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Development of Nordihydroguaiaretic Acid Derivatives as Potential Multidrug-Resistant Selective Agents for Cancer Treatment

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In this research, we designed and synthesized a new series of nordihydroguaiaretic acid (NDGA) derivatives for multidrug resistance (MDR) research. A methylsulfonyl NDGA derivative, ((2*R*,3*S*)-2,3-dimethylbutane-1,4-diyl)bis(benzene-4,1,2-triyl) tetramethanesulfonate (**5d**), was found to inhibit MDR1 gene expression and suppress drug resistant MES-SA/Dx5 cells. Moreover, the combination of **5d** and doxorubicin/terameprocol (M4N) showed a profound synergistic effect on inhibition of the drug resistant cancer cells, suggesting that **5d** is a potential adjuvant applied with doxorubicin or terameprocol in cancer treatment.

Cancer is a major health problem worldwide that claims millions of lives annually. Although there have been great advances in the diagnosis and treatment of cancer, many people still succumb to cancer death because the chemotherapeutics adopted do not respond well to the spreading cancer. A major reason for the resistance of cancer cells to anticancer drugs is multidrug resistance (MDR), which is an intrinsic survival program deploying a variety of mechanisms that desensitize cancer cells to anticancer drugs, and increase the endurance of malignancies to therapeutic cytotoxic agents. Of these mechanisms, the most commonly encountered is mediated by ATP-binding cassette (ABC) transporters, which extrude a broad range of hydrophobic cytotoxic drugs that have already invaded the tumor cell.¹ P-glycoprotein (P-gp) [ABCB1/MDR1] is an important member of the ABC transporter family, and acts as a vacuum cleaner of the plasma membrane to expel hundreds of chemically unrelated toxins, preventing the cellular entry of xenobiotics and keeping the concentration of cytotoxic drugs below a cytotoxic threshold.²⁻³ Overexpression of P-gp has been found in various drug resistant tumor cells, and is a notorious wire-puller that causes nearly 90% of treatment failure.³ Therefore, P-gp has become a well-known and feasible drug

target for moderating the drug resistance of cancer cells.⁴

In this research, we have designed and synthesized a new series of nordihydroguaiaretic acid (NDGA **1**, Figure 1) derivatives for MDR research. NDGA is a lignin isolated from creosote bush (*Larrea tridentate*), which has been used as an herbal medicine by ethnic groups in America for the treatment of many diseases for a long time.⁵ Known as Masoprocol, NDGA was approved in the United States as an antineoplastic drug of topical treatment for actinic keratosis.⁶

In terms of biological activities, NDGA has very important and well-known antioxidant properties, such as being a scavenger of reactive oxygen species (ROS) and an inhibitor of lipoxygenase (LOX). Interestingly in non-tumor cells, NDGA displays cytoprotective effects, which are related to modulating the

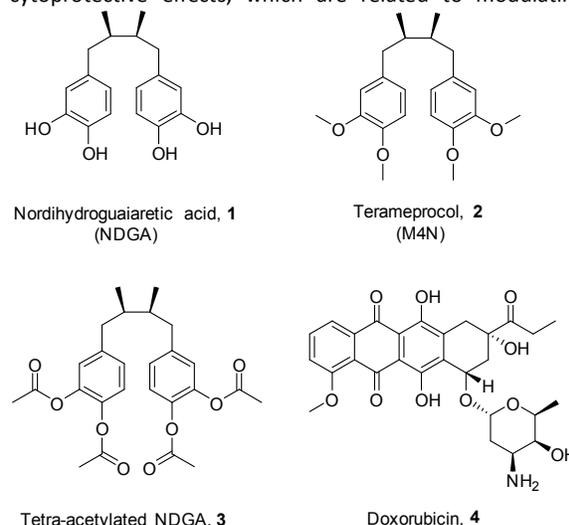


Figure 1. Nordihydroguaiaretic acid (NDGA) derivatives and chemotherapeutic agents.

