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## Induction of Mitochondrial Pathways and Endoplasmic Reticulum Stress for Increasing Apoptosis in Ectopic and Orthotopic Neuroblastoma Xenografts

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### Abstract

Cancers are characterized by dysregulation of multiple signaling pathways and thus monotherapies are hardly effective. Neuroblastoma, which often occurs in adrenal glands, is the most common childhood malignancy. Malignant neuroblastoma resists traditional treatments and further studies are needed for effective therapeutic interventions. We evaluated synergistic efficacy of N-(4-hydroxyphenyl) retinamide (4-HPR) and genistein (GST) for induction of apoptosis in human malignant neuroblastoma SH-SY5Y and SK-N-BE2 cells in culture and activation of multiple pathways for increasing apoptosis in ectopic and orthotopic neuroblastoma xenografts in nude mice. Combination of 4-HPR and GST synergistically reduced cell viability, caused subG1 accumulation, increased caspase-3 activity for apoptosis *in vitro* and reduced tumor growth *in vivo*. Western blotting indicated that combination therapy down regulated Id2 to induce differentiation, increased pro-apoptotic Bax and decreased anti-apoptotic Bcl-2 leading to an increase in Bax:Bcl-2 ratio, increased mitochondrial Bax level, caused mitochondrial release of Smac/Diablo, down regulation of the baculovirus inhibitor-of-apoptosis repeat containing (BIRC) proteins such as BIRC-2 and BIRC-3, and activation of calpain and caspase-3 in SH-SY5Y xenografts. Accumulation of apoptosis-inducing-factor (AIF) in cytosol and increase in caspase-4 activation suggested involvement of mitochondrial pathway and endoplasmic reticulum (ER) stress, respectively, for apoptosis in SH-SY5Y xenografts. *In situ* immunofluorescent labelings of SH-SY5Y and SK-N-BE2 xenograft sections showed overexpression of calpain, caspase-12, and caspase-3, and AIF, suggesting induction of mitochondrial caspase-dependent and caspase-independent pathways for apoptosis. Collectively, synergistic effects of 4-HPR and GST induced mitochondrial pathways and also ER stress for increasing apoptosis in ectopic and orthotopic neuroblastoma xenografts in nude mice.

### Keywords

apoptosis; endoplasmic reticulum stress; genistein; N-(4-hydroxyphenyl) retinamide; neuroblastoma

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## 1. Introduction

Retinoid controls a wide range of biological phenomena including differentiation, proliferation, and apoptosis in a variety of cancers [1]. Fenretinide or N-(4-hydroxyphenyl) retinamide (4-HPR), a synthetic retinoid, has shown the most promising broad spectrum anti-cancer efficacy against a variety of *in vitro* and *in vivo* animal studies [2], and chemopreventive clinical trials [3]. Notably, 4-HPR showed low toxicity and anti-tumor efficacy even in all-*trans* retinoic acid (ATRA) or 13-*cis* retinoic acid (13-CRA) resistant cancer cells; and several *in vitro* studies reported that dose-dependent anti-tumor efficacy of 4-HPR was mainly due to growth arrest and apoptosis in neuroblastoma [3, 4]. Treatment of neuroblastoma patients with high dose of 4-HPR accumulated a plasma 4-HPR concentration that was able to induce apoptosis in neuroblastoma cells [5]. Interestingly, 4-HPR in comparatively low concentrations induces cellular differentiation due to differential expression of a variety of key proteins [6]. Thus, therapeutic potential of low dose of 4-HPR in combination with other therapeutic agent needs to be explored for induction of apoptosis in various cancers including neuroblastoma.

Development of therapeutic agents from dietary phytochemicals is very promising to fight various human cancers [7]. Genistein (GST), an isoflavonoid primarily derived from soybean, has been reported to inhibit the growth of various cancer cells through altering cell signaling pathways, cell cycle, and apoptosis [8]. GST induced apoptosis through endoplasmic reticulum stress and mitochondrial damage in human hepatoma Hep3B cells [9] and Ca<sup>2+</sup>-mediated calpain/caspase-12-dependent apoptosis in breast cancer MCF-7 cells [10]. GST potentially inhibited cellular proliferation in 5 neuroblastoma N2A, JC, SK-N-SH, MSN, and Lan5 cell lines through induction of apoptosis [11]. We previously reported that GST induced activation of calpain and caspases for apoptosis in human neuroblastoma SH-SY5Y cells [12]. These studies suggest that GST could be a strong anti-cancer agent for treating neuroblastoma *in vivo* as well.

Neuroblastoma is the most frequent extracranial childhood solid tumor derived from sympathetic nervous system mostly affecting adrenal gland and it also usually metastasizes in other body parts including chest, neck, lymph nodes, pelvis, liver, and bone [13, 14]. The current therapy for this childhood malignancy comprises surgery, radiation, and chemotherapy; but in most cases of neuroblastoma, therapeutic inefficacy leading to poor clinical outcome may render in excess of 15% of all pediatric cancer related deaths in children [15, 16]. Thus, innovative therapeutic approach is urgently warranted for successful treatment of neuroblastoma. Neuroblastoma shows a complex clinical as well as biological heterogeneity [13]. In this investigation, we explored the efficacy of the combination of 4-HPR (for induction of differentiation) and GST (for induction of apoptosis) in human neuroblastoma cells and in ectopic SH-SY5Y and orthotopic SK-N-BE2 xenografts in athymic nude mice. Our data showed that combination of 4-HPR and GST caused more anti-tumor efficacy than monotherapy and they worked synergistically to activate multiple molecular mechanisms for increasing apoptosis in human neuroblastoma cells and in pre-clinical ectopic SH-SY5Y and orthotopic SK-N-BE2 xenografts in athymic nude mice.

## 2. Materials and Methods

### 2.1. Human Neuroblastoma Cell Lines and Culture Conditions

Human neuroblastoma SH-SY5Y and SK-N-BE2 cell lines were purchased from the American Type Culture Collection (ATCC, Manassas, VA). Cells were grown in 75-cm<sup>2</sup> flasks containing 10 ml of 1xRPMI 1640 supplemented with 10% fetal bovine serum (FBS) and 1% penicillin and 1% streptomycin in a fully-humidified incubator containing 5% CO<sub>2</sub>

at 37°C. Cell lines were serially passaged following trypsinization using a trypsin/EDTA solution.

## 2.2. [3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide] (MTT) Assay

The MTT assay was used to determine the sensitivities of human neuroblastoma SK-N-BE2 and SH-SY5Y cell lines to 4-HPR (0.25, 0.5, and 1 µM for 72 h) and GST (50, 100, and 200 µM for 24 h) alone or in combination. Cell viability data were analyzed on the basis of the multiple drug-effect equation (classical isobologram) using the Compusyn software (ComboSyn, Paramus, NJ). The effect (synergistic, additive, or antagonistic) of the combination of drugs (4-HPR and GST) was expressed as the combination index (CI), where  $CI < 1$  indicated synergism,  $CI = 1$  indicated additive effect, and  $CI > 1$  indicated antagonism [17].

## 2.3. Flow Cytometric Analysis of Cell Cycle

Following treatment, cells were collected by trypsinization and centrifugation, then washed with PBS, and fixed with 70% ethanol. Cells were labeled with propidium iodide (PI) solution (0.05 mg/ml PI, 2 mg/ml RNase A, 0.01% Triton X-100 in PBS) and incubated for 30 min at room temperature in darkness. Cell cycle (DNA content) was analyzed using an Epics XL-MCL Flow Cytometer (Beckman Coulter, Fullerton, CA) [17].

