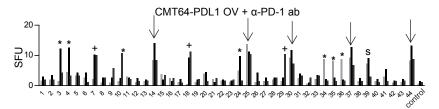
## Figure S3

Tumor-associated epitopes of CMT64 induced by PD-1 blockade and virotherapy (≥50% of responsive mice)					
gene i.d.	gene name	mutation	minimal sequence	score	MHC-type
NM_010872	NLR family, apoptosis inhibitory protein 2 (Naip2)	Y540N	CPLL <u>N</u> RFQL	22	H2-Db
NM_172967	Lung adenoma susceptibility 2 (Las2)	194T	H <u>T</u> PDNAFVNL	26	H2-Db
NM_016701	Nestin (Nes)	S570L	SIEEN <u>L</u> GTV	22	H2-Db

Fig. S3a A table listing the epitopes that were specifically triggered by combination therapy in at least 50% of responding mice.



**Fig. S3b** PD-L1-expression by CMT64 tumors does not affect epitope spreading upon combined treatment. The same experimental setup as in Fig. 4B for the combined treatment was applied in mice bearing CMT64-PDL1 tumors (n=8 treated animals). In total, strong expansion of mutanome-specific responses was detected in 9 out of 16 treated animals (8 animals bearing CMT64-tumors, another 8 animals bearing CMT64-PDL1-tumors).