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Ethanol Enhances Estrogen Mediated Angiogenesis in Breast Cancer

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Abstract

Angiogenesis, a highly regulated process, is exploited by tumors like breast cancer to ensure a constant supply of oxygen and nutrients and is key for tumor survival and progression. Estrogen and alcohol independently have been observed to contribute to angiogenesis in breast cancer but their combinatorial effects have never been evaluated. The exact mechanism by which estrogen and alcohol contribute to breast cancer angiogenesis remains to be elucidated. In this study, we defined the in vitro effects of the combination of estrogen and alcohol in breast cancer angiogenesis using the tubulogenesis and scratch wound assays. Conditioned media, generated by culturing the murine mammary cancer cell line, TG1-1, in estrogen and ethanol, enhanced tubule formation and migration as well as modulated the MAP Kinase pathway in the murine endothelial cell line, SVEC4-10. Additionally, estrogen and ethanol in combination enhanced the expression of the pro-angiogenic factors VEGF, MMP-9, and eNOS, and modulated Akt activation. These observations suggest that TG1-1 cells secrete pro-angiogenic molecules in response to the combination of estrogen and ethanol that modulate the morphological and migratory properties of endothelial cells. The data presented in this study, is the first in attempting to link the cooperative activity between estrogen and ethanol in breast cancer progression, underscoring correlations first made by epidemiological observations linking the two.

Key words: breast cancer, angiogenesis, estrogen, alcohol, ethanol

Introduction

Breast cancer remains one of the most commonly occurring cancers in women. The estimated breast cancer cases and deaths in 2018 stand at 268,670 and 41,400 respectively; making it second only to lung cancer in the number of cancer related deaths in the United States of America. Factors known to influence the development of breast cancer include genetics, evidenced by the positive association between breast cancer and mutated BRCA1 and BRCA2 genes, and the female sex hormone, estrogen. In addition to its essential physiological roles, estrogen is known to sustain breast cancer development and progression by enhancing cell viability and motility. The actions of estrogen are primarily initiated by its binding to estrogen receptor alpha (ERα), resulting in the activation of key cellular signaling pathways including the MAPK, NFκB and AP-1 pathways.

Other long-known risk factors for breast cancer include early menarche, late menopause, age at first birth, intervals between births as well as number of births, all of which implicate estrogen exposure as a key factor for breast cancer development. This led to the development of anti-estrogens like tamoxifen as a mainstay treatment strategy for breast cancer. Anti-estrogens like tamoxifen, in addition to inhibiting the various cellular effects of estrogen, have been demonstrated to inhibit angiogenesis, promoting a better prognosis in treated patients, and enhanced cell death in animal models.

Interestingly, in addition to the aforementioned risk factors, epidemiological observations positively correlate breast cancer incidence with alcohol consumption. A meta-analysis of 53 epidemiological studies, including 58,515 women with invasive breast
cancer and 95,067 healthy controls demonstrated a consistently higher relative risk of breast cancer in the alcohol consuming group. Moreover, the relative risk increased with increasing levels of alcohol intake. Another meta-analysis spanning 33 years revealed a similar result and estimated that an average of one drink per day leads to a monotonic 10% rise in breast cancer risk. On the molecular level, experimental studies have suggested that by-products of ethanol metabolism, namely acetaldehyde, play a role in increasing oxidative stress, epigenetic changes, and DNA damage, thus promoting tumor development. A study by Williams et al evaluating risk of invasive breast cancer in women who consume alcohol demonstrated an increased risk in invasive breast cancer in certain populations that drank more than 7 alcoholic beverages per week. Although epidemiological studies link alcohol and increased breast cancer incidence, a concrete mechanism is yet to be established.

A significant observation was made wherein increased risk and incidence of breast cancer was positively correlated in women with ER positive breast cancer who reported increased alcohol consumption. This observation was further corroborated in a study, that included 38,454 women, and observed a positive correlation between breast cancer risk and moderate alcohol consumption in postmenopausal women on hormone replacement therapy. Together, these studies provide anecdotal evidence suggesting a relationship between alcohol and estrogen that promotes breast cancer; however, the mechanisms regulating this risk have not been fully elucidated.

