



POSTER PRESENTATION

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“Immune B Cells know it better”: tumorimmunological panel assay to define tumor-associated antigen binding antibodies in patients with metastatic melanomas

Beatrix Kotlan^{1,2*}, Timea Balatoni³, Katalin Csirbesz³, Judit Olasz⁴, Orsolya Csuka^{4,6}, Laszlo Toth⁵, Emil Farkas⁵, Akos Savolt⁵, Andras Szollar⁵, Mihaly Ujhelyi⁵, Szabolcs Horvath², Klara Eles², Miklos Kasler⁶, Francesco M Marincola⁷, Gabriella Liszky^{3,6}

From Melanoma Bridge Meeting 2014
Naples, Italy. 03-06 December 2014

Background

Revealing novel cancer targeting biomarkers is a great challenge, and especially urging in cancer types with a more pronounced metastatic feature. We focus on potential anti-tumor immune reactions of the host. In order to harness the natural humoral immune response a novel immunological and molecular genetic panel assay has been developed for the investigation of patients with melanomas.

Materials and methods

Punch biopsies were taken of surgically removed fresh cancerous tissues and peripheral blood was gathered from the patients involved into the study (n= 125). Ethical permission was provided by the Scientific and Research Ethics Committee of the Medical Research Council of the Hungarian Ministry of Health (ETT TUKEB 16462- 02/2010). We established and standardized two experimental strategies (Epstein Barr virus transformation and cloning with limiting dilution assay (LDA) and tumor infiltrating B cell (TIL-B) antibody phage display technology) and started basic processes for a detailed immunoglobulin repertoire analysis at DNA level. We set up a novel native tumor cell membrane preparation technique extremely useful for specific detection of tumor reacting antibodies or antibody fragments (eg: immunoblotting, scFv phagemid ELISA). Defining

essential tumor-associated antigens on the cancerous tissue specimen by immunohistochemistry became the other part of the tumorimmunological panel assay. Results: We claim that this complex quantitative and qualitative analysis of antibodies in sera and in the tumor microenvironment results in revealing tumorspecific antibodies of human origin. Our antibody profile analysis revealed glycoprotein and sialylated glycolipid based tumor-associated antigen-specific antibody-variable regions in various patterns.

Conclusions

The present technological developments enable the specific detection of cancer associated sialylated glycolipid and glycoprotein antigens with unique characteristics. The study helps to understand the question of “abnormal glycosylation” and its role in cancer. We conclude that the complex tumorimmunological assay has important potentials in evaluating the host’s anti tumor immune status.

Acknowledgement

Harry J. Lloyd Charitable Trust Melanoma Research Award (2010), previous Fulbright No1206103, present Fulbright No1214104 and OTKA T038488 Grants.

Authors’ details

¹Molecular Immunology and Toxicology, National Institute of Oncology, Budapest, Hungary. ²Center of Surgical and Molecular Tumorpathology, National Institute of Oncology, Budapest, Hungary. ³Oncodermatology, National Institute of Oncology, Budapest, Hungary. ⁴Pathogenetics, National Institute of Oncology, Budapest, Hungary. ⁵Oncosurgery, National Institute of Oncology, Budapest, Hungary. ⁶Board of Directors, National Institute of

¹Molecular Immunology and Toxicology, National Institute of Oncology, Budapest, Hungary

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

Oncology, Budapest, Hungary. ⁷SIDRA Medical and Research Centre, Doha, Qatar.

Published: 15 January 2015

doi:10.1186/1479-5876-13-S1-P16

Cite this article as: Kotlan *et al.*: "Immune B Cells know it better": tumorimmunological panel assay to define tumor-associated antigen binding antibodies in patients with metastatic melanomas. *Journal of Translational Medicine* 2015 13(Suppl 1):P16.

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