Statins affect ETS1-overexpressing triple-negative breast cancer cells by restoring DUSP4 deficiency

Hae Hyun Jung1,*, Soo-Hyeon Lee2,*, Ji-Yeon Kim, Jin Seok Ahn2, Yeon Hee Park1,2,3 & Young-Hyuck Im2,3

We investigated the molecular mechanisms underlying statin-induced growth suppression of triple-negative breast cancer (TNBC) that overexpress the transcription factor ets proto-oncogene 1 (ets-1) and downregulate dual specific protein phosphatase 4 (dusp4) expression. We examined the gene expression of BC cell lines using the nCounter expression assay, MTT viability assay, cell proliferation assay and Western blot to evaluate the effects of simvastatin. Finally, we performed cell viability testing in TNBC cell line-transfected DUSP4. We demonstrated that ETS1 mRNA and protein were overexpressed in TNBC cells compared with other BC cell lines (P < 0.001) and DUSP4 mRNA was downregulated (P < 0.001). MTT viability assay showed that simvastatin had significant antitumor activity (P = 0.002 in 0.1 μM). In addition, simvastatin could restore dusp4 deficiency and suppress ets-1 expression in TNBC. Lastly, we found that si-DUSP4 RNA transfection overcame the antitumor activity of statins. MAPK pathway inhibitor, U0126 and PI3KCA inhibitor LY294002 also decreased levels of ets-1, phosphor-ERK and phosphor-AKT on Western blot assay. Accordingly, our study indicates that simvastatin potentially affects the activity of transcriptional factors such as ets-1 and dusp4 through the MAPK pathway. In conclusion, statins might be potential candidates for TNBC therapy reducing ets-1 expression via overexpression of dusp4.

Triple-negative breast cancer (TNBC) is defined by a lack of expression of estrogen receptor (ER), progesterone receptor (PgR) and overexpression of human epidermal growth factor receptor-2 (HER-2). This cancer accounts for approximately 15–20% of all breast cancers (BCs)1,2. Although many advances have been made recently in BC treatment, TNBC represents a significant challenge since it is associated with dismal prognosis and no effective targeted therapy has yet been identified. Cytotoxic chemotherapy is currently the only treatment option for TNBC; thus, there is an urgent unmet need to improve treatment outcomes3,4.

Many studies have endeavored to determine the molecular mechanisms underlying the aggressiveness of TNBC. This aggressiveness was recently found to be associated with expression of dual specificity phosphatase 4 (DUSP4), an extracellular-regulated kinase (ERK) phosphatase known to be a negative regulator of ERK. Specifically, decreased dusp4 expression resulted in a loss of negative feedback to the Ras-ERK pathway, which facilitated tumor cell proliferation and survival in TNBCs5. Moreover, patients with basal-like BC who received neoadjuvant chemotherapy and had low levels of dusp4 expression achieved poorer pathologic complete remission rates and shorter recurrence-free survival periods compared with patients with high levels of dusp4 expression.

The ETS family of transcription factors regulates the expression of genes involved in normal cell development, proliferation, and differentiation6. Therefore, the dysregulation of these transcription factors facilitates cell proliferation in many cancers, and several ETS members have been shown to participate in invasion and metastasis by activating gene transcription7. V-ets avian erythroblastosis virus E26 oncogene homology 1 (ETS-1) is the founding member of the ETS family and regulates transcription and DNA binding by binding to ETS-domain proteins8,9. Previous have demonstrated that ETS1 is required for activation of the Ras/ERK pathway and migration of Ras/ERK-activated cells by transcriptional activation through ETS/AP-1 sites10,11. In invasive BC, upregulation

1 Department of Health Sciences and Technology, SAIHST, Sungkyunkwan University, Seoul 06351, Korea. 2Division of Hematology-Oncology, Department of Medicine, Samsung Medical Center, Sungkyunkwan University School of Medicine, Seoul 06351, Korea. 3Biomedical Research Institute, Samsung Medical Center, Seoul 06351, Korea. * These authors contributed equally to this work. Correspondence and requests for materials should be addressed to Y.H.P. (email: yhparkhmo@skku.edu) or Y.-H.I. (email: imyh00@skku.edu)
of \textit{ETS1} is associated with high aggressiveness and poor prognosis. Consistent with this observation, \textit{ETS1} regulates the expression of important angiogenetic and extracellular matrix remodeling factors such as \textit{VEGF}, \textit{MMP2}, and \textit{MMP9}\textsuperscript{12–15}. Furthermore, \textit{ETS1} also represses genes such as \textit{DUSP4}, \textit{DUSP6}, and \textit{SPRY4}, all of which provide negative feedback to the RAS/ERK pathway\textsuperscript{16–18}.

