Dopamine Blocks Stress-Mediated Ovarian Carcinoma Growth

Myrthala Moreno-Smith1, Chunhua Lu1, Mian M.K. Shahzad7, Guillermo N. Armaiz Pena1, Julie K. Allen1, Rebecca L. Stone1, Lingegowda S. Mangala8, Hee Dong Han1, Hye Sun Kim1, Donna Farley9, Gabriel Lopez Berestein2,3,4, Steve W. Cole5, Susan K. Lutgendorf6, and Anil K. Sood1,2,4

Abstract

**Purpose:** Increased adrenergic activity in response to chronic stress is known to promote tumor growth by stimulating the tumor microenvironment. The focus of the current study was to determine whether dopamine, an inhibitory catecholamine, could block the effects of chronic stress on tumor growth.

**Experimental Design:** Expression of dopamine receptors (DR1–DR5) was analyzed by reverse transcriptase-PCR and by Western blotting. *In vitro* effects of dopamine on cell viability, apoptosis, and migration were examined. For *in vivo* therapy, murine and human DR2-siRNAs were incorporated into chitosan nanoparticles (CH-NP).

**Results:** In this model of chronic stress, tumoral norepinephrine levels remained elevated whereas dopamine levels were significantly decreased compared with nonstressed animals. Daily restraint stress resulted in significantly increased tumor growth in both immunodeficient (SKOV3ip1 and HeyA8) and immunocompetent (ID8) ovarian cancer models. This increase was completely blocked with daily dopamine treatment. Dopamine treatment also blocked the stress-induced increase in angiogenesis. Endothelial and ovarian cancer cells expressed all dopamine receptors except for the lack of DR3 expression in ovarian cancer cells. DR2 was responsible for the inhibitory effects of dopamine on tumor growth and microvessel density as well as the stimulatory effect on apoptosis, as the DR2 antagonist eticlopride reversed these effects. Dopamine significantly inhibited cell viability and stimulated apoptosis *in vitro*. Moreover, dopamine reduced cyclic AMP levels and inhibited norepinephrine and vascular permeability factor/VEGF-induced Src kinase activation.

**Conclusions:** Dopamine depletion under chronic stress conditions creates a permissive microenvironment for tumor growth that can be reversed by dopamine replacement. *Clin Cancer Res;* 17(11); 3649–59. ©2011 AACR.

Introduction

The stress response is a complex process arising from interactions between environmental contexts and the organism’s evaluation of potential threat and its capacity to respond. These factors initiate a cascade of information processing in both central and peripheral nervous systems as well as hormonal cascades (1). This results in activation of the sympathetic nervous system (SNS) and the hypothalamic-pituitary-adrenal (HPA) axis (2, 3). Norepinephrine and epinephrine levels are known to be elevated in the plasma and tumor microenvironment of individuals with acute and chronic stress (4, 5). We have recently shown that both norepinephrine and epinephrine levels are elevated in a sustained fashion in ovarian and other peritoneal tissues in preclinical models of chronic stress (6). These hormonal increases were related to greater tumor burden, which was mediated by increased angiogenesis. Recent evidence suggests that the third catecholamine dopamine has the opposite effect with regard to effects on tumor angiogenesis, growth, and development of ascites (7, 8). *In vivo* and *in vitro* studies have shown that dopamine, via its specific DR2 receptors, inhibits tumor growth by suppressing the actions of vascular permeability factor (VPF)/vascular endothelial growth factor-A (VEGF-A) on both tumor endothelial cells and bone marrow–derived endothelial progenitor cells (9). Dopamine inhibits VEGF-induced angiogenesis by suppressing VEGFR-2 phosphorylation (10–12) and inhibits mitogen-activated protein kinase and focal adhesion kinase activation (12). Dopamine can also inhibit mobilization...
Translational Relevance

Epidemiologic and clinical studies have provided growing evidence for links between chronic stress, depression, social isolation, and cancer progression. In the present work, our data provide a new understanding of the mechanisms by which chronic stress contributes to tumour progression. Specifically, our findings indicate that tumoral dopamine levels are depleted under chronic stress. Dopamine replacement blocked the growth stimulatory effects of chronic stress. On the basis of these findings, dopamine analogues may represent a novel therapeutic strategy.

of endothelial progenitor cells (EPC) from the bone marrow (13). It is known that dopamine levels are increased in the brain during acute stress (14). In contrast, under chronic stress conditions, dopamine levels are lower as a consequence of decreased release of dopamine (15). However, it is not known whether dopamine levels are depleted in the tumor microenvironment in response to chronic stress. Moreover, it is not known whether dopamine can counteract the stimulatory effects of norepinephrine on tumor growth. These unanswered questions along with underlying mechanisms are addressed in this article.

