

Supplementary Material

Table S1. List of CMV T-cell epitopes used in the quantiferon technique.

Epitope Sequence	HLA Restriction	Epitope Code	Antigen
IMREFNSYK	A3	IMR	gB
VLEETSVML	A2	VLE	IE-1
QIKVRVDMV	B8	QIK	IE-1
AYAQKIFKIL	A23&A24	AYA	IE-1
FEQPTETPP	B41	FEQ	IE-2
VTEHDTLLY	A1	VTE	pp50
TPRVTGGGAM	B7	TPR	pp65
QYDPVAALF	A24	QYD	pp65
QMWQARLTV	B52	QMW	pp65
QEFFWDANDI	B44	QEF	pp65
QAIRETVEL	B57&B58	QAI	pp65
PFTSQYRIQGKL	A24	FTS	pp65
NLVPMVATV	A2	NLV	pp65
KMQVIGDQY	B40/60	KMQ	pp65
GPISGHVLK	A11	GPI	pp65
FPTKDVAL	B35	FPT	pp65
DIYRIFAEL	A26	DIY	pp65
DALPGPCI	B51	DAL	pp65
CPSQEPMSIYVY	B35	CPS	pp65
CEDVPSGKL	B40/60	CED	pp65
ARVYEIKCR	B27	ARV	pp65

Current immunoassays use a broad spectrum of CMV-antigen types (synthetic peptides, viral lysates, recombinant antigens). However, such materials have, to date, not been issued as an official WHO standard. Of note, in quantiferon analysis, most of the epitopic peptides are derived from **pp65** (i.e. 15 to highlight presented in the bold font), but only one was epitope from immunodominant surface glycoprotein B (**gB**), sometimes used in the pentamer technique. Most notably, HLA specificities are not uniformly distributed e.g., only one **HLA-A3**-restricted epitope (**blue**), contrary to 3 **-HLA-A24**-restricted (**red**). It may be a source of inter-individual variation. Under the assumption of the quantiferon technique, such HLA-restricted CMV epitopes acts via TCR to activate T cell immune response and cytokine release. Limitations of CMV-interferon gamma release assay (IGRA) and a high rate of intermediate results [11,13,14,32] were described, but IL-2 as a T cell specific cytokine yielded inferior results and was negligible as a surrogate marker [32]. The IFN γ release was significantly higher (1324 or 5871 pg/mL) than IL-2 (44 or 883 pg/mL) under the influence of recombinant or synthetic pp65 peptides [32].