SUPPLEMENTARY MATERIAL

Supplementary Table 1: Clinical characteristics of endometrial cancer patients' cohort.

Supplementary Figure 1. Characterization of endometrial cancer cell lines. (**A**) Restriction map of the pmCherry-N1 vector. (**B**) ANXA2 and Vimentin levels in H-Ctrl and H-ANXA2 cells analyzed by qPCR (Mean±SEM; Mann-Whitney test; **p*=0.029). (**C**) Representation of western blot analysis shown in Figure 1. Protein levels of IK-Cherry-Ctrl and IK-Cherry-ANXA2 cell lines by original western blot. Actin (42KDa); Vimentin (57KDa); ANXA2 (38KDa) and Cherry-ANXA2 (67KDa).

Supplementary Figure 2. *High Throughput Screening (HTS)* assay. (**A**) Dose-response curves comparing the cytotoxic activity of Novobiocin in IK-pLKO (IC₅₀=1,310x10⁻⁷M) and IK-shANXA2 (IC₅₀=2,004x10⁻⁶M) cells and (**B**) in IK-Ctrl (IC₅₀=1,589x10⁻⁶M) and IK-ANXA2 (IC₅₀=2,888x10⁻⁶M) cells for 5 days in a 96 well plate colony formation assay.

Supplementary Figure 3. Absence of cytotoxic effect of Daunorubicin on EAhy926-GFP endothelial cells. (**A**) Quantification of the total area of fluorescence signal from EAhy926-GFP endothelial cells treated or not with Daunorubicin $(1x10^{-6}\text{M})$ for 7 days in 3D transendothelial migration culture using Operetta® High Content Imaging System (Mean±SEM; Mann-Whitney test; p=ns). (**B**) Alamar blue-based cell viability assay showing dose-response curve of Daunorubicin in EAhy926-GFP endothelial cells (IC₅₀=1.69x10⁻⁶M). These results evidence the absence of cytotoxic effect of Daunorubicin on endothelial cells at concentrations showing a deleterious impact on ANXA2-expressing endometrial tumor cells.