## 2.4. Colorimetric Assay for Determination of Caspase-3 Activity

Caspase-3 activity in the lysates of cells or xenografts was measured using the commercially available caspase-3 colorimetric assay kit (Sigma, St. Louis, MO).

## 2.5. Annexin V-FITC/PI Staining and Flow Cytometry for Determination of Apoptosis

Cells from single and combination treatments were harvested and processed with an Annexin V-FITC/PI staining kit (BD Biosciences, San Jose, CA) for flow cytometric analysis for quantitative determination of apoptosis using an Epics XL-MCL Flow Cytometer (Beckman Coulter, Fullerton, CA, USA). Further, we used 100 µM z-DEVD-fmk (caspase-3-specific inhibitor) to determine the involvement of caspase-3 in apoptosis in cells following combination therapy [17].

## 2.6. Human Neuroblastoma Ectopic and Orthotopic Xenografts in Athymic Nude Mice

Six-week-old female athymic nude (nu/nu) mice were obtained from Charles River Laboratories (Wilmington, MA). All animal studies were conducted following guidelines of the NIH Animal Advisory Committee and also approved by our Institutional Animal Care and Use Committee (IACUC). For developing ectopic human neuroblastoma xenografts, SH-SY5Y cells ( $6 \times 10^6$ ) in 100 µl of (1:1) mixture with Matrigel (BD Biosciences, San Jose, CA) were implanted by subcutaneous (s.c.) injection on the flank of each mouse under isoflurane anesthesia condition. Palpable xenografts were developed within 6–8 days. Tumors were measured using an external caliper and volume was calculated by the formula:  $4\pi/3 \times (\text{length}/2) \times (\text{width}/2)^2$ . For developing orthotopic human neuroblastoma xenografts, nude mice were anaesthetized using ketamine (50 mg/kg) and xylazine (35 mg/kg) intraperitoneally (i.p.). After a midline incision, a total of  $6 \times 10^5$  SK-N-BE2 neuroblastoma cells in 15 µl of 1xRPMI 1640 were injected in the capsule of the left adrenal gland using a 22 gauge needle attached to a Hamilton syringe. After implantation of cells, abdominal wall and skin were closed with a 5-0 LOOK® suture (Surgical Specialties, Reading, PA) and antiseptic-analgesic ointments were applied and animals were kept in a pathogen-free animal facility [18]. Animals with 3 weeks-old neuroblastoma xenografts were randomly divided into four groups: control (CTL), 4-HPR, GST, and 4-HPR + GST. Animals in control group did not receive any therapy. Each animal in other group received i.p. a daily dose of 4-HPR

(20 µg/kg/day), GST (2 mg/kg/day), or 4-HPR (20 µg/kg/day) + 4 h later GST (2 mg/kg/day) for 8 days.

### 2.7. Histological Examination of Tumor Xenografts

After completion of treatment schedule, mice were sacrificed, both ectopic and orthotopic neuroblastoma xenografts were surgically collected. After washing with PBS, pH 7.4, tumors were either immediately frozen in liquid nitrogen and stored at  $-80^{\circ}\text{C}$  for future use or immediately frozen ( $-80^{\circ}\text{C}$ ) in Optima Cutting Temperature media (Fisher Scientific, Suwanee, GA) and 5 µm sections were cut with a cryostat. These sections were stained for routine histological examinations [19].

### 2.8. *In Situ* Immunofluorescent Labeling for Detection of Pro-apoptotic Protein Expression

For *in situ* immunofluorescent labelings, samples were fixed to slides using 95% (v/v) ethanol for 10 min and rinsed twice for 10 min each in PBS, pH 7.4. Fixation and all subsequent steps were conducted at room temperature. Sections were blocked with 2% (v/v) horse and goat sera (Sigma, St. Louis, MO) for 30 min. Samples were probed with (1:100) primary IgG antibody for 1 h and slides were rinsed twice in PBS, pH 7.4, for 5 min each followed by incubation with (1:75) secondary Texas Red-conjugated anti-rabbit IgG antibody (Vector Laboratories, Burlingame, CA) for 30 min. Slides were then rinsed twice in PBS, pH7.4, for 5 min and in distilled water for 5 min. The slides were immediately mounted with VectaShield (Vector Laboratories) and viewed promptly under a fluorescence microscope at 200x magnification (Olympus, Japan), and digital pictures were taken with Image-Pro Plus software (Media Cybernetics, Silver Spring, MD), as we described recently [20].

### 2.9. Combined Terminal dUTP Nick-End Labeling (TUNEL) and Double Immunofluorescent Stainings for Detection of Apoptosis and Upregulation of a Pro-apoptotic Protein

Briefly, xenograft tissues were sectioned, fixed in 95% (v/v) ethanol for 10 min and then postfixed in 4% methanol-free formaldehyde for 10 min. The sections were saturated with TdT equilibration buffer (Promega, Madison, WI, USA) (50 µL/slide) for 5 min and then replaced with TUNEL reaction mixture (50 µL/slide) and were incubated for 1 h at  $37^{\circ}\text{C}$  in an OmniSlide Thermal Cycler (Hypaid, UK). Following the TUNEL reaction, slides were immuno-stained with primary IgG antibody for specific protein and then incubated with fluoresceinated secondary antibodies and images were taken [20].

### 2.10. Western Blotting and Detection of Specific Protein Bands

Western blotting was used for analyzing specific proteins [20]. Briefly, total protein samples from tumor tissue were extracted, measured spectrophotometrically and denatured. Samples were loaded onto the SDS-polyacrylamide gradient (4–20%) gels (Bio-Rad, Hercules, CA), resolved by electrophoresis, and electroblotted to membranes. The blots were incubated with a primary IgG antibody followed by incubation with an alkaline horseradish peroxidase (HRP)-conjugated secondary IgG antibody. Specific protein bands on the blots were detected by HRP/ $\text{H}_2\text{O}_2$  catalyzed oxidation of luminol in alkaline condition using enhanced chemiluminescence (ECL) system (GE Healthcare Bio-Sciences, Piscataway, NJ) followed by autoradiography. Autoradiograms were scanned using Photoshop software (Adobe Systems, Seattle, WA), and OD of each band was determined using NIH Image software.

### 2.11. Colorimetric Assay for Determination of Caspase-4 Activity

Caspase-4 activity in tumor lysates was measured using the commercially available caspase-4 activity colorimetric assay kit (BioVision, Mountain View, CA).

## 2.12. Statistical Analysis

Data were analyzed using the StatView software (Abacus Concepts, Berkeley, CA), expressed as arbitrary units  $\pm$  standard error of mean (SEM) of separate experiments ( $n \geq 3$ ), and compared by one-way analysis of variance (ANOVA) followed by the Fisher post hoc test. A significant difference between control and treatment was indicated by \* ( $p < 0.05$ ) or \*\* ( $p < 0.001$ ).