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Nrf2/antioxidant response element (ARE) antioxidant pathway and preventing mitochondrial damage. In the case of tumor cells, NDGA displays pro-apoptotic activity and anti-tumor effects.⁷ In the case of tumor cells, NDGA displays pro-apoptotic activity and anti-tumor effects.⁸ Also, NDGA prevents Alzheimer's pathology by interfering with the amyloid- β aggregation pathway.⁹ Recently, NDGA was found to have the capability to extend murine lifespan,¹⁰ inhibiting replication of dengue virus,¹¹ and reducing secondary damage after spinal cord injury in rats via anti-inflammatory effects.¹²

The derivatives of NDGA, especially *O*-methylated NDGA, terameprocol (**2**, Figure 1), have antitumor activity based on the selective inhibition of proteins regulating specificity protein 1 (Sp1), including cyclin-dependent kinase 1, survivin and VEGF. Using this mechanism, terameprocol potentially inhibits the cell cycle, triggers apoptosis, and decreases angiogenesis.¹² Furthermore, terameprocol was found to have the capability to circumvent the drug resistance of the tumor cells in a xenograft model, and concomitantly strengthen the toxicity of doxorubicin and paclitaxel to the drug resistant cancer cells in combination treatment.¹⁴ It is also reported that tetra-acetylated NDGA (**3**, Figure 1) enhanced the toxicity of doxorubicin and several cytotoxic agents against drug resistant cancer cells as an adjuvant agent.¹⁵ In the case of uterine sarcoma, a major barrier in improving treatment outcomes is the intrinsic or acquired resistance to cytotoxic drugs.¹⁶ The major mechanisms of drug resistance in uterine sarcoma involve overexpression of P-gp, which is encoded by the MDR1 gene.¹⁷ Our approach to overcoming MDR1/P-gp-mediated MDR is unique, as we adopted NDGA, which itself is a potent anticancer agent, as the core scaffold.

In this research, we designed and synthesized a new series of NDGA derivatives for the inhibition of MDR gene expression in cancer cells. Several studies on NDGA structure modification have been carried out over the past few decades.¹⁷⁻²⁵ The central linker (7, 7', 8, 8', 9, and 9' positions of NDGA, Figure 2) of potential NDGA derivatives tend to be hydrophobic, and adding any polar functional group would diminish the bioactivities. Stereoisomers of NDGA (8 and 8' positions, Figure 2) were shown to give almost the same level of activity.²⁴ The phenyl group of NDGA has undergone modification of several functional

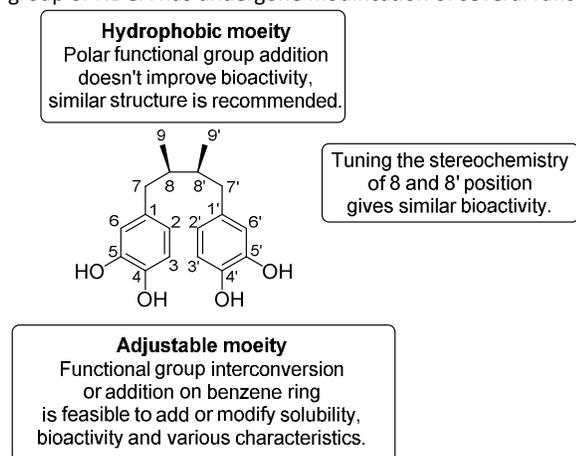
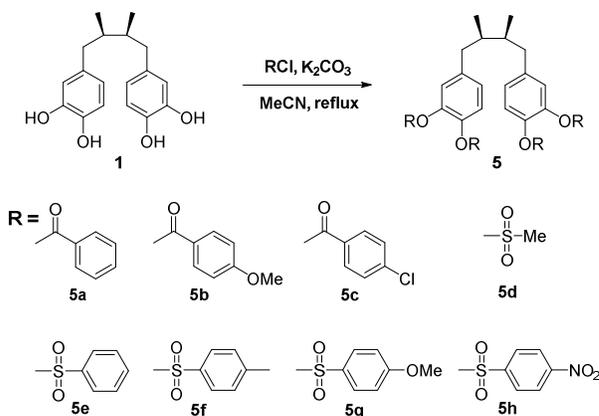


Figure 2. Synthetic strategy for modifying NDGA.



Scheme 1. Synthesis of benzoyl and sulfonyl NDGA derivatives.

Table 1. The lipophilicity and water solubility of synthesized NDGA derivatives.

Compound	clogP ^a	clogS ^a
1	3.44	-4.35
2	5.77	-6.27
5a	8.30	-7.53
5b	7.91	-7.29
5c	9.24	-8.25
5d	1.17	-4.57
5e	3.54	-6.23
5f	3.90	-6.46
5g	3.84	-6.11
5h	3.97	-7.45

^a clogP and clogS values were calculated using AlogPS 2.1 software. (<http://www.vcclab.org/lab/alogps/>)

groups, which have led to compounds with potent anti-cancer activity, *i.e.*, terameprocol. Here, we attached various sulfonyl and benzoyl groups to the hydroxyl terminal of NDGA with corresponding sulfonyl and benzoyl chlorides (Scheme 1), to increase water solubility and lipid solubility, respectively.

