E-cadherin-Fc chimera protein matrix enhances cancer stem-like properties and induces mesenchymal features in colon cancer cells

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Abstract
Cancer stem cells (CSC) are a subpopulation of tumor cells with properties of high tumorigenicity and drug resistance, which lead to recurrence and poor prognosis. Although a better understanding of CSC is essential for developing cancer therapies, scarcity of the CSC population has hindered such analyses. The aim of the present study was to elucidate whether the E-cadherin-Fc chimera protein (E-cad-Fc) enhances cancer stem-like properties because studies show that soluble E-cadherin stimulates human epithelial growth factor receptor (EGFR) and downstream signaling pathways that are reported to play a crucial role in CSC. For this purpose, we used ornithine decarboxylase (ODC)-degron–transduced (Degron(+)) KM12SM cells as a CSC model that retains relatively low CSC properties. Compared to cultures without E-cad-Fc treatment, we found that E-cad-Fc treatment further suppressed proteasome activity and largely enhanced cancer stem-like properties of ODC-degron–transduced KM12SM cells. These results include increased expression of stem cell markers Lgr5, Bmi-1, SOX9, CD44, and CD44v9, aldehyde dehydrogenase (ALDH), and enhancement of robust spheroid formation, and chemoresistance to 5-fluorouracil (5-FU) and oxaliplatin (L-OHP). These effects could be attributed to activation of the EGFR pathway as identified by extensive phosphorylation of EGFR, ERK, PI3K, AKT, and mTOR. In SW480 cells, E-cad-Fc matrix induced some CSC markers such as CD44v9 and ALDH. We also found that E-cad-Fc matrix showed high efficiency of inducing mesenchymal changes in colon cancer cells. Our data suggest that the E-cad-Fc matrix may enhance CSC properties such as enhancement of chemoresistance and sphere formation.

KEYWORDS
cancer stem cell, colon cancer, E-cadherin-Fc, epithelial-mesenchymal transition, extracellular matrix
1 | INTRODUCTION

Epithelial-mesenchymal transition (EMT) is a cellular process first discovered in embryogenesis and is increasingly recognized as the main factor in cancer progression. Cancer cells with mesenchymal-like phenotype present a loss of tight cell-cell adhesion and show cellular polarity reversal with enhanced expression of EMT-activating transcription factors such as SNAIL, TWIST and ZEB families. The process of EMT facilitates invasiveness and migration capability and is thus considered a trigger of cancer metastasis and recurrence.

Cumulative evidence suggests that EMT is linked to cancer stem cells (CSC). EMT process is considered to be involved in altering the microenvironment and regulating CSC initiation. CSC undergo self-renewal and produce differentiated cells by asymmetric division. Because CSC have properties such as drug resistance and high tumorigenicity, they are associated with cancer recurrence and poor prognosis. Decades of research have shown that the CSC niche microenvironment plays a pivotal role in their development, offering the possibility of supporting their stem cell status. Several studies have indicated that CSC stem cell properties could be enhanced through the extracellular matrix (ECM). For instance, endothelial-derived ECM propagates the population of mesenchymal stem cells (MSC) by providing MSC with a perivascular niche. Cancer cells cultured on hepatocyte-derived ECM show increased expression of stem cell markers Lgr5 and CD133. Additionally, feeder cells such as fibroblasts or laminin-511-coated matrix support growth of human pluripotent embryonic stem cells (hPSC). These findings indicate an essential role of the ECM in supporting stem cells by cell-cell or cell-matrix interactions.

Recently, E-cadherin Fc chimera protein (E-cad-Fc) has been noted as cell culture material for stem cell maintenance. E-cadherin is an epithelial cell-cell adhesion molecule that is frequently downregulated in many cancers. Its expression is inversely correlated with EMT. Culturing with the E-cad-Fc matrix satisfactorily retains the undifferentiated state of embryonic stem (ES) cells and maintains isolation of adhered cells without colony formation. It is notable that E-cad-Fc matrix promotes adhesion and proliferation of MSC compared with Matrigel. Studies have also shown that cell adherence and junctions between cancer cells are disturbed by addition of the soluble fragment of E-cadherin, leading to malignancy in skin squamous cell carcinoma and breast cancer, possibly by activation of EGFR and its downstream signaling pathways.

