

Supporting Information

The interplay between oxidation process and cytotoxicity effects of antimonene nanomaterials

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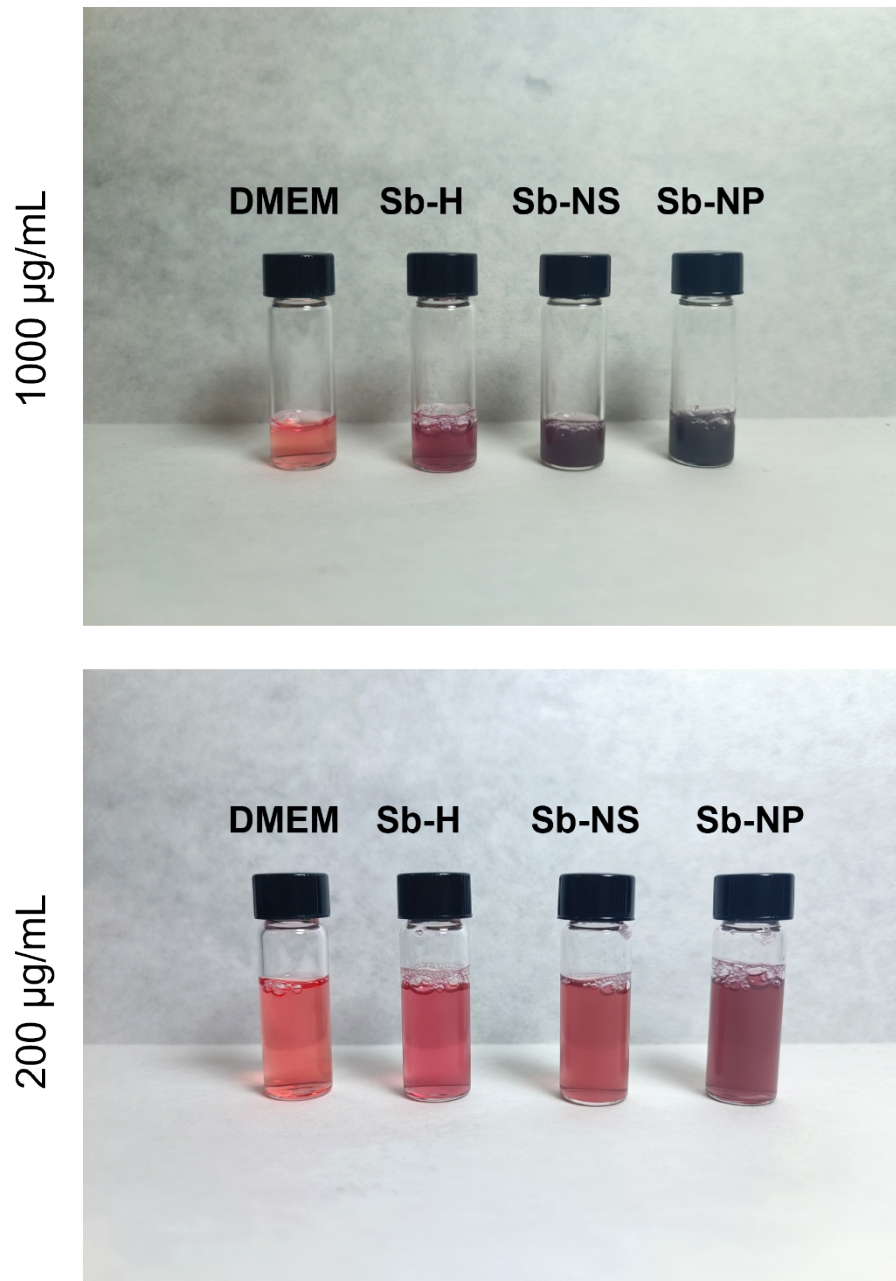


Figure S1: Photographs of the of the DLS dispersions at 1 mg/ml (resuspended concentration) and at 200 g/ml.