Statins, small molecule inhibitors of 3-hydroxy-3-methylglutaryl-coenzyme A reductase (HMG-CoA\textsubscript{R}), exert antitumor effects by altering the RAS/MEK/ERK and RAS/PIK3CA/AKT signaling pathways\textsuperscript{19,20}. Lipophilic statins inhibit the growth and proliferation of BC cells, especially hormone receptor-negative, basal-like BC cells\textsuperscript{21–23}.

Based on these studies, we hypothesized that statins suppress TNBC growth by altering the expression of \textit{DUSP4} and \textit{ETS1}.

**Results**

**Expression of \textit{DUSP4} and \textit{ETS} family genes in 22 BC cell lines using Western blot, RT-PCR, and nCounter expression assay.** To compare the levels of \textit{dusp4} and \textit{ets-1} expression in different BC subtypes, we performed Western blot and RT-PCR analyses of \textit{ER}, \textit{PgR}, and \textit{HER-2} expression in 22 BC cell lines representing various BC subtypes (Fig. 1A,B). These results were used to classify each of the 22 BC cell lines into TNBC or non-TNBC subtypes.

Western blot analysis showed that \textit{dusp4} expression was downregulated in TNBC cell lines compared with non-TNBC cell lines; the opposite effect was observed for \textit{ets-1} (Fig. 1A).

We also examined the mRNA expression levels of \textit{ETS1} and \textit{DUSP4} in the 22 BC cell lines using quantitative RT–PCR. The relative \textit{ETS1} mRNA expression levels are shown in Fig. 1B. Consistent with the Western blot analysis, RT-PCR revealed that \textit{ets-1} mRNA expression was higher in most TNBC cell lines than in the non-TNBC cell lines. Similarly, \textit{DUSP4} expression was decreased in TNBC cell lines compared with non-TNBC cell lines.

To gain a more comprehensive picture of the differences between TNBC versus non-TNBC cells, we next analyzed the expression of 70 additional genes in the 22 BC cell lines (Supplementary Table 1). Of the 27 \textit{ETS} genes analyzed, \textit{ETS1}, \textit{ETS2}, \textit{ETV1}, \textit{ETV5}, \textit{ELK3}, and \textit{FLI1} were overexpressed in TNBC cell lines [TNBC vs. non-TNBC; false discovery rate (FDR) = < 0.001, 0.014, 0.003, 0.002, < 0.001, and < 0.001, respectively] (Table 1).
and Fig. 1C). In contrast, SPDEF genes were expressed at lower levels in TNBC cell lines than in non-TNBC cell lines (FDR < 0.001). We also observed reduced DUSP4 expression in TNBC cell lines (FDR = 0.008).

Of the 18 cell cycle-related genes examined, CCNA1, CCND2, and CDK6 exhibited significantly higher expression in the TNBC cell lines. Generally, cell cycle-related genes were upregulated in TNBC cell lines.

Antitumor effects of simvastatin on BC cells in vitro. To determine whether statin inhibits cell proliferation, we performed MTT assays. We found that TNBC cells were sensitized to simvastatin compared with non-TNBC cell lines, as indicated by the IC50 values. Six of the 10 TNBC lines tested had an IC50 < 20 μmol/L, whereas all non-TNBC cell lines had an IC50 ≥ 20 μmol/L (Fig. 2A). A colony-forming assay was performed by seeding equal numbers of MDA-MB-231 cells and MCF-7 cells in triplicate on soft agar in 6-well tissue culture plates. After treating the cells with 0.1 μM simvastatin, the MDA-MB-231 cells formed fewer colonies than the MCF-7 cells (Fig. 2B). Thus, stimulation with 0.1 μM simvastatin significantly inhibited colony formation in MDA-MB-231 cells (P = 0.002).