## 3. Results

### 3.1. Changes in Cell Viability

We used human malignant neuroblastoma SK-N-BE2 and SH-SY5Y cells in cultures to determine the effects of treatments (Figure 1). Changes in cell viability were determined by the MTT assay (Figure 1A). We used 4-HPR or GST alone as monotherapy and also 4-HPR + GST as combination therapy. Notably, 4-HPR (0.5  $\mu\text{M}$  for 72 h) + GST (100  $\mu\text{M}$  for 24 h) and 4-HPR (1  $\mu\text{M}$  for 72 h) + GST (200  $\mu\text{M}$  for 24 h) caused most significant decrease in cell viability when compared with CTL and other treatment groups (Figure 1A). Data were analyzed to generate the combination index (CI) of the drugs. The effect of 4-HPR + GST was found to produce  $\text{CI} < 1$  in neuroblastoma SK-N-BE2 ( $\text{CI} = 0.78$ ) and SH-SY5Y ( $\text{CI} = 0.95$ ) cells, indicating synergistic action of the combination of drugs. Finally, 4-HPR (0.5  $\mu\text{M}$  for 72 h), GST (100  $\mu\text{M}$  for 24 h), and 4-HPR (0.50  $\mu\text{M}$  for 72 h) + GST (100  $\mu\text{M}$  for 24 h) were selected as effective treatments and used in all subsequent cell culture studies.

### 3.2. Cell Cycle Analysis

We found treatments (4-HPR, GST, and 4-HPR + GST) significantly increased apoptotic subG1 phase (10, 18, and 28%, respectively) in SK-N-BE2 cells (Figure 1B). In case of SH-SY5Y cells (Figure 1B), 4-HPR treatment showed non-significant change, while GST (35%) and 4-HPR + GST (51%) treatments exhibited significant increases in apoptotic subG1 phase, compared with untreated cells (Figure 1B).

### 3.3. Increase in Caspase-3 Activity

We determined the caspase-3 activity in apoptosis and amounts of apoptotic cells following treatments (Figure 2). Increases in caspase-3 activity in course of apoptosis in both neuroblastoma cell lines were measured by a colorimetric assay (Figure 2A). The caspase-3 activity in cell lysate was most significantly found in both cell lines after treatment with 4-HPR + GST, compared with untreated cells (Figure 2A).

### 3.4. Increase in Apoptosis was Associated with Upregulation of Caspase-3

Our Annexin V-FITC/PI staining and flow cytometric studies showed the percentages of Annexin V-positive/PI-negative populations (early apoptotic) in neuroblastoma SK-N-BE2 (CTL, 2%; 4-HPR, 8%; GST, 20%; and 4-HPR + GST, 32%) and SH-SY5Y (CTL, 2%; 4-HPR, 3%; GST, 32%; and 4-HPR + GST, 47%) cells (Figure 2B). Pre-treatment with 100  $\mu\text{M}$  z-DEVD-fmk (a caspase-3-specific inhibitor) significantly reduced the capability of combination therapy for induction of apoptosis to 5.28% in SK-N-BE2 cells and to 18.45% in SH-SY5Y cells, indicating involvement of caspase-3 in induction of apoptosis in both cell lines (Figure 2B).

### 3.5. Histological Evaluation of Efficacy of Combination Therapy

We treated ectopic SH-SY5Y xenografts in nude mice (Figure 3). As seen in the animals (Figure 3A), treatments caused tumor regression. Following treatments, the tumors were collected surgically to measure tumor volume (Figure 3B). Treatment with 4-HPR + GST caused more regression of SH-SY5Y tumors than either drug used alone. Further, we

performed hematoxylin and eosin (H&E) staining of tumor sections and histological examination at 200x magnification under the light microscope (Figure 3C), showing maximum cell death after combination therapy.

Similarly, the efficacy of combination therapy was evaluated in orthotopic SK-N-BE2 xenografts in nude mice (Figure 4). Following 4-HPR or GST monotherapy and 4-HPR + GST combination therapy of orthotopic SK-N-BE2 xenografts (Figure 4A), tumor sections from each group were subjected to H&E staining followed by histological examination under the light microscope at 200x magnification (Figure 4B). In orthotopic xenografts also, combination therapy caused maximum cell death.

Examination of sections of all neuroblastoma ectopic SH-SY5Y xenografts and orthotopic SK-N-BE2 xenografts after H&E staining revealed that untreated tumors maintained characteristic growth of neuroblastoma, 4-HPR alone inhibited tumor cell proliferation and caused neuronal differentiation, GST alone induced apoptosis, and 4-HPR + GST combination therapy dramatically increased the amounts of apoptosis.

### **3.6. Combination Therapy Down regulated Id2 and Increased Bax:Bcl-2 Ratio in SH-SY5Y Xenografts**

We performed Western blotting (Figure 5) and found that treatment with 4-HPR + GST down regulated inhibitor-of-differentiation 2 (Id2), and changed the levels of expression of total Bax and Bcl-2 (Figure 5A) resulting in significant increase in the Bax:Bcl-2 ratio, compared with CTL xenograft (Figure 5B). The elevation of Bax:Bcl-2 ratio could make cells fully committed to apoptosis due to mitochondrial release of several pro-apoptotic molecules. Further, we isolated the mitochondrial fractions to analyze the change in mitochondrial Bax level and monitored the COX-4 level as mitochondrial internal control (Figure 5C). We found that treatment with 4-HPR + GST caused more increase in mitochondrial Bax level in the SH-SY5Y xenografts than either therapy alone (Figure 5D).

### **3.7. Combination Therapy Caused Mitochondrial Release of Pro-apoptotic Molecules and Down regulation of BIRC Proteins for Activation of Cysteine Proteases in SH-SY5Y Xenografts**

Therapeutic treatment may cause mitochondrial release of pro-apoptotic molecules [21]. We performed Western blotting to examine mitochondrial release of Smac/Diablo and levels of BIRC proteins and cysteine proteases in SH-SY5Y xenografts (Figure 6). Compared with CTL or 4-HPR alone or GST alone, 4-HPR + GST highly induced mitochondrial release of pro-apoptotic protein 25 kDa Smac/Diablo into the cytosol (Figure 6A) and levels of the anti-apoptotic proteins 72 kDa BIRC-2 (cIAP-1), 68 kDa BIRC-3 (cIAP-2) in the cytosol (Figure 6A). We also found that combination therapy increased expression of 80 kDa calpain, 20 kDa caspase-3, and mitochondrial 67 kDa apoptosis-inducing factor (AIF) into the cytosol (Figure 6A). Further, we used Western blotting and colorimetric assay and found increased caspase-4 activation and total caspase-4 activity, respectively, indicating increase in ER stress, for apoptosis in the SH-SY5Y xenografts after treatment with 4-HPR + GST (Figure 6B).

### **3.8. Combination Therapy Caused Upregulation of Calpain and Caspase-12 for Apoptosis in Ectopic and Orthotopic Neuroblastoma Xenografts**

We performed the single immunofluorescent (SIF) and double immunofluorescent (DIF) stainings of the xenograft sections after the treatments (Figure 7). The SIF staining showed that compared with CTL or monotherapy, 4-HPR + GST combination therapy very significantly increased expression of calpain and caspase-12 in ectopic SH-SY5Y xenografts (Figure 7A) as well as in orthotopic SK-N-BE2 xenografts (Figure 7B). Subsequently, the

DIF staining showed highly remarkable overexpression of calpain and caspase-12 in association with increase in DNA fragmentation in ectopic SH-SY5Y xenografts (Figure 7A) and orthotopic SK-N-BE2 xenografts (Figure 7B) following 4-HPR + GST combination therapy.

### 3.9. Combination Therapy Induced Activation of Caspase-dependent and Caspase-independent Pathways for Apoptosis

To explore the downstream executioner steps, we examined the caspase-3 and AIF levels in all treatment groups from both SH-SY5Y and SK-N-BE2 xenografts (Figure 8). Following the SIF staining, we found significant increases in caspase-3 and AIF expression in ectopic SH-SY5Y xenografts (Figure 8A) as well as in orthotopic SK-N-BE2 xenografts (Figure 8B) due to 4-HPR + GST combination therapy. The DIF staining showed that significant increases in caspase-3 and AIF expressions were associated with DNA fragmentation in both ectopic and orthotopic xenografts following the 4-HPR + GST combination therapy (Figure 8B). These studies confirmed the involvement of caspase-dependent pathway along with non-caspase pathway for induction of apoptosis in both ectopic SH-SY5Y and orthotopic SK-N-BE2 xenografts after the 4-HPR + GST combination therapy.