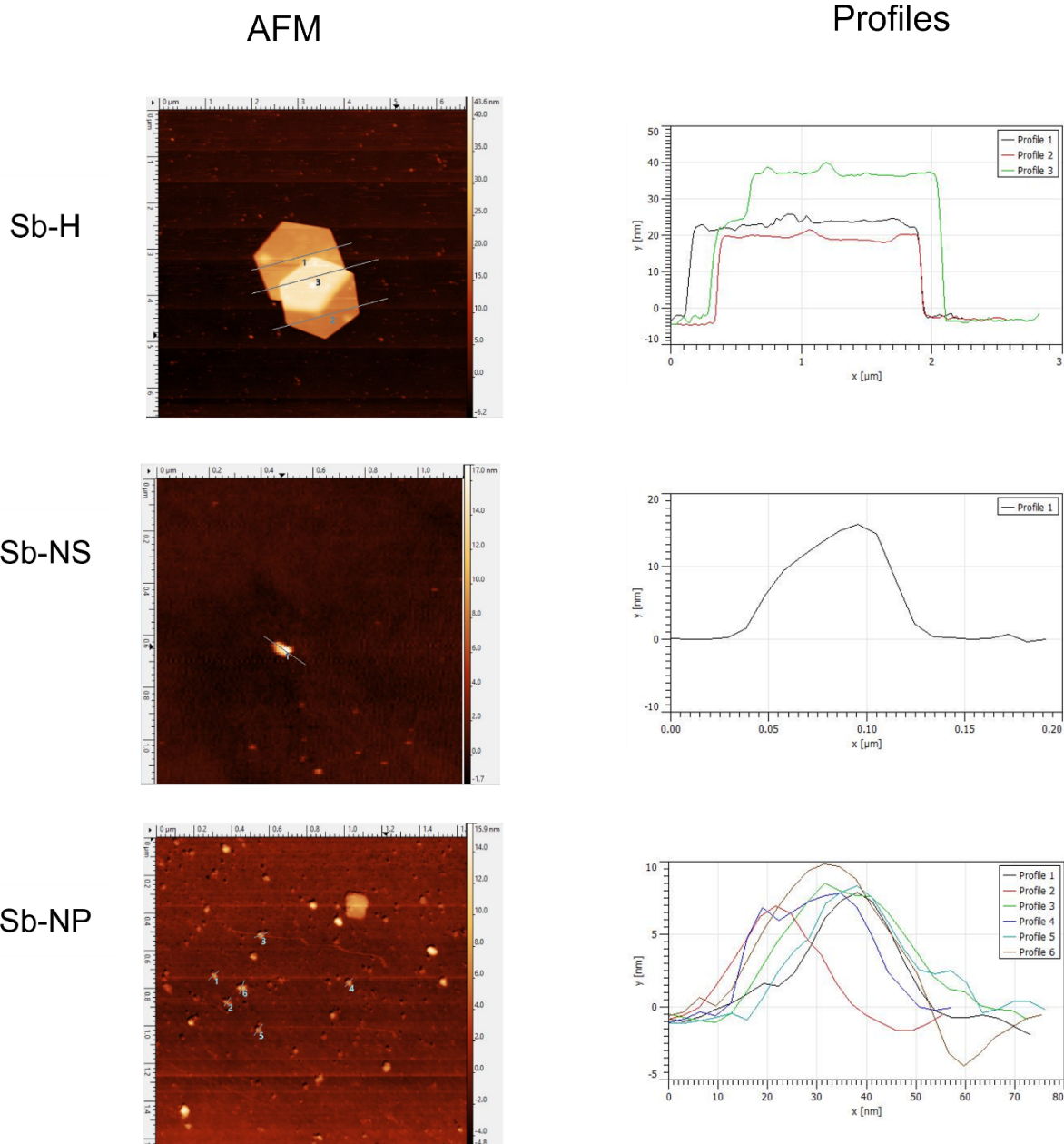


Figure S2: AFM images of the obtained Sb-Nanomaterials. AFM profiles showed on left correspond to the lines indicated in the images.

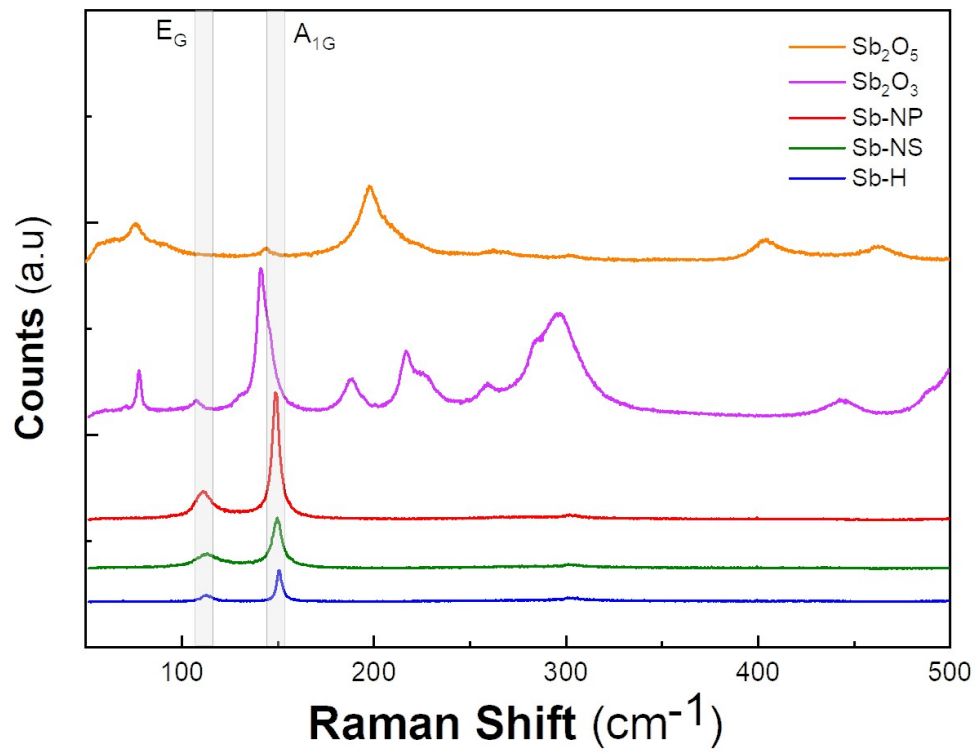


Figure S3: Comparison of the Raman spectra of the Sb-nanomaterials and Sb oxides.