The synthetic route of the new NDGA derivatives is shown in Scheme 1. NDGA (**1**) was treated with various benzoyl and sulfonyl chlorides in the presence of potassium carbonate in acetonitrile at 80–85°C. After cooling down the reaction mass to room temperature, the inorganic residue was filtered off, and the filtrate was concentrated under reduced pressure to afford a residue. The residue was extracted by ethyl acetate, and washed with brine followed by purification by column chromatography or MPLC to give the yield of 67–78%. All these products were purified with 95% purity by chromatography and recrystallization was conducted for bioactivity assays.

The lipophilicity and water solubility of synthesized compounds were evaluated with the ALOGPS 2.1 program, developed by the virtual computational chemistry laboratory.²⁶ Calculated *n*-octanol/water partition coefficients (clogP) are shown in Table 1. NDGA and terameprocol clogP is 3.44 and 5.77 respectively, and both of them are current antineoplastic agents. The benzoyl derivatives of NDGA, **5a**, **5b**, and **5c** were found to be highly lipophilic with clogP value of 7.91–9.24. On the other

hand, phenylsulfonyl derivatives **5e–5h** were found to be less lipophilic than the benzoyl derivatives with a clogP value of 3.54–3.97. These results show that the sulfonyl group possess higher hydrophilicity than the benzoyl groups and the methoxy group of terameprocol. Whereas **5d** had a clogP of 1.17, indicating the methylsulfonyl derivative is much more hydrophilic than other synthesized NDGA derivatives, terameprocol and NDGA. The result of clogS (Table 1) shows that the sulfonyl derivatives have better water solubility, especially **5d**, which has a clogS of -4.57 , a higher level of water solubility than terameprocol.

To determine the biological activity of synthesized compounds, we applied MTT assay to evaluate the cytotoxicity of synthesized NDGA derivatives towards ovarian carcinoma cells, OVCAR-3 and SKOV3 (supplementary data, Figure S1-S4). After treatment for 48 and 72 hours, no significant toxicity was observed in either cancer cell line. However, terameprocol showed a certain level of suppression of tumor growth, and the cell viability was decreased to below 60% after 72 hours. This preliminary result suggested that sulfonyl and benzoyl NDGA derivatives did not have good anticancer potency.

When we investigated the toxic concentration (TC_{50}) of the test compounds on MDR cell lines; however, we found a different outcome compared to the cytotoxicity result of the tumor cells described above. Previously, we reported that

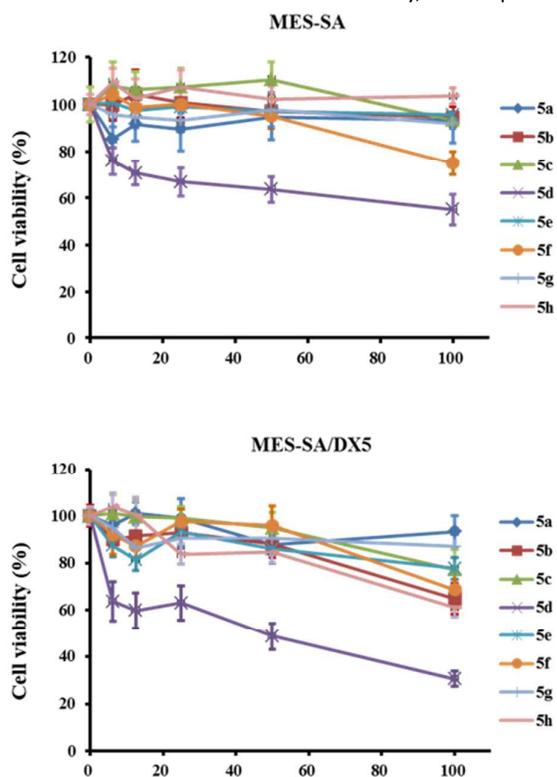


Figure 3. Dose-dependent inhibition of human uterine sarcoma cell line (MES-SA) and multidrug resistant uterine sarcoma cell line (MES-SA/Dx5) by **5a–5h** in culture. The data are expressed as the mean and the standard deviation of the mean ($n = 4$).

Table 2. Effect of **5a–5h** on cytotoxicity of MES-SA and MES-SA/Dx5 cells.