Materials and Methods

Reagents

Dopamine (DA), bromocriptine [dopamine receptor 2 (DR2) agonist], eticlopride (DR2-antagonist), SKF 38393 (DR1 agonist), butaclamol (DR1 antagonist), and norepinephrine (NE) were obtained from Sigma Aldrich; recombinant human VEGF was from R&D Systems. Annexin V–(fluorescein isothiocyanate FITC) and (terminal deoxynucleotidyl transferase–mediated dUTP nick end labeling TUNEL) staining kits were purchased from BD Pharmingen and Promega, respectively.

Cell lines and culture conditions

The ovarian nontransformed (HIO-180) and cancer cells (SKOV3ip1, HeyA8, A2780, RMG2, and IGROV) were maintained in RPMI 1640, 15% FBS, and 0.1% gentamicin sulfate (16, 17). Endothelial cells isolated from the mesentery or ovary of the immortomouse (MOEC) were a kind gift from Dr. Robert Langley (18) and were maintained in DMEM, 10% FBS. All materials and methods were conducted at 60% to 80% confluence, unless otherwise specified. For in vivo experiments, all in vitro experiments were conducted at 4°C. Ovarian cancer cells were trypsinized and centrifuged at 1,000 rpm for 7 minutes at 4°C, washed twice, and reconstituted in Hank’s balanced salt solution (Gibco).

Determination of dopamine concentration in tumor and normal tissue

Dopamine levels were determined by high-pressure liquid chromatography with electrochemical (HPLC–EC) detection in the College of Pharmacy at the University of Iowa. The method uses electrochemical detection to quantitate dopamine levels; HPLC is used to separate one catecholamine from another. Values were calculated by comparing the peak height of the unknown (sample) to that of a pure standard of known concentration, these were expressed in picograms of dopamine per milligram of wet tissue.

RT-PCR analysis of dopamine receptors

Total RNA was isolated by using the Qiagen RNeasy Kit. cDNA was synthesized by using the SuperScript First-Strand Kit (Invitrogen) as per the manufacturer’s instructions. cDNA was subjected to PCR using specific primer sequences for human dopamine receptors (DR1–DR5) previously reported (19). Specific primer sequences for murine dopamine receptors (DR1–DR5) were designed on the basis of the reported NCBI nucleotide sequences using the Oligo Perfect Software (Invitrogen). β-Actin was used as a housekeeping gene.

siRNA preparation

Specific siRNA sequences targeted against murine dopamine receptor 2 (DR2; duplex of 5′-GAGUUGUUGUGUAGAGUG-3′ and 5′-GAUUGAAGUAGUAGUGU-3′) and human DR2 (duplex of 5′-CACACAUCCCCAGACACGA and 5′-UGUCAUUCAGGAACUGACUG-3′) were obtained from Sigma Aldrich. These sequences were incorporated into chitosan nanoparticles (CH-NP) by using a gelation method of anionic tripolyphosphate (TPP; ref. 20). Briefly, predetermined TPP (0.25% w/v) and siRNA (1 μg/μL) were added in CH solution, resulting in siRNA-CH-NP generated under constant stirring at room temperature. After incubating at 4°C for 40 minutes, the siRNA-CH1 was collected by centrifugation (Thermo Biofuge) at 12,000 rpm for 40 minutes at 4°C. The CH-NPs were purified 3 times and stored at 4°C until used.

Chronic stress model and treatment schema

Female athymic nude and immunocompetent (C57BL6) mice (10- to 12-week-old) were obtained from the U.S. National Cancer Institute. All experiments were approved by the Institutional Animal Care and Use Committee of the University of Texas MD Anderson Cancer Center. The animals were experimentally stressed using a well-characterized restraint system, in which periodic immobilization induces high levels of HPA and SNS activity characteristic of chronic stress (6). Ovarian cancer cells were injected intraperitoneally into mice 7 days after starting daily stress applied for 2 hours. Then, mice were further divided into treatment groups (10 animals per group) as follows: (i) control PBS; (ii) dopamine (75 mg/kg daily intraperitoneal (i.p.)); (iii) dopamine + eticlopride (10 mg/kg daily i.p.); (iv) control siRNA-CH; (v) dopamine + murine dopamine receptor 2-siRNA (mDR2-siRNA-CH (5 μg per injection)); or (vi) dopamine + human dopamine receptor 2-siRNA (hDRD2-siRNA; 5 μg per injection). siRNA treatments (150 μg/kg) were...
given through the i.v. route twice per week. Following 3 weeks of treatment, mice were euthanized and tumor weight and nodules were recorded. Tumor and relevant tissue samples were collected.