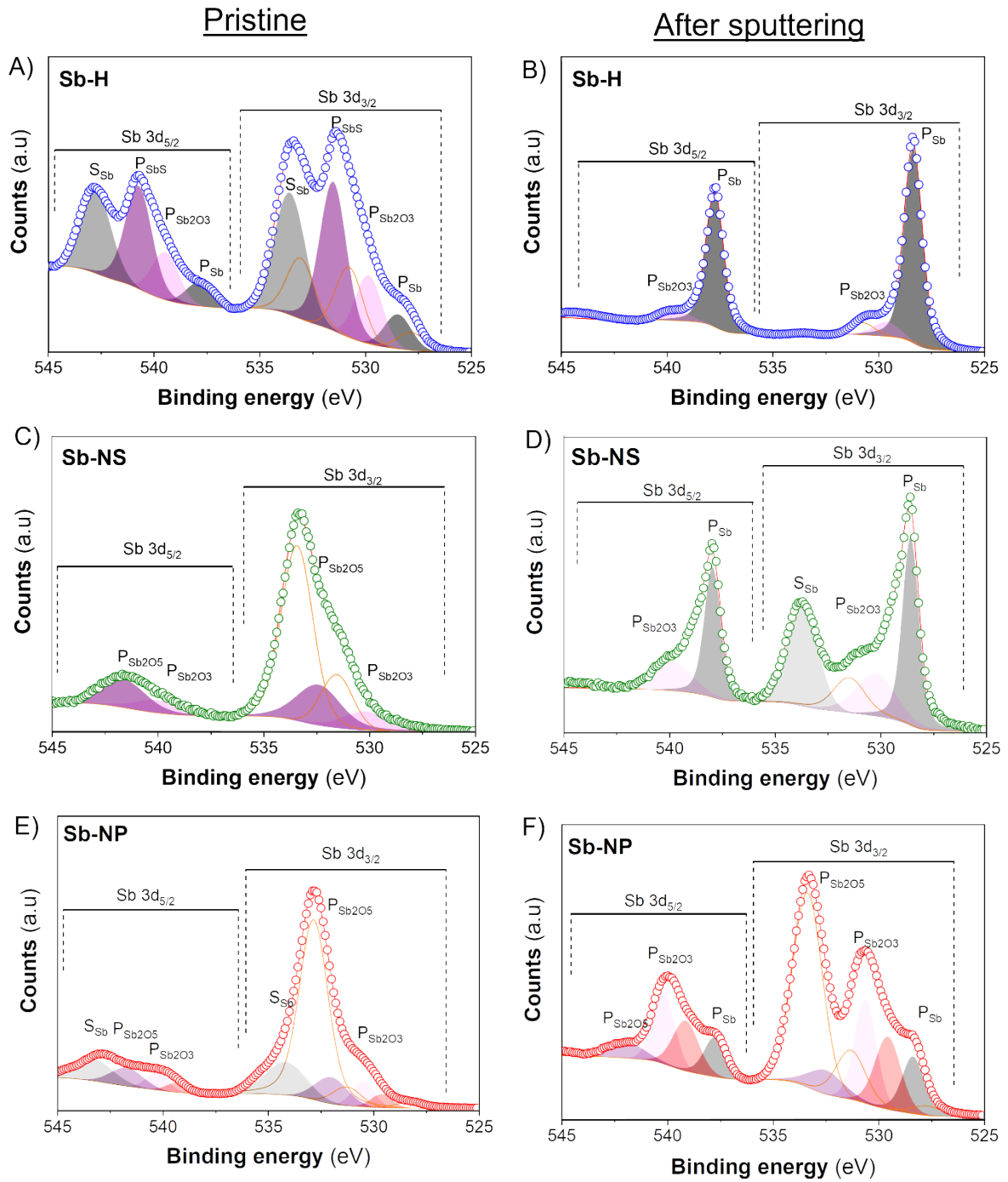


Figure S4: Deconvoluted XPS spectra of the nanomaterials before A, C, E) and after B, D, F) the sputtering process of the three Sb-Nanomaterials. P indicates the peak of the indicated atom and S the satellite.

