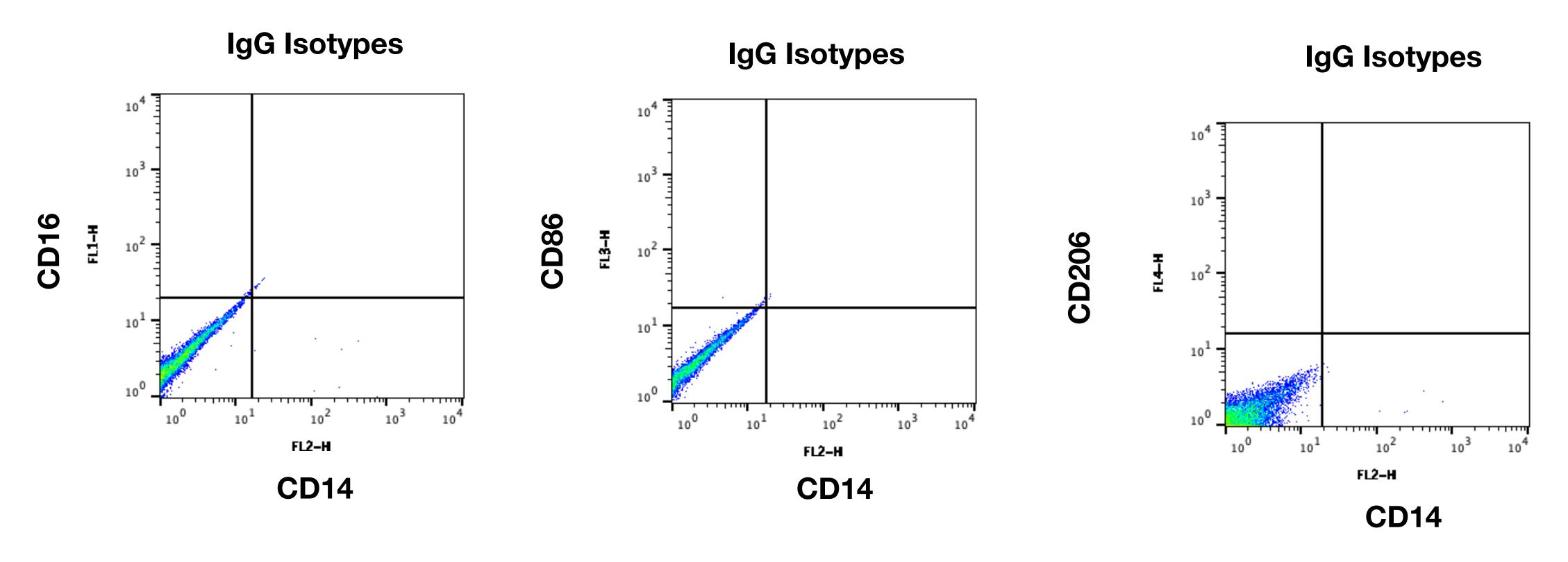
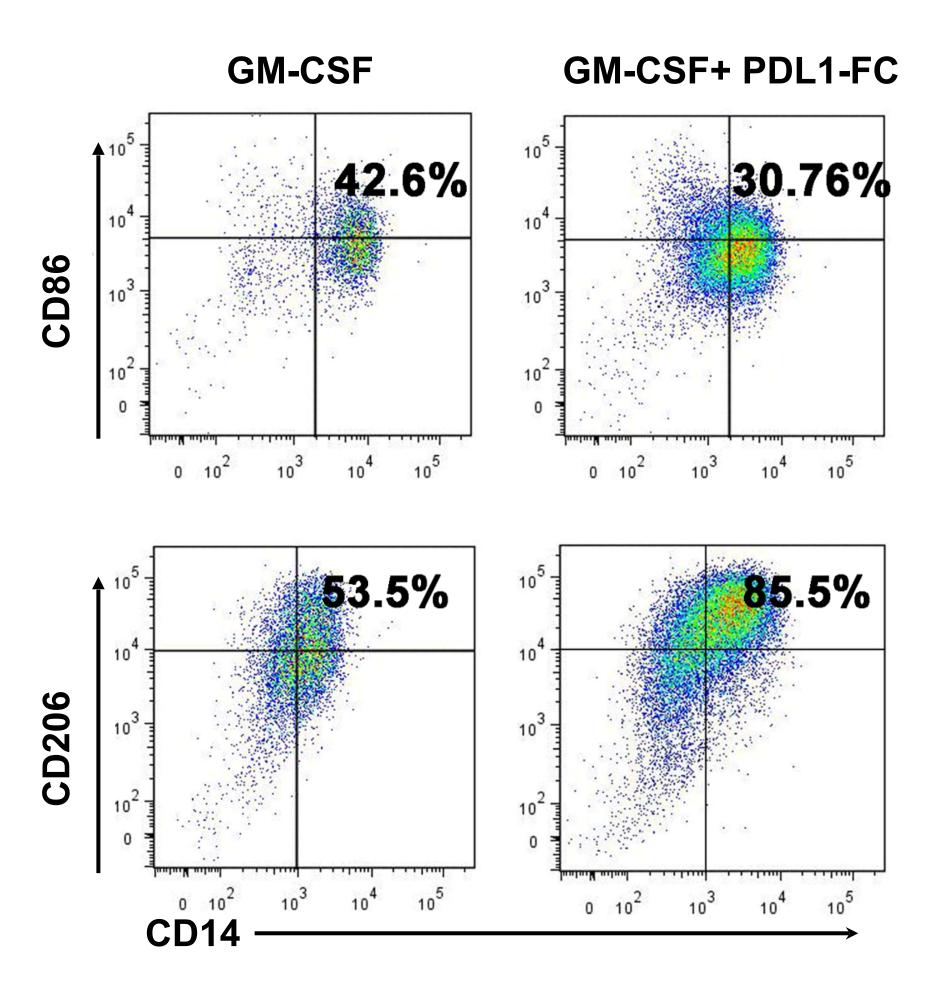
Sup Fig 1