One biological mechanism that is targeted by both alcohol and estrogen in tumors is angiogenesis; a process indispensable for tumor growth and progression. Ethanol consumption has been demonstrated to promote mammary tumor growth, leading to increased cellular proliferation and enhanced expression of VEGF, an essential protein involved in angiogenesis. Estrogen, over and above increasing viability and proliferation of breast cancer cells via the ER-alpha pathway, is also known to promote endothelial cell activity; consequently neo-vascularization and angiogenesis. We reported earlier that estrogen mobilizes endothelial progenitor cells (EPCs) from the bone marrow to the tumor site thus increasing breast tumor tissue neo-vascularization.

Based on our current knowledge we speculate that a cooperative mechanism exists between estrogen and alcohol which acts to increase angiogenesis. To date, studies addressing this interaction between estrogen and alcohol on neo-vascularization in breast cancer have not been undertaken. This formed the basis for this study; an effort to explain a mechanism for this potential relationship between alcohol and estrogen in enhancing angiogenesis. We hypothesized that both alcohol and estrogen play complementary roles to enhance angiogenesis in breast cancer, thereby promoting it. To that end, we used in vitro model systems to explore the effects of ethanol, estrogen and a combination of the two on both a murine mammary cancer cell line, TG1-1, and a murine endothelial cell line, SVEC4-10. Treated TG1-1 cells were assayed for the effect of either estrogen, ethanol, or their combination on expression of various pro-angiogenic factors, as well as the effects on the PI3K/Akt cell-signaling pathway. In addition, conditioned media from treated TG1-1 cells was harvested and used to determine the effects of TG1-1 secretory factors on SVEC4-10 cells. To that end, SVEC4-10 cells were cultured in TG1-1 conditioned media after which tubulogenesis, migration, proliferation and activation of relevant signaling pathways was assayed for. This study, the first of its kind, demonstrates an enhanced expression of angiogenic markers when cells are treated with a combination of alcohol and estrogen.

**Materials and Methods**

**Cell Culture and Treatments**

The TG1-1 murine mammary cancer cell line, a kind gift from Dr. Rakesh Jain (Harvard University), was developed from spontaneous tumors in transgenic FVB mice expressing the c-neu oncogene under the control of the mouse mammary tumor virus promoter. SVEC4-10, a murine endothelial cell line, was purchased from ATCC (CRL-2181). Both cell lines were maintained in complete 1X DMEM (Corning #10-013-CV) supplemented with 10% (v/v) Fetal Bovine Serum (FBS) (BioTC #FBS-02), 2mM L-glutamine (Corning #25-005-CI), 1X penicillin/streptomycin (P/S) (Corning #30-002-CI) and 25μg/ml Plasmocin prophylactic (Invivogen #ant-mp). For ethanol treatments, TG1-1 cells were incubated with 200 proof absolute ethanol (Fisher Scientific #BP2818-100) at a concentration of 0.2% and 0.5%, since physiologically relevant doses are set to be at 0.3%. For estrogen treatment, TG1-1 cells were incubated with β-estradiol (Sigma #E8875-1G) at a concentration of 10⁻⁸ M. Treatments were carried out for 24 hours either in combination or as stand-alone treatments. SVEC4-10 cells were treated for 24 hours with the conditioned media generated from alcohol and/or estrogen treated TG1-1 cells.

**Conditioned Media Experiments**

Conditioned media was generated by treating...
TG1-1 cells with 0.2% or 0.5% ethanol and/or 10⁻⁸M estrogen in starvation media; phenol red free RPMI (Corning #17-105-CV) containing 5% charcoal stripped FBS (Sigma #F6765) and 1X P/S. The media was collected at the end of 24 hours, centrifuged to remove cell debris, and stored at -80°C until further use. SVEC4-10 cells were plated in complete DMEM. Once 70% confluence was reached, the SVEC4-10 cells were incubated in starvation media overnight. The following day, starvation media was removed and TG1-1 conditioned media added to the SVEC4-10 cells and incubated for 24 hours. At the end of 24 hours, cells were lysed and whole cell lysates analyzed for protein expression by western blots.

