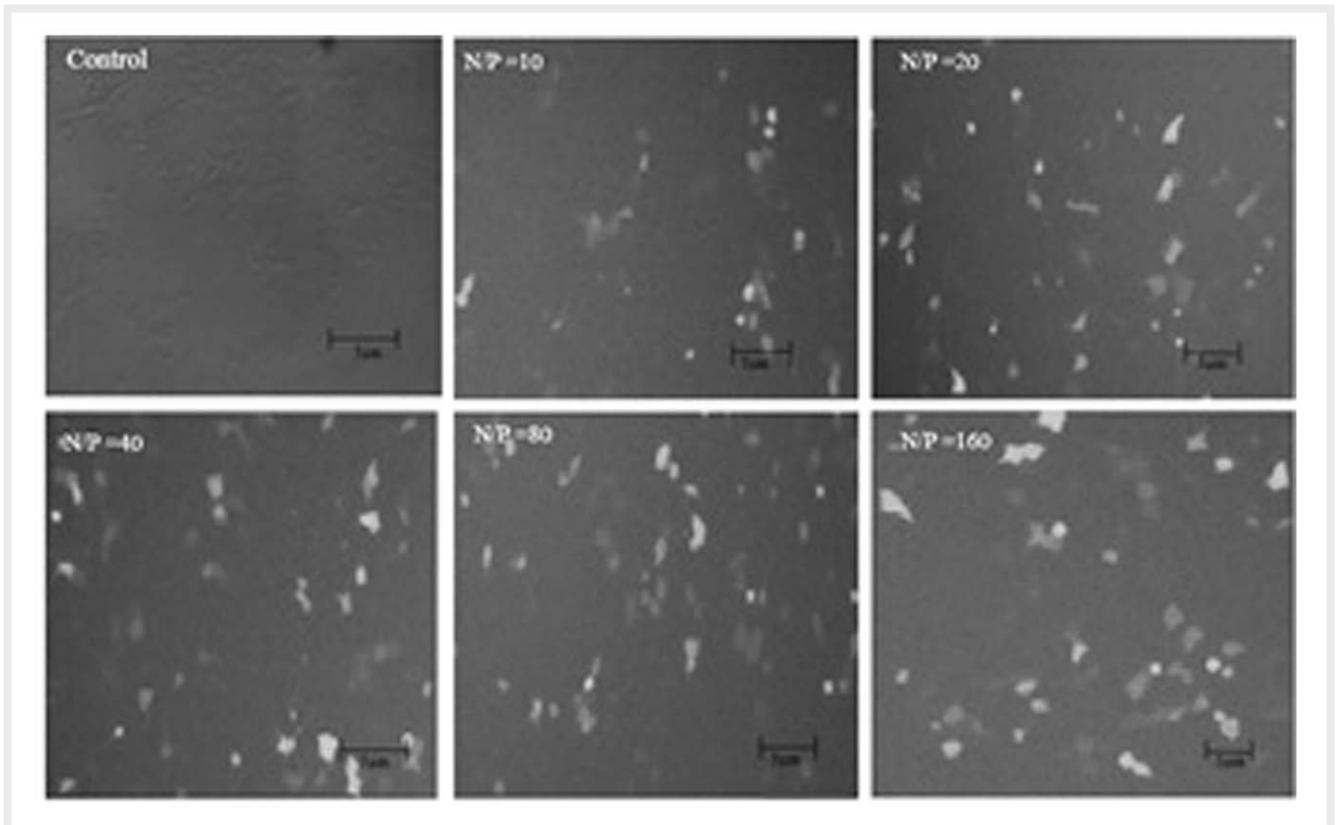
To determine the effect of cell supernatant of transfected cells on prodrug, concentrations of 33 % and 66 % of cell supernatant were added on non-transfected cells and treated with different concentrations of prodrug. Toxicity in cells treated with cell supernatant was confirmed (► **Fig. 11**).