Effects of statin treatment on gene expression in TNBC cell lines. We next sought to determine the effect of simvastatin on the MAP kinase signaling pathway. To this end, we performed a PCR array of 45 genes associated with the MAP kinase signaling pathway using HCC1806 and MDA-MB-231 cells (Table 2). This analysis showed that simvastatin significantly suppressed ETS-1 expression in TNBC cell lines. In addition, we found that the mRNA expression levels of 13 cell cycle-related genes and of the MAP kinase genes MAP2K3 and MAP2K6 were also downregulated by statin treatment.

In addition, we evaluated the effects of statins on dusp4 and ets-1 expression using RT-PCR and Western blot analyses (Fig. 3A,B). In HCC1806 cells, simvastatin treatment (>5 μM) resulted in decreased ets-1 mRNA expression.
expression and increased dusp4 expression (Fig. 3A). MDA-MD-231 cells were more sensitive to simvastatin than HCC1806 cells; 0.4 μM simvastatin was sufficient to affect ets-1 and dusp4 mRNA expression. Western blot analysis also revealed that the effects of simvastatin on ets-1 and dusp4 expression were dose-dependent (Fig. 3B).
Additionally, we evaluated the effects of statin on the gene expression profile in TNBC cells using the nCounter expression assay. Statin treatment restored dusp4 and suppressed ets-1 expression to levels that were not significantly different from those in non-TNBC cells (both $P = 0.068$) (Supplementary Fig. 1).

Effect of ETS1 and DUSP4 expression on the viability of TNBC cells. To determine the effects of ets-1 and dusp4 expression on cell viability, we transfected TNBC cells with ETS1 siRNA. In MTT assay, after transfecting si-ETS1 into HCC1806 and MDA-MB-231 cells, we found that cell viability of both HCC1806 and MDA-MB-231 cells were reduced ($P < 0.05$) (Fig. 4A). We also performed DUSP4 transfection into TNBC cells. We also analyzed cell viability and ets-1 expression after transfection of Hs578t and MDA-MB-231 cells. Transfection of DUSP4 resulted in decreased ets-1 expression in TNBC cells. Moreover, ectopic expression of DUSP4 in the Hs 579t and MDA-MB-231 TNBC cell lines resulted in significantly decreased cell viability compared with empty vector (pcDNA) transfection ($P < 0.05$) (Fig. 4B).

Inhibition of dusp4 expression blunts simvastatin-induced apoptosis. To investigate the mechanism of simvastatin-induced apoptosis, we transfected HCC1806 and MDA-MB-231 cells with DUSP4-targeting siRNA and analyzed apoptosis by FACS (Fig. 5). As expected, simvastatin induced apoptosis in cells transfected with control siRNA. In contrast, cells transfected with DUSP4 siRNA exhibited decreased simvastatin-induced apoptosis. In MDA-MB-231 cells, DUSP4 siRNA transfection inhibited apoptosis in the presence of 0.5, 1, and 5 μM simvastatin ($P < 0.05$), whereas DUSP4 siRNA transfection suppressed apoptosis in HCC1806 cells induced by 1 and 5 μM simvastatin ($P < 0.05$).

Dusp4 and ets-1 expression in TNBC cells after in vitro treatment with simvastatin, a MEK inhibitor, and a PIK3CA inhibitor. To investigate the downstream signaling pathways activated by simvastatin, the expression levels of mitogen-activated protein kinase (MEK) and phosphatidylinositol 3-kinase (PI3K) were evaluated after simvastatin treatment. TNBC cells were treated with 0, 1, 5, or 10 μM simvastatin, and the expression levels of total and phosphorylated Erk and Akt were assessed by Western blot analysis (Supplementary Fig. 2A). Simvastatin treatment decreased the levels of both phosphorylated Erk and Akt in a dose-dependent manner. We next used the specific pharmacological inhibitors U0126 and LY294002 to inhibit the MEK-Erk and PI3K-Akt pathways, respectively (Supplementary Fig. 2B). U0126-mediated inhibition of MEK abrogated ets-1 overexpression and suppressed Erk phosphorylation in a dose-dependent manner. In a similar pattern, LY294002-mediated PI3K/Akt inhibition decreased ets-1 expression in a dose-dependent manner.