Based on the above findings, in the present study, we hypothesized that the E-cad-Fc matrix might serve as potential material for enhancement or maintenance of CSC properties. In parallel, we investigated the effect of E-cad-Fc matrix on EMT. For this purpose, we used colon cancer SW480 cells because the cells readily undergo EMT with transforming growth factor-beta (TGF-β)1 and EGF cocktail. We also used ornithine decarboxylase (ODC)-degron–transduced KM12SM colon cancer cells that show relatively weak CSC properties in this system. Low proteasome activity (LPA) is considered a hallmark of CSC in human cancers, including gastrointestinal cancer, cervical cancer, and osteosarcoma. By using the ZsGreen-labeled degron (Gdeg) proteasome reporter system, we and other groups showed that compared with non-LPA cells, colon and pancreatic cancer cells with LPA showed strong stem cell potential, enhanced chemo- or radio-resistance, and upregulated expression of stem cell markers such as CD44, CD133, or ALDH.

2 | MATERIALS AND METHODS

2.1 Cell culture

KM12SM cells acquire highly metastatic ability through mouse serial transplantation of primary colon cancer KM12C cells. KM12SM was a kind gift from Professor T. Minamoto (Cancer Research Institute, Kanazawa University, Ishikawa Japan). Human colon cancer cell line SW480 was purchased from ATCC. Cells were maintained in DMEM supplemented with 10% FBS and Myco-Zap Plus CL (Lonza) at 37°C in a humidified 5% CO₂ atmosphere. The E-cad-Fc fusion protein was purchased from Somar Corporation, Ltd and partially donated by Prof. T. Akaike (FAIS, Ibaraki, Japan).

2.2 Epithelial-mesenchymal transition induction

Epithelial-mesenchymal transition induction assay was conducted according to a previous study. Briefly, after cells were seeded, they were treated with TGF-β1 (2.5 ng/mL; Sigma-Aldrich) and epithelial growth factor (EGF) (10 ng/mL; Sigma-Aldrich) for 48 hours. To form the E-cad-Fc matrix, culture dishes were coated with the E-cad-Fc fusion protein at a concentration of 10 μg/mL for 2 hours at room temperature. Dishes were washed gently with PBS twice, then cells were seeded and grown for the indicated time.

2.3 Transduction of the degron reporter

The degron sequence of ODC is recognized directly by proteasomes, which leads to immediate destruction of the involved protein. The retroviral expression vector pQCXIN-ZsGreen-cODC, containing green fluorescence Gdeg, was kindly provided by Dr Frank Pajonk (Jonsson Comprehensive Cancer Center, UCLA, CA, USA) to S. Tanaka. The vector was transfected into platinum retroviral packaging cells, and the retrovirus collected from the supernatant was used to infect KM12SM cells. Stable transfectants were selected with G418 solution (Roche). The top 0.86% of cell populations of the EGF channel were enriched by sorting three times using flow cytometry (Cell Sorter SH800; SONY) and maintained in 0.1 mg/mL G418 solution.

2.4 Cell morphology

Cells were cultured in polystyrene tissue culture (TC) dishes with a TC-treated surface (IWAKI) or in E-cad-Fc coated dishes. After treatment for 24, 48, and 72 hours, microscopy observation was carried out using BZ-X700 (Keyence).
2.5 Western blot analysis

Cell lysates were extracted by RIPA buffer with 1% Halt Protease Inhibitor Cocktail Kit (Thermo Fisher Scientific). For detection of phosphorylation, 1% Halt Phosphatase Inhibitor Cocktail (Thermo Fisher Scientific) was added to the lysates. Cell extracts were loaded onto Mini-Protean TGX 4%-15% gels (Bio-Rad) and transferred using the Trans-Blot Turbo Blotting System (Bio-Rad). Primary antibodies used in this study are shown in Table S1. Secondary antibodies were incubated with ECL substrate (Bio-Rad), and bands were visualized using the ChemiDoc Touch Imaging System (Bio-Rad). Images were processed with Image Lab 5.2.1 software.