## RT-PCR of CD gene expression into transfected 4T1 cell

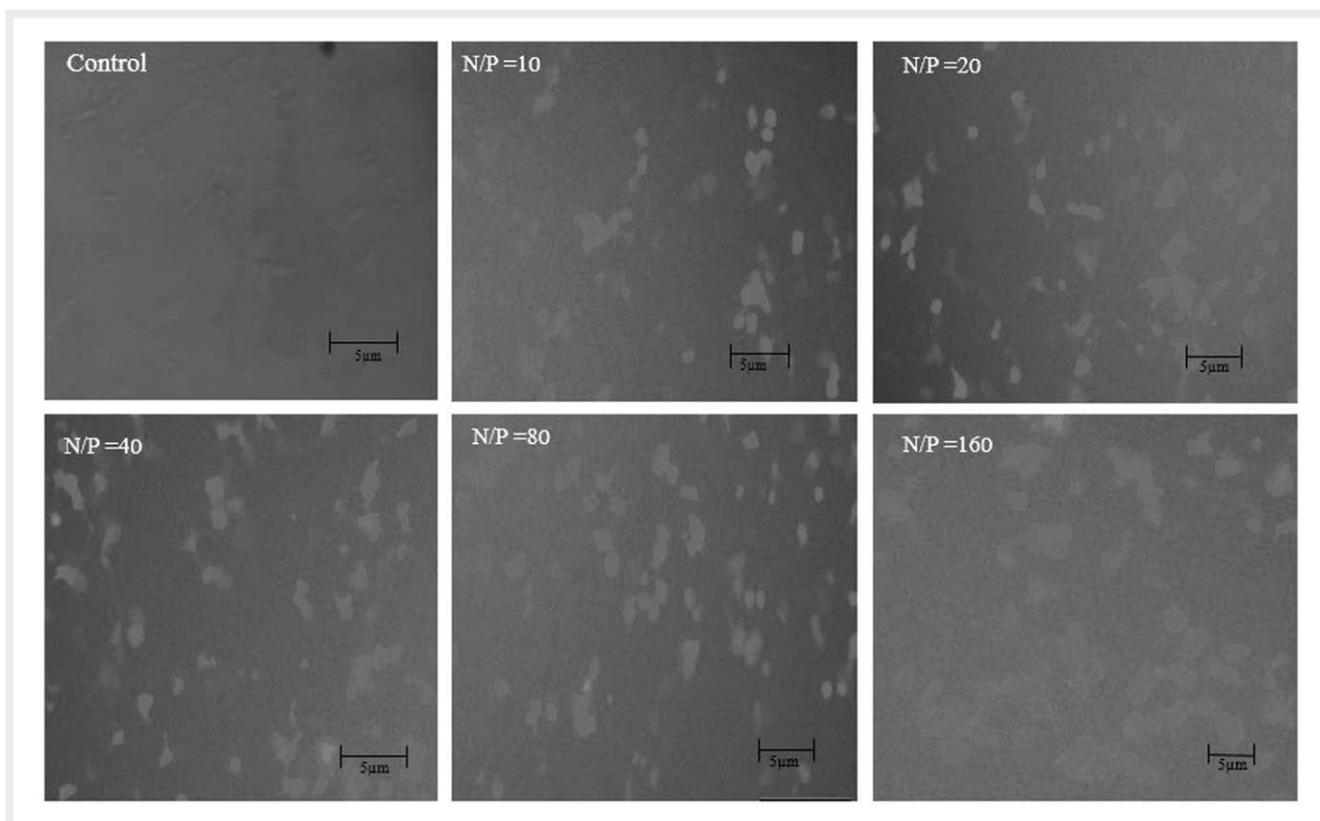
We used RT-PCR to detect the expression of the CD gene 1300-bp fragment and expression rate comparison between of VEGF-CD-pEGFP-N1 and CD - pEGFP-N1 treated group cells. The transcriptional activity of the CMV promoter (positive control) in each cell line was considered as 100 %. The expression of CD gene by the VEGF promoter was approximately from 36 to 72 % of positive control (CD - pEGFP-N1) (► **Fig. 12**).