Finally, we performed MTT assays to determine the effect of statin treatment in combination with LY294002 on the viability of BT-549, Hs578t, MDA-MB-231, and MCC 1806 cells. Treatment with either simvastatin or LY294002 alone each resulted in decreased cell viability; moreover, simvastatin and LY294002 acted synergistically to inhibit cell proliferation when given together (Supplementary Fig. 2C).
Discussion

Previous studies have shown that ETS family members are significantly upregulated in TNBC cells compared with other cells, whereas DUSP4 is downregulated in TNBC cells\(^5,24\). Our results indicate that statins suppress proliferation of TNBC cell lines by restoring dusp4 expression and suppressing ets-1 upregulation; thus, statins are potential candidate molecules for treating TNBCs.

Our research has several clinical implications for the treatment of TNBCs. ETS1 might have a role in TNBC pathogenesis, especially for phenotypes resistant to conventional treatment, and could be a biomarker for TNBC cells. Increased ets-1 expression is associated with aberrant transcription of multiple cancer-associated genes, which can result in enhanced energy metabolism, cell survival, matrix degradation, cell growth, angiogenesis, migration, and invasion\(^13,25\). ETS1 contains a Ras-responsive phosphorylation site at threonine-38, phosphorylation of which strongly increases the transcriptional activity of ets-1. In a complex signal transduction network, ets-1 is an effector of oncogenic Ras via the MAPK pathway\(^26,27\), a main mediator of Ras signaling that is dysregulated in many cancers, particularly BC. In addition, loss of DUSP4 also promotes MARK pathway activation and induces the cancer stem cell-like phenotype of TNBC\(^24\). Accordingly, the negative correlation of ets-1 and dusp4 expression in TNBC could plausibly contribute to the aggressive nature of TNBC. In this study, we found that ETS1 and other ETS family members such as ETS2, ETV1, ETV5, ELK3, and FLI1 were overexpressed in TNBC cell lines compared with other BC cell lines. Therefore, the elevated expression of ETS family members (e.g., ETS1) might contribute to chemotherapy resistance as well as tumor invasion and aggressiveness in TNBC cells\(^28–30\).

Our results imply that statins, which are small molecule inhibitors of HMG-CoAR, might be able to potently inhibit TNBC growth by restoring dusp4 expression and downregulating ets-1 expression. Dusp4 expression is consistently lost in TNBC cells\(^31\), which has been reported to promote resistance to treatment\(^7\). In addition, statin treatment has been shown to alter Dusp4 expression\(^32\). We previously reported that simvastatin is a promising candidate for treating TNBCs, especially those with wild-type pten expression\(^33\). In the present study, we used a PCR array and an nCounter expression assay to demonstrate that dusp4 expression is lost in TNBC cells. We propose that statin effectively suppresses ets-1 expression by restoring dusp4 levels in TNBC cells.

Finally, our results indicate that ets-1 overexpression and dusp4 deficiency, a combination that might contribute to treatment failure in refractory TNBCs, can be overcome by statin treatment with mitogen-activated protein kinase (MEK) inhibitors. Since ETS family members can affect the MAPK pathway\(^28\), MEKs are potential therapeutic targets for treating TNBC.

In summary, our results suggest that the loss of dusp4, a potential biomarker of treatment-resistant TNBC, is associated with ets-1 overexpression via the PI3K and MAPK pathways. Statin, a small inhibitor of HMG-CoAR, is a potential therapeutic candidate for treatment-resistant TNBC because it can reverse ets-1 overexpression by restoring dusp4 expression.
Materials and Methods

Human TNBC cell lines and culture. Twenty-two BC cell lines were purchased from the American Type Culture Collection (Manassas, VA, USA). Each cell line was phenotypically characterized as either TNBC or non-TNBC, BT-549, Hs578T, MDA-MB-157, MDA-MB-231, MDA-MB-468, HCC1806, HCC38, HCC70, HCC1395, and HCC1937 were TNBC cell lines, whereas SK-BR-3, UACC-893, ZR-75-30, HCC1419, MDA-MB-361, BT-474, CAMA-1, HCC1500, MCF-7, T-47D, and ZR-75-1 were non-TNBC cell lines. All cells were grown in RPMI-1640 medium supplemented with 10% fetal bovine serum (FBS), 100 mU/mL penicillin, and 100 μg/mL streptomycin under 5% CO₂ in humidified air at 37 °C. All media and supplements were purchased from Invitrogen Corp. (Carlsbad, CA, USA).