2.6 RNA extraction and qRT-PCR

Total RNA was collected from cultured cells using TRIzol Reagent (Thermo Fisher Scientific), and complementary DNA was synthesized from 1.0 μg total RNA using oligo dT primer and a Reverse Transcription System (Promega) according to the manufacturer’s instructions. Quantitative real-time PCR (q-PCR) was carried out using LightCycler FastStart DNA Master SYBR Green I (Roche) or LightCycler TaqMan Master (Roche) on a LightCycler 2.0 II (Roche).

Expression of the target gene was normalized relative to β2M mRNA expression using the 2−ΔΔCt method. Primers are shown in Table S2.

2.7 Flow cytometric analysis

Cells were suspended in FluoroBrite DMEM (Thermo Fisher Scientific) with 10% FBS. For analysis of CSC markers, rat antihuman CD44v9/RV3 (LKG-M001; Cosmo Bio Co., Ltd) was used as the primary antibody and phycoerythrin (PE)-conjugated mouse antirat IgG2a (Becton Dickinson) as the secondary antibody. BV421-conjugated mouse antihuman CD44 (Becton Dickinson) and PE-conjugated mouse antihuman CD24 (Becton Dickinson) were used as controls. Data were collected from three independent experiments. Stem cell marker ALDH was detected using ALDEFLUOR Kit (STEMCELL Technologies) according to the manufacturer’s protocol. All samples included a test group and negative control treated with ALDH inhibitor, diethylaminobenzaldehyde (DEAB). Immunofluorescent intensity was measured by flow cytometry.

2.8 Spheroid culture

Serum-free DMEM/F12 medium was added to ultra-low attachment 96-well plates. Cells were then suspended in DMEM and added to wells at a density of 1000 cells per well. In the E-cad-Fc–treated group, cells were transferred to ultra-low attachment plates 48 hours after incubation. Cells were cultured for 18 days. Images were acquired digitally using BX-Z analysis software.

2.9 Proliferation assay

Cells were suspended with DMEM, and 5000 cells of each group were plated into a 96-well plate. Medium was changed every 2 days. Cell proliferation was measured every 24 hours with CCK-8 (Dojindo) according to the manufacturer’s protocol. Absorbance (450 nm) was measured using a spectrophotometer.

2.10 Chemosensitivity assay

A total of 8000 cells per group was seeded onto a 96-well plate and treated with chemotherapeutic drugs at concentrations of 0.5-256 μmol/L for 5-FU and 0.25-100 μmol/L for oxaliplatin (L-OHP). Cells were treated for 72 hours, then measured with CCK-8 (Dojindo).

2.11 Proteasome activity assay

One million cells of each group were suspended with 500 μL of 0.5% NP-40, then centrifuged at 15 300 g for 15 minutes at 4°C to remove insoluble cellular debris. Cell lysates were collected to detect proteasome activity according to the manufacturer’s protocol.