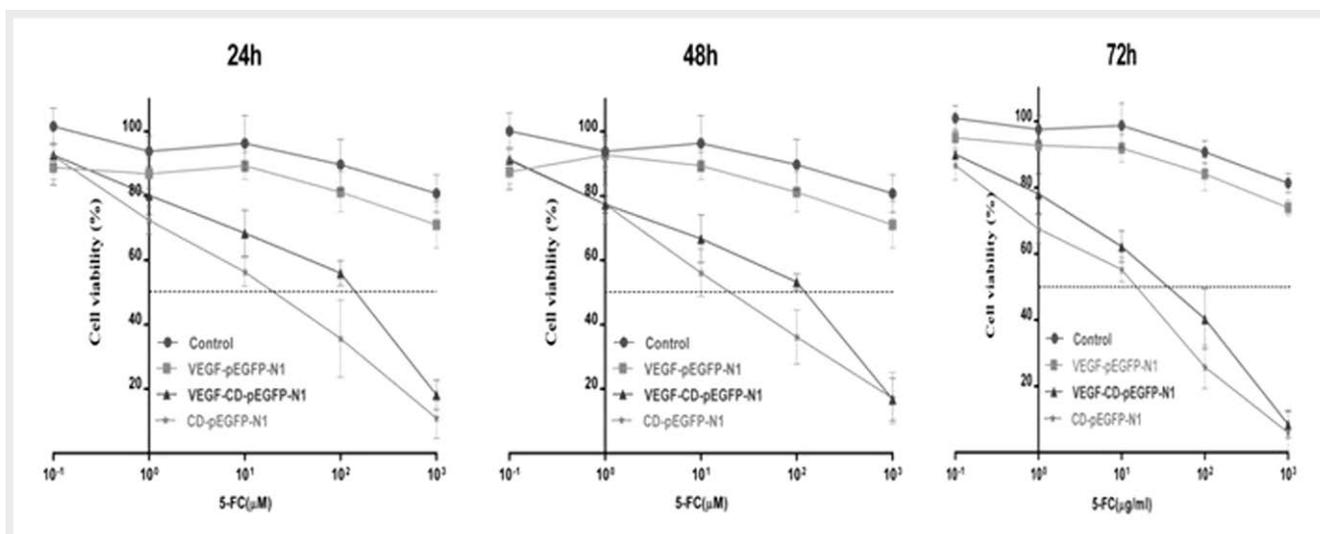
► **Fig. 6** Characterization of the PAMAM /pEGFP-N1 (dendriplex). **a** Mean diameter and surface charge of the polyplex versus various N/P ratios using DLS (data expressed as mean + S.D. of three-independent measurements). **b** cellular cytotoxicity and transfection efficiency of dendriplex versus various N/P ratios; **c** and **d** Polyplex sizes and zeta potentials measured by Zetasizer for example.



► **Fig. 7** GFP fluorescence imaging: 4T1 cells were transfected by pEGFP-N1 after 48h at different N/P ratios (Control, and: N/P= 10, N/P= 20, N/P= 40, N/P= 80, and N/P= 160).



► **Fig. 8** Flowcytometry imaging: 4T1 cells were transfected by pEGFP-N1 after 48 h at different N/P ratios: **a** Control, **b** N/P = 10 (% transfection; 7.9), **c** N/P = 20 (%transfect; 17.6), **d** N/P = 40 (%transfect; 25.9), **e** N/P = 80 (%transfect; 35.2), **f** N/P = 160, (%transfect; 45.8)

