

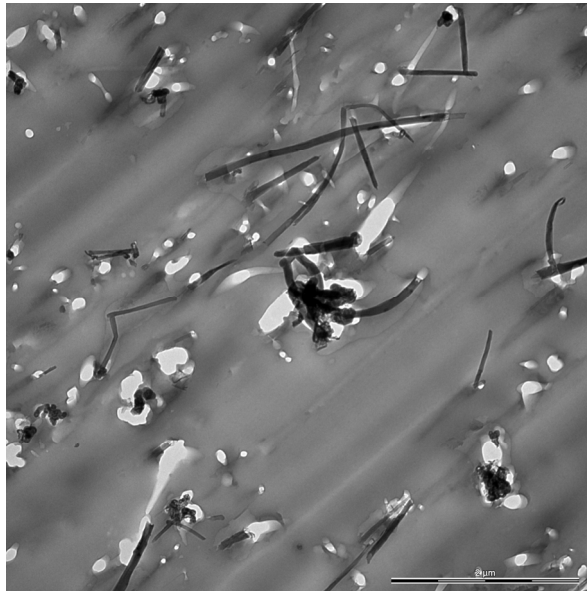
Table S1. Physicochemical properties of NMs from the JRC nanomaterial repository.

Nanomaterial (NM)	Diameter (TEM) (nm)	Hydrodynamic diameter (DLS) (nm)	Crystalline size (XRD) (nm)	Specific surface area (BET) (m <sup>2</sup> /g)	Elemental composition (XPS/ICP-MS)
<sup>1</sup> NM104 (TiO <sub>2</sub> )	26 ± 10	128.3 ± 0.8	21–27	41.216	O,C,Ti,Al
