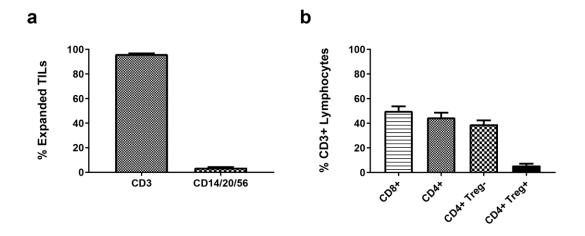
## Supplementary data

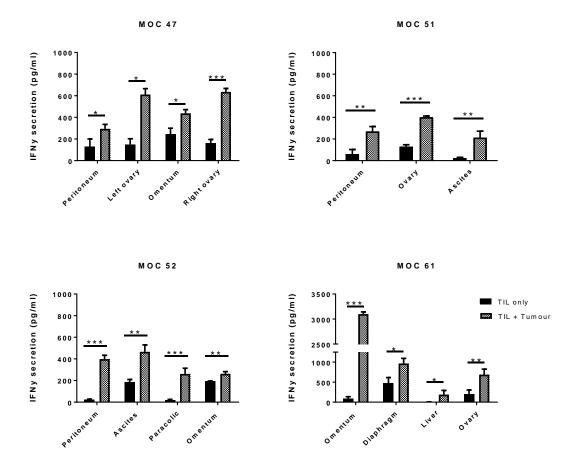
## **Supplementary Table 1. Flow cytometry antibodies**

Name	Fluorophore	Clone	Company	Panel used
Anti-CCR7	APC	REA108	Miltenyi, Germany	Differentiation
Anti-CD45RA	PE	HI100	BD Biosciences, UK	
Anti-CD45RO	PE	UCHL1		
Anti-CD28	PE	CD28.2		
Anti-CD27	PE	M-T271		
Anti-CD25	PE	M-A251		
Anti-CD8	FITC	HIT8A		
Anti-CD62L	Pe/Cy7	DREG-56	eBioscience, UK	
Live/Dead	BV395	N/A	Invitrogen	Activation, Treg,
Anti-CD4	BV785	OKT4	Biolegend, UK	Exhaustion & Functional Panels
Anti-CD14	Pe-Cy7	HCD14		
Anti-CD20	Pe-Cy7	2H7		
Anti-CD56	Pe-Cy7	HCD56		
Anti-CD45	BV510	UCHL1		
Anti-CD3	APC	UCHT1		
Anti-PD-1	PE	EH12.2H 7		
Anti-TIM3	BV421	F38-2E2		
Anti-CD69	BV421	FN50		
Anti-CD25	BV711	BC96		
Anti-OX40	FITC	ACT35		
Anti-Foxp3*	PE	206D		
Anti-CD127	BV421	A019D5		
Anti- TNFα*	BV650	MAb11		
Anti- IFNγ*	BV711	4S.B3		
GranzymeB	FITC	GB11		
Anti-ICOS	BV650	Dx29	BD Biosciences, UK	
Anti-CD8	APC-H7	SK1		
Anti-IL2*	BV421	5344.111		
Anti-LAG3	FITC	17B4	Enzo Lifesciences	

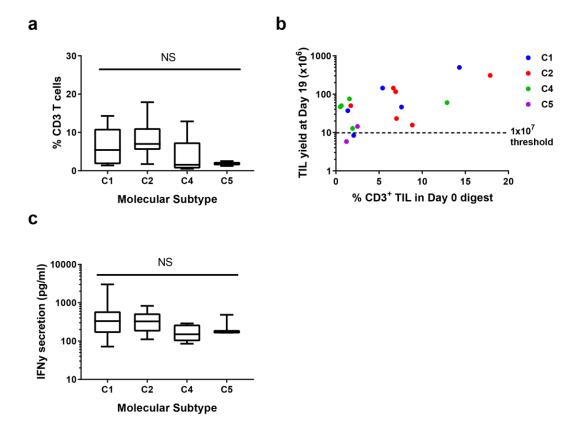
<sup>\*</sup>Denotes intracellular antigens



**Supplementary Figure 1. Phenotype of expanded ovarian TILs.** (a) Frequency of CD3<sup>+</sup> T cells and other lineages (CD14<sup>+</sup>/CD20<sup>+</sup>/CD56<sup>+</sup>) in the expanded TIL cultures. (b) Percentage CD8<sup>+</sup>, CD4<sup>+</sup> and Treg cells in the expanded TILs cultures. Data are reported as mean % ± SEM.



Supplementary Figure 2. Comparison of functional activity of TILs expanded from different anatomical sites. In four patients, TILs were successfully expanded from a minimum of three different anatomical sites. Expanded TILs were co-cultured against autologous tumour from cognate sites, to determine whether expanded TILs retained functional activity. TILs were considered to show functional activity if significantly higher levels of IFNγ were produced by the co-culture compared to TILs alone. Mean and SD of three replicates are shown. \*P<0.05, \*\*P<0.01, \*\*\*P<0.001, paired student T-test.



Supplementary Figure 3. Comparison of CD3 T cell infiltration, expansion and IFNγ secretion by the different molecular subtypes of ovarian cancer. (a) Percentage of CD3+ T cells in the tumour digest was determined using flow cytometry and compared between molecular subtypes of ovarian cancer (C1/mesenchymal; C2/immune; C4/differentiated; C5/proliferative). (b) Final TIL yield at day 19 compared against percentage of CD3+ TIL in tumour digest. Dashed line illustrates 1x10<sup>7</sup> threshold for a positive TIL culture. (c)IFNγ produced by expanded TILs in response to autologous tumour cells was compared between molecular subtypes. Data represents the median, upper and lower quartiles and range. Kruskal-Wallis test; P<0.05, NS, not significant.