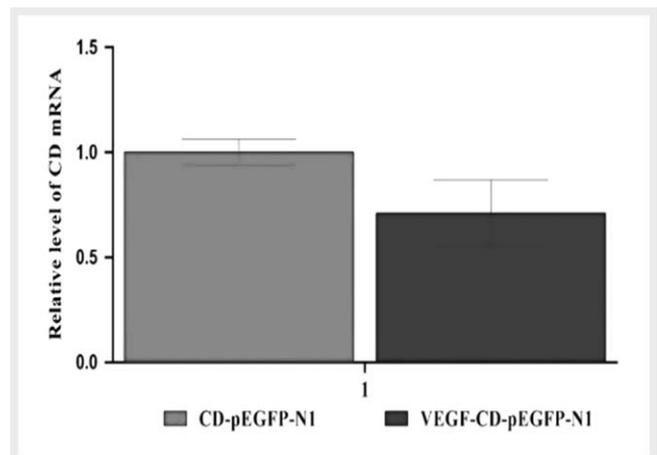
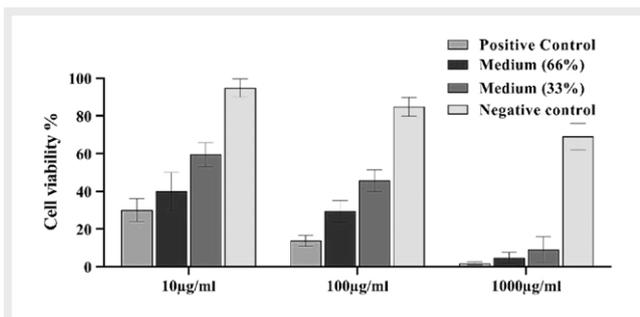
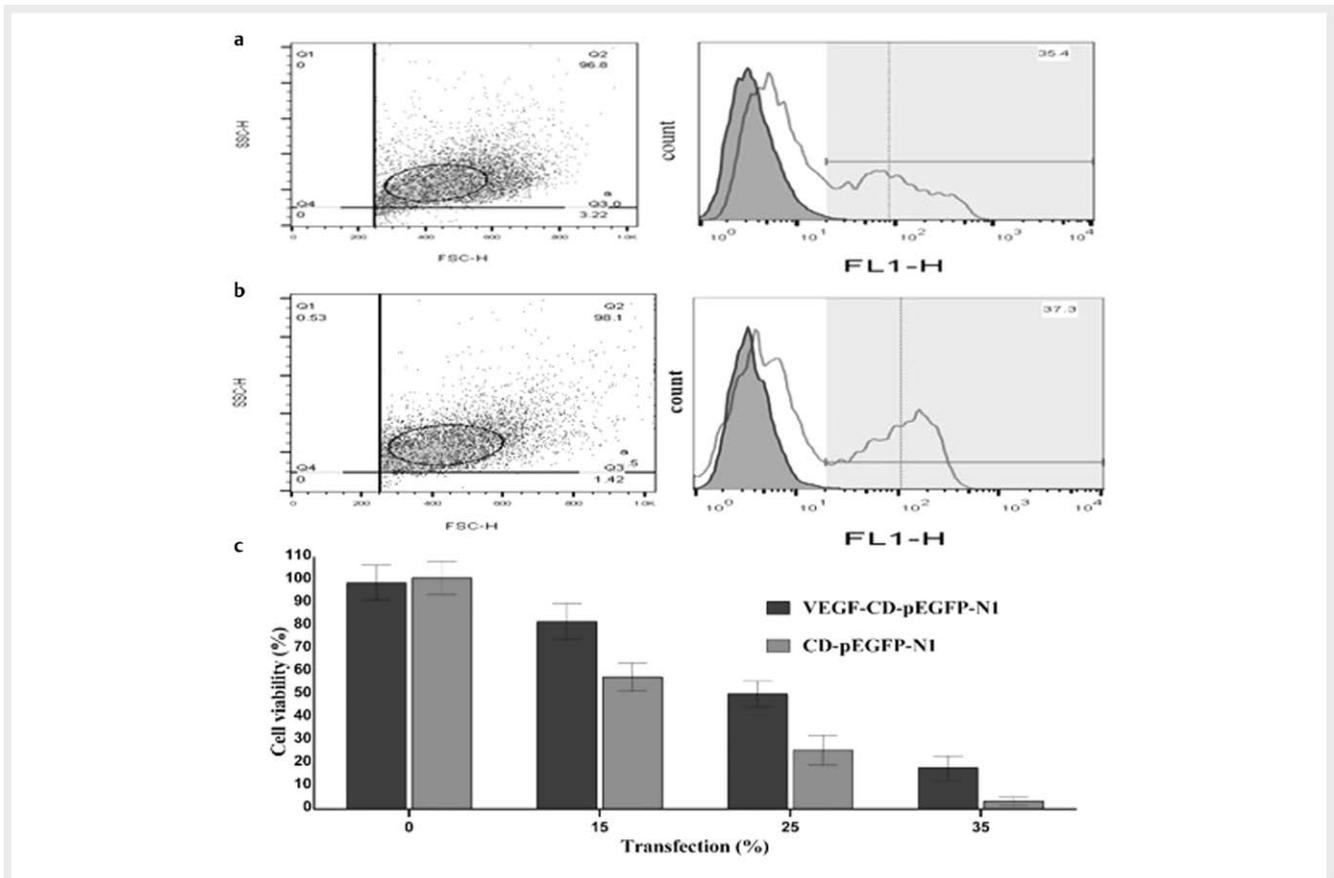