<sup>2</sup> NM300K (Ag)	8–50 (average: 17.6)	50-70	NA	NA	NA
<sup>3</sup> NM401 (MWCNT)	60–70 (length: 4048 ± 2371)	NA	10–30	140.46	Na, Fe, Al, Ni, Mg

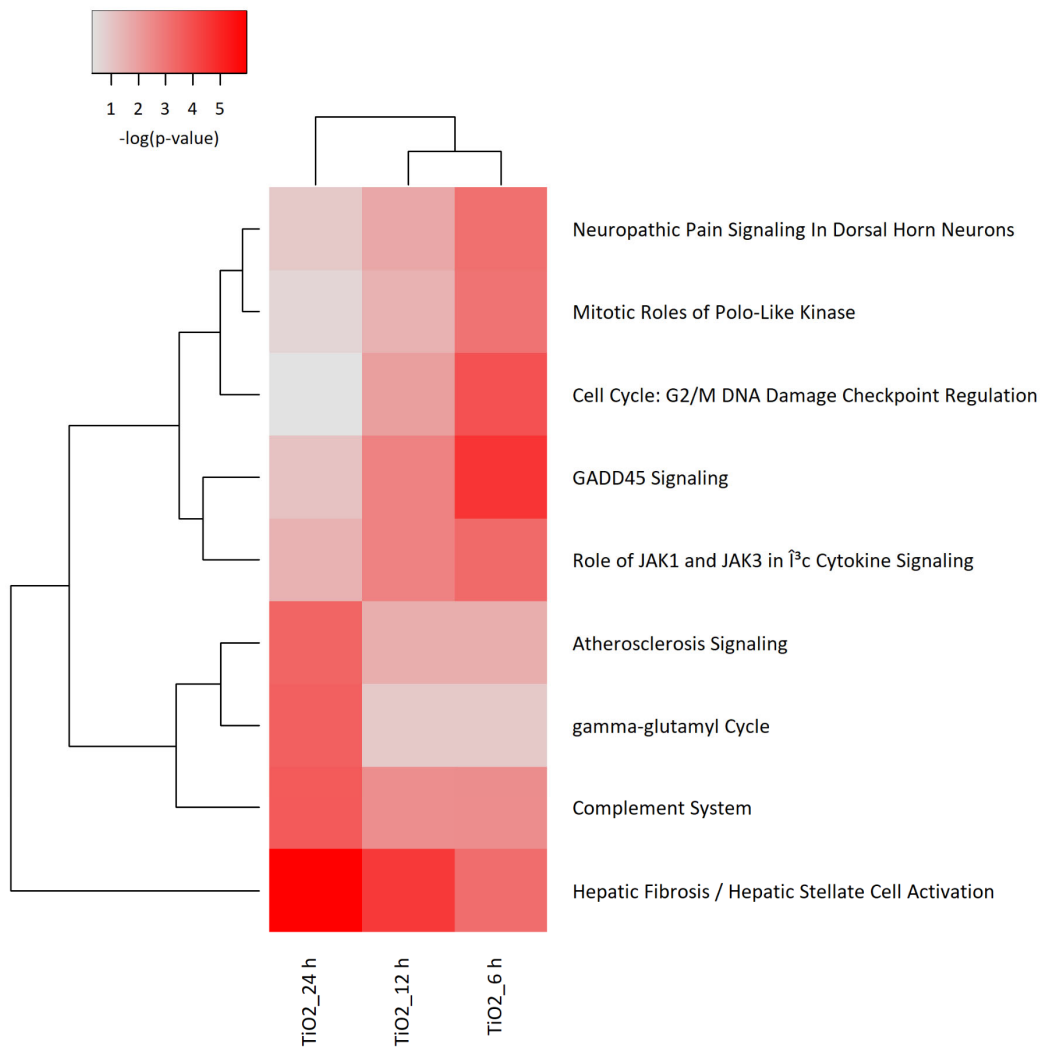
<sup>1</sup>Rasmussen K. et al. Scientific and Technical Research Reports – Titanium Dioxide, NM-100, NM-101, NM-102, NM-103, NM-104, NM-105: Characterisation and Physico-Chemical Properties. EUR 26637. (2014).