Western Blot
Whole cell lysates were made from TG1-1 and SVEC4-10 cells using RIPA buffer (50 mM Tris-HCl, pH 7.4, 150 mM NaCl, 0.2% sodium deoxycholate, 0.1% SDS, 0.5% NP-40) supplemented with HALT protease/phosphatase inhibitor cocktail (Thermofisher #78440). Samples were placed on ice for 45 minutes with intermittent vortexing. The lysates were then centrifuged for 20 minutes at 14000 rpm at 4°C. Protein concentrations were determined using the Bio-rad Protein Assay Dye Reagent Concentrate (Bio-Rad #5000006). Fifteen micrograms of total protein were loaded per sample and resolved on 10% SDS gels under reducing conditions and transferred onto Immobilon-P membranes (Millipore #IPVH00010) at 220mA for 2 hours at 4°C. Membranes were blocked using 5% dried milk in 1X TBST (200mM Tris-HCl, 150mM NaCl, pH 7.4, and 0.1% Tween-20 added fresh/liter of 1XTBS (TBS-T)) for 2 hours on a shaker at room temperature. They were incubated with primary antibodies (VEGF, eNOS, MAPK, AKT, MEK, FAK) overnight at a 1:1000 dilution (unless indicated otherwise) in 2% bovine serum albumin and transferred onto Immobilon-P membranes (Millipore #IPVH00010). Fifteen micrograms of total protein were loaded per sample and resolved on 10% SDS gels under reducing conditions and transferred onto Immobilon-P membranes (Millipore #IPVH00010) at 220mA for 2 hours at 4°C. Membranes were blocked using 5% dried milk in 1X TBST (200mM Tris-HCl, 150mM NaCl, pH 7.4, and 0.1% Tween-20 added fresh/liter of 1XTBS (TBS-T)) for 2 hours on a shaker at room temperature. They were incubated with primary antibodies (VEGF, eNOS, MAPK, AKT, MEK, FAK) overnight at a 1:1000 dilution (unless indicated otherwise) in 2% bovine serum albumin made in 1X TBST at 4°C. The next day, membranes were washed three times with 1X TBST, followed by incubation with horse radish peroxidase conjugated secondary antibodies, made at a 1:10,000 dilution in 2% dried milk in 1X TBST, for 2 hours, on a shaker, at room temperature. After 4 washes in 1XTBST, the membrane was developed using Pierce ECL western blotting substrate (Thermofisher #32106), and detected on CL-Xposure Film (Thermofisher #34091). Films were scanned and band intensity analyzed using the ImageJ Software.

Tubulogenesis Assay
Twenty four well plates were coated with 100µl/well of growth factor reduced matrigel (Corning #356231) and incubated for 30 minutes at 37°C to enable polymerization. Overnight starved SVEC4-10 cells were re-suspended in treated or untreated TG1-1 conditioned media as well as unconditioned starvation media and complete DMEM as controls, and plated at 50,000 cells in 500µl/well. The plates were incubated overnight. The following day, excess media was carefully removed and tubulogenesis observed under the microscope in bright field at 500X magnification. Four fields of view were counted per well (2 wells/treatment). Using Image J’s Angiogenesis Analyzer toolset, the number of master segments and total segment length were determined for each field of view and averaged.

XTT Assay
SVEC4-10 cells were plated at 10,000 cells/well in a 96 well plate and allowed to adhere overnight. The next day, starvation media was added and the cells were starved for 14-18 hours after which the starvation media was replaced by TG1-1 conditioned media as well as unconditioned starvation media and complete DMEM serving as controls. Twenty four hours post treatment, conditioned media was removed and 100µl serum free clear DMEM containing 25µl of the XTT (Thermo Fisher #X6493) reagent (1mg/ml) with 25mM Phenazine Methosulfate (PMS) (Fisher Scientific #AC130160010) was added to each well. The plate was incubated at 37°C for 4 hours and OD was measured at 450nm with a reference wavelength of 630nm.

