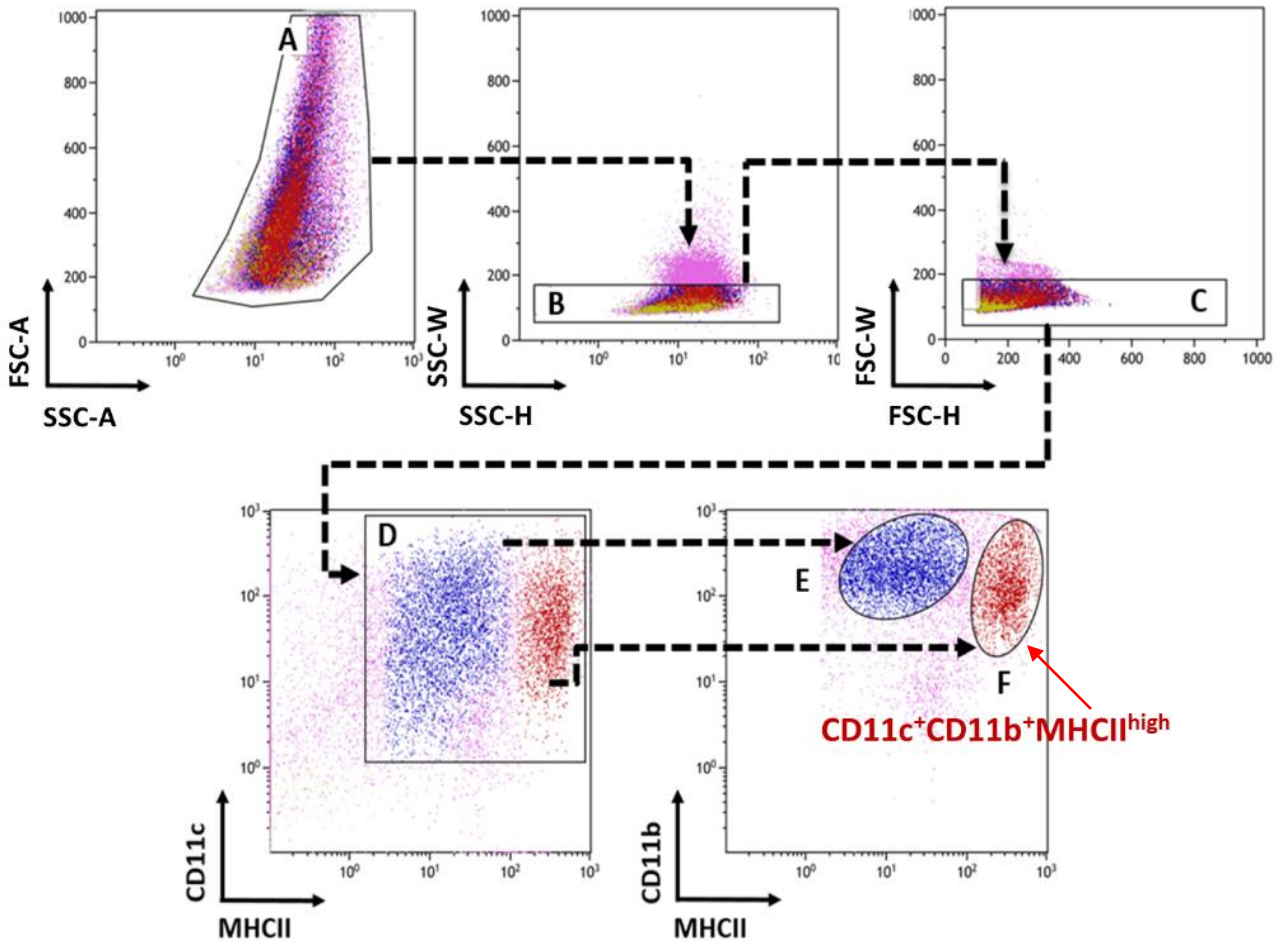


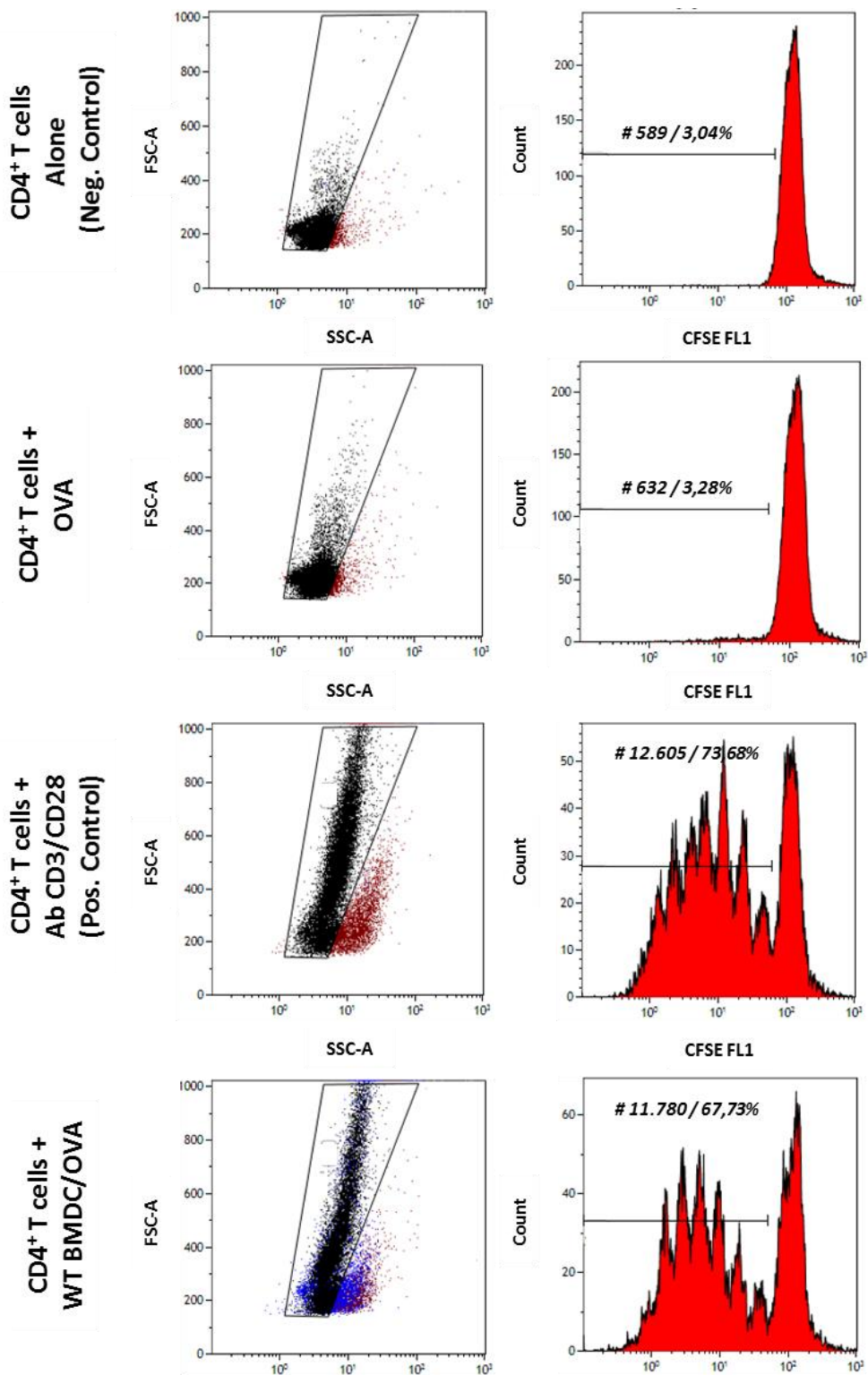
Supplementary Table 1. Primers sets for the different genes amplified by PCR.

	Forward primer	Reverse primer
<i>hPARP1</i>	5'-GGAAGCTGGAGGAGTGACAG-3'	5'-ACACCCCTTGCACGTACTIONTC-3'
<i>hiNOS</i>	5'-TCCTACACCACACCAAAC-3'	5'-CTCCAATCTCTGCCTATCC-3'
<i>hGAPDH</i>	5'-TCCCTGAGCTGAACGGGAAG-3'	5'-GGAGGAGTGGGTGTCGCTGT-3'
<i>mVCAM-1</i>	5'-AGTTGGGGATTTCGGTTGTTCT-3'	5'-CCCCTCATTCCCTTACCACCC-3'
<i>mβ-actin</i>	5'-ACCGTGAAAAGATGACCCAGATC-3'	5'-TAGTTTTCATGGATGCCACAGG-3'