<sup>2</sup>Klein C.L. et al. NM-series of Representative Manufactured Nanomaterials. NM-300 Silver. Characterisation, Stability, Homogeneity. JRC60709. (2011).

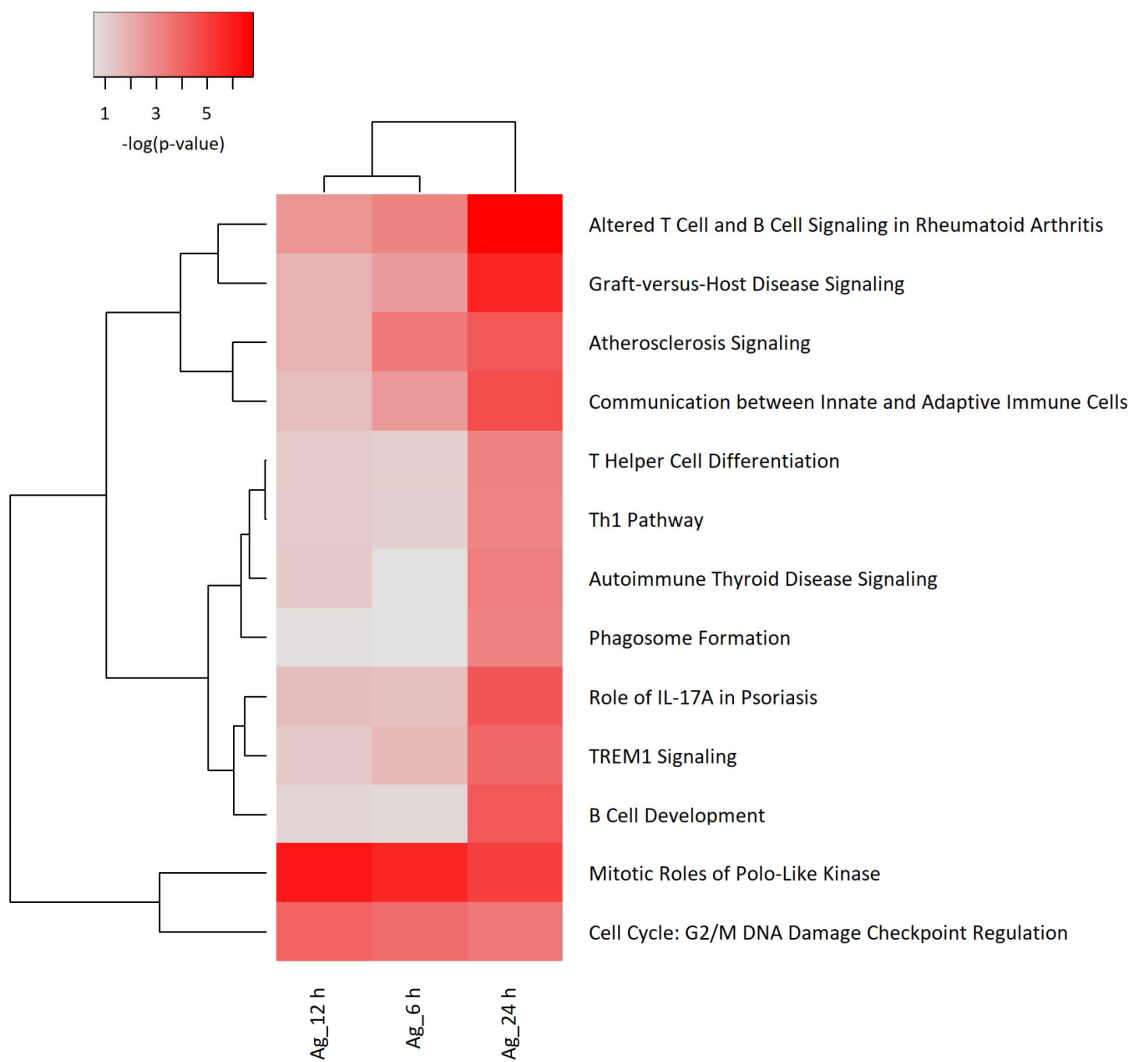
<sup>3</sup>Rasmussen K. et al. Scientific and Technical Research Reports – Multi-Walled Carbon Nanotubes, NM-400, NM-401, NM-402, NM-403: Characterisation and Physico-Chemical Properties. EUR 26796. (2014).