Antibodies and reagents. Anti-ER antibodies were purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA). Anti-ets-1 antibodies were purchased from Abcam (Cambridge, UK). Anti-phospho-ERK1/2 (Thr202/Thr204), anti-ERK1/2, anti-Akt (Ser473), and anti-Akt antibodies were purchased from Cell Signaling Technology (Beverly, MA, USA). Anti-β-actin antibodies and simvastatin were purchased from Sigma-Aldrich (St. Louis, MO, USA); U0126 and LY294002 were from Cell Signaling Technology.

nCounter expression assay (NanoString). The expression levels of 70 genes previously reported as belonging to the ETS family or associated with the ETS family (Table 1) were tested in 22 cell lines using the NanoString nCounter Analysis System (NanoString Technologies, Seattle, WA, USA)34.
The effects of simvastatin treatment (10 μmol, 24 hr) on the expression profiles of these 70 genes in TNBC cell lines were also investigated using the nCounter expression assay (NanoString). NanoString nCounter gene expression data were filtered using quality control (QC) criteria according to the manufacturer’s recommendations. Raw counts of QC-passed samples were normalized using five reference genes as internal controls (GUSB, PUM1, TBP, TFFC, and TUBB). All QC and normalization procedures were performed using nSolver Analysis Software v2.0 (NanoString Technologies); all data were log2-transformed before further analysis. The student's t-test was used to compare normalized expression values between TNBC and non-TNBC cell lines. P-values were adjusted using the false discovery rate (FDR) method for multiple comparisons35. Genes with FDRs below 0.001 and absolute log fold change greater than 2 were considered to be differentially expressed and are displayed on the volcano plot. Among the genes related to the ETS family, the cell cycle, and signaling pathways, a hierarchically clustered heat map was generated using genes whose adjusted p-values were less than 0.1. The paired t-test was applied to identify genes that were differentially expressed before versus after simvastatin treatment. Differential expression was defined as a p-value less than 0.05 and an absolute log2 fold change greater than 1.

Intrinsic subtype classification was performed using the PAM50 predictor, as described in Parker et al.36. R version 3.0.2 (http://www.R-project.org/) was used to perform all statistical tests, generate all plots, and predict all PAM50 subtypes.

Isolation of RNA and reverse transcription polymerase chain reaction (RT–PCR). Total cellular RNA was isolated using Trizol (GIBCO BRL/Life Technologies, Carlsbad, CA, USA). For RT–PCR, 2 μg of RNA was treated with RNase-free DNase and cDNA was synthesized with the Omniscript Reverse Transcription kit (QIAGEN). The levels of DUSP4, ets-1, and glyceraldehyde 3-phosphate dehydrogenase (GAPDH; loading control) cDNA were quantified with an ABI Prism 7900 Real-Time PCR system (Applied Biosystems, Foster City, CA, USA). All primers and probes (GAPDH cat. no. Hs99999905-m1; DUSP4 cat. no. Hs01027785; ets-1 cat. no. Hs00428293) were obtained commercially and are proprietary; thus, sequences are not available (TaqMan Gene Expression Assay, Applied Biosystems). The following amplification conditions were used: 50 °C for 2 min and 95 °C for 10 min, followed by 40 cycles of 94 °C for 15 sec and 60 °C for 1 min. Data were analyzed with ABI Prism 7900 SDS 2.3 software (Applied Biosystems).