Compound	TC_{50} (μ M)	
	MES-SA	MES-SA/Dx5
5a	>100	>100
5b	>100	>100
5c	>100	>100
5d	>100	44.38
5e	>100	>100
5f	>100	>100
5g	>100	>100
5h	>100	>100

terameprocol²⁷ effectively inhibited the growth of drug resistant human cancer cell lines ovarian carcinoma NCI/ADR-RES in culture, with $TC_{50} \leq 10 \mu$ M.¹⁴ In this study, we further examined the effect of synthesized NDGA derivatives **5a–5h**, on human uterine sarcoma (MES-SA, P-gp deficient) and the drug resistant uterine sarcoma cell lines (MES-SA/Dx5, P-gp proficient) by following previous reported protocols.²⁸ Table 2 shows the cytotoxicity of synthesized NDGA derivatives on MES-SA and MES/Dx5 cell lines. The results demonstrated that **5a**, **5b**, **5c**, **5e**, **5f**, **5g**, and **5h** show low inhibition activity to human tumor cells. However, the methylsulfonyl derivative **5d** was able to inhibit the growth of drug resistant uterine sarcoma MES-SA/Dx5 cells in culture (Figure 3 and Table 2), indicating that **5d** suppresses the growth of P-gp overexpressing cancer cells selectively to a certain extent.

The effect of NDGA derivative **5d** on MDR1 promoter activity was further examined by luciferase reporter assay (Figure 4A). The MDR1 promoter gene modified vector containing luciferase reporter gene, pMetLuc-MDR1, was transfected into drug resistant MES-SA/DX5 cells. The control counterpart luciferase containing vector without MDR1 promoter gene, pMetLu, was also transfected into MES-SA/DX5 cells. Two kinds of transfected cell were cultured with and without **5d** treatment for 48 hours. MDR1 promoter activity was reported by the luciferase reporter system, depicted in Figure 4A, indicating that **5d** has the capability to abolish MDR1 promoter activity, and further obstruct P-gp expression. According to results of luciferase activity, we further examined the effect of **5d** on MDR1 gene expression. To examine this possibility, MES-SA/Dx5 cells were exposed to 0, 5, 10, and 20 μ M **5d** for 2-day, after which total protein were examined for levels of P-gp. After treatment with 20 μ M **5d**, the amount of P-gp was reduced with its abundance decreasing to 31.2% of the control amount after a 2-day exposure to 20 μ M **5d** (Figure. 4B). Even a 2-day exposure to 5 μ M **5d** resulted in a 12.8% reduction in P-gp. The levels of P-gp were normalized to β -actin.

A combination of chemotherapeutics and **5d** has a more profound effect on MDR1 promoter activity than single drug treatment. The pMetLuc-MDR1 transfected MES-SA/DX5 cells were treated with various combinations of doxorubicin, terameprocol (M4N) and **5d**, as demonstrated in Figure 5. After incubation for 48 hours, cells treated only with doxorubicin showed full activity of MDR1 promoter, whereas cells treated with terameprocol alone showed apparent lower level activity of

