

1 **Supplementary Material**

2 **MTT assay**

3 The cells were seeded on 96-well microplates (5000 cells/100 μ L/well) and
4 cultured in DMEM/F-12 medium for 24 h, then in serum-free medium for 18 h and in
5 Met-free medium for 6 h. Finally, the cells were cultured in the Met-free medium
6 (Met starvation group) or the medium containing different concentrations of Met (0.5,
7 1, 2 and 5 mM) for 24 h. At the end of the treatment, 10 μ L of a 5 mg/mL solution of
8 MTT in PBS was added to each well and the plates were incubated at 37°C for an
9 additional 4 h. The supernatant was then discarded, and the purple formazan crystals
10 formed were solubilized with 100 μ L/well of DMSO (Sigma-Aldrich, St. Louis, MO,
11 USA). The absorbance of the 96-well plate was determined using an automatic
12 microplate spectrophotometer at 570 nm. Cell viability was calculated as the percent
13 ratio of absorbance of the samples against the control medium.