Western blot analysis. For Western blot analysis, cells were lysed with RIPA buffer (0.5% sodium deoxycholate, 1% Nonidet P-40, 150 mM NaCl, 50 mM Tris pH 7.5, 0.1% sodium dodecyl sulfate [SDS], and 1 mM PMSF). The protein concentrations of the supernatants were determined using the BCA protein assay (Pierce, Rockford, IL, USA). Equal amounts of protein were loaded onto SDS–polyacrylamide gels and transferred to nitrocellulose membranes. Nonspecific binding sites were blocked by incubating the membranes in blocking solution containing 5% nonfat dry milk for 2 h, after which membranes were incubated with primary antibodies. After incubation with horseradish peroxidase (HRP)-conjugated secondary antibodies, immunoreactive bands were visualized using enhanced chemiluminescent (ECL) reagents (Amersham). To confirm equal protein loading, blots were stripped and reprobed with anti-beta-actin antibodies.

Cell proliferation and colony formation assays. The effect of simvastatin on BC cell growth in vitro was determined by measuring the absorbance of 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) in living cells. Briefly, cells were seeded (3 × 10^4 cells per well) on 96-well microtiter plates (Nunc, Roskilde, Denmark). After 72 h of drug exposure, 5 μg/mL MTT (Sigma-Aldrich) solution was added to the culture medium (10 μL per 100 μL of medium), and the plates were incubated for an additional 4 h at 37 °C. Excess MTT solution was aspirated. Formazan crystals in viable cells were dissolved in 100 μL of acid-isopropanol (0.04 N HCl in isopropanol), and the absorbance at 540 nM was measured. The half-maximal inhibitory concentration (IC50) values were estimated using Prism software (Version 4.0, GraphPad Software Inc., San Diego, CA, USA).

For colony formation assays, cells were plated in 6-well plates (Nunc; 2 × 10^4 cells/well for MCF7 cells and 1 × 10^5 cells/well for MDA-MB-231 cells) and treated with 0.1 μM simvastatin for 3 days. Triple cultures of each cell type were maintained at 37 °C for 14 days under 5% CO2 in humidified air. Fresh medium was added after 7 days, and cells were stained with 0.1% crystal violet. Colonies, defined as groups of at least 50 cells, were counted under an inverted phase contrast microscope (CK × 31SF; Olympus Biosystems, Hamburg, Germany). The percentage of relative cell proliferation is expressed as (number of colonies from treated cells/number of colonies from controls) × 100. Each assay was repeated three times.

Ectopic expression of DUSP4 in TNBC cell lines. The DUSP4-pCMV6-Entry expression plasmid was purchased from Origene. Cells were transfected with 4 μg of the DUSP4-pCMV6-Entry plasmid using the Neon Transfection System (Invitrogen). Briefly, cells (2 × 10^4 per transfection) were harvested, washed with PBS, resuspended in 100 μl of resuspension buffer, and electroporated according to the manufacturer's instructions.

siRNA transfection. TNBC cells were transfected with siGENOME SMARTpool silencing (si)RNA targeting human ETS1 or DUSP4. Alternatively, cells were transfected with control siRNA. All siRNAs were purchased from Dharmacon (Lafayette, CO, USA). Transfections were performed using the Neon Transfection System according to the manufacturer's instructions.

Apoptosis assay. TNBC cells were transfected with DUSP4 siRNA. At 24 h post-transfection, cells were treated with simvastatin for an additional 48 h. For FACS analysis of cell death, cells were stained with Annexin V-FITC (BD Biosciences) and propidium iodide (PI, Sigma) following the manufacturer's protocol. Briefly, 100 μl of cell suspension (1 × 10^5 cells in 1X binding buffer) was stained with 2.5 μl of Annexin V-FITC and 5 μl of PI [50 μg/mL]. The suspension was gently mixed and incubated for 15 min at room temperature in the dark.
after which 200 μl of 1X Annexin binding buffer was added. Cells were immediately analyzed on a FACS VERSE flow cytometer. Data from 10,000 cells were collected at a low flow rate and analyzed using BD FACSuite (BD Biosciences).

**Statistical analysis.** Comparisons between experimental groups and the relevant controls were performed using the student’s t-test. P values < 0.05 were considered to indicate significant differences. All experiments were performed at least three times.

**References**

1. Perou, C. M. et al. Molecular portraits of human breast tumours. *Nature* **406**, 747–752 (2000).

2. Kreike, B. et al. Gene expression profiling and histopathological characterization of triple-negative/basal-like breast carcinomas. *Breast cancer research: BCR* **9**, R65 (2007).