Cell viability assay
To examine the effect of dopamine on cell viability, MTT assay was carried out as previously described (21).

Cyclic AMP determination
The effect of dopamine on cyclic AMP (cAMP) accumulation was examined exposing cells to 0, 10, and 50 μmol/L dopamine for 30 minutes at 37°C. cAMP levels were measured in total cell lysates using an enzyme immunoassay kit (BioMol).

Cell invasion assay
Invasion through human-defined matrix was assessed using the Membrane Invasion Culture System (MICS), as previously described (22). A total of 1.5 × 10⁵ SKOV3ip1 cells were loaded into the upper chamber in media only or in media containing the stimulant of interest. Agonist (DR1, SKF 38393; DR2, bromocriptine) and antagonist (DR1, butaclamol; DR2, eticlopride) were used at 50 μmol/L, norepinephrine at 10 μmol/L, and VEGF at 10 ng/mL. Cells were allowed to invade in a humidified incubator for 24 hours. Cells that had invaded through the basement membrane were collected, stained, and counted by light microscopy in 5 random fields (original magnification ×200) per sample.

Western blot and immunoprecipitation analyses
Western blot analysis was carried out as previously described (21). Briefly, lysates from cultured cells were prepared using modified radioimmunoprecipitation assay (RIPA) buffer, protein concentrations were determined using a BCA Protein Assay Reagent kit (Pierce Biotechnology). Briefly, lysates from cultured cells were prepared using modified radioimmunoprecipitation assay described (21). Lysates from cultured cells were incubated with anti-DR2 antibody at 4°C overnight. Samples were washed in PBS and incubated with blocking solution for 10 minutes and then with a goat anti-rabbit Alexa 488 antibody (1:1,000) for 1 hour. Afterward, tissues were subjected to CD31 and TUNEL staining, as previously described. (18, 21, 23, 24).

Western blot analysis was carried out as previously described (21). Briefly, lysates from cultured cells were prepared using modified radioimmunoprecipitation assay (RIPA) buffer, protein concentrations were determined using a BCA Protein Assay Reagent kit (Pierce Biotechnology). Protein lysates were subjected to 10% SDS–PAGE separation and transferred to a nitrocellulose membrane. Blots were probed with primary antibodies DR2 (Santa Cruz Biotechnology), Src and phosphorylated Src (pSrc) kinase (Cell Signaling Biotechnology), and horseradish peroxidase–conjugated secondary antibody (Amersham) and developed with enhanced chemiluminescence detection kit (ECL; Pierce Biotechnology). Equal protein loading was confirmed reprobing membranes with an anti-β-actin antibody (Sigma Aldrich). To examine the association between Goi1-protein and DR2, MOEC cells were exposed to 10 μmol/L dopamine for 0, 10, and 30 minutes at 37°C. Cell lysates were then prepared and immunoprecipitated with DR2 antibody at 4°C for 2 hours. Immunocomplexes were captured with 2% protein A–agarose beads (Upstate). Protein was eluted in reducing sample Laemmli buffer, subjected to 10% SDS–PAGE separation, and transferred to a nitrocellulose membrane. Anti-Goi1 (Abcam) was used as primary antibody. Immunodetection of Goi1-protein was carried out as described earlier.

Immunohistochemistry
Analysis of microvessel density (MVD) and assessment of tumor and endothelial cell apoptosis were conducted following procedures described previously (18, 21, 23, 24). Double immunofluorescence for DR2/CD31 and for DR2/TUNEL was carried out in frozen tissue sections as follows: after acetone fixation and blocking with gelatin (4%), tissues were incubated with rabbit DR2 antibody (Santa Cruz Biotechnology; 1:100) at 4°C overnight. Samples were washed in PBS and incubated with blocking solution for 10 minutes and then with a goat anti-rabbit Alexa 488 antibody (1:1,000) for 1 hour. Afterward, tissues were subjected to CD31 and TUNEL staining, as previously described. (18, 21, 23, 24).