Scratch Wound (Migration) Assay
SVEC4-10 cells were plated at 500,000 cells/well in a 6 well plate and incubated until a 60% confluent monolayer was observed. Subsequently, the cells were starved for 14-18 hours in starvation media followed by treatment with TG1-1 conditioned media for 24 hours. Prior to start of the treatment, two vertical wounds were created using a 20µl sterile pipette tip. Horizontal lines were made, using a sharpie, on the underside of the plate to allow for visualization at the same spots at 0 hours and 24 hours post treatment. Pictures were taken in the bright field phase contrast at 100X magnification, and migration distance calculated using the scale on the microscope.

Statistical Analysis
Data was normalized to estrogen treatment groups. The ethanol and estrogen combination treatments were compared to the estrogen stand-alone treatment using a Student’s T Test where possible; P<0.05 was considered to be statistically significant. Error bars on all figures represent standard error of mean.
Results

**SVEC4-10 cells demonstrate enhanced tubulogenesis when cultured in conditioned media obtained from estrogen and alcohol treated TG1-1 cells.**

Previous observations by our laboratory demonstrated that estrogen stimulates bone marrow derived endothelial progenitor cell (BM-EPCs) migration from the bone marrow thus resulting in enhanced neo-vascularization in breast cancer. Based on our observations, and other groups stating that ethanol can influence angiogenesis, we sought to determine if estrogen and ethanol in combination act to enhance angiogenesis in breast cancer, by using an \textit{in vitro} tubulogenesis assay. In addition to recruitment from the bone marrow, angiogenesis is a complex process determined by the ability of endothelial cells to differentiate and form stable tubules in response to pro-angiogenic factors. In an effort to mimic the tumor microenvironment, conditioned media from TG1-1 cells treated with ethanol, in the presence and absence of estrogen, as well as estrogen alone, was added onto SVEC4-10 cells for 24 hours. Enhanced tubule formation was observed in SVEC4-10 cells when treated with conditioned media generated (Fig 1A) in presence of a combination of ethanol at a concentration of 0.2% or 0.5% with estrogen as compared to estrogen alone. This was further quantified; number of master segments formed (Fig 1B) and the total segment length (Fig 1C) determined. There was an increase in the number of master segments as well as total segment length, commonly used to represent the extent of tubulogenesis. Our results provide evidence that TG1-1 cells, treated with estrogen and alcohol in combination presumably secrete pro-angiogenic factors which result in increased SVEC4-10 endothelial cell differentiation and formation of stable tubules on a basement membrane matrix.

**Ethanol enhances estrogen mediated expression of pro-angiogenic markers in TG1-1 breast cancer cells.**

Angiogenesis is a dynamic process that requires the interplay between various pro-angiogenic factors, which are required both \textit{in vivo} and \textit{in vitro}, such as that observed during tubule formation. Vascular endothelial growth factor (VEGF), is considered to be the most important pro-angiogenic factor, playing a critical role in angiogenesis. In order to determine the levels of VEGF protein expression that could have contributed to the enhanced tubulogenesis observed, TG1-1 whole cell lysates were used and levels of VEGF determined by a western blot analysis (Fig 2A). We observed an increase in the expression

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**Figure 1. Enhanced Tubulogenesis in SVEC4-10 cells in presence of alcohol and estrogen**

A) Representative brightfield images (50X) of SVEC4-10 cells cultured in alcohol (Ethanol/EtOH) and/or estrogen (10^{-8}M) treated TG1-1 conditioned media; B) Number of master segments, and C) Total Segment Length in pixels, analyzed using angiogenesis analyzer by Image J. (n=2) Error bars represent ±SEM.
levels of VEGF protein when TG1-1 cells were treated with a combination of ethanol at 0.2% and 0.5% and estrogen, in a dose dependent manner, compared with estrogen treatment alone. Another molecule that plays an important role in angiogenesis is endothelial nitric oxide synthase (eNOS)\textsuperscript{45–49}. eNOS on converting L-arginine to L-citrulline produces nitric oxide (NO) which plays a crucial role in vascular remodeling and maintaining a vasodilatory tone\textsuperscript{50–52}. We assessed the expression of eNOS in treated Tg1-1 cells, which was similar to that observed in VEGF. Enhanced eNOS expression was observed when TG1-1 cells were treated with a combination of ethanol at 0.2% and 0.5% and estrogen compared to estrogen alone (Fig 2B). In addition to VEGF and eNOS, we assessed the expression of MMP-9, which belongs to a family of proteins collectively known as matrix metalloproteinases. Matrix metalloproteinases play a role in the proteolytic degradation of the extracellular matrix leading to tumor invasion, metastasis, as well as tissue remodeling that occurs during angiogenesis as observed in many cancers\textsuperscript{53–56}. Expression of MMP-9 in TG1-1 cells (Fig 2C) was greatly enhanced upon treating the cells with a combination of ethanol, at 0.2%, 0.5% as well as 1.0%, and estrogen, in a dose dependent manner, when compared to estrogen alone. Taken together these results demonstrate that ethanol and estrogen enhance the expression of pro-angiogenic and ECM remodeling factors in tumor cells, which are in part responsible for the stimulated proliferation and differentiation of endothelial cells and thus, angiogenesis.