► **Fig. 9** 4T1 cells transfected with different plasmid shown in ► **Fig. 10**. The MTT assay was applied for each cell groups at the different concentration of prodrug (5-FC) at 24 h, 48 h, and 72 h of incubation. Viability percentages and the error bars are expressed as mean  $\pm$  SD and represent the result of five experiments. No cytotoxicity from PAMAM -G4 dendrimer and 5-FC was observed on this cell line.

## Discussion

The purpose of this study was evaluation of the effectiveness of VEGF promoter and bystander effect on the efficiency of the CD/5-FC enzyme/prodrug system using a dendriplex-based non-viral

gene delivery system. Non-viral gene delivery systems are highly regarded for their many advantages compared to viral gene delivery systems [45]. However, their low efficiency in cell transfection has restricted their applications in clinic. Therefore, researchers are



looking for a way to compensate this shortcoming of non-viral systems and replace non-viral systems with today's high-risk viral systems. One potential therapeutic strategy for cancer treatment is the technique of inserting suicide genes that activate prodrugs to produce cytotoxicity in tumor cells. Suicide gene therapy, utilizing the cytosine deaminase/5-fluorocytosine (CD/5-FC) system, is an efficient procedure for targeted therapy in cancer research with promising results in previously reported studies [46, 47]. It is based on the introduction of a viral or bacterial gene (and or fungal gene)

that encodes a metabolic enzyme into target cells, which allows the conversion of a non-toxic compound into a lethal drug, causing the death of target cells. The 'bystander effect' is a phenomenon whereby the transfection or transduction of a small fraction of

target cells with the suicide gene may result in widespread target cell death, including the non-transfected cells [48]. Among many GDEPT, the most frequently used suicide gene therapy for clinical trials approved by the FDA, are two major systems as: the CD/5-FC system along or with the (HSV-TK)/GCV system [46]. Cytosine Deaminase, which normally catalyzes the deamination of cytosine to uracil during RNA biosynthesis, is found in many bacteria and fungi but not in mammalian cells [49]. However, the drug has no or low toxicity for normal mammalian cells. 5-FC is an antifungal drug compound that is non-cytotoxic but is transformed into 5-FU following deamination by the CD gene, which is more cytotoxic and may act to inhibit cellular proliferation [48]. Because of these many advantages of the CD/5-FC system and having a suitable bystander effect, we decided to deliver this system with a dendrimer-based non-viral gene delivery system. We can have a comparison between the high-risk viral method at literature and the low-risk non-viral method [50], and we hoped that the low efficiency of the non-viral gene delivery system would be offset by the bystander effect.