Immunofluorescence and confocal microscopy
Immunofluorescence microscopy was carried out using a Zeiss Axioplan fluorescence microscope (Carl Zeiss, Inc.) equipped with a 100-W Hg lamp and narrow band-pass excitation filters. Representative images were obtained using a cooled charge-coupled device Hamamatsu C5810 camera (Hamamatsu Photonics) and Optimas software (Media Cybernetics). Confocal fluorescence images were collected using 20× objectives on a Zeiss LSM 510 laser scanning microscopy system (Carl Zeiss, Inc.) equipped with a motorized Axioplan microscope, argon laser (458/477/488/514 nm, 30 mW), HeNe laser (413 nm, 1 mW, and 633 nm, 5 mW), LSM 510 control and image acquisition software, and appropriate filters (ChromaTechnology Corp.). Composite images were constructed with Photoshop software (Adobe Systems, Inc.).

In vitro assessment of apoptosis
MOEC and SKOV3ip1 cells were exposed to different dopamine concentrations (0–50 μmol/L) for 48 hours. The relative percentage of apoptotic cells was assessed using the Annexin V-FITC Apoptosis Detection Kit-1 (BD Pharmingen) according to the manufacturer’s protocol.

Statistical analysis
Continuous variables were compared with the Student t test (between 2 groups) or analysis of variance (for all groups) if normally distributed and the Mann–Whitney rank-sum test if distributions were nonparametric. For in vivo therapy experiments, 10 mice in each group were used, and a P < 0.05 on 2-tailed testing was considered significant. To control for the effects of multiple comparisons, a Bonferroni adjustment was made; for the analysis, a value of P ≤ 0.017 was considered statistically significant.

Results
Dopamine levels during chronic stress
We determined dopamine and norepinephrine concentrations in tumors and peritoneal tissues including ovary, liver, and omentum from mice subjected to daily stress for 1, 3, 7, or 14 days. Norepinephrine levels increased after 1 day of stress and remained elevated until day 14 in all
tissues samples. In contrast, dopamine concentrations increased significantly after 1 day of stress but declined after 3 to 14 days of stress in tumor, ovarian, and omental tissues (Supplementary Fig. S1).

**Dopamine dose-response in vivo**

We have previously shown that chronic stress establishes favorable conditions for promoting angiogenesis in the tumor microenvironment by SNS activation (6). Because the levels of dopamine, the antiangiogenic catecholamine, are decreased in the tumor microenvironment under chronic stress, we asked whether dopamine could block the stimulatory effects of SNS on ovarian cancer growth. Before conducting such blocking experiments, we first tested several doses of dopamine to identify the lowest dose required for inhibiting cancer growth in nonstressed or stressed mice. SKOV3ip1 tumor-bearing nude mice (n = 10 per group) were treated with (i) control PBS, (ii) dopamine at 50 mg/kg, (iii) dopamine at 75 mg/kg, or (iv) dopamine at 100 mg/kg. Treatment was started 1 week after injection of SKOV3ip1 ovarian cancer cells. In stressed mice, doses of 75 and 100 mg/kg significantly reduced tumor growth (P < 0.05), with the greatest decrease noted at the 75 mg/kg dose. All 3 doses significantly decreased the number of tumor nodules (P < 0.05) compared with controls (Supplementary Fig. S2A). In contrast, in nonstressed mice, no significant changes in tumor weight or tumor nodules were observed at any of the dopamine doses tested. Given the known effects of dopamine on angiogenesis (7), we also assessed MVD. There was a significant reduction in MVD at all dopamine doses tested (P < 0.05; Supplementary Fig. S2B). Dopamine at 75 mg/kg was selected for all subsequent in vivo experiments because of its inhibitory effect on cancer growth.

To assess the longitudinal effects of dopamine, we carried out an in vivo experiment in SKOV3ip1 tumor-bearing mice. Tumor growth curves in nonstressed and stressed mice are presented in Supplementary Figure S2D. Although there was no significant effect of dopamine in the nonstress setting, dopamine completely blocked the growth stimulatory effects of daily restraint stress (Supplementary Fig. S2D).

**Dopamine blocks stress-induced ovarian cancer growth via DR2**

As expected, daily stress increased tumor growth by 2.4-fold (Fig. 1). In stressed mice, daily dopamine treatments resulted in 68% reduction in tumor growth compared with control mice injected with PBS (P < 0.05). We also tested the effects of dopamine in an immunocompetent syngeneic mouse model of ovarian cancer (ID8-C57BL6 model). Consistent with the other models, daily restraint stress increased tumor growth by 50% (P < 0.01; Supplementary Fig. S3) in the ID8 model. Moreover, dopamine treatment significantly reduced tumor growth (47%; P < 0.02) in the nonstressed group and blocked the stress-mediated increase in tumor growth (78%, P < 0.001; Supplementary Fig. S3).

