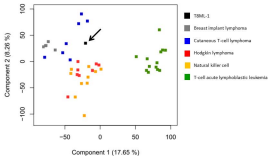


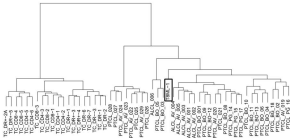
Peripheral T-cell lymphoma cell line T8ML-1 highlights conspicuous targeting of PVRL2 by t(14;19)(q11.2;q13.3)

Stefan Ehrentraut,^{1*} Stefan Nagel,¹ Claudia Pommerenke,¹ Wilhelm G. Dirks,¹ Hilmar Quentmeier,¹ Maren Kaufmann,¹ Corinna Meyer,¹ Margarete Zaborski,¹ Robert Geffers,² Hiroshi Fujiwara,³ Hans G. Drexler¹ and Roderick A. F. MacLeod¹

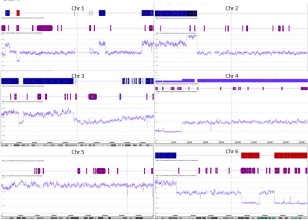
¹DSMZ - German Collection of Microorganisms and Cell Cultures, Department of Human and Animal Cell Lines, Braunschweig, Germany; ²Genome Analytics Research Group, Helmholtz Centre for Infection Research, Braunschweig, Germany and ³First Department of Internal Medicine, Ehime University Hospital, Ehime, Japan

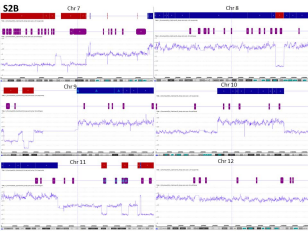
Correspondence: rmf@dsmz.de
doi:10.3324/haematol.2017.168203





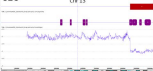
S2A





S2C

Chr 13



Chr 14



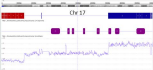
Chr 15



Chr 16



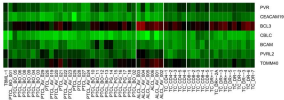
Chr 17



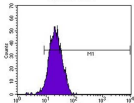
Chr 18



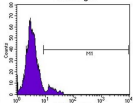
S2D**Chr 19****Chr 20****Chr 21****Chr 22****Chr X****Chr Y**



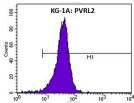
T8ML-1: PVRL2



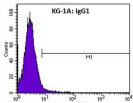
T8ML1: IgG1

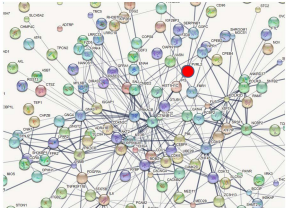


KG-1A: PVRL2



KG-1A: IgG1





Supplementary Figure Legends

Supplementary Figure S1: Gene expression profiling (GEP) of T-cell lymphoma cell lines. Principal component analysis was performed on GEP data. Note isolation of T8ML-1. Names of cell lines are available on request. **B: Cluster diagram of T8ML-1 and PTCL patients:** GEP clustering shows that T8ML-1 consorts with PTCL patient profiles. Primary lymphoma and normal T cells expression data were obtained from the public GEO dataset GSE3668 (9). After RMA-background correction and quantile normalization of the spot intensities via R/Bioconductor using *limma* and *affy* packages (<http://www.bioconductor.org>), hierarchical clustering was calculated on basis of Ward's method.

Supplementary Figure S2: Copy number alterations and zygoty in T8ML-1. Figure summarizes genomic array data for chromosomes 1-6 (A), 7-12 (B), 13-18 (C), and 19-22, X and Y (D). Note gross changes in copy number associated with cytogenetic breakpoints (**Fig. 1A**). For technical details see legend to Fig. 1.

Supplementary Figure S3: PTCL patient gene expression at 19q13.3: Shows GEP incorporating public GEP data (GEO-GSE6338) from ref. 9. Note moderate upregulation of *PVRL2* in PTCL patient PTCL-026 only and in two ALCL patients, while *BCL3* is usually downregulated in PTCL patients, when compared to primary T-cells. For data source see Supplementary Figure 1.

Supplementary Figure S4: Surface expression of PVRL2 in T8ML-1 cells. Flow cytometric plots for PVRL2 and (reference) IgG1 on T8ML-1 and (positive) control KG-1A cells. Cell surface expression of PVRL2 closely matches that of positive control cells. Cells were analyzed by flow cytometry on a FACSCalibur (BD Biosciences, Heidelberg/Germany) using CellQuest Pro software.

Supplementary Figure S5: Protein interactogram. Shows STRING analysis results of interactions predicted among top 500 expressed protein coding genes in T8ML-1 versus congener T-cell lymphoma cell lines. PVRL2 (shown red) is connected to CD96, CTNNB1, GOPC based on co-citation evidence. Functional associations remain to be determined.