**FIGURE 1** Effects of E-cadherin-Fc chimera protein (E-cad-Fc) on epithelial-mesenchymal transition (EMT) and cancer stem cell markers in SW480 colon cancer cells. SW480 cells were divided into three groups, control group, E-cad-Fc(+) group, and epithelial growth factor (EGF) + transforming growth factor (TGF)β1(+) group. A, Time course study of cell morphology (24, 48, and 72 h). Cells were cultured on normal tissue culture plate (left), cultured on a non-treated plate coated with E-cad-Fc matrix (middle), or cultured on normal tissue culture plate with medium supplemented by EGF and TGF-β1 (right). Mesenchymal-like cells that showed a spindle-like shape appeared as time passed in the E-cad-Fc(+) and EGF + TGF-β1(+) groups. B, Expressions of epithelial or mesenchymal markers at the protein level when treated for 48 h with E-cad-Fc matrix or EGF and TGF-β1. Quantitative analysis in three independent experiments is shown in the right panel. N-cadherin expression was significantly upregulated in the E-cad-Fc–treated group (P < 0.05). C, Expression of epithelial or mesenchymal markers at the mRNA level was detected after 24 h treatment using quantitative real-time PCR assay. Expression of the target gene was normalized relative to β2M mRNA expression using the 2−ΔΔCt method. Data were collected from two independent experiments. **P < 0.001. D, Expression of cancer stem cell markers. Left panel: CD44v9, CD44s, and CD24 were detected by immunostaining using flow cytometry after 48 h treatment. Results are shown from two independent experiments as a ratio of immunofluorescence intensity of Degron(+) E-cad-Fc(+) group to that of Degron(−) control group. ***P < .001. Right panel: Stem cell marker aldehyde dehydrogenase (ALDH) expression. Cells of each group were divided into test group and negative control group blocked by DEAB (ALDH inhibitor). ALDH positive rate: Degron(−) control, 15.55%; Degron(+) E-cad-Fc(+), 19.99%
Each reaction well (96-well plate [opaque white; Thermo Fisher Scientific]) was mixed with 30 μL sample lysate and 70 μL assay buffer provided with the kit, and negative control cell lysate was measured in the presence of proteasome inhibitor. Proteasome activity (mU) per one million cells was calculated.

2.12 Statistical analysis

Data are presented as mean ± SEM. Statistical significance of differences was calculated using one-way ANOVA followed by Bonferroni’s correction for multiple comparisons. Statistical analyses were carried
We first examined EMT-related changes in morphology and related molecular events in SW480 cells. Because EGF plus TGF-β1 cocktail is widely recognized as an EMT inducer, we used this treatment as a positive control. We found that compared with control cells, both E-cad-Fc matrix-treated cells and EGF + TGF-β1-treated cells showed spindle-like morphology, a hallmark of EMT and this change became evident as the time passed (Figure 1A). Western blot analyses showed that EGF + TGF-β1-treated cells showed increased expression of mesenchymal markers, N-cadherin, Snail1, and Vimentin to some extent (Figure 1B). Because SW480 cells expressed a relatively low level of E-cadherin, apparent reduction in E-cadherin was not noted by E-cad-Fc treatment. Quantitative analysis of band intensity in three independent experiments indicated that N-cadherin protein expression was significantly upregulated in the E-cad-Fc–treated group (P < .05, Figure 1B). RT-PCR assay in two independent experiments indicated concordant results to those obtained by western blotting. Thus, N-cadherin mRNA was significantly upregulated by E-cad-Fc treatment (P < .001, Figure 1C).

We next examined expression of CSC-associated cell surface markers including CD44v9, CD44s,36,37 and CD24.38 Flow cytometric analysis showed that the E-cad-Fc–treated group showed an increase in CD44v9 expression (P < .001) and ALDH activity (from 15.55% to 19.99%), another hallmark of CSC.39

As the second cell line, we used ODC-degron–transduced KM12SM cells because these cells acquire insufficient CSC properties even with the ODC-degron system. Time-course study showed that only the Degron(+) E-cad-Fc(+) group showed spindle-like morphology (Figure 2A). Western blot analysis indicated that the Degron(+) E-cad-Fc(+) group, compared to the other groups, highly expressed mesenchymal markers, including N-cadherin, Vimentin, ZEB1, Snai1, Slug, and Twist1 while E-cadherin expression was mostly reduced (Figure 2B). Notably, both latent (45 kDa) and active (26 kDa) forms of TGF-β1 increased in the Degron(+) E-cad-Fc(+) group. Quantitative analysis in two independent experiments indicated that there was >twofold increase in protein expression of Vimentin, Snai1, Twist1, TGF-β1 in the Degron(+) E-cad-Fc(+) group (Figure 2B). These changes of mesenchymal markers were also noted in the Degron(−) E-cad-Fc(+) group (Figure S1). The RNA study showed concordant results except for vimentin and TGF-β1 (Figure 2C). Time-course study showed that Vimentin mRNA increased at 48 hours, but TGF-β1 mRNA did not increase throughout the time point examined (Figure S2).