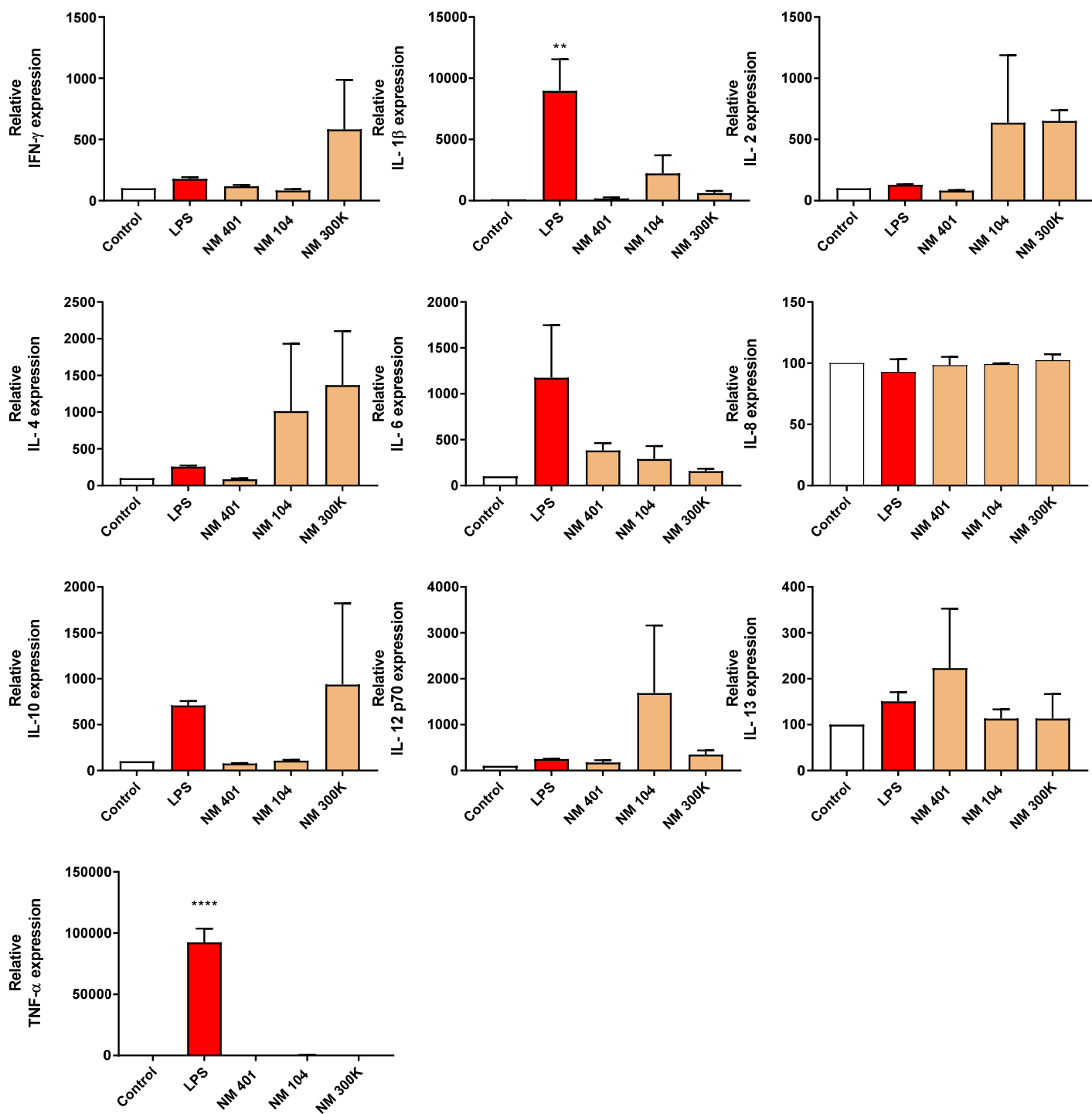
Suppl. Fig. S1. TEM imaging of THP-1 cells exposed to NM401 failed because the microtome blade used to cut the ultrathin sections was damaged by the NM401 material. Nevertheless, some remnants of the rigid, needle-like MWCNTs can be seen.