**Ethanol and estrogen induced TG1-1 secretory factors enhance activation of MAP kinase cell survival pathway in SVEC4-10 cells**

The MAPK pathway has a wide array of downstream effector molecules that affect cellular functions, such as cell proliferation, differentiation, and cell survival, which has been observed to play a role in angiogenesis\textsuperscript{57–59}. In order to evaluate if secretory factors from treated TG1-1 cells would lead to activation of the MAPK pathway in SVEC4-10 cells, we treated the cells with conditioned media harvested from TG1-1 cell cultures followed by western blot analyses. The MAPK pathway targets assayed for were Mek/pMek and MAPK/pMAPK (Fig 3A and B). There was an upregulation of activation of MEK1/2, as denoted by the p-MEK/MEK ratio (Fig 3A), in SVEC4-10 cells cultured in conditioned media generated from TG1-1 cells treated with a combination of 0.2%, 0.5% and 1.0% ethanol and estrogen as compared to estrogen alone. Surprisingly, this increase in MEK1/2 activation did not carry on downstream to its effector, MAPK, as no major differences were noted in expression of MAPK/pMAPK (Fig 3B). One plausible explanation is that there may be a regulating factor playing a role, which may be a result of an inhibitory phosphorylation of MEK1 by cdk5\textsuperscript{60}.

**Ethanol and estrogen induced TG1-1 secretory factors enhance expression of the cellular migration mediator FAK in SVEC4-10 Cells**

Endothelial focal adhesion kinase (FAK), induced by VEGF, plays a critical role during tumor angiogenesis, as it is involved in cell signaling via the...
PI3 Kinase/Akt signaling pathway and maintaining endothelial cell adhesion during angiogenesis\textsuperscript{61–64}. The importance of FAK in angiogenesis was also observed in experiments using FAK knockout mice in which angiogenesis was inhibited\textsuperscript{65}. We evaluated the expression of FAK in SVEC4-10 cells upon treatment with TG1-1 conditioned media. FAK expression in SVEC4-10 cells was enhanced in a dose dependent manner upon culturing the cells with CM derived from TG1-1 cells treated with a combination of ethanol and estrogen as compared to estrogen alone (Fig 4). These data indicate that enhanced FAK expression could play a role in endothelial cell migration by modulating cytoskeletal proteins, such as actin and microtubules, which are both directly involved in cell motility\textsuperscript{66,67}.

Figure 3. Enhanced MEK Activation in SVEC4-10 cells in the presence of alcohol and estrogen Western blot analysis from whole cell lysates of SVEC4-10 cells cultured in conditioned media from ethanol and estrogen treated TG1-1 cells. Whole cell lysates were assayed for expression of A) MEK Activation; and B) MAPK Activation. Expression was determined by normalizing to SVEC4-10 protein lysates treated with TG1-1 conditioned media cultured in the presence of estrogen alone (n=2) Error bars represent ±SEM.