We then examined the expression of the CSC markers Lgr5, Bmi-1,40,41 and SOX942 by RT-PCR. ODC-degron–transduced cells increased SOX9 mRNA expression (Figure 3A; P < .05), whereas Degron(+) E-cad-Fc(+) cells showed a significant increase in Lgr5, Bmi-1, and SOX9 mRNA expression compared with the Degron(−) control cells (Figure 3A; *P < .05, **P < .01). Flow cytometric analysis showed that the Degron(+) group showed no increase in CSC surface markers; however, the Degron(+) E-cad-Fc(+) group had significantly higher expression of CD44v9 and CD44s (but not CD24) (Figure 3B, left panel, *P < .05, **P < .01). Flow cytometric analysis indicated that ALDH activity increased from Degron(−) control cells to ODC-degron–transduced cells to Degron(+) E-cad-Fc(+) cells in a stepwise method (6.79% ~ 11.65% ~ 22.50%; Figure 3B, right panel).

RESULTS

3.1 Effects of E-cad-Fc on EMT and cancer stem cell markers in SW480 colon cancer cells

3.2 Effects of E-cad-Fc on EMT and cancer stem cell markers in ODC-degron–transduced KM12SM cells

3.3 E-cadherin-Fc chimera protein enhanced stem cell properties

3.4 E-cadherin-Fc chimera protein activated the EGFR signaling pathway

Out with GraphPad Prism version 6.00 for Mac (GraphPad Software). P < .05 was considered to indicate significance.

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FIGURE 2  Mesenchymal-like characteristics in human colon cancer cell line KM12SM. A, Cells were cultured for 24, 48, and 72 h. Degron(−) group was cultured on normal tissue culture plate, Degron(+) group was cultured on normal tissue culture plate, and Degron(+) E-cad-Fc(+) group was cultured on a non-treated plate coated with E-cad-Fc matrix. Degron(+) E-cad-Fc(+) group cells showed a spindle-like shape as time passed. B, Degron(+) E-cad-Fc(+) group showed upregulation of mesenchymal markers at the protein level after 48 h treatment. Latent (45 kDa) and active (26 kDa) forms of transforming growth factor (TGF)-β1 protein also increased. Quantitative analysis in two independent experiments are shown. C, RNA study was done after 24 h treatment. *P < .05, **P < .01, ***P < .001. E-cad-Fc, E-cadherin-Fc chimera protein.
3.5 | E-cadherin-Fc chimera protein matrix suppressed proteasome activity

Because earlier studies showed that LPA is a hallmark of CSC, we measured proteasome activity in KM12SM cells. Our results showed that the Degron(+) group significantly inhibited proteasome activity compared to Degron(−) control cells (Figure 6; *P < .05, **P < .001). We also found that the proteasome activity tended to be further suppressed when Degron(+) cells were cultured on the E-cad-Fc matrix (Degron(+) E-cad-Fc(+) compared with the Degron(+) group (Figure 6, P = .068).
Despite rigorous studies, an effective treatment targeting CSC has not yet been established. One reason is that CSC are relatively rare among tumor cells and readily lose their properties through dilution with rapidly expanding differentiated daughter cells. This scarcity hinders analyses involving CSC. One solution is development of tools that enhance CSC properties and allow for efficient analysis of these cells.

Homeostatic proteasome activity plays a critical role in regulating the physiological activities of cancer cells and contributes to CSC initiation and development. During experiments, we were aware that the ODC-degron system worked well to show CSC properties in some cells such as colon cancer cell lines HCT116 and DLD1 and pancreatic cancer cell line Panc-1, but did not work in other cancer cell types. Therefore, it is assumed that the success of the ODC-degron system may depend on the degree to which LPA in the cell affects CSC properties. In the present study, we used KM12SM cells because ODC-degron-transduced KM12SM cells did not acquire sufficient CSC properties as shown in the present study. Therefore, we viewed these cells as suitable for evaluating an enhanced effect of cancer stemness.