## 4. Discussion

In this investigation, we found that 4-HPR + GST worked synergistically to significantly reduce the viability of neuroblastoma SH-SY5Y cells (N-Myc expression without N-myc gene amplification) and SK-N-BE2 cells (N-Myc overexpression with N-myc gene amplification) [22] and induce caspase-3 activity for apoptosis. Previous studies reported that 4-HPR [3, 23] or GST [12] induced apoptosis in a variety of cell lines including neuroblastoma [4, 12] with activation of caspase-3. The synergistic efficacy of 4-HPR + GST activated multiple pathways for increasing apoptosis, as suggested by molecular and histological observations, in two preclinical neuroblastoma animal models such as ectopic SH-SY5Y and orthotopic SK-N-BE2 xenografts.

We found that 4-HPR + GST caused down regulation of Id2, a biochemical marker well known to antagonize differentiation [24], indicating induction of molecular mechanism of differentiation in SH-SY5Y xenografts. The increases in Bax:Bcl-2 ratio and translocation of Bax to mitochondria could be the key events for apoptosis in neuroblastoma cells [12, 25] and also well correlated following combination therapy with 4-HPR and 1-threo-dihydrosphingosine (safingol) in SK-N-MC cells [26]. Similarly, GST induced apoptosis through alterations in Bax and Bcl-2 levels in breast [27] and head and neck squamous cell carcinoma cell lines [28]. Thus, it would be highly plausible that combination of 4-HPR and GST increased Bax:Bcl-2 ratio to trigger the mitochondrial release of pro-apoptotic molecules Smac/Diablo to down regulate the anti-apoptotic BIRC proteins and activate the downstream apoptotic pathways.

Levels of expression of BIRC-2 (cIAP-1) and BIRC-3 (cIAP-2) are often stimulated in cancers and cause suppression of caspase activation for inhibition of apoptosis [29] and can act as oncogenes leading to cell survival, transformation, and resistance to radiation and chemotherapies [30]. Smac/Diablo acts as an indirect activator of caspases by inhibition of the BIRC proteins [21] to facilitate the caspase-dependent pathway of apoptosis following treatment with combination of 4-HPR and GST in ectopic SH-SY5Y xenografts. Treatment with this combination therapy increased activation of the Ca<sup>2+</sup>-dependent cysteine protease calpain, which could play a significant role in cell death with alternations in expression of the Bcl-2 family members leading to an increase in Bax:Bcl-2 ratio [31] and Bax translocation to mitochondria for release of Smac/Diablo [32]. Calpain can also activate caspase-3 for cytoskeletal breakdown and apoptosis in cancers including neuroblastoma [12,

25]. Our finding is consistent with previous studies demonstrating that increases in calpain [33] and caspase-3 [34] activation play important role in induction of apoptosis in SH-SY5Y xenografts after treatment with 4-HPR + GST.

Loss of mitochondrial membrane integrity can release AIF to induce caspase-independent apoptosis [35] also we found following 4-HPR + GST therapy in ectopic SH-SY5Y xenografts. Calpain has been strongly associated with mitochondrial release of AIF and translocation to the nucleus [36] to cause caspase-independent nuclear DNA fragmentation. The increase in caspase-4 activation and activity following 4-HPR + GST combination therapy indicated involvement of ER stress in induction of apoptosis that could be due to 4-HPR [37] or GST [9] monotherapy or due to their combined effects. Recent studies demonstrate that 4-HPR induces the generation of reactive oxygen species (ROS) for mitochondrial damage leading to apoptosis in six neuroblastoma cell lines [38] and also ER stress induces cell death in neuroectodermal tumor cells [39]. Studies in oral squamous cell carcinoma (OSCC) cell line demonstrated that combination of cetuximab and genistein caused dose-dependent significant decrease in cell proliferation when compared with either drug alone [40]. Our observations suggested that 4-HPR + GST combination therapy triggered mitochondrial caspase-dependent and caspase-independent pathways and also ER stress for apoptosis in SH-SY5Y xenografts. In line with our observation, a recent investigation reported that combination of bortezomib and 4-HPR increased apoptosis through activation of the ER stress in neuroblastoma cells when compared with either monotherapy [41]. Also, increased survival time in tumor-bearing mice and histological evaluation proved that combination of bortezomib and 4-HPR increased anti-tumor and anti-angiogenic mechanisms [41]. Similarly, combination of cetuximab and genistein significantly inhibited tumor growth and caused a substantial delay in tumor growth in animal models of OSCC while no delay in tumor growth was observed using either monotherapy alone [40]. Thus, understanding the complex regulation of cell death pathways through mitochondrial damage or ER stress- in response to 4-HPR plus GST combination therapy has the potential to reveal novel therapeutic targets in neuroblastoma as observed in present preclinical models.

We used the SIF staining to examine a specific protein expression and the DIF staining to examine a specific protein expression in association with DNA fragmentation in course of apoptosis in both neuroblastoma ectopic SH-SY5Y and orthotopic SK-N-BE2 xenografts. Our results indicated significant involvement of both calpain and caspase-12 overexpression in apoptosis in both neuroblastoma xenografts. We also found significant upregulation of caspase-3 and AIF for apoptosis in both neuroblastoma xenografts. Calpain is known to cause ER stress and activation of caspase-12 for apoptosis [42]. Active caspase-12 is able to directly activate caspase-9 and caspase-3 for inducing apoptosis [43, 44]. Our observation is correlated with previous studies showing that GST increased intracellular  $\text{Ca}^{2+}$  concentration due to release from ER  $\text{Ca}^{2+}$  storage to activate the  $\text{Ca}^{2+}$ -dependent cysteine protease calpain and the ER protease caspase-12 for apoptosis in breast cancer cells [10]. We previously reported that GST caused increase in intracellular free  $\text{Ca}^{2+}$  for activation of calpain and caspase-12 leading to activation of caspase-3 in SH-SY5Y cells for apoptosis [12]. Activation of calpain could cause mitochondrial release of AIF to promote caspase-independent apoptosis.

In conclusion, our investigation showed that 4-HPR + GST worked synergistically for increasing apoptosis in human malignant neuroblastoma SH-SY5Y and SK-N-BE2 cells and induced multiple molecular mechanisms for increasing apoptosis in ectopic SH-SY5Y xenografts as well as in orthotopic SK-N-BE2 xenografts. The results from our investigation strongly suggest that combination of 4-HPR and GST should be explored as a promising



therapeutic strategy for the treatment of malignant neuroblastoma in humans in the near future.

