Figure legends

Figure S1. Isolated normal primary ovarian surface epithelial cells (OSE) has epithelial morphology and express epithelial markers. OSE cells were isolated as described in material and methods. **A.** Photos showed cobble-stone epithelial morphology of OSE cells (a and b) compared to spindle-like morphology of fibroblasts in c. **B.** RT-PCR showed that OSE cells express epithelial markers (CDH1, EpCAM, KRT18 and KRT7) and to a lesser extent mesenchymal marker vimentin (VIM). Beta-2-macroglobulin (B2M) was used as positive control. Scale bar=100 μm in a and c. Scale bar=200 μm in b.

Figure S2. Exogenous Gal-3 increase SKOV-3 cell proliferation and cell survival. A. SKOV-

3 MCTS were treated with rhGal-3 (30 μ M) for 48h followed by BCL-2 and CCDN1 expression levels assessment by qRT-PCR. Results were normalized related to ACTB as internal control. **B.** Cell death analysis was performed in the presence or absence of rhGal-3 for 48h. Mean \pm SD, n=3

Figure S3. Differential expression levels of alpha integrin subunits in BLSOC, LGSOC and

HGSOC human specimens. The extracted RNA from fresh serous type EOC tumors (n = 46) were divided into three groups borderline serous ovarian cancer (BLSOC, n=12); low-grade serous ovarian cancer (LGSOC, n=12) and high-grade serous ovarian cancer (HGSOC, n=22) as well as fresh normal ovarian tissue (n =10) were analyzed by qRT-PCR for the following mRNA expression levels: (A) ITGA2, (B) ITGA4, (C) ITGA6, (D) ITGA5, (E) ITGAv. Dot plot comparing the distribution of normalized qRT-PCR analysis for afore-mentioned gene expression level in tumor specimens related to normal healthy ovaries. Values were normalized relative to ACTB expression levels used as internal control. *: $P \le .05$; ***: $P \le .001$ compared to control.

Figure S4. Differential expression levels of beta integrin subunits in BLSOC, LGSOC and HGSOC human specimens. The extracted RNA from fresh serous type EOC tumors (n = 46) were divided into three groups borderline serous ovarian cancer (BLSOC, n=12); low-grade serous ovarian cancer (LGSOC, n=12) and high-grade serous ovarian cancer (HGSOC, n=22) as well as fresh normal ovarian tissue (n =10) were analyzed by qRT-PCR for the following mRNA expression levels: (A) ITGB1, (B) ITGB2, (C) ITGB3, (D) ITGB4, (E) ITGB6. Dot plot comparing the distribution of normalized qRT-PCR analysis for afore-mentioned gene expression level in tumor specimens related to normal healthy ovaries. Values were normalized relative to ACTB expression levels used as internal control. *: $P \le .05$; **: $P \le .01$; ***: $P \le .001$ compared to control.

Tumor grade	n	Median age	Treatment
Normal ovary	10	53	-
BLSOC	12	42	None
LGSOC (Grade I)	7	55	None
LGSOC (Grade II)	5	47	None
HGSOC (Grade III)	12	56	None
HGSOC (Grade IV)	10	61	None

Table S1. Clinicopathological characteristics of patients

Borderline serous ovarian cancer (BLSOC), Low-grade serous ovarian cancer (LGSOC), Highgrade serous ovarian cancer (HGSOC).

Genes	Primer sequences
ACTB	F: 5'- CTTCCTTCCTGGGCATG-3' R: 5'- GTCTTTGCGGATGTCCAC-3'
B2M	<pre>F: 5´- CCTGAATTGCTATGTGTCTGGG -3 ' R: 5´-TGATGCTGCTTACATGTCTCGA-3'</pre>
LGALS3	<pre>F: 5´CCTCGCATGCTGATAACAATTCT-3' R: 5´TGACTCTCCTGTTGTTCTCATTGAA-3'</pre>
ITGB1	F: 5'- GTGGGTGGTGCACAAATTC-3' R: 5'-GGTCAATGGGATAGTCTTCAGC-3'
ITGB2	<pre>F: 5'-TTCGGGTCCTTCGTGGACA-3' R: 5'- ACTGGTTGGAGTTGTTGGTCA-3'</pre>
ITGB3	<pre>F: 5'- AGCCAACAACCCACTGTA-3' R: 5'- CTGACATTCTCCCAACCTAC-3'</pre>
ITGB4	<pre>F: 5'-TGGAAGTACTGTGCCTGCTG-3' R: 5'-TGCATGTTGTTGGTGACCTT-3'</pre>
ITGB6	<pre>F: 5'- TCCATCTGGAGTTGGCGAAAG-3' R: 5'- TCTGTCTGCCTACACTGAGAG-3'</pre>
ITGA2	<pre>F: 5' - TAGCGCTCAGTCAAGGCATT-3' R: 5' - GCACTGCATAGCCAAACTGT-3'</pre>
ITGA4	<pre>F: 5'- AGCCCTAATGGAGAACCTTGT-3' R: 5'- CCAGTGGGGAGCTTATTTTCAT-3'</pre>
ITGA5	F: 5'- TTTATCGGTCTCGGGAGTTG-3' R: 5'-CTTCAACTTAGACGCGGAGG -3'
ITGA6	F: 5'- TTTATCGGTCTCGGGAGTTG-3' R: 5'- GGCCACTGAATGTTCAAGGT-3'
ITGAV	F: 5'- GCAACAGGCAATAGAGAT-3' R: 5'- TGCTGAATCCTCCTTGACAA-3'
EPCAM	<pre>F: 5'- CCATGTGCTGGTGTGTGAAC -3' R: 5'-CCTTCTGAAGTGCAGTCCGC -3'</pre>
CDH1	F: 5'- CAGGAGTCATCAGTGTGGT -3' R: 5'- GGAGGATTATCGTTGGTGTCAG -3'
KRT7	F: 5'- TCCGCGAGGTCACCATTAAC-3' R: 5'- GCTCTGTCAACTCCGTCTCAT-3'
KRT18	F: 5' - GGAGGCATCCAGAACGAGAA- 3' R: 5' - CCAGCTGCAGTCGTGTGATA- 3'
VIM	F: 5' - GGCTCGTCACCTTCGTGAAT - 3' R: 5' - GAGAAATCCTGCTCTCCTCGC - 3'

Table S2. Primer sequences used for qRT-PCR in this study

Figure S1

A















4

11650C

LGSOC





Figure S4

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0.

BLSOC