Because dopamine can signal through multiple receptors, we examined the RNA expression of DR1–DR5 in ovarian and endothelial cell lines. Figure 1A shows the RNA expression in ovarian nontransformed epithelial (HIO-180), cancer (A2780, HeyA8, and SKOV3ip1), and endothelial cells [human umbilical vein endothelial cells (HUVEC)]. Although all dopamine receptors are expressed in HUVEC, DR3 was not detected in any of the ovarian cell lines tested. Protein expression of DR1 and DR2 was also confirmed by Western blotting in these and other ovarian cancer cell lines (RMG2 and IGROV) as well as MOEC cells (Fig. 1A).

Given the probable role of DR2 in mediating the antiangiogenic effects of dopamine (7), we next examined whether the suppression of tumor growth by dopamine was specifically mediated by DR2. In stressed mice, the addition of a DR2 antagonist (eticlopride) to dopamine treatment reversed the protective effects of dopamine (Fig. 1B). These data suggest that dopamine indeed acted through DR2 to block stress-mediated ovarian cancer growth.

**Dopamine targets murine endothelial and human cancer cells**

To further examine whether the inhibitory effects of dopamine were mediated by targeting (host) endothelial and/or (human) tumor cells through their corresponding DR2 receptors, we utilized siRNA incorporated into CH-NPs. The specificity of siRNA sequences for mouse and human DR2 was confirmed using RT-PCR (Supplementary Fig. S4). SKOV3ip1 tumor-bearing mice were divided into 4 experimental groups: (i) control siRNA, (ii) dopamine alone, (iii) dopamine + mDR2-siRNA, and (iv) dopamine + hDR2-siRNA; each group was used for both nonstress and stress conditions (Fig. 1C). In daily stressed mice, dopamine treatment significantly blocked tumor weight (67%) and the number of tumor nodules (65%). This growth inhibitory effect of dopamine was significantly abrogated by dopamine/mDR2-siRNA-CH but not by dopamine/hDR2-siRNA-CH combined treatment (P = 0.02; Fig. 1C). These results suggest that in the SKOV3ip1 model, dopamine-mediated effects occur primarily through DR2 on endothelial cells. The effects of dopamine were also tested in a second ovarian cancer model, HeyA8 (Fig. 1D). Compared with mice injected with control siRNA, dopamine treatment blocked tumor growth by 84% (P = 0.007) and resulted in a 63% decrease in the number of nodules (P = 0.02). Interestingly, in this model, both dopamine/mDR2-siRNA-CH and dopamine/hDR2-siRNA-CH treatments significantly blocked the inhibitory effects of dopamine.

**Effect of dopamine on angiogenesis and cell viability**

Given the known effects of catecholamines on angiogenesis, we examined MVD in the SKOV3ip1 tumors harvested from dopamine-treated animals. In stressed mice, dopamine treatment resulted in a significant reduction (61%) in MVD compared with controls (Fig. 2A; P < 0.01). In
contrast, dopamine/eticlopride treatment led to a significant increase (Fig. 2A; $P < 0.01$) in MVD compared with dopamine-treated mice. We also examined effects on tumor cell apoptosis, as it is known that norepinephrine can reduce sensitivity of cancer cells to apoptosis (19). Tissues from dopamine-treated mice revealed a 2.2-fold increase ($P < 0.01$) in tumor cell apoptosis compared with control tissues (Fig. 2A). The combined dopamine/eticlopride treatment abrogated the dopamine-mediated effects on cell apoptosis. In addition, the combined treatment of dopamine/mDR2-siRNA and dopamine/hDR2-siRNA-CH reversed the dopamine-blocking effect on tumor growth. Data are the mean ± SE; n = 10 mice per group. *, $P < 0.05$ compared with stress controls; †, $P < 0.005$ compared with nonstress controls. C and D, tumor weight and tumor nodules in stressed mice bearing SKOV3ip1 and HeyA8 tumors treated with control siRNA, DA, DA/mDR2-siRNA-CH, and DA/hDR2-siRNA-CH. DA reduced significantly tumor growth and tumor nodules in the SKOV3ip1 ($P < 0.05$) and HeyA8 models ($P < 0.01$). The combined treatment of DA/mDR2-siRNA significantly abrogated the DA inhibitory effect on tumor growth in the SKOV3ip1 model. In the HeyA8 model, both combined treatments (DA/mDR2-siRNA-CH and DA/hDR2) reversed the DA-blocking effect on tumor growth. Data are the mean ± SE; n = 10 mice per group. *, $P < 0.05$; **, $P < 0.01$, compared with controls; †, $P < 0.05$, compared with DA-treated mice.