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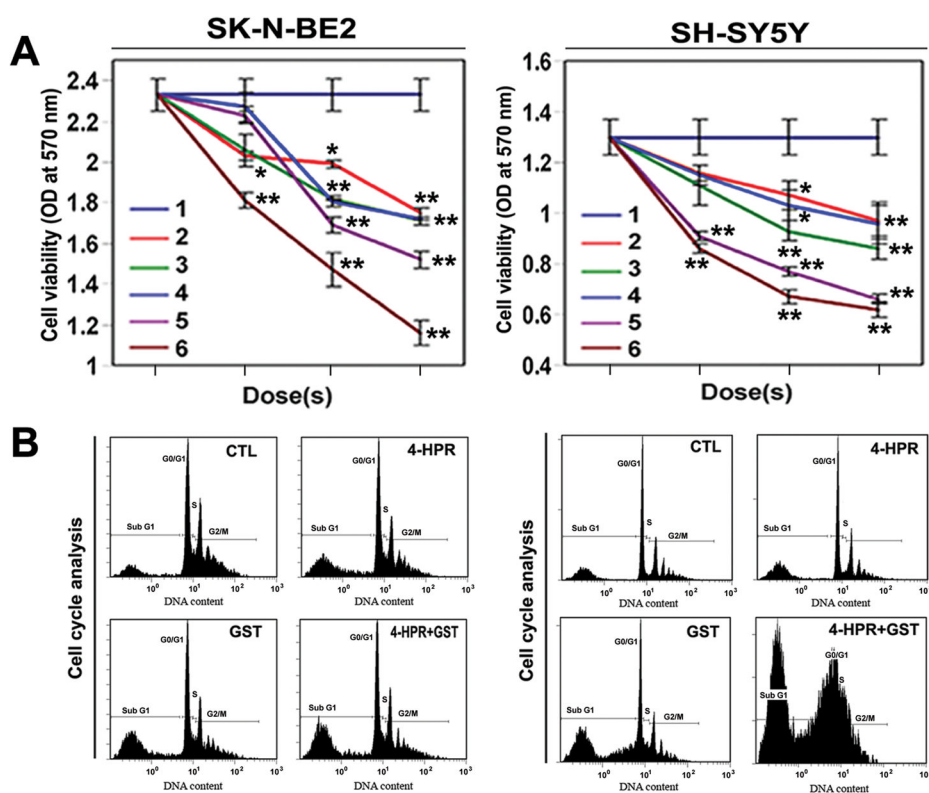
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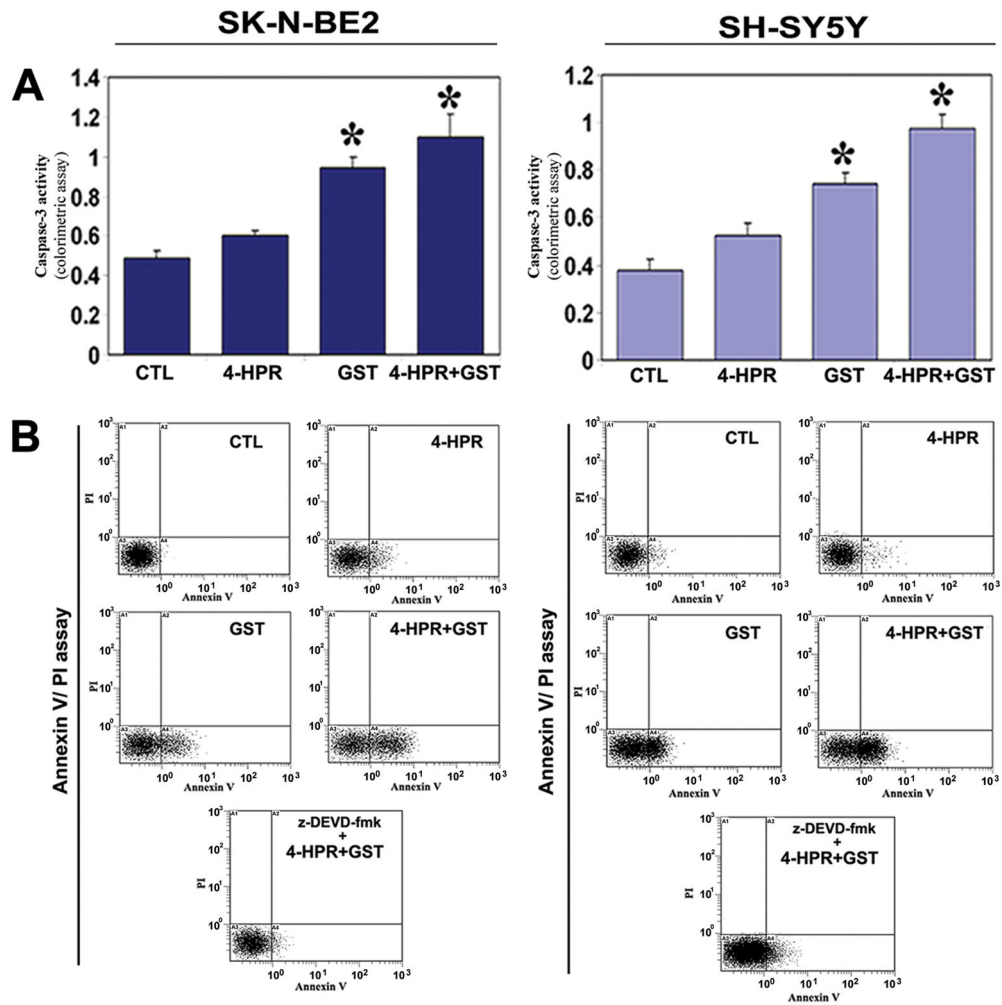
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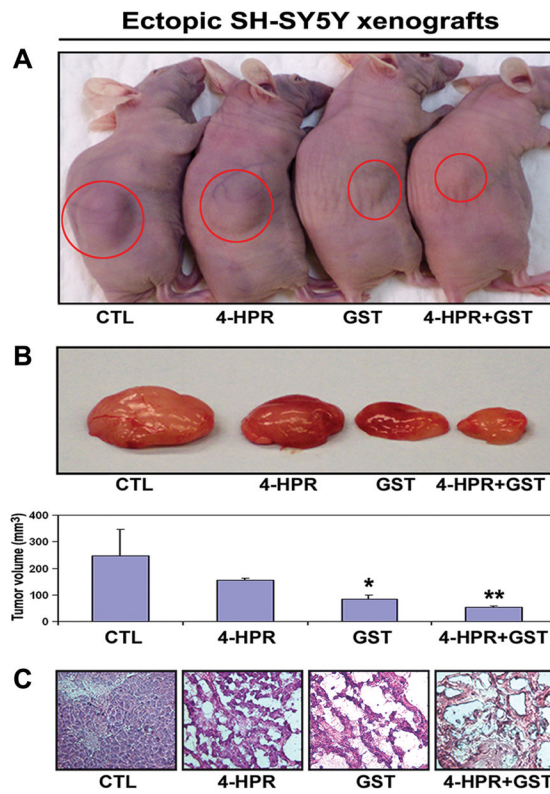
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**Figure 1.** Effects of treatments on human malignant neuroblastoma SK-N-BE2 and SH-SY5Y cells. (A) Changes in cell viability in neuroblastoma cells as determined by MTT assay. Treatments: 1, control (CTL); 2, 4-HPR (0.25, 0.5, and 1  $\mu$ M for 72 h); 3, GST (50, 100, and 200  $\mu$ M for 24 h); 4, 4-HPR (0.25  $\mu$ M for 72 h) + GST (50  $\mu$ M for 24 h); 5, 4-HPR (0.5  $\mu$ M for 72 h) + GST (100  $\mu$ M for 24 h); and 6, 4-HPR (1  $\mu$ M for 72 h) + GST (200  $\mu$ M for 24 h). Finally, 4-HPR (0.5  $\mu$ M for 72 h) + GST (100  $\mu$ M for 24 h) was selected for all other cell culture studies. (B) Cell cycle analysis by flow cytometry in SK-N-BE2 and SH-SY5Y cells. Combination therapy induced apoptotic subG1 accumulations.