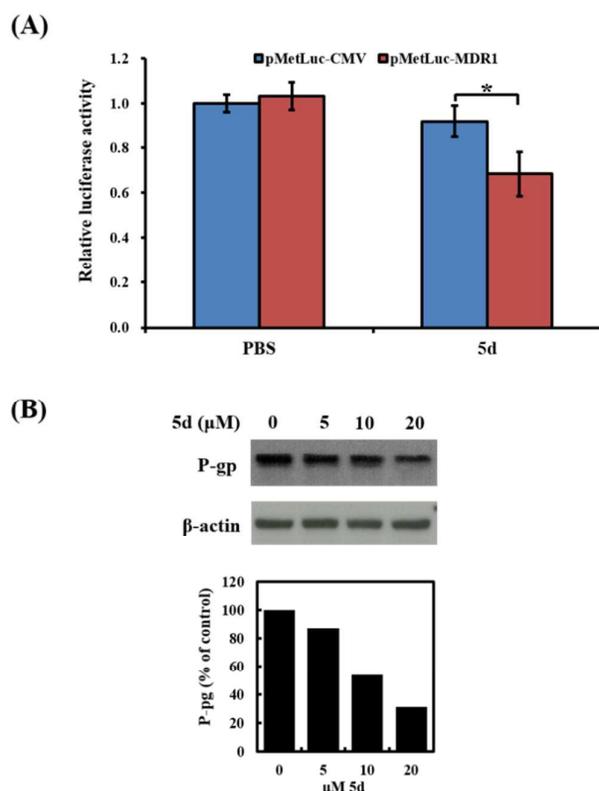


Figure 4. Effect of compound **5d** on MDR1 promoter activity and P-glycoprotein levels (P-gp) in MES-SA/Dx5 cells. **(A)** Luciferase activity in MES-SA/Dx5 stable transfected with MDR1 promoter (pMetLuc-MDR1), or control vector (pMetLuc-CMV) after treatment with compound **5d**. *Significant difference ($p < 0.05$) for the comparison is indicated. Data are expressed as the mean \pm SEM. **(B)** Western blot analysis of P-gp and β -actin (normalization control) protein levels in cells treated for 3 days with 0, 5, 10, and 20 μM compound **5d**. Results in bar graph form normalized to β -actin.

MDR1 promoter, which was indicated by nearly 50% of luciferase activity. Cells treated with **5d** showed 70% luciferase activity, higher than the cells treated only with terameprocol; however, when cells were treated with the combination of **5d** with doxorubicin or terameprocol, the activity of luciferase was suppressed to nearly 20%, revealing that **5d** intensifies attenuation of MDR1 promoter. Moreover, to confirm **5d** and combination treatments on MDR1 promoter activity in Figure 5, we measured viability of MES-SA/Dx5 cells incubated with the **5d**, Dox, M_4N and their combination for 2-day respectively by MTT assay. After incubation for 2-day, cells treated only with doxorubicin showed almost 100% viability, whereas cells treated with M_4N and **5d** alone showed 75% to 80% viability. However, when cells were treated with the combination of **5d** with doxorubicin or M_4N , the cell viability was suppressed to 50–60% (Figure 6). According to these results, formulae of **5d** combined with doxorubicin or terameprocol have a synergistic effect to control MDR1 promoter activity. Moreover, these results also

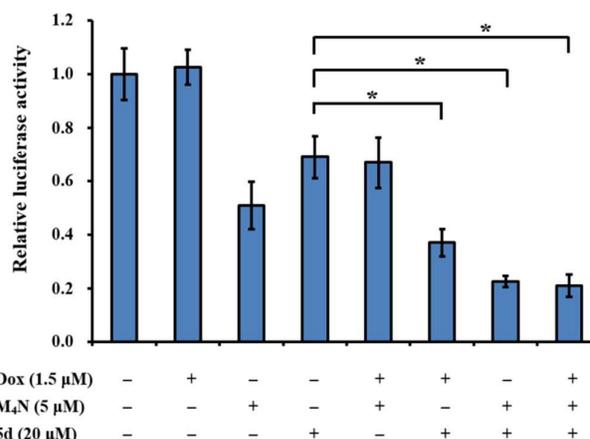


Figure 5. Inhibition of MDR1 promoter activity in MES-SA/DX5 treated with doxorubicin (Dox), M_4N (**2**), **5d** and combination formula. Luciferase activity was used to measure promoter response of in MES-SA/DX5 stable transfected with MDR1 promoter reporter vector (PMetLuc-MDR1). Relative luciferase activity are expressed as mean \pm SEM. Student's t-test was used to compare the difference between **5d** alone and each combination treated group, and $P < 0.05$ (*) are shown to be statistically significant.

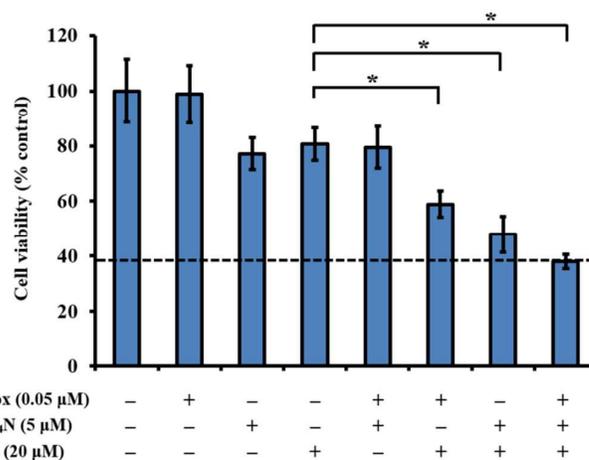


Figure 6. Effect of doxorubicin (Dox), M_4N (**2**) and **5d** and their combination on the cell viability of MES-SA/DX5 drug resistant cell lines. The number of viable cells after 72 hours of treatment with Dox, M_4N (**2**) and **5d** alone and in combination. The horizontal line in graph represents the number of viable at the initiation of drugs exposure. Cell viability are expressed as mean \pm SEM. Student's t-test was used to compare the difference between **5d** alone and each combination of treatment groups, and $P < 0.05$ (*) are shown. $P < 0.05$ (*) are considered to be statistically significant.

indicated a combination of chemotherapeutics and **5d** has a more extreme effect on multi-drugs resistant cancer cells than single drug treatment.