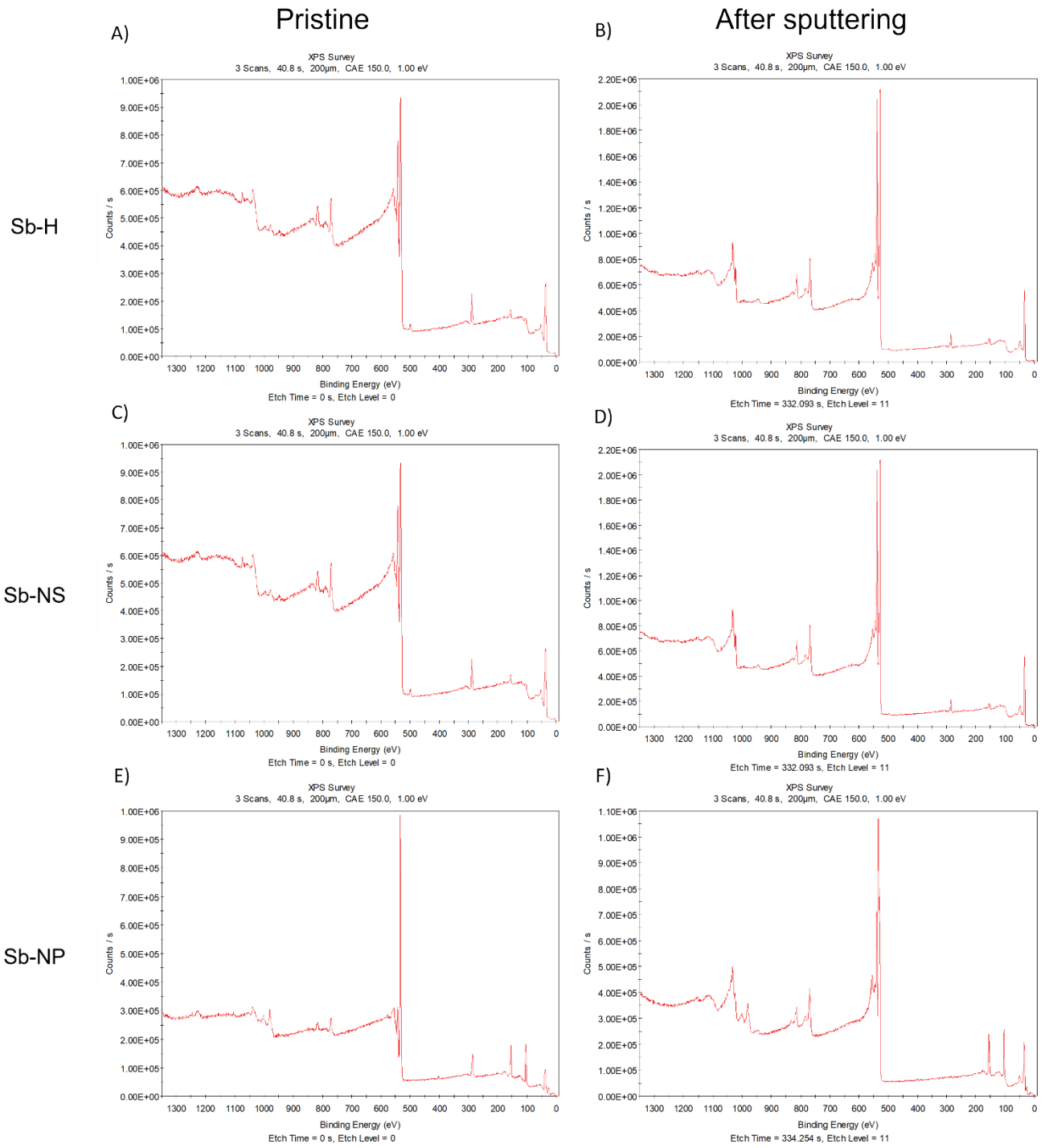


Figure S5: XPS surveys of the nanomaterials before A, C, E) and after B, D, F) the sputtering process of the three Sb-Nanomaterials.