**Figure 2.**

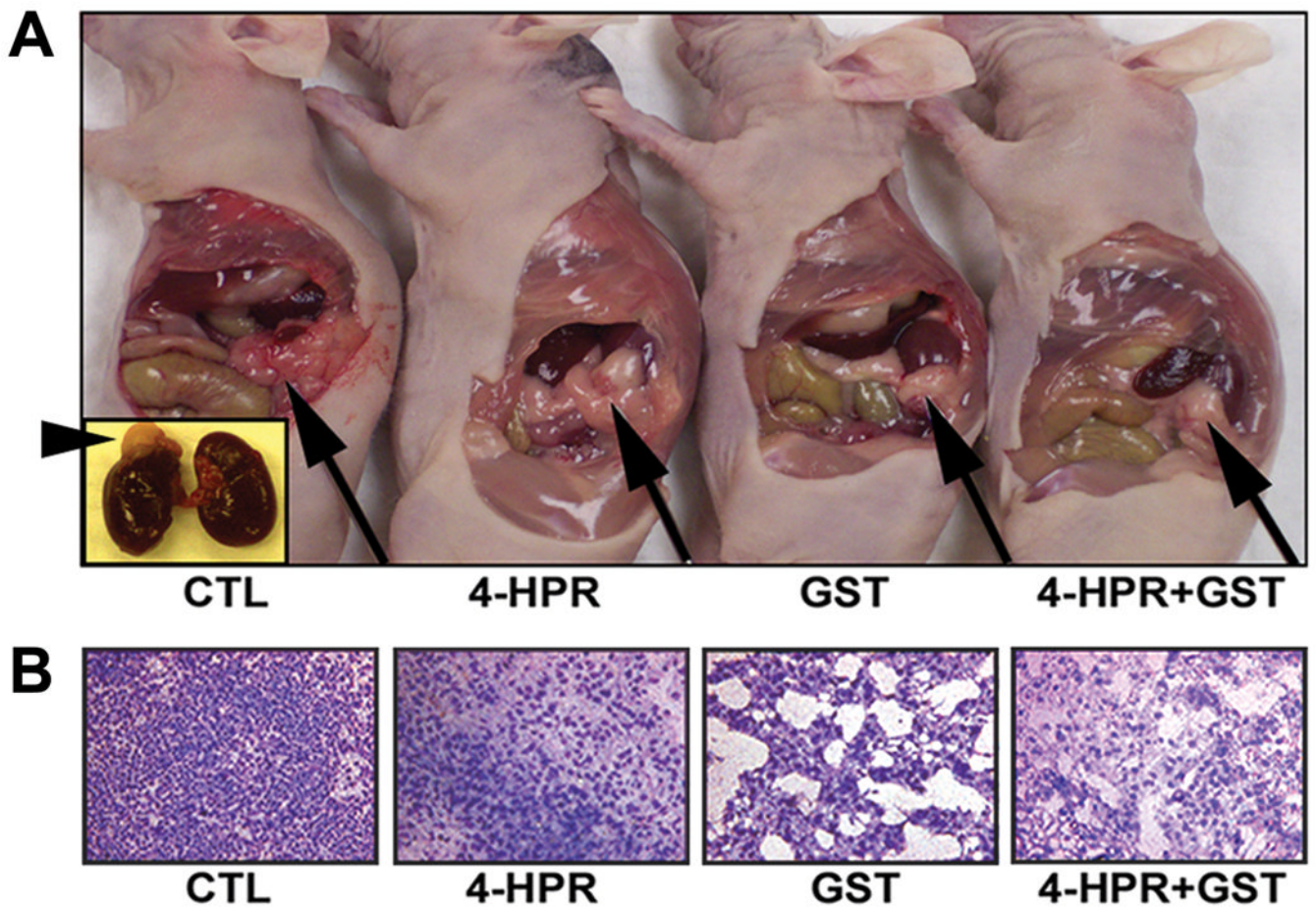
Increases in caspase-3 activity and apoptotic populations in SK-N-BE2 and SH-SY5Y cells. Treatments: CTL, 4-HPR (0.5  $\mu$ M for 72 h), GST (100  $\mu$ M for 24 h), and 4-HPR (0.5  $\mu$ M for 72 h) + GST (100  $\mu$ M for 24 h). (A) Determination of caspase-3 activity using a colorimetric assay. (B) Annexin V-FITC/PI staining and flow cytometric determination of apoptosis. Treatment with 4-HPR + GST effectively controlled growth of neuroblastoma cells due to increase in caspase-3 activity for apoptosis.



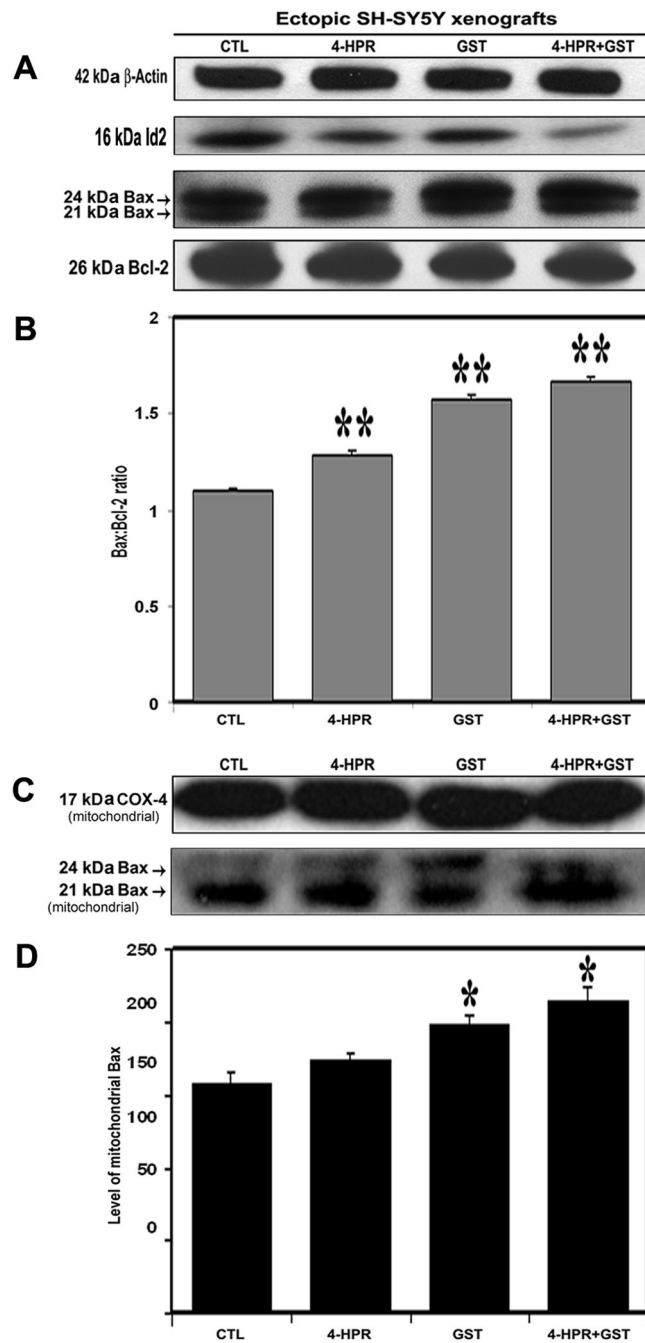
**Figure 3.**

Efficacy of combination therapy in ectopic SH-SY5Y xenografts. (A) Ectopic SH-SY5Y xenografts in nude mice. Treatments (8 days): control (CTL), 4-HPR (20  $\mu\text{g}/\text{kg}/\text{day}$ ), GST (2  $\text{mg}/\text{kg}/\text{day}$ ), and 4-HPR (20  $\mu\text{g}/\text{kg}/\text{day}$ ) plus 4 h later GST (2  $\text{mg}/\text{kg}/\text{day}$ ). (B) The representative tumors after treatments. (C) Histological features (at 200x magnification) of SH-SY5Y xenografts after treatments.