Methylsulfonyl functionalized NDGA, **5d**, has better water solubility than commercial NDGA-type antineoplastic agents **1** and **2**, and much more hydrophilic than other synthesized NDGA derivatives in this research, as described in Table 1. With more hydrophilic clogP value, **5d** itself has certain suppression effect on doxorubicin resistant uterine sarcoma cells, MES-SA/DX5. Having more than 100 μM of TC_{50} value in normal uterine sarcoma cells (MES-SA), **5d** has 44.38 μM of TC_{50} in drug resistant counterpart (Table 2); although **5d** has similar viability in both cell lines, the suppression of MES-SA/DX5 is more obvious with increasing amount of **5d** (Figure 3). Such a mild toxicity might make **5d** a suitable adjuvant agent in cancer treatment. **5d** could suppress MDR1 promoter to a certain extent, and further interrupt the expression of P-gp, one of the cruxes that caused multidrug-resistant cancer (Figure 4). Moreover, combining the chemotherapeutics Dox **4** and M_4N **2** with **5d** shows a remarkable synergistic effect on inhibition of MDR1 promoter activity, which is much better than single-agent treatment with Dox and M_4N (Figure 5). This shows that **5d** could prevent the early development of MDR in cancer treatment, i.e., **5d** inhibits doxorubicin mediated induction of MDR1 gene expression. Furthermore, combination of **5d** with doxorubicin and/or terameprocol (M_4N , **2**) has much improved suppression effect on the growth of MES-SA/DX5 cells, especially compared with the single-drug treatments with Dox or M_4N , i.e., **5d** has a good synergistic effect on inhibition of dox-resistant uterine sarcoma cell growth drug-resistant (Figure 6). Also, good water solubility allows **5d** to work very well with water-soluble doxorubicin and M_4N ; the consumption of doxorubicin would decrease massively with **5d** as an adjuvant agent in regular cancer treatment.

Conclusions

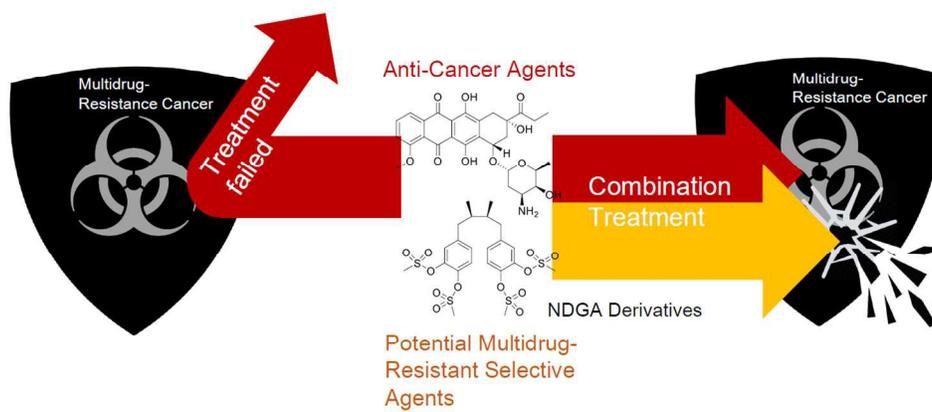
In conclusion, we synthesized 8 new NDGA derivatives and evaluated their potential as multidrug resistance selective agents for cancer treatment. We found one of the derivatives, the methylsulfonyl functionalized NDGA derivative, **5d**, can inhibit MDR1 gene expression, decrease the P-gp protein formation, and then suppress drug resistant MES-SA/Dx5 cells (resistant uterine sarcoma cells). Moreover, the combination of **5d** and doxorubicin/terameprocol (M_4N) has much better efficacy of suppressing the drug-resistant cancer cells than single-agent treatment. From the clinical point of view, this good synergistic effect shows that **5d** has great potential adjuvant drug with doxorubicin or terameprocol for cancer treatment.

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