3. Hudis, C. A. & Gianni, L. Triple-negative breast cancer: an unmet medical need. *The oncologist* **16** Suppl 1, 1–11 (2011).

4. Bauer, K. R., Brown, M., Cress, R. D., Parise, C. A. & Caggiano, V. Descriptive analysis of estrogen receptor (ER)-negative, progesterone receptor (PR)-negative, and HER2-negative invasive breast cancer, the so-called triple-negative phenotype: a population-based study from the California cancer Registry. *Cancer* **109**, 1721–1728 (2007).

5. Balko, J. M. et al. Profiling of residual breast cancers after neoadjuvant chemotherapy identifies DUSP4 deficiency as a mechanism of drug resistance. *Nature medicine* **18**, 1052–1059 (2012).

6. Sharrocks, A. D. The ETS-domain transcription factor family. *Nature reviews Molecular cell biology* **2**, 827–837 (2001).

7. Okawa, T. & Yamada, T. Molecular biology of the Ets family of transcription factors. *Gene* **303**, 11–34 (2003).

8. Goetz, T. L., Gu, T. L., Speck, N. A. & Graves, B. J. Auto-inhibition of Ets-1 is counteracted by DNA binding cooperativity with core-binding factor alpha2. *Molecular and cellular biology* **20**, 81–90 (2000).

9. Kim, W. Y. et al. Mutual activation of Ets-1 and AML1 DNA binding by direct interaction of their autoinhibitory domains. *The EMBO journal* **18**, 1609–1620 (1999).

10. Chang, F. et al. Signal transduction mediated by the Ras/Raf/MEK/ERK pathway from cytokine receptors to transcription factors: potential targeting for therapeutic intervention. *Leukemia* **17**, 1263–1293 (2003).

11. Hollenhorst, P. C., McIntosh, L. P. & Graves, B. J. Genomic and biochemical insights into the specificity of ETS transcription factors. *Annual review of biochemistry* **80**, 437–471 (2011).

12. Span, P. N. et al. Expression of the transcription factor Ets-1 is an independent prognostic marker for relapse-free survival in breast cancer. *Oncogene* **21**, 8506–8509 (2002).

13. Park, Y. H., Jung, H. H., Ahn, J. S. & Im, Y. H. Ets1 regulates HER2-induced MMP-1 expression in breast cancer cells. *Biochemical and biophysical research communications* **377**, 389–394 (2008).

14. Hollenhorst, P. C. et al. Oncogenic ETS proteins mimic activated RAS/MAPK signaling in prostate cells. *Genes & development* **25**, 2147–2157 (2011).

15. Selvaraj, N., Budka, J. A., Ferris, M. W., Jerde, T. J. & Hollenhorst, P. C. Prostate cancer ETS rearrangements switch a cell migration gene expression program from RAS/ERK to PI3K/ AKT regulation. *Molecular cancer* **13**, 61 (2014).

16. Plotnik, J. P., Budka, J. A., Ferris, M. W. & Hollenhorst, P. C. ETS1 is a genome-wide effector of RAS/ERK signaling in epithelial cells. *Nucleic acids research* **42**, 11928–11940 (2014).

17. Yanae, M. et al. Statin-induced apoptosis via the suppression of ERK1/2 and Akt activation by inhibition of the geranylgeranyl-pyrophosphate biosynthesis in glioblastoma. *Journal of experimental & clinical cancer research: CR* **30**, 74 (2011).

18. Tsubaki, M. et al. Blockade of the Ras/MEK/ERK and Ras/PI3K/Akt pathways by statins reduces the expression of bFGF, HGF, and TGF-beta as angiogenic factors in mouse osteosarcoma. *Cytokine* **54**, 100–107 (2011).

19. Goard, C. A. et al. Identifying molecular features that distinguish flavustatin-sensitive breast tumor cells. *Breast cancer research and treatment* **113**, 301–312 (2014).

20. Campbell, M. J. et al. Breast cancer growth prevention by statins. *Cancer research* **66**, 8707–8714 (2006).

21. Garwood, E. R. et al. Fluvastatin reduces proliferation and increases apoptosis in women with high grade breast cancer. *Breast cancer research and treatment* **119**, 137–144 (2010).