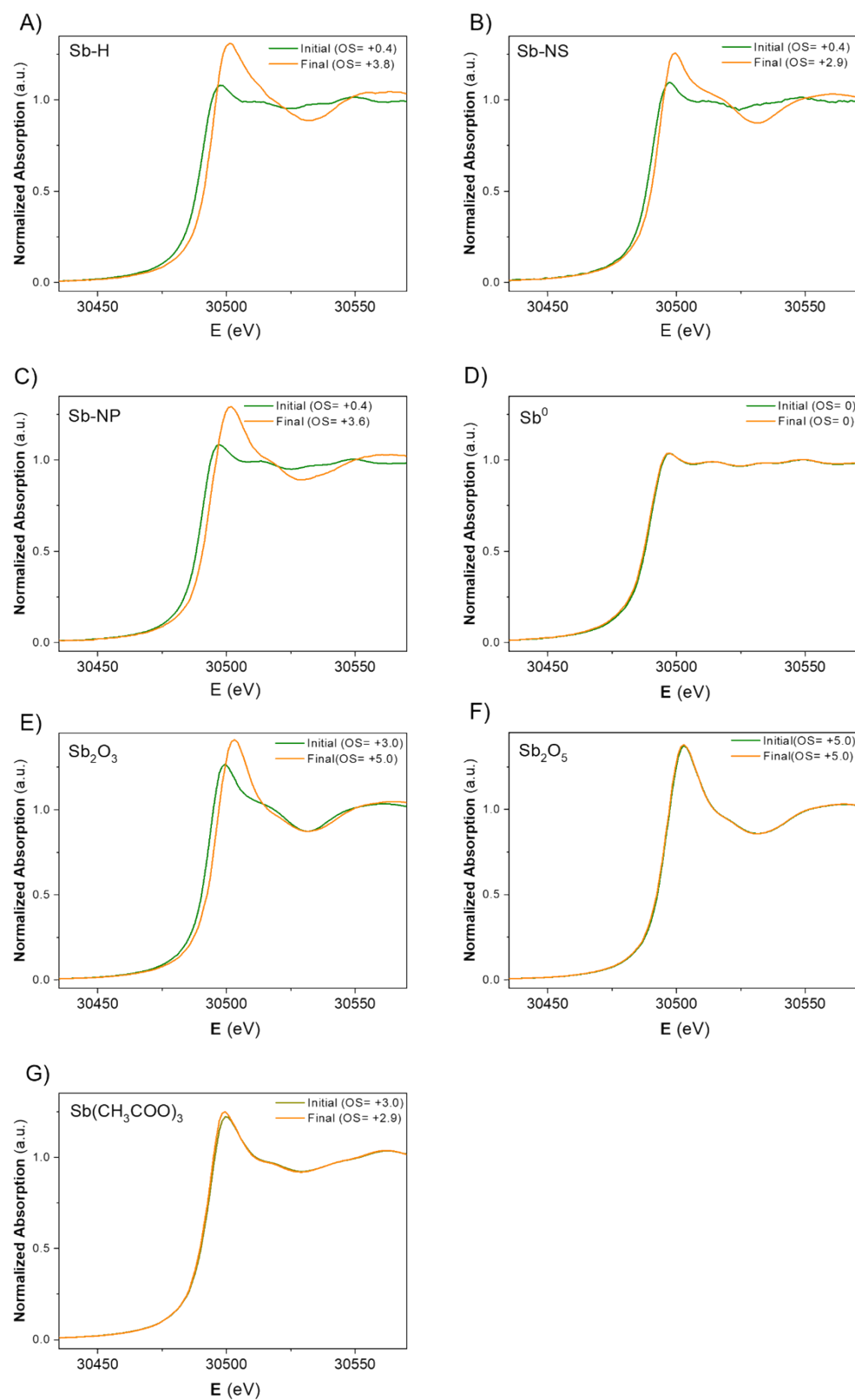


Figure S6: XANES spectra of the ex-situ measurements on the Sb reference species before (initial) and after (final) incubating 1 week in DMEM cell culture media.

