

POSTER PRESENTATION

Open Access

Tetramer based approach for efficient identification and isolation of neo-antigen specific CD8 T cells from peripheral blood (PBL) of patients with metastatic cancers

Mini Bharathan¹, Katarina Trebska-McGowan², Pasetto Anna², Drew C Deniger¹, Ken-ichi Hanada², Jared J Gartner², James C Yang³, Steven A Rosenberg¹, Paul F Robbins^{3*}

From 30th Annual Meeting and Associated Programs of the Society for Immunotherapy of Cancer (SITC 2015) National Harbor, MD, USA. 4-8 November 2015

Background

Adoptive cell therapy with T cells bearing mutation specific T cell receptors (TCR) can be an effective method for treating metastatic cancers. The objective of this study was to identify mutation reactive T cells in the circulation of patients with different types of metastatic cancer.

Methods

The strategy utilized whole exome sequencing data to identify somatic non-synonymous mutations and then in-silico algorithms to predict minimal epitopes encoding mutated amino acids for each patient specific HLA-allele. CD8-enriched PBL from each patient were stained with tetramers generated in house by a UV-exchangeable technique as previously described for A*02:01, A*03:01, A*11:01, B*07:02, and a commercial tetramer was acquired for B*57:01. Based on the initial staining frequency (+tetramer⁺ T cells recognizing 7 unique neo-antigens from the PBL of 4 patients (ranging from 1 to 4 per patient). We enriched the frequencies of CD8⁺tetramer⁺ cells from 0.5 to >85%, 0.3 to >65% and 0.01 to 3% from the PBL of patients with colorectal (3971-A*02:01), lung (4014-B*57:01), and ovarian (4067-B*07:02) cancers respectively, using individual tetramers. Populations reactive with three HLA-A*11:0-restricted and one HLA-A*03:01-restricted neoantigens were also isolated from the PBL of lung cancer patients 4014 and 4037, respectively, using a pooled tetramer approach.

Results

Overall, the isolated T cells recognized mutated epitopes when co-cultured with autologous CD14⁺ monocytes pulsed with mutated peptides in the context of appropriate MHC-I alleles including HLA-A*02:01, HLA-A*03:01, HLA-A*11:01, HLA-B*07:02, and HLA-B*57:01, with reactivity detected using IFN- γ ELISA. Using single cell PCR, we could clone the TCRs reactive with an HLA-*02:01-presented colon cancer neoantigen and an HLA-B*57:01-presented lung cancer neoantigen. Evaluation of PBL retrovirally-transduced with these TCRs demonstrated that they bound to tetramers and secreted IFN- γ when co-cultured with CD14⁺ monocytes pulsed with appropriate mutated peptides.

Conclusions

To conclude, tetramers offer a sensitive, fast and reliable methodology to isolate mutation specific tumor reactive T cells from PBL of cancer patients. Furthermore, this method facilitates the identification, and cloning of mutation reactive TCR with which to construct receptor-engineered T cells for adoptive T cell therapy.

Authors' details

¹NIH/NCI, Bethesda, MD, USA. ²NIH/NCI Surgery Branch, Bethesda, MD, USA. ³National Institutes of Health, Bethesda, MD, USA.

³National Institutes of Health, Bethesda, MD, USA

Full list of author information is available at the end of the article

Published: 4 November 2015

doi:10.1186/2051-1426-3-S2-P47

Cite this article as: Bharathan *et al.*: Tetramer based approach for efficient identification and isolation of neo-antigen specific CD8 T cells from peripheral blood (PBL) of patients with metastatic cancers. *Journal for ImmunoTherapy of Cancer* 2015 **3**(Suppl 2):P47.

**Submit your next manuscript to BioMed Central
and take full advantage of:**

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at
www.biomedcentral.com/submit