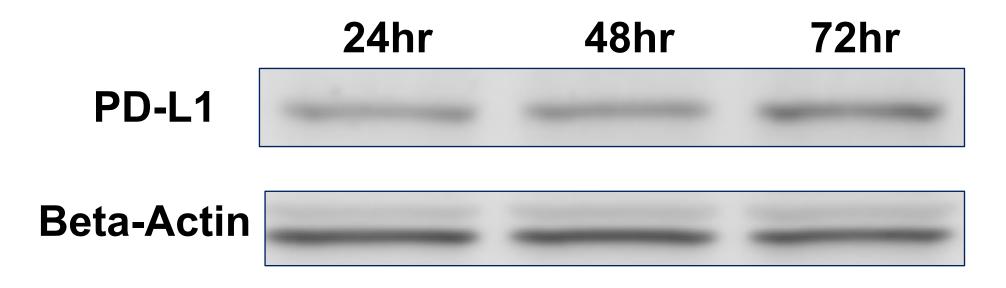
Sup. Fig. 1. Flow Cytometry for macrophage differentiation markers. Isotype antibodies used for each marker.

Sup Fig 2

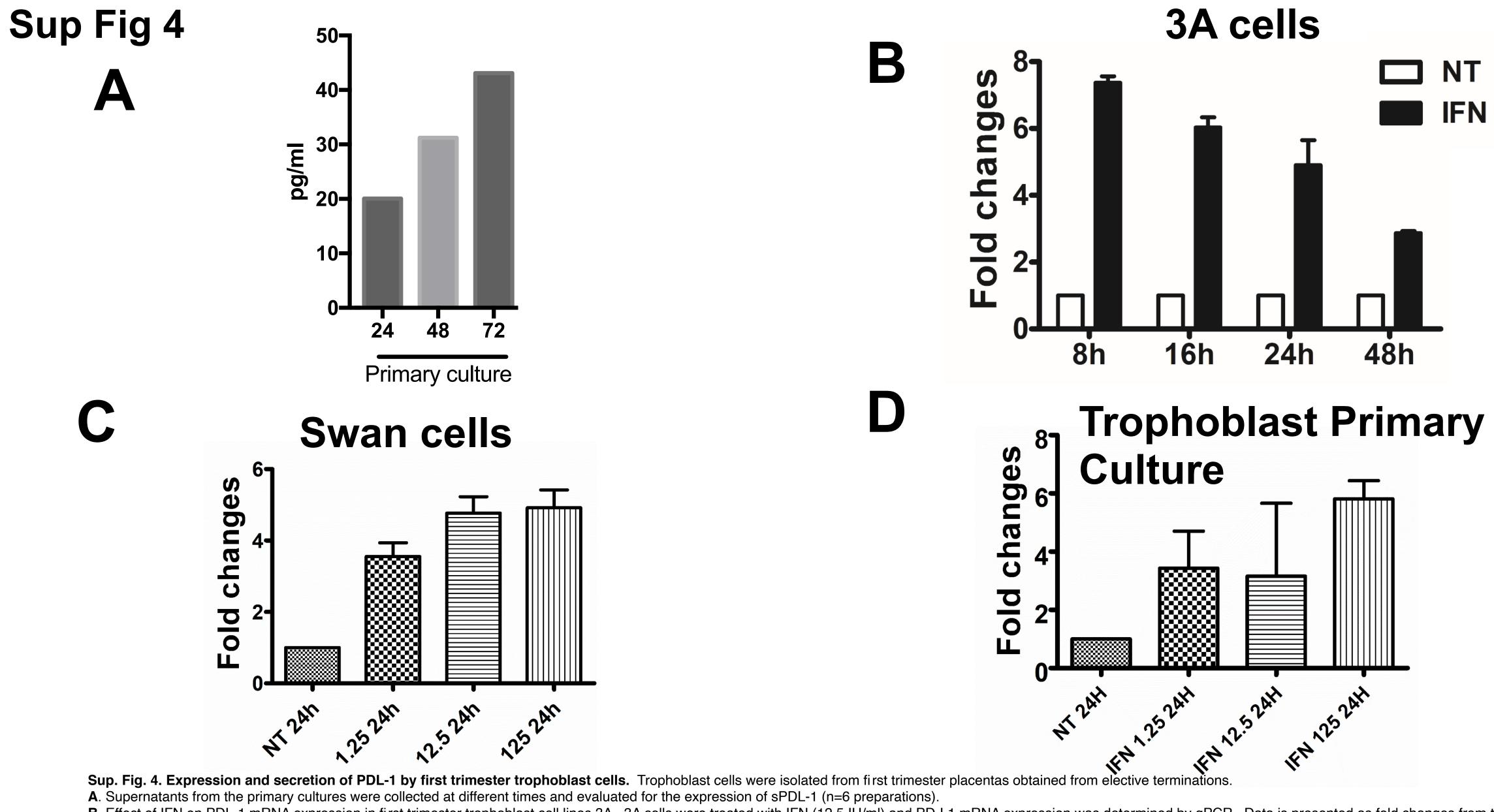


Sup. Fig. 2. Effect of stimulating the PD-1 signaling pathway in GM-CSF M1 differentiated macrophages. Peripheral CD14+ macrophages were differentiated with GM-CSF in the presence or absence of the recombinant PD-L1 agonist (PD-L1-FC) and their phenotype evaluated by Flow Cytometry.

Sup Fig 3



Sup. Fig. 3. Expression of PD-L1 in first trimester trophoblast Swan 71 cell line. Cell lysate was prepared from trophoblast Swan 71 cells every 24h and PD-L1 protein expression determined by western blot analysis. Beta Actin was housed as housekeeping gene.



B. Effect of IFN on PDL-1 mRNA expression in first trimester trophoblast cell lines 3A. 3A cells were treated with IFN (12.5 IU/ml) and PD-L1 mRNA expression was determined by qPCR. Data is presented as fold changes from the control group treated with vehicle (NT).

C. Effect of IFN on PDL-1 mRNA expression in first trimester trophoblast cell Swan 71 cells. Time dependent induction of PDL-1 mRNA expression by trophoblast cells incubated in the presence of increasing concentrations of IFN. n=3 independent experiments

D. Effect of IFN on PDL-1 mRNA expression in primary cultures of first trimester trophoblast cells. Time dependent induction of PDL-1 mRNA expression by trophoblast cells incubated in the presence of increasing concentrations of IFN. n=6 independent preparations