Ethanol and estrogen induced TG1-1 secretory factors enhance migration but not proliferation in SVEC4-10 cells

In order to evaluate the migratory potential of SVEC4-10 cells in presence of ethanol and estrogen treated TG1-1 conditioned media, we performed a scratch wound assay. Treating SVEC4-10 cells with secretory factors of ethanol and estrogen treated TG1-1 cells demonstrated an enhanced migration compared to ethanol or estrogen alone, as observed by the scratch wound results. In presence of conditioned media generated with 0.2% and 0.5% ethanol in combination with estrogen, the wound closed by 300µm and 400µm respectively. This migration was higher as compared to conditioned media generated by treating TG1-1 cells with ethanol alone, where the wound closed by 100µm for both 0.2% and 0.5% and estrogen alone, where the wound closed by 200µm (Fig 5A). Migration and proliferation form the two foundation pillars for successful angiogenesis and frequently occur simultaneously during angiogenesis\textsuperscript{68–70}. To evaluate the effects of TG1-1 secretory factors on proliferation of SVEC4-10 cells, we performed an XTT Proliferation assay on SVEC4-10 cells cultured in the presence of Tg1-1 conditioned media. Surprisingly, no striking differences in cell proliferation were observed when comparing the different treatment conditions (Fig 5B). These data indicate that TG1-1 secretory factors enhance migration of endothelial cells in presence of ethanol and estrogen in combination.

Figure 4. Alcohol and estrogen enhance expression of FAK in SVEC4-10 cells. Western blot analysis from whole cell lysates of SVEC4-10 cells cultured in conditioned media from ethanol and estrogen treated TG1-1 cells. Whole cell lysates were assayed for expression of FAK, a marker for migration. Expression was determined by normalizing to SVEC4-10 protein lysates treated with TG1-1 conditioned media cultured in the presence of estrogen alone (n=2) Error bars represent ±SEM.

Ethanol and estrogen enhance activation of the pro-survival PI3K/AKT signaling pathway in TG1-1 breast cancer cells

We have provided evidence that treating TG1-1 breast cancer cells with a combination of ethanol and estrogen leads to an increase in the expression of pro-angiogenic factors. Interestingly, the pro-angiogenic factors (VEGF, MMP-9, and eNOS) assayed for in this study require the PI3K/Akt signaling pathway, amongst other signaling pathways, to control their expression\textsuperscript{71–75}. Additionally, AKT phosphorylation leads to the activation of
pro-survival pathways leading to enhanced cancer progression\(^8\,76,\,77\). We thus wanted to evaluate the effects of ethanol and estrogen on activation of Akt in TG1-1 cells when treated with a combination of ethanol and estrogen. Following treatment of TG1-1 cells, whole cell lysates were prepared followed by western blot analyses in order to determine the expression of AKT and phosphorylated Akt (Fig 6). Treatment of TG1-1 cells with a combination of 0.5% and 1.0% ethanol and estrogen led to enhanced activation of AKT compared to estrogen alone.

**Discussion**

Angiogenesis is a highly regulated process exploited by tumors in order to sustain their growth and development, ultimately providing the necessary vasculature needed for metastasis\(^30,\,78,\,79\). Various factors, such as estrogen and ethanol have been shown to modulate angiogenesis. Our laboratory has demonstrated that estrogen mobilizes bone marrow derived endothelial progenitor cells (BM-EPCs) thus resulting in increased neo-vascularization of mammary tumor tissue in mice\(^32\). Studies have demonstrated that ethanol as well is directly linked to the process of angiogenesis by means of increasing the expression of VEGF, monocyte chemoattractant protein 1 (MCP-1), and hypoxia inducible factor 1 alpha (HIF-1\(\alpha\))\(^29,\,38\). Interestingly, despite the available literature, no studies have been undertaken to address the possible correlation between estrogen and ethanol, in combination, on the process of angiogenesis. This comes as a surprise being that literature exists in which epidemiological observations have been made linking estrogen and alcohol in increasing breast cancer initiation and progression\(^7,\,20,\,21,\,80,\,81\).