To determine whether the effects of dopamine on tumor growth are direct or indirect, we carried out a series of in vitro experiments. First, effects of dopamine on in vitro viability of ovarian cancer or endothelial cells were tested. Dopamine did not significantly affect viability of the non-transformed (HIO-180) or ovarian cancer cells (A2780 and SKOV3ip1; Fig. 3). However, cell viability of HeyA8 ovarian cancer cells was significantly decreased at dopamine doses from 12.5 to 50 μmol/L. This is consistent with the in vivo inhibitory effect of dopamine on HeyA8 tumor growth. In MOEC cells, there was a dose-dependent decrease in cell viability with increasing doses of dopamine ($P < 0.01$; Fig. 3).

We also examined the effects of dopamine on apoptosis in MOEC and SKOV3ip1 cells after treatment for 48 hours. The percentage of early apoptosis increased proportionally to dopamine concentrations used in MOEC (Fig. 3),
whereas late apoptotic rates did not vary. Total apoptotic rates in these cells were significantly elevated at 25 μmol/L (51%) and 50 μmol/L (75%) compared with untreated cells (P<0.01). SKOV3ip1 cells showed significantly higher percentages of late apoptotic cells at 50 μmol/L dopamine (P<0.01).

Figure 2. Dopamine (DA) decreases stress-stimulated angiogenesis and stimulates tumor cell apoptosis via its DR2. A, MVD was significantly decreased by DA treatment in SKOV3ip1 tumor tissues of stressed mice (*, P<0.01). DA/eticlopride treatment significantly †, (P<0.01) reversed DA inhibitory effect on MVD, which was evaluated by immunohistochemical analysis of CD31 [3,3'-diaminobenzidine (DAB) staining]. Microvessels were counted in 5 fields at 100× of each tissue sample (n=5). Values are means ± SE. DA treatment caused significantly (*, P<0.005) higher apoptotic rates than control mice. This effect was also significantly †, (P<0.005) reversed by DA/eticlopride combined treatment. Confocal images (200×) showing colocalization of DR2 (red fluorescence) in CD31-positive tumor endothelial cells (green fluorescence). Apoptotic cells (green fluorescence) were detected by TUNEL staining (Promega kit). Percentages of apoptotic cells were calculated in 5 fields at 200× of each tissue sample (n=5). Values are mean ± SE. Detection of DR2 (red fluorescence) in tumor apoptotic cells (green fluorescence). Confocal images 200× B, DA/mDR2-siRNA and DA/hDR2-siRNA-CH combined treatments induced significant increase †, (P<0.005) in MVD in stressed mice, reverting the DA inhibitory effect on MVD (*, P<0.005). Values are mean ± SE. Detection of DR2 (red fluorescence) in tumor CD31-positive endothelial cells (green fluorescence). Confocal images 200×. DA stimulatory effect on tumor cell apoptosis was also significantly abrogated by the DA/mDR2-siRNA and DA/hDR2-siRNA treatments †, (P<0.005). Values are mean ± SE. Confocal images (200×) revealing colocalization of DR2 (red fluorescence) in tumor apoptotic cells (green fluorescence).
Dopamine counteracts the stimulatory effect of VEGF and norepinephrine on tumor cell invasion

Although dopamine did not affect cell viability of all ovarian cancer cells tested, we explored whether it could affect other steps in the metastatic cascade. We examined the effect of dopamine on cell invasion. We analyzed the effect of dopamine and various dopamine agonists and antagonists on the invasive potential of SKVO3ip1 cells. Dopamine at 10 μmol/L inhibited VPF/VEGF-induced cell invasion (25) by 59% (P < 0.05; Fig. 4), as did the DR2 agonist bromocriptine (56%; P < 0.05). However, the DR1 agonist SKF 38393 (50 μmol/L) had no effect. Eticlopride (50 μmol/L), a specific DR2 receptor antagonist, significantly abrogated the dopamine-mediated inhibition of cancer cell invasion (P < 0.01). These results further suggest that the inhibitory action of dopamine on cell invasion was mediated specifically through DR2.