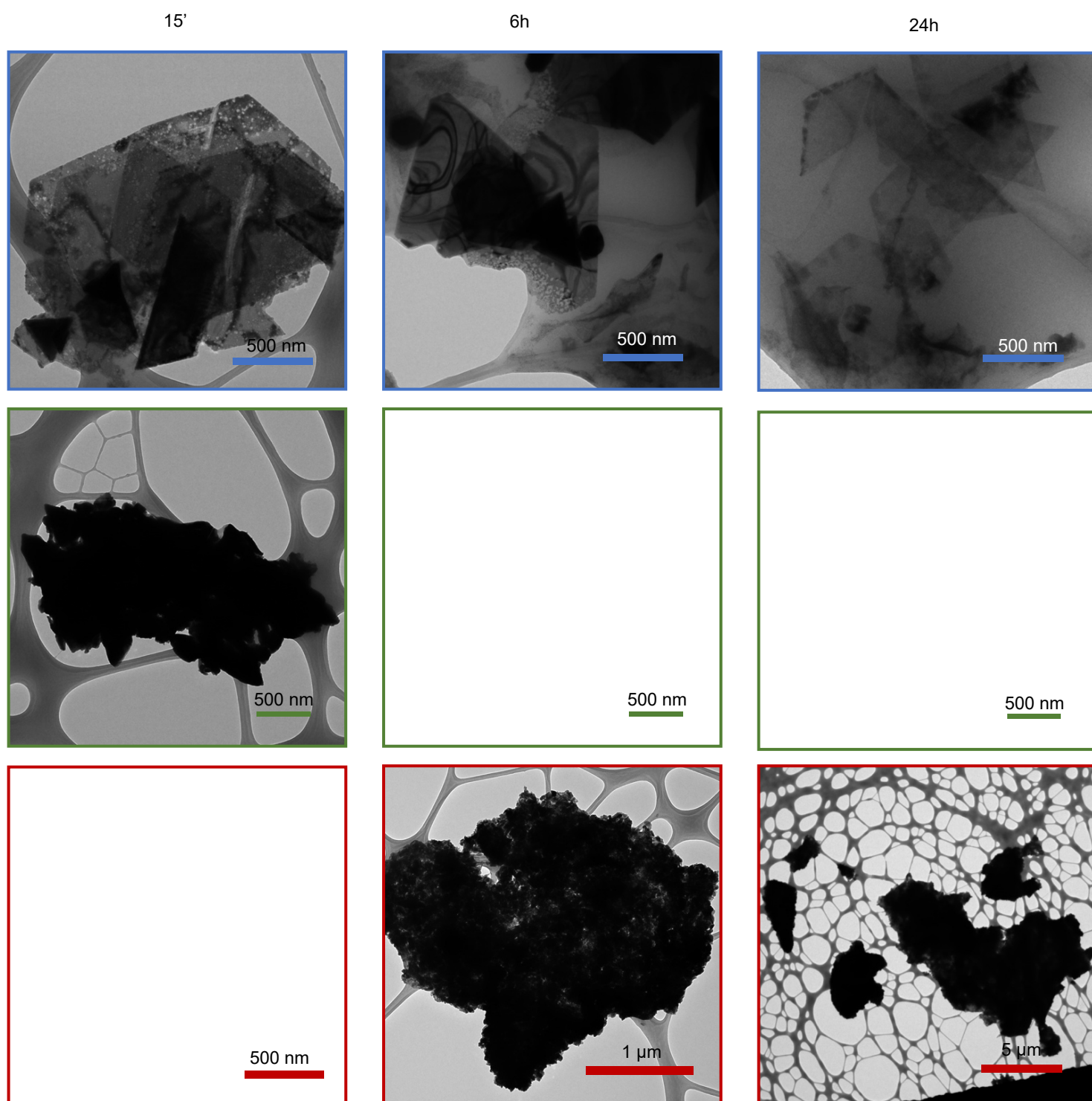


Figure S7: TEM images of the Sb-nanomaterials incubated at 37 °C in DMEM during the indicated times for the Sb-H, Sb-NS and Sb-NP samples highlighted in blue, green and red colors, respectively.

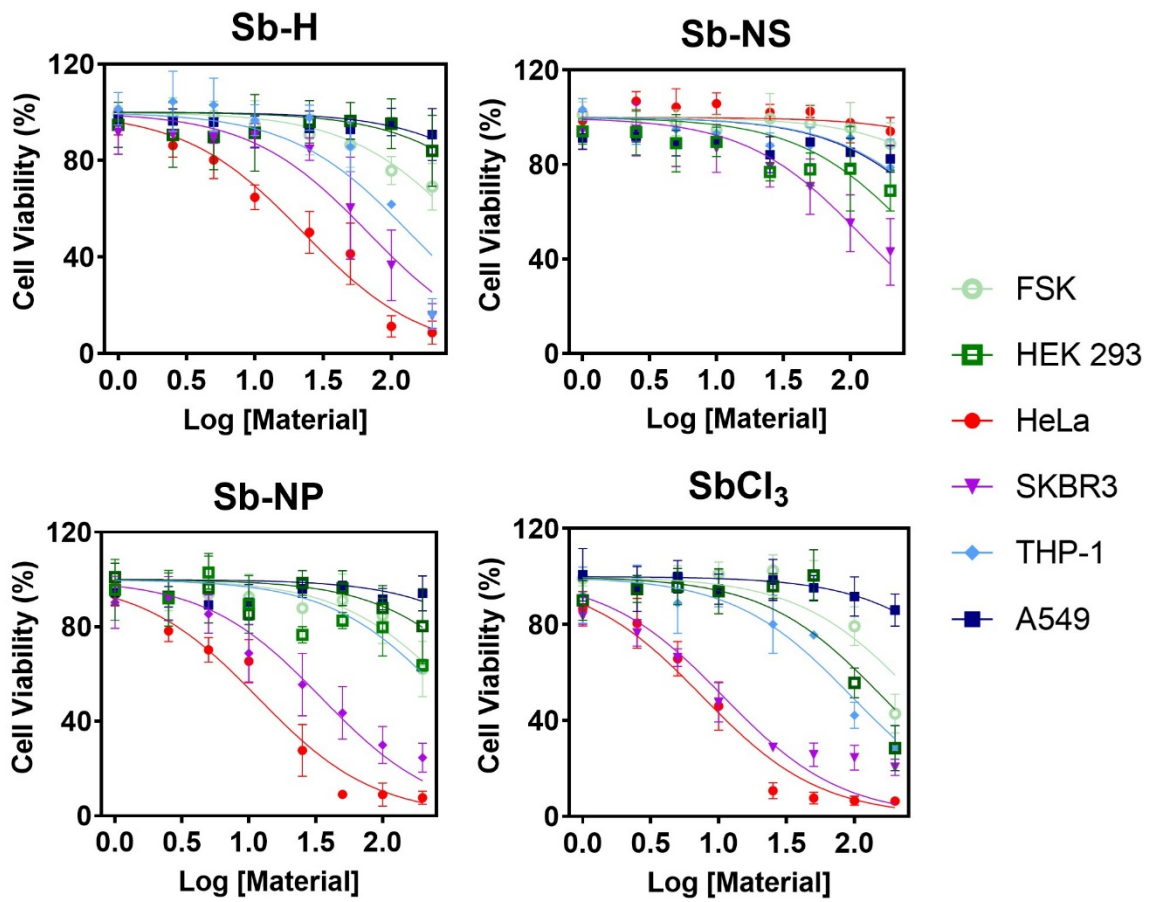


Figure S8: Dose-response curves of the tested compounds, obtained from the MTT viability assays. Obtained using the GraphPad Prism 8 software.