![Figure 5. SVEC4-10 cell migration is enhanced in the presence of alcohol and estrogen. A) Representative bright field (100X) image of a scratch wound assay using SVEC4-10 cells in presence of conditioned media from ethanol and estrogen treated TG1-1 cells for 24 hours, demonstrating enhanced migration in presence of ethanol and estrogen. (n=2) B) XTT assay using SVEC4-10 cells cultured in presence of conditioned media from ethanol and estrogen treated TG1-1 cells for 24 hours (n=3) Error bars represent ±SEM.](http://www.jcancer.org)
In this study we demonstrate, for the first time, that ethanol enhances estrogen-mediated angiogenesis in breast cancer using the estrogen responsive murine mammary cancer cell line, TG1-1, and the murine endothelial cell line, SVEC4-10. Expression of pro-angiogenic molecules is regulated by multiple cell signaling pathways, including the PI3K/Akt pathway, which is known to play a prominent role in angiogenesis\textsuperscript{72,77,82}. We observed an enhanced activation of the PI3K/Akt pathway, as determined by the phosphorylation status of Akt in TG1-1 when treated with estrogen and ethanol. The enhanced expression of Akt/pAkt observed in response to both ethanol and estrogen correlates with the enhanced expression of pro-angiogenic factors observed in figure 2. Activation of the PI3K/Akt pathway in tumor cells is known in literature to modulate expression of various pro-angiogenic factors\textsuperscript{72,73,77}. Treatment of TG1-1 cells in the combination of estrogen and ethanol resulted in an enhanced expression of VEGF, eNOS, and MMP-9 compared to treatment with either estrogen or ethanol alone.

Vascular endothelial growth factor (VEGF) plays a pivotal role in tumor angiogenesis\textsuperscript{43,44,46,83}. VEGF is also known to function in an autocrine signaling loop in hormone responsive cancers\textsuperscript{84–87} and activate endothelial nitric oxide synthase (eNOS).\textsuperscript{83} Expression of eNOS is classically restricted to the vascular endothelium but has been shown to be expressed in cancers and is positively correlated with cancer aggressiveness\textsuperscript{88,89}. The enhanced expression of eNOS that we observed in response to both estrogen and ethanol could be caused, in part, by the autocrine function of VEGF. Lastly, we observed a substantial increase in the expression of MMP-9 in TG1-1 cells upon treatment with the combination of estrogen and ethanol, compared to either estrogen or ethanol alone. Tumor derived matrix metalloproteinase 9 (MMP-9) has been known to play a crucial role in tumor progression, tissue remodeling, and angiogenesis\textsuperscript{90–92}. The Williams study demonstrated an increased invasive breast cancer risk in women who drink more than 7 alcoholic drinks per week\textsuperscript{24}. MMP-9 has been shown to lead to proteolytic activation of TGFβ, consequently leading to increased invasion as well as angiogenesis\textsuperscript{90,93}, and release of VEGF from ovarian cancer cells\textsuperscript{94}. Based on these observations, we conclude that ethanol and estrogen, in combination, result in activation of the PI3K/Akt pathway, which results in enhanced expression of pro-angiogenic molecules. We speculate that increased MMP9 as a result of the combination of estrogen and ethanol points to increased invasiveness; however this needs to be evaluated further.

Having assayed for the expression of various pro-angiogenic factors, we focused our attention on characterization of SVEC4-10 endothelial cells in response to conditioned media generated by culturing TG1-1 in estrogen and ethanol. The two parameters that we assayed for were tubulogenesis, a direct measure of angiogenesis in vitro, and SVEC4-10 migration and proliferation. We observed increased tubule formation in SVEC4-10 endothelial cells treated with conditioned media generated from TG1-1 cells cultured in a combination of ethanol and estrogen, when compared to TG1-1 cells cultured in either estrogen or ethanol alone. We evaluated migration using the classical scratch wound assay. As with tubulogenesis, enhanced migration of SVEC4-10 cells was observed when cultured in conditioned media obtained from TG1-1 cultured in both ethanol and estrogen together but, no differences in SVEC4-10 proliferation were observed under the same experimental conditions.