We also examined whether dopamine can block the stimulatory effect of norepinephrine on SKOV3ip1 cell invasion, previously described by our group (22). Dopamine blocked the norepinephrine-mediated effect by 61% (P < 0.05) in SKOV3ip1 cells exposed to norepinephrine plus dopamine. In addition, the combined treatment of dopamine/norepinephrine/VEGF led to a 52% reduction (P < 0.001) in cell invasion compared with norepinephrine/VEGF treatment, indicating the ability of dopamine to counteract the stimulatory effects of norepinephrine and VEGF on cell invasion (Fig. 4).

Dopamine decreases cAMP levels in MOEC and SKOV3ip1 cells

Dopamine, acting through G-protein–coupled receptors, exerts stimulatory (DR1, DR5) or inhibitory (DR2–DR4) effects on adenylate cyclase leading to increased and decreased intracellular cAMP levels, respectively. We examined the effects of dopamine on cAMP levels in MOEC and SKOV3ip1 cells. Dopamine at 10 μmol/L resulted in a reduction in intracellular cAMP levels by 31% and 70% in MOEC and SKOV3ip1 cells, respectively (Fig. 5A). Further decreases in cAMP levels were noted at higher doses (50 μmol/L) of dopamine.
Dopamine inhibits VEGF- and norepinephrine-induced cell signaling

The effect of dopamine on phosphorylation of several key effectors of the VEGF angiogenic pathway was analyzed in MOEC and SKOV3ip1 cells. In MOEC cells, VEGF (10 ng/mL) for 5 minutes resulted in increased phosphorylation of Src(Y416) (Fig. 5B). This effect was blocked by a 10-minute preincubation of cells with dopamine at 10 μmol/L. Similarly, SKOV3ip1 cells stimulated with norepinephrine (10 μmol/L) for 5 minutes revealed a significant increase (2.4-fold) of pSrc(Y419) (Fig. 5B), which was blocked by a combined treatment of dopamine + norepinephrine for 5 and 15 minutes.

Our in vitro data confirmed that dopamine decreases intracellular cAMP levels in MOEC and SKOV3ip1 cells. This strongly suggested that dopamine could be acting through the DR2–cAMP signaling pathway to exert its inhibitory effects in vivo on tumor growth. To address this question, we examined the association of DR2 to Gαi1-protein in MOEC cells exposed to dopamine (10 μmol/L) for 10 and 30 minutes. There was a 1.6-fold increase in the interaction between DR2 and Gαi1-protein after 10 minutes of dopamine exposure compared with untreated cells (Fig. 5C).

Discussion

The main findings of this study are that dopamine significantly reduces stress-mediated cancer growth in ovarian carcinoma. Our data strongly suggest that dopamine retards tumor growth by inhibiting tumor angiogenesis and stimulating tumor cell apoptosis. In addition, we provide the first evidence that dopamine can block the stimulatory effects of chronic stress on cancer growth.

The physiologic actions of dopamine are mediated by at least 5 distinct G-protein–coupled receptor subtypes. (26, 27). Two DR1-like receptor subtypes (DR1 and DR5) couple to the G-protein Gs, activate adenylate cyclase, and increase cAMP levels. The other receptor subtypes belong to the DR2-like subfamily (DR2, DR3, and DR4) and are prototypic of G-protein–coupled receptors that inhibit adenylate cyclase and decrease cAMP production. The ovarian nontransformed cancer and endothelial cells tested in this work showed expression of DR1- and DR2-like dopamine receptors, indicating that dopamine might regulate stimulatory and/or inhibitory processes in these cells.

Dysfunction of dopaminergic system is known to be associated with various disorders, including schizophrenia and Parkinson's disease (28). The consequences of dopamine dysfunction indicate the importance of maintaining dopamine functionality through homeostatic mechanisms based on the delicate balance between synthesis, storage, release, metabolism, and reuptake (28). A decrease in dopamine in the brain has been implicated as the cause of Parkinson's disease. In contrast, it is argued that a functional excess of dopamine or oversensitivity of certain DRs is one of the causal factors in schizophrenia (29). Interestingly, the incidence of cancer in patients with schizophrenia has been reported to be lower than in the general population (30–32). Whether this reduced incidence is related to hyperactive dopaminergic system is not known.