## Orthotopic SK-N-BE2 xenografts



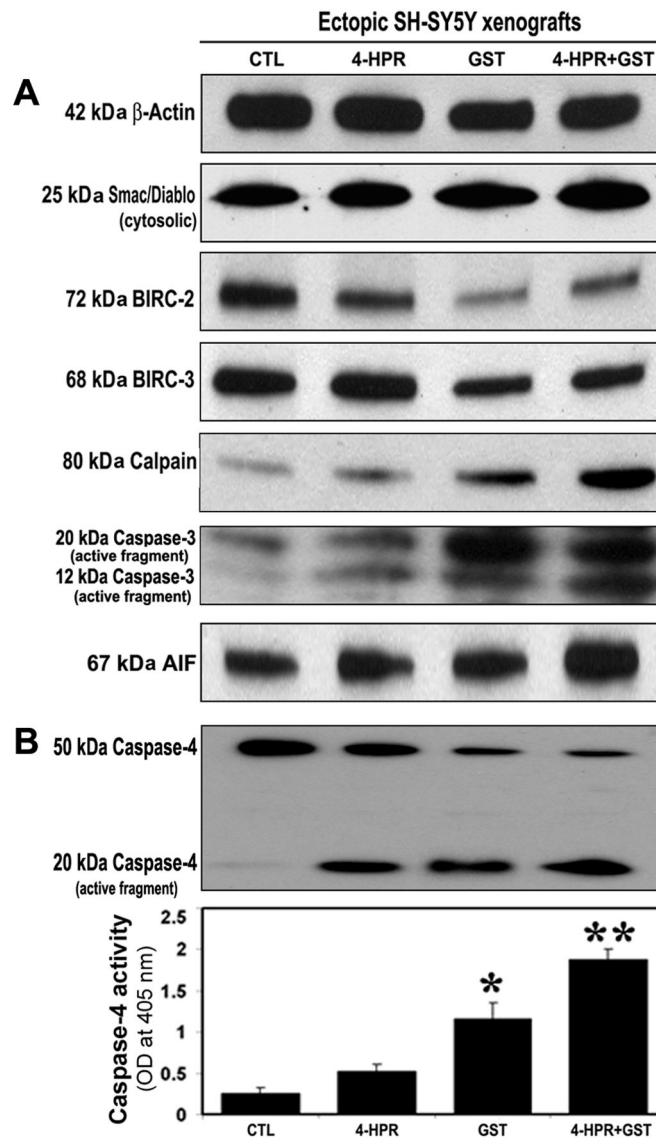
**Figure 4.** Efficacy of combination therapy in orthotopic SK-N-BE2 xenografts. (A) Orthotopic SK-N-BE2 xenografts in nude mice. Treatments (8 days): CTL, 4-HPR (20  $\mu\text{g}/\text{kg}/\text{day}$ ), GST (2  $\text{mg}/\text{kg}/\text{day}$ ), and 4-HPR (20  $\mu\text{g}/\text{kg}/\text{day}$ ) + 4 h later GST (2  $\text{mg}/\text{kg}/\text{day}$ ). Inset: arrowhead indicated location of orthotopic SK-N-BE xenograft on the adrenal gland. Arrow indicated location of tumor on the adrenal gland in animal. (B) Histological features (at 200x magnification) of orthotopic SK-N-BE2 xenografts following treatments.



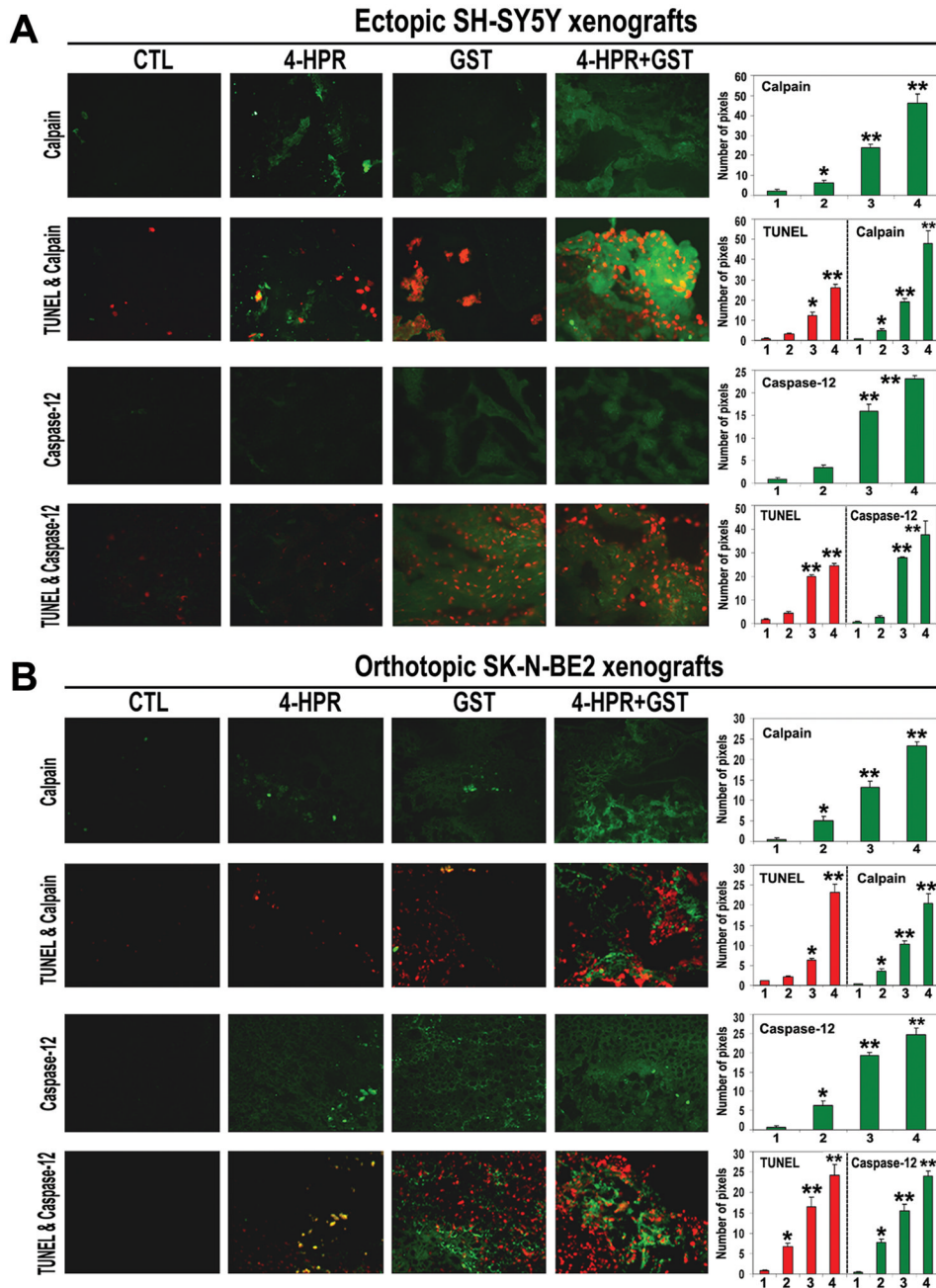
**Figure 5.**

Western blotting for biochemical feature of differentiation and determination of Bax:Bcl-2 ratio and mitochondrial Bax level in SH-SY5Y xenografts. Treatments (8 days): CTL, 4-HPR (20  $\mu$ g/kg/day), GST (2 mg/kg/day), and 4-HPR (20  $\mu$ g/kg/day) + 4 h later GST (2 mg/kg/day). (A) Representative Western blots ( $n \geq 3$ ) to show expression of 42 kDa  $\beta$ -actin, 16 kDa Id2, 21 and 24 kDa Bax, 26 kDa Bcl-2. (B) Densitometric analysis to show change in the Bax:Bcl-2 ratio. (C) Representative Western blots ( $n \geq 3$ ) to show expression of 17 kDa COX-4, 21 and 24 kDa Bax in mitochondrial fraction. (D) Densitometric analysis to show relative changes in the mitochondrial Bax level following treatments.