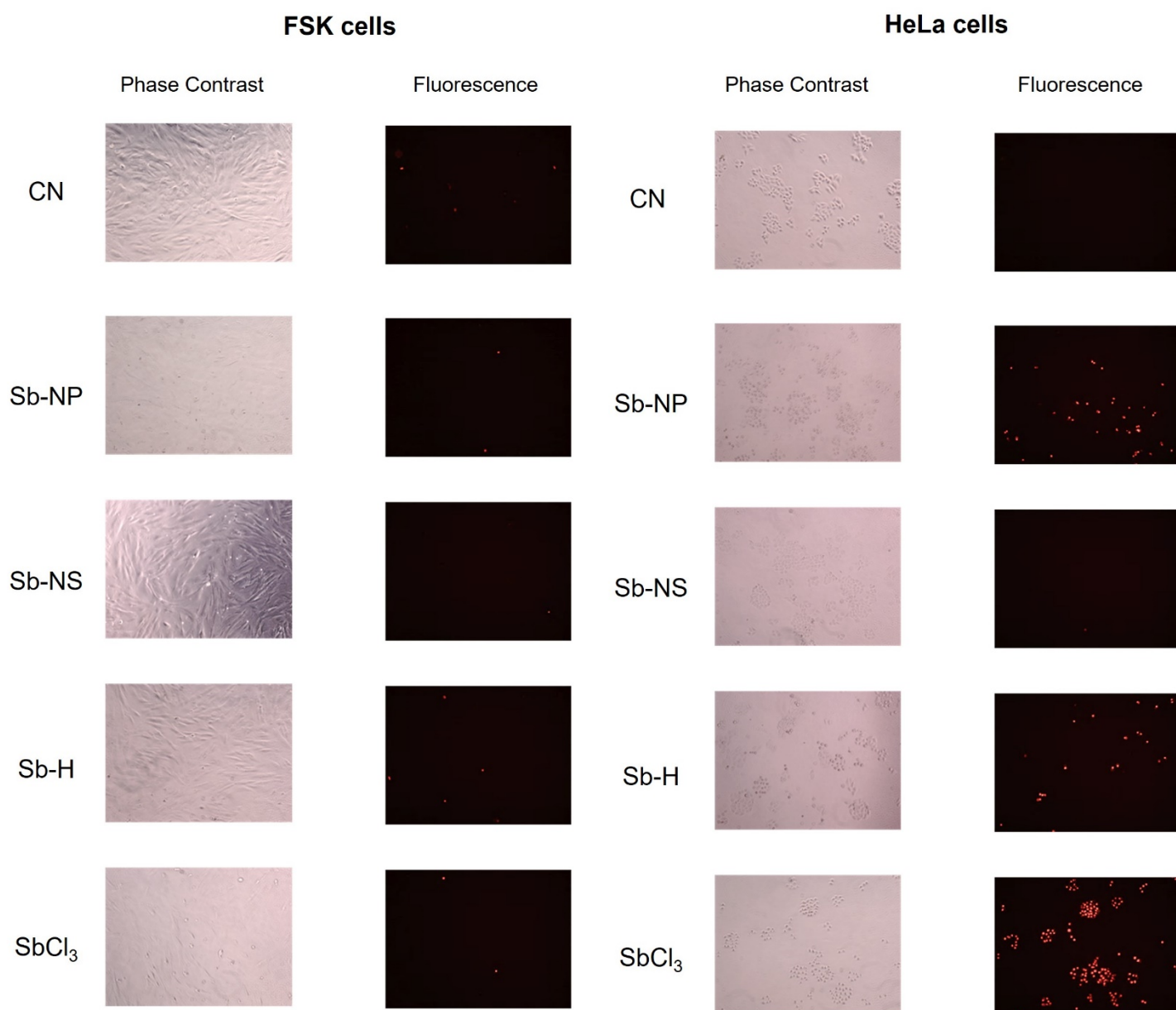


Figure S9: Optical microscope images and fluorescent images ($\lambda_{exc}= 545nm$) of the Propidium iodine test after treatment with the different nanomaterials at 25 $\mu g/mL$.

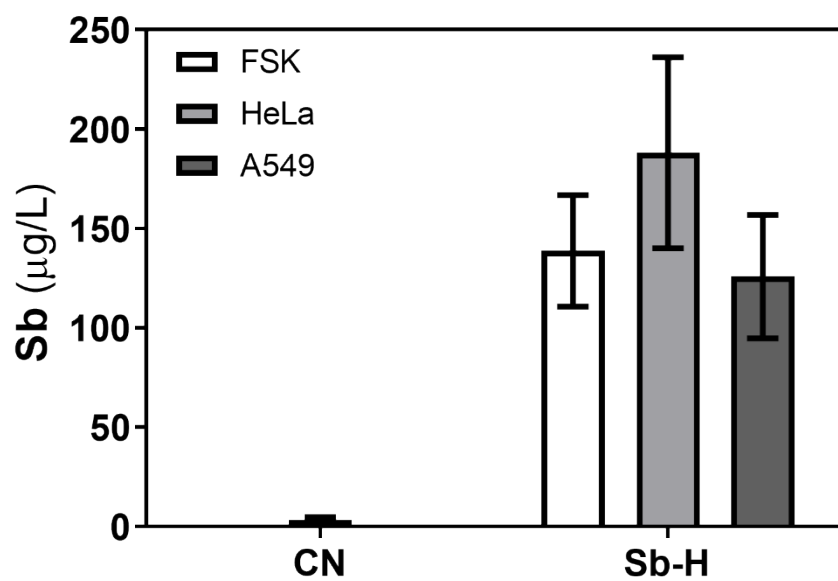


Figure S10: ICP-MS analysis of the intracellular content of the cells treated with the Sb-H at 25 µg/mL.