It has been shown that dopamine concentrations are lower in the tumor microenvironment than in normal tissues (11, 33). These findings prompted us to consider whether the increases in tumor growth and angiogenesis may result from a permissive microenvironment created by a relative shift toward growth-promoting catecholamines. In the present study, we show that dopamine blocks stress-induced tumor growth by activating DR2. The central role of DR2 was confirmed with the DR2 antagonist eticlopride in combination therapy with dopamine, as well as with siRNA targeted against murine or
human DR2. The dopamine-suppressing effect on tumor growth in our stress models was significantly blunted by DR2 gene silencing. These findings indicate that dopamine targets both host murine endothelial cells and human cancer cells through DR2 to exert its growth-suppressive effects. Our results are supported by the reported growth inhibitory effects of dopamine under nonstress conditions (7, 9, 34).

Our data also indicate that dopamine, via DR2, blocked the VPF/VEGF or norepinephrine-mediated invasion of ovarian cancer cells. The inhibitory effects of dopamine on cell invasion would explain, in part, the in vivo blocking effect of dopamine on tumor progression and metastasis under chronic stress. Our in vitro studies show that dopamine decreases viability not only of endothelial cells, as previously described (7, 12) but also of ovarian cancer cells. These results were further confirmed using in vivo models of ovarian carcinoma. The dual cell targeting (endothelial and tumor cells) of dopamine in the tumor microenvironment underlines the significance of dopamine as a potential modality to block the growth stimulatory effects of chronic stress.

Dopamine and DR2 agonists have been used for many years for the treatment of Parkinson’s disease and hyperprolactinemia (35, 36) as well as for the management of cardiovascular disorders and renal dysfunction. (37, 38).
Dopamine agonists such as bromocriptine mesylate (Parlodel, oral) have also been used clinically to treat hyperprolactinemia and are well tolerated (39). Furthermore, effective shrinkage of prolactinomas has been observed after injection of long-acting form of Parlodel (40). Such agents may represent a new strategy for blocking the effects of chronic stress on tumor growth. With such therapies, however, some adverse reactions such as nausea, hallucinations, and orthostatic hypotension have been reported and would require careful monitoring.

To identify the signaling pathways by which dopamine affects endothelial and ovarian cancer cell function, we examined the effects of dopamine on phosphorylation and activation of various effectors in the metastatic cascade. In MOEC cells, dopamine inhibited VEGF-induced phosphorylation and activator of Src kinase. In addition, dopamine reversed norepinephrine-stimulated Src phosphorylation. Src is a key mediator in multiple signaling pathways that regulate critical cellular functions (41). In ovarian cancer, Src has been shown to play a functional role in cancer cell invasion and angiogenesis (42). Src is also a regulator of VEGF-mediated vascular permeability (43). Recent studies in HUVECs have shown that dopamine reduces VEGF-mediated permeability by inhibiting VEGF-induced Src activation (44). These studies coupled with our data implicate blockade of Src activation as a key mediator of the inhibitory effects of dopamine on ovarian cancer growth.

Overall, our data suggest that dopamine inhibits stress hormone-stimulated Src activation in endothelial and ovarian cancer cells. Moreover, we conclude that dopamine replacement effectively counteracts the stimulatory effects of norepinephrine on ovarian cancer growth during chronic stress. Considering the stimulatory effects of chronic stress on cancer growth (6), our findings implicate dopamine as a potential therapeutic strategy for blocking the deleterious effects of chronic stress.

Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

Grant Support

Portions of this work were supported by the NIH (CA110793, CA109298, P50 CA 083639, CA128797, RC2GM092599; U54 CA151668), the Ovarian Cancer Research Fund, Inc. (Program Project Development Grant), the DOD (OC093146), the Zarrow Foundation, the Marcus Foundation, the Blanton-Davis Ovarian Cancer Research Program, the Laura and John Arnold Foundation, and the Betty Anne Asche Murray Distinguished Professorship. R.L. Stone is supported by NCI-DRHS-NIH T32 Training Grant (T32 CA101642). M.M.K. Shahzad is supported by the NIH/NICHD WRHR Grant (HD050128) and the CCF-Molly Cade Ovarian Cancer Research Grant.

The costs of publication of this article were defrayed in part by the payment of page charges. This article must therefore be hereby marked advertisement in accordance with 18 U.S.C. Section 1734 solely to indicate this fact.

Received September 13, 2010; revised March 25, 2011; accepted March 27, 2011; published OnlineFirst April 29, 2011.

References