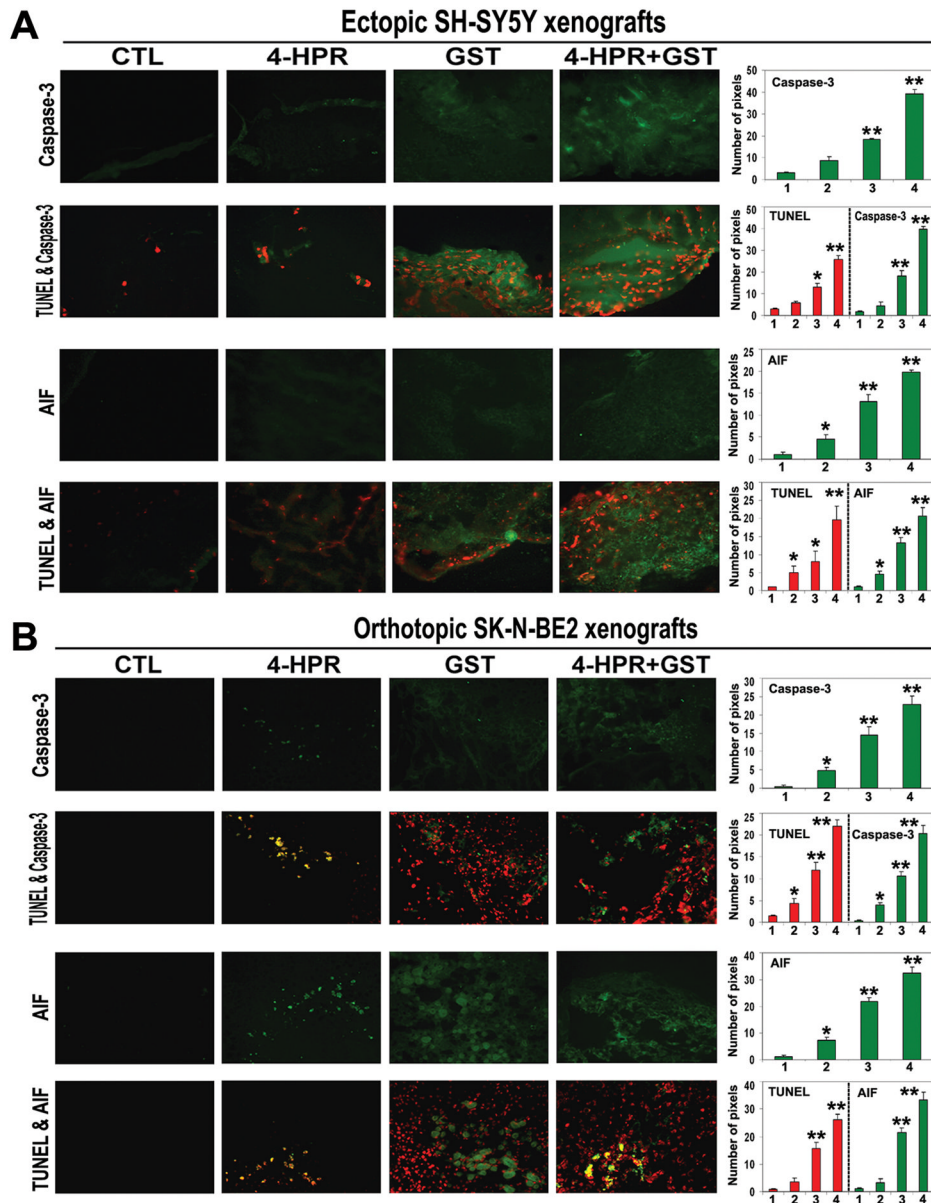




**Figure 6.** Activation of caspase-dependent and caspase-independent pathways for apoptosis in SH-SY5Y xenografts. Treatments (8 days): CTL, 4-HPR (20  $\mu$ g/kg/day), GST (2 mg/kg/day), and 4-HPR (20  $\mu$ g/kg/day) + 4 h later GST (2 mg/kg/day). (A) Representative Western blots ( $n \geq 3$ ) to show expression of 42 kDa  $\beta$ -actin, 25 kDa Smac/Diablo, 72 kDa BIRC-2, 68 kDa BIRC-3, 80 kDa Calpain, 20 kDa and 12 kDa active caspase-3, and 67 kDa AIF. (B) Western blotting showed active 20 kDa caspase-4 fragment. Bar diagram to show changes in caspase-4 activity as determined by a colorimetric assay.

**Figure 7.**

*In situ* stainings for examination of upregulation of calpain and caspase-12 in association with DNA fragmentation in the sections of neuroblastoma xenografts. Treatments (8 days): CTL, 4-HPR (20  $\mu\text{g}/\text{kg}/\text{day}$ ), GST (2  $\text{mg}/\text{kg}/\text{day}$ ), and 4-HPR (20  $\mu\text{g}/\text{kg}/\text{day}$ ) + 4 h later GST (2  $\text{mg}/\text{kg}/\text{day}$ ). (A) The SIF and DIF stainings of the sections of ectopic SH-SY5Y xenografts. (B) The SIF and DIF stainings of the sections of orthotopic SK-N-BE2 xenografts. In the sections (at 200x magnification), calpain or caspase-12 is stained in green and DNA fragmentation (TUNEL labeling) is stained in red. In the bar diagrams, x-axis denotes 1 = CTL, 2 = 4-HPR, 3 = GST, and 4 = 4-HPR + GST.

**Figure 8.**

*In situ* stainings for examination of upregulation of caspase-3 and AIF in association with DNA fragmentation in the sections of neuroblastoma xenografts. Treatments (8 days): CTL, 4-HPR (20  $\mu\text{g}/\text{kg}/\text{day}$ ), GST (2  $\text{mg}/\text{kg}/\text{day}$ ), and 4-HPR (20  $\mu\text{g}/\text{kg}/\text{day}$ ) + 4 h later GST (2  $\text{mg}/\text{kg}/\text{day}$ ). (A) The SIF and DIF stainings of ectopic SH-SY5Y xenografts. (B) The SIF and DIF stainings of the sections of orthotopic SK-N-BE2 xenografts. In the sections (at 200x magnification), caspase-3 or AIF is stained in green and DNA fragmentation (TUNEL labeling) is stained in red. In the bar diagrams, x-axis denotes 1 = CTL, 2 = 4-HPR, 3 = GST, and 4 = 4-HPR + GST.