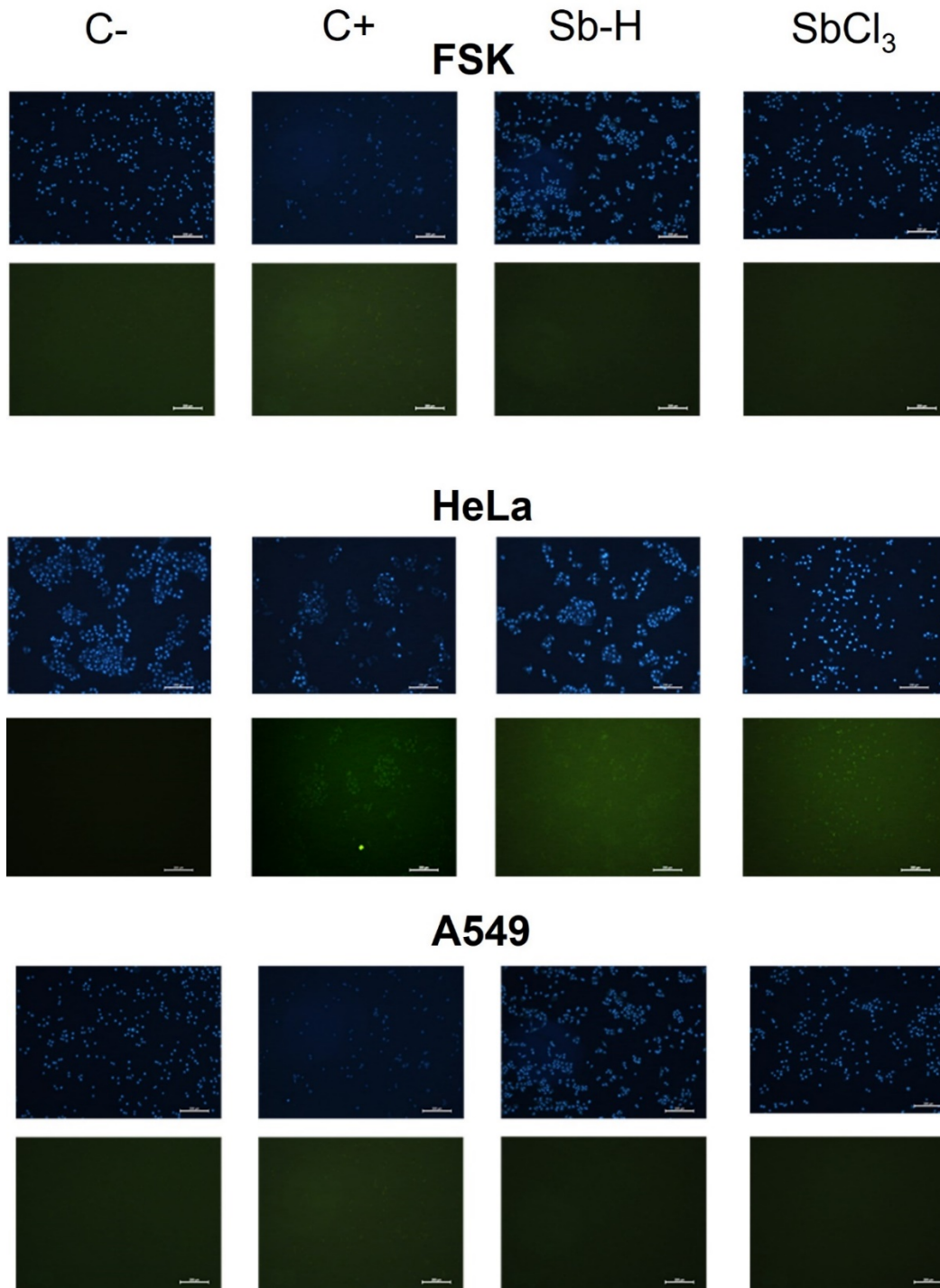


Figure S11: Representative fluorescence microscopy images of ROS experiments. Cell lines (HeLa, FSK and A549) were seeded on 24-well plates and incubate in the presence of Sb-nanomaterials at 25 $\mu\text{g}/\text{mL}$ for 24h. Next day, cells were treated with 25 μM of 6-carboxy-2',7'-dichlorodihydrofluorescein diacetate (carboxy- $\text{H}_2\text{DCF-DA}$) to detect ROS formation (Fluorescein FITC filter) and Hoechst 33342 to visualize nuclei (DAPI filter). C-: non treated cells, C+: cells treated with tert-butyl hydroperoxide (100 μM). SB-H: cells treated with Sb nanomaterial. SbCl₃: cells treated with antimony trichloride. Scalebar 200 μm .