We showed that culturing with E-cad-Fc-coated plates raised CD44v9 expression significantly and ALDH expression to some extent in SW480 cells and enhanced various CSC-related molecular in the ODC-degron-transduced KM12SM cells. In the latter cells, Lgr5, Bmi-1, and SOX9 mRNA were increased by E-cad-Fc treatment. They are reported to play an important role in regulating Wnt signaling and are associated with stem cell maintenance. Cell surface markers for CSC, CD44v9 and CD44s but not CD24 were also increased. These changes could lead to cancer stem-like properties such as sphere formation and chemoresistance to 5-FU and L-OHP. In sphere-formation assays, Degron(+) E-cad-Fc(+) cells showed enhanced formation of smooth and round robust spheres. This finding is consistent with a report by Frankel et al. showing that smooth and round spheroids have a higher spherical quality, whereas rough spheroids begin to disintegrate and lose cell junctions. Tumorigenicity assay is a valid method to estimate CSC activity and partially activated the EGFR downstream pathway, which was insufficient to induce a variety of CSC cell surface markers and chemoresistance, although it may contribute to some increase in SOX9 mRNA and ALDH expression (Figure 3A,B). In contrast, when ODC-degron–transduced KM12SM cells were cultured on E-cad-Fc-coated plates, proteasome activity was further lowered and the EGFR signaling pathway was markedly activated as identified by intense phosphorylation of the constituent molecules. It is assumed that these drastic changes led to induction of the broad range of CSC markers (Lgr5, Bmi-1, SOX9, CD44v9, CD44s, and ALDH), increased chemoresistance to 5-FU and oxaliplatin and smooth-edged, robust spheroid formation. These findings suggest that E-cad-Fc, like soluble E-cadherin, could affect EGFR and enhanced cancer stem-like properties.

During the experiments, we found that E-cad-Fc-treated cells had a spindle-like cell morphology, a characteristic of EMT. Analyses at protein and RNA levels showed the underlying molecular events. Thus, N-cadherin was highly induced by E-cad-Fc in SW480 cells. In particular, KM12SM cells showed a large increase in a variety of mesenchymal markers with E-cad-Fc treatment although E-cadherin, an epithelial marker, was not reduced much (Figure 2B). KM12SM cells acquire highly metastatic ability through mouse serial transplantation of primary colon cancer cells. Therefore, we postulate that this cell line may have potential to transform in the mesenchymal direction. RT-PCR analyses indicated that most mesenchymal molecules were upregulated by E-cad-Fc at the mRNA level. However, TGF-β1, a typical inducer of EMT, did not increase at the mRNA level whereas both latent and active forms of TGF-β1 were highly induced at the protein level, suggesting the existence of certain post-translational mechanisms.

Several studies have shown that low proteasome activity enhances EMT through stabilization of the EMT-related transcription factor Snail1 in gastric cancer and mammary epithelial cells. However, this is not likely to be the current case because ODC-degron–transduced cells without E-cad-Fc treatment showed no change in EMT-related protein expression despite significant down-regulation of proteasome activity. However, considering that various mesenchymal markers were similarly induced by E-cad-Fc even in control KM12SM cells (Figure S1), it is assumed that E-cad-Fc directly facilitates mesenchymal changes, irrespective of cancer stemness. Because currently used E-cad-Fc is a fusion protein consisting of human E-cadherin and mouse Fc domain, we examined whether a fusion protein of human E-cadherin and human Fc domain would directly induce mesenchymal changes in ODC-degron-transduced KM12SM cells. As shown in Figure S3, recombinant human E-cadherin-human Fc chimera protein caused spindle-like morphology at 48 hours and highly induced mesenchymal markers, Snail1, Slug, Twist1 expression and TGF-β1. These findings emphasize that human E-cadherin fragment is essentially important to induce mesenchymal changes and that this unique effect is not likely to be a non-specific reaction caused by the mouse Fc domain.