The plasmid (pEGFPN1) contains of the cytosine deaminase (CDase) suicide gene, under the VEGF promoter (VEGF-CD-pEGFP-N1), was used to transfect 4T1 cells with the G4-PAMAM dendriplex. The highest efficiency after optimization of the most factors for our non-viral system was between 40–50%, and we investigated if cells transfection was 50%, it is enough to kill all of the transfected cells and nontransfected cells? For this purpose, we organize the bystander effect test [43] performed on on transfected cells, with non-viral gene delivery systems. The results showed that in 35% transfection, cells transfected with plasmid VEGF-CD-pEGFP-N1 and CD-pEGFP-N1 inhibited the growth of cancer cells by 80 and 90%, respectively.

Another aim of this project was to confirm the expression of CD gene by transfected cells. For this purpose, three methods were used. First, because we know that both the CD and GFP genes are under the same promoter, the expression of GFP will also express the expression of the CD gene in a way that the expression of GFP gene was quantitatively and qualitatively confirmed. Second, unlike transfected cells with CD-plasmids, cell transfected with CD<sup>+</sup> plasmids showed toxicity at a certain concentration of prodrug, which could be due to the function of CD enzyme and conversion of prodrug to a more toxic drug (5-fluorouracil). Third, the expression of the CD gene at the transcriptional level was confirmed by real-time PCR method the presence of CD gene mRNA in total RNA of transfected cells with CD<sup>+</sup> plasmids.

In this study, we created a plasmid controlled by a cancer specific promoter (VEGF). The VEGF promoter was obtained from 4T1 cell lines and inserted into the pEGFP-N1 expression plasmid. Another plasmid was constructed under the control of the CMV promoter, as a standard promoter [51], to make an appropriate comparison between the function of the VEGF promoter and the standard CMV promoter. The results of bystander effect test and real-time PCR showed that the VEGF promoter function can act close to the CMV promoter of the standard pEGFP-N1 plasmid. The mRNA content of the CD gene produced by VEGF-CD - pEGFP-N1 in the 4T1 cells was 70% of mRNA content of the CD gene under control of CMV promoter (CD - pEGFP-N1) in the 4T1 cells.

In summary, it demonstrated that the G4-PAMAM dendrimer-mediated suicide gene CD/5-FC system can inhibit 4T1 cell proliferation, and exerted a 'bystander effect'. This approach may be providing a novel framework for suicide gene therapy to treat breast cancer and or other kinds of cancers (especially gliomas). 4T1-specific delivery by G4-PAMAM-dendrimer-mediated suicide CD gene with the addition of specific promoters upstream of the suicide gene may be provide a therapeutic potential for treatment of breast cancer in future.

## Conclusion

In this study we have attempted to improve the antitumor efficacy in 4T1 metastasizing breast carcinoma mice by using expressional shuttle vector to transfer *cytosine deaminase* gene under control of VEGF promoter in combination with non-viral PAMAM dendrimer, followed by 5-FC treatment. The results show inducing apoptosis in 4T1-transfected cell and a marked inhibition of tumor cell growth, suggesting CD gene transfer driven by VEGF promoter plus a safe non-viral vector followed by 5-FC treatment, may be an alternative therapeutic approach for the breast cancer treatment. Hence; Future studies in animals and clinical trials remain to follow the *in vivo* safety and efficiency of this therapeutic platform.

## Authors' Contributions

M. E. and A.A. contributed to methodology, conceptualization, software, and writing of original draft; T. S contributed supervision; A.S. and A.B contributed project administration, supervision, review, and editing.

## Informed Consent

All authors have read the manuscript and approved it for submission to your journal. All authors have agreed to authorship and order of authorship for this manuscript. Additionally, all authors have the appropriate permissions and rights to the reported data. Authors declare that they have no competing interests.

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## Conflict of Interest

There is no conflict of interest.

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