22. Balko, J. M. et al. Activation of MAPK pathways due to DUSP4 loss promotes cancer stem cell-like phenotypes in basal-like breast cancer. *Cancer research* **73**, 6346–6358 (2013).

23. Verschoor, M. L., Wilson, L. A., Verschoor, C. P. & Singh, G. Ets-1 regulates energy metabolism in cancer cells. *PloS one* **5**, e13565 (2010).

24. Yang, B. S. et al. Ras-mediated phosphorylation of a conserved threonine residue enhances the transactivation activities of c-Ets1 and c-Ets2. *Molecular and cellular biology* **16**, 538–547 (1996).

25. Wasylyk, C., Bradford, A. P., Gutierrez-Hartmann, A. & Wasylyk, B. Conserved mechanisms of Ras regulation: evolutionarily related transcription factors, Ets1 and pointed P2. *Oncogene* **14**, 899–913 (1997).

26. Wallace, J. A. et al. Ets2 in tumor fibroblasts promotes angiogenesis in breast cancer. *PloS one* **8**, e71533 (2013).

27. Yuan, Z. Y. et al. Overexpression of ETV4 protein in triple-negative breast cancer is associated with a higher risk of distant metastasis. *OncoTargets and therapy* **7**, 1733–1742 (2014).

28. Scheiber, M. N. et al. FLI1 expression is correlated with breast cancer cellular growth, migration, and invasion and altered gene expression. *Neoplasia (New York, NY)* **16**, 801–813 (2014).

29. Manzano, R. G., Martinez-Navarro, E. M., Forteza, J. & Brugarolas, A. Microarray phosphatome profiling of breast cancer patients unveils a complex phosphatase regulatory role of the MAPK and PI3K pathways in estrogen receptor-negative breast cancers. *International journal of oncology* **45**, 2250–2266 (2014).

30. Bjarnadottir, O. et al. Global Transcriptional Changes Following Statin Treatment in Breast Cancer. *Clin Cancer Res* **21**, 3402–3411 (2015).

31. Park, Y. H., Jung, H. H., Ahn, J. S. & Im, Y. H. Statin induces inhibition of triple negative breast cancer (TNBC) cells via PI3K pathway. *Biochemical and biophysical research communications* **439**, 275–279 (2013).

32. Park, Y. H. et al. A seven-gene signature can predict distant recurrence in patients with triple-negative breast cancers who receive adjuvant chemotherapy following surgery. *International journal of cancer journal international du cancer* **136**, 1976–1984 (2015).

33. Benjamin, Y. & Hochberg, Y. Controlling the False Discovery Rate - a Practical and Powerful Approach to Multiple Testing. *J Roy Stat Soc B Met* **57**, 289–300 (1995).

34. Fisher, B. et al. Supervised risk predictor of breast cancer based on intrinsic subtypes. *Journal of clinical oncology: official journal of the American Society of Clinical Oncology* **27**, 1160–1167 (2009).
Acknowledgements
This study was supported by a grant from Samsung Biomedical Research (SBRI C-A9-234-3).

Author Contributions
Y.-H.I. and Y.H.P. conceived the study design and oversaw its conduct. Y.H.P. wrote the first draft of the paper and Y.H.P. and J.-Y.K. wrote and reviewed the final draft of the paper. H.H.J. and S.-H.L. conducted experiments and analyzed data. H.H.J. and J.-Y.K. contributed to data interpretation, statistical analysis, and preparation of the figures and tables. J.S.A. and Y.-H.I. contributed to the study design and conduct.

Additional Information
Supplementary information accompanies this paper at http://www.nature.com/srep

Competing financial interests: The authors declare no competing financial interests.

How to cite this article: Jung, H. H. et al. Statins affect ETS1-overexpressing triple-negative breast cancer cells by restoring DUSP4 deficiency. Sci. Rep. 6, 33035; doi: 10.1038/srep33035 (2016).

This work is licensed under a Creative Commons Attribution 4.0 International License. The images or other third party material in this article are included in the article’s Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit http://creativecommons.org/licenses/by/4.0/

© The Author(s) 2016