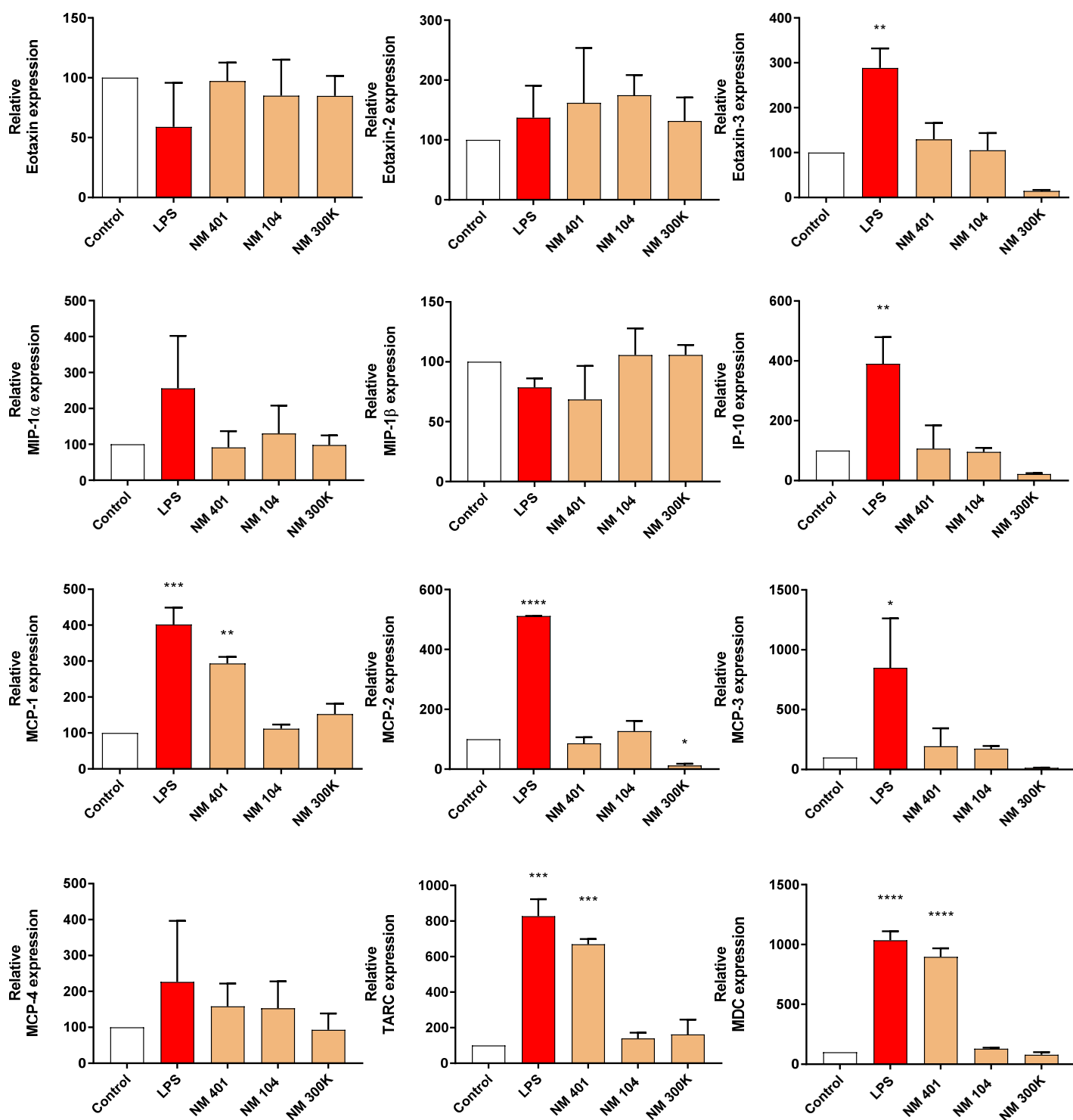
Suppl. Fig. S2. Canonical pathway analysis of transcriptomics data obtained from macrophage-like THP-1 cells exposed to TiO<sub>2</sub> (NM104) at 25 µg/mL. The significance values indicate the probability of association of differentially expressed genes (DEGs) with the respective pathway. The cut-off for the *p*-value was *p*<0.001 for at least one of the conditions.



Suppl. Fig. S3. Canonical pathway analysis of transcriptomics data obtained from macrophage-like THP-1 cells exposed to Ag (NM300K) at 25  $\mu\text{g}/\text{mL}$ . The significance values indicate the probability of association of differentially expressed genes (DEGs) with the respective pathway. The cut-off for the  $p$ -value was  $p < 0.001$  for at least one of the conditions.



Suppl. Fig. S4. Cytokine release in macrophage-differentiated THP-1 cells exposed for 24 h to NM401, NM104, and NM300K. LPS (0.1 μg/mL) was included as a positive control. The data shown are mean values  $\pm$  S.D. derived from three independent experiments. \* $p$ <0.05, \*\* $p$ <0.01, \*\*\* $p$ <0.001, \*\*\*\* $p$ <0.0001 (One-way ANOVA with Dunnett's multiple comparisons test).



Suppl. Fig. S5. Chemokine release in macrophage-differentiated THP-1 cells exposed to NM401, NM104, and NM300K. LPS (0.1 μg/mL) was included as a positive control. The data shown are mean values ± S.D. derived from three independent experiments. \* $p$ <0.05, \*\* $p$ <0.01, \*\*\* $p$ <0.001, \*\*\*\* $p$ <0.0001 (One-way ANOVA with Dunnett's multiple comparisons test).