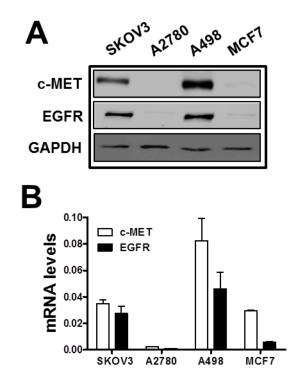
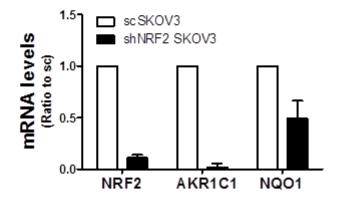
## NFE2L2/NRF2 silencing-inducible miR-206 targets c-MET/EGFR and suppresses BCRP/ABCG in cancer cells

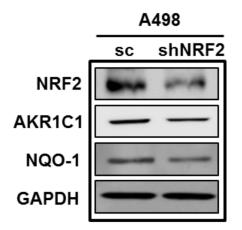
## SUPPLEMENTARY MATERIALS



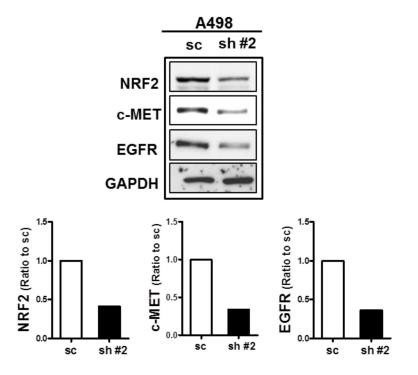
**Supplementary Figure 1: (A)** Protein levels for c-MET and EGFR in four cell lines: ovarian carcinoma SKOV3 and A2780, renal carcinoma A498, and breast carcinoma MCF7. **(B)** Transcript levels for c-MET and EGFR were assessed in four cell lines. Transcript levels of each gene were normalized with the level of HPRT.



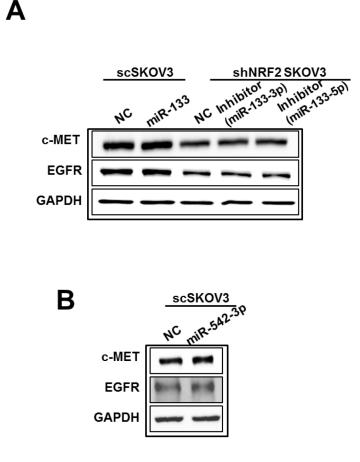
Supplementary Figure 2: *NRF2*-silending was verified by measuring transcript levels for *NRF2*, *AKR1C1* and *NQO-1* in established scSKOV3 and shNRF2-SKOV3 cells. Values are means ± SD from 3-4 experiments.



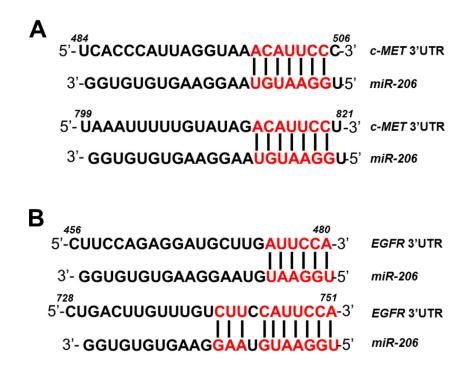
Supplementary Figure 3: NRF2, AKR1C1, and NQO-1 protein levels were determined by Western blotting in scA498 and shNRF2-A498 cells.



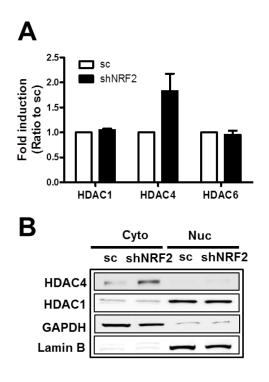
Supplementary Figure 4: Additional *NRF2*-specific shRNA (shNRF2 <sup>#</sup>2) was transiently transduced into A498, and protein levels of NRF2, c-MET and EGFR were assessed.



**Supplementary Figure 5: (A)** The effect of the miR-133 mimic or miR-133 inhibitor on c-MET and EGFR protein levels. The scSKOV3 cells were transfected with the miR-133 mimic (100 nM) and the shNRF2 cells were transfected with the miR-133 inhibitor (100 nM). After 18 h-incubation, protein levels for c-MET and EGFR were monitored by Western blotting. **(B)** After transfection of the scSKOV3 with the miR-542 mimic (100 nM), c-MET and EGFR protein level was measured.



**Supplementary Figure 6:** Target analysis with TargetScan (http://www.targetscan.org/vert\_71/) and Diana Tools (http:// diana.imis.athena-innovation.gr/DianaTools/index.php?r=tarbase/index) showed that miR-206 can bind to the 3'-UTR of the *c-MET* (A) and *EGFR* (B) gene.



**Supplementary Figure 7: (A)** The transcript levels for HDAC1, HDAC4, and HDAC6 were measured in sc and shNRF2-SKOV3 cells. **(B)** Cytoplasmic (Cyto) and nuclear (Nuc) protein levels of HDAC1 and HDAC4 were assessed by Western blotting.