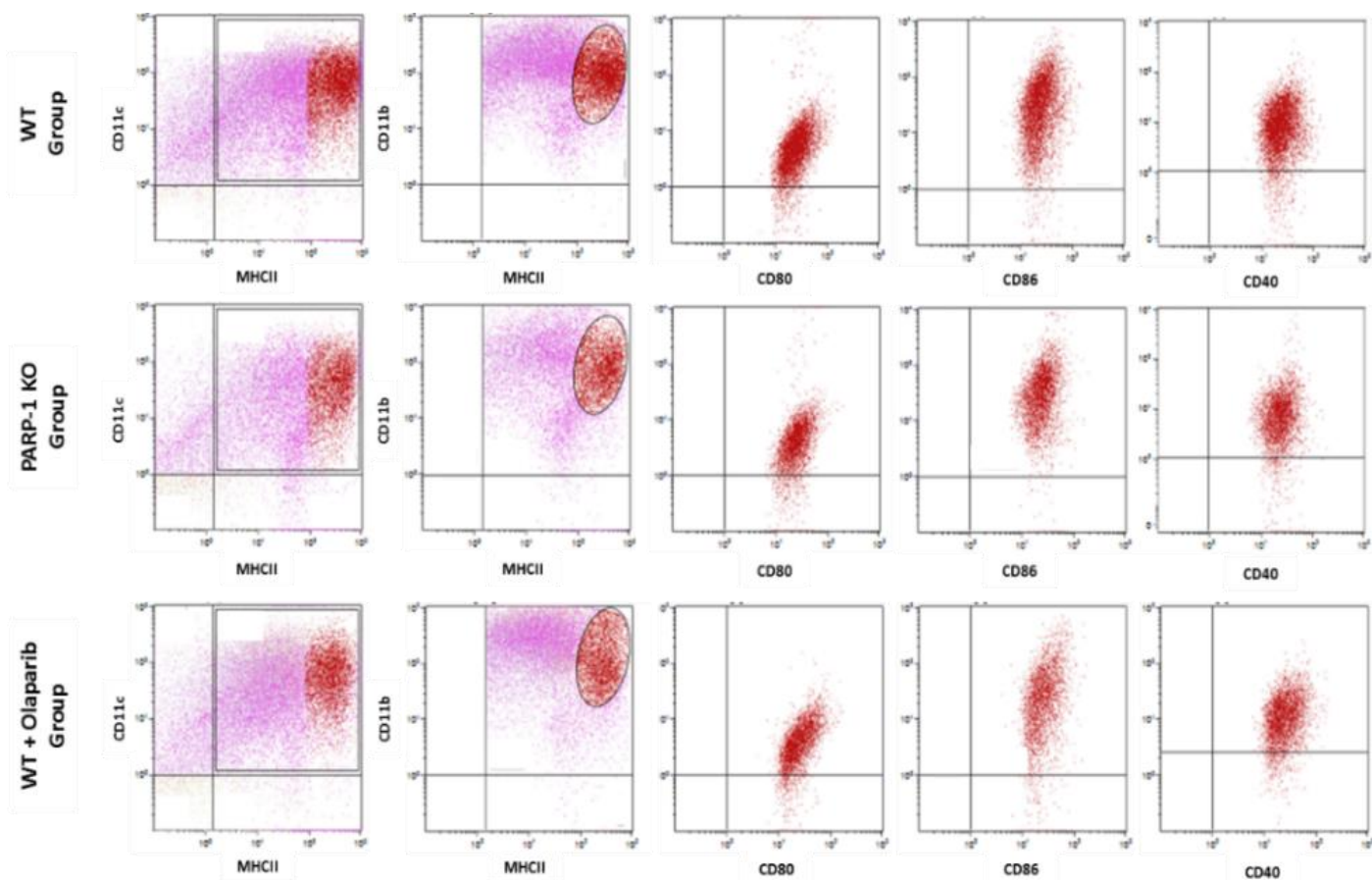
Supplementary Fig. S1. Gating strategy for DC identification and phenotyping.



Supplementary Fig. S2. Gating strategy and representative histograms pertinent to the determination of T cell proliferation.



Supplementary Fig. S3. Representative dot plots for the data displayed in Fig. 2.



Supplementary Fig. S4. Percent of CD11b⁺/CD11c⁺/MHCII^{interm} cells in the different experimental groups; this is related to results displayed in Fig.2B and 2C.