The observations made with respect to enhanced tubulogenesis and migration of SVEC4-10 cells prompted us to explore the MAP-Kinase pathway, as is it known to have numerous downstream effectors that target cellular processes including proliferation, migration and survival.\textsuperscript{57} Surprisingly, our data indicate that the media generated from TG1-1 cells treated with a combination of ethanol and estrogen, while demonstrating enhanced MEK activation, failed to translate into activation of its downstream effector MAPK. We believe that this is a phenomenon observed due to the role of cdk5 in angiogenesis. Cdk5 is known to mediate endothelial cell migration\textsuperscript{85}. Phosphorylation of MEK1 by Cdk5, has been demonstrated to lead to its inactivation and thus a downregulation of MAPK activation, pushing cells towards a more migratory phenotype\textsuperscript{60}. Keeping this in mind, it was not surprising when we observed an enhanced migratory capacity in the SVEC4-10 cell plated in ethanol and estrogen treated TG1-1.
conditioned media, but no proliferation differences. Lastly, to validate enhanced migratory capacity of SVEC4-10 cells, we assayed for the expression of focal adhesion kinase (FAK) in response to TG1-1 conditioned media. Enhanced expression of FAK was also observed in cells cultured in conditioned media generated from both ethanol and estrogen treated TG1-1 cells. This is significant as FAK associates with paxillin and cellular integrins leading to enhanced endothelial cell migration.63,96,97

The possible mechanism we propose in outlined in Figure 7, in which the enhanced angiogenic events are initiated by activated AKT in response to both estrogen and ethanol, thus leading to the secretion of pro-angiogenic factors from TG1-1 murine breast cancer cells. In turn, these pro-angiogenic factors act upon SVEC-4-10 murine endothelial cells, resulting in enhanced activation of the MAP kinase pro-survival pathway and FAK expression, consequently enhancing cellular migration and tubulogenesis.

Thus, the experimental observations made in this study support various epidemiological observations in that increased alcohol consumption in post-menopausal women on estrogen replacement therapy have an increased risk of developing breast cancer. We demonstrate that alcohol acts as a promoter of estrogen driven biological processes observed in breast cancer. Specifically, we propose a mechanism by which estrogen and alcohol, in combination, can lead to enhanced angiogenesis in breast cancer. The mechanism reported in this study may indirectly be responsible for the increased risk of breast cancer associated with these women, supporting the later stages that involve a sustained blood supply via angiogenesis. This study does not limit itself to post-menopausal women; in premenopausal women, increased alcohol intake could also enhance various estrogen mediated processes as well as enhance the risk of developing estrogen responsive cancers.

A potentially different pro angiogenic mechanism may be via the NMDA glutamate receptor. The NMDA receptor, vital for cancer survival and growth is known to be expressed on various cancer cells including the estrogen responsive MCF-7 breast cancer cells.96–101. Estrogen is known to upregulate the expression and activity of this receptor in the central nervous tissue. One can speculate that it may have a similar effect on estrogen responsive breast cancer cells. Ethanol has also been demonstrated to upregulate NMDA receptor expression.104 This study opens up multiple avenues for further exploration. Alcohol metabolism begins with the conversion of alcohol to acetaldehyde by the enzymatic action of alcohol dehydrogenase.105 Alcohol dehydrogenase activity is known to be lower in women than in men, leading to a higher blood alcohol level in women. This prolonged exposure could potentiate the pro-tumorigenic effect seen in breast cancer patients. Additionally, the effects of acetaldehyde in combination with estrogen remain to be studied. This study does not include the kinetics and effects of alcohol metabolites in combination with estrogen, an important future direction to pursue.

Our lab has established an in vivo model linking estrogen to enhanced angiogenesis using the estrogen responsive TG1-1 breast cancer cell line. Based on our previous study, we are now planning on performing animal studies aimed at addressing the systemic effects of estrogen and ethanol on breast cancer angiogenesis; the concentration of ethanol in serum and tumor tissue and the effect of both on angiogenesis in breast cancer. Our study opens up a new area of exploration into other components of the tumor microenvironment, such as macrophages. Anti-inflammatory tumor associated M2 macrophages play a role in angiogenesis and tumor progression,107–109 and, can be studied in presence of both ethanol.
and estrogen. This will provide novel intervention strategies that target alcohol metabolism pathways in an effort to dampen angiogenesis and improve patient prognosis.

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Competing Interests

The authors have declared that no competing interest exists.

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