Cumulative evidence further emphasizes the links between EMT and CSC including their similar characteristics, such as tumorigenesis, enhanced survival signals, and metastatic ability.
(A) Whole view  Magnified view

Degron(−)

Degron(+)

Degron(+) E-cad-Fc(+)

(B) Sphere-formation assay

|                | Degron(−) | Degron(+) | Degron(+) E-cad-Fc(+) |
|----------------|-----------|-----------|-----------------------|
| Sphere-counting per well | [Bar graph showing data] | [Bar graph showing data] | [Bar graph showing data] |

* P < .05
ns P > .05

(C) Proliferation

Absorbance (492 nm) vs. Time (day)

5-FU

|                | Degron(+) E-cad-Fc(+) | Degron(+) | Degron(−) |
|----------------|-----------------------|-----------|-----------|
| Cell viability (%) | [Graph showing data] | [Graph showing data] | [Graph showing data] |

* P < .05

L-OHP

|                | Degron(+) E-cad-Fc(+) | Degron(+) | Degron(−) |
|----------------|-----------------------|-----------|-----------|
| Cell viability (%) | [Graph showing data] | [Graph showing data] | [Graph showing data] |

* P < .05
Mesenchymal markers including ZEB1, Snail1, and Slug are thought to be involved in CSC maintenance and tumorigenesis. SOX9 is stabilized through combination with the transcription factor Slug, promoting CSC properties and metastasis. Mesenchymal change may also contribute to spheroid formation and chemoresistance. Recent research on ovarian cancer cells showed that N-cadherin-positive mesenchymal-type cells form stable, highly cohesive solid spheroids with a smooth shape, and that their adhesive, migratory, and invasive cell properties were promoted. It is also reported that CSC markers such as CD44, CD44v9, and ALDH, along with EMT markers including Snail1, Slug, and ZEB1, contribute to resistance for chemotherapy and radiotherapy.

In conclusion, we showed that the E-cad-Fc matrix facilitated CSC properties and induced mesenchymal changes. Our data suggest that the E-cad-Fc matrix may be efficient for providing better access to CSC research as well as EMT investigation.

**FIGURE 4** Stem cell properties endowed by E-cadherin-Fc chimera protein (E-cad-Fc). A, Sphere formation of each group seeded at a density of 1000 cells per well cultured for 18 days. In the Degron(+) E-cad-Fc(+) group, cells were cultured with E-cad-Fc-coated plates for 48 h and then transferred to ultra-low attachment plates. Spheroid morphology is shown in whole view (left) and magnified view (right). Rough-edged spheres are marked by black arrows, and smooth spheres by white arrowheads. Scale bar, 200 μm. B-a, Sphere size ≥100 μm in diameter was counted to assess sphere-formation ability. *P < .05. B-b, Sphere size ≥100 μm in diameter was counted to assess sphere-formation ability, and smooth-edged spheres and rough-edged spheres were separately counted. *P < .05, **P < .01. C-a, Proliferation assay and chemoresistance assay. In the Degron(+) E-cad-Fc(+) group, cells were cultured with E-cad-Fc-coated plates for 48 h and then transferred to plates at a density of 5000 cells per well, and cultured with DMEM supplemented with 10% FBS. Proliferation was measured by CCK-8 every 24 h. C-b, Cells of each group were treated with fluorouracil (5-FU) or oxaliplatin (L-OHP) for 72 h. After drug treatment, cell survival rates were measured. *P < .05

**FIGURE 5** Activation of the epithelial growth factor receptor (EGFR) signaling pathway with E-cadherin-Fc chimera protein (E-cad-Fc) treatment. Upper panel: Quantitative analysis in two independent western blotting experiments. E-cad-Fc treatment of ornithine decarboxylase-degron-transduced cells markedly enhanced phosphorylation of EGFR, AKT, ERK1/2, mTOR, and PI3K, although total expression of each molecule was not affected. *P < .05, **P < .01, ***P < .001. Lower panel: Representative blots are shown.
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DISCLOSURE

Authors declare no conflicts of interest for this article.

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SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section at the end of the article.

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