

POSTER PRESENTATION

Open Access

Association between the CCR5 Δ 32 polymorphism and preeclampsia

Bianca de Paula Telini*, Tiago Degani Veit, Priscila Vianna, José Artur Bogo Chies

From 5th Congress of the Brazilian Biotechnology Society (SBBIOTEC)
Florianópolis, Brazil. 10-14 November 2013

Background

Preeclampsia (PE) is a condition that occurs in up to 7% of all pregnancies. Its exact pathophysiology is not known yet, but there is the involvement of genetic and immune factors maternal and fetal, with occurrence of hypertension and proteinuria. There is evidence of increased systemic inflammation during the first trimester of pregnancy in women with preeclampsia. The PE usually develops in the second half of pregnancy and is characterized by events of endothelial dysfunction and inflammation in its pathogenesis. Chemokines (proinflammatory cytokines) are considered the main determinants of the inflammatory response and its action by binding to specific receptors can be directly related to the development of PE. The chemokine receptor type 5 (CCR5) is a protein encoded by the CCR5 gene which is located on chromosome 3p21.3 p24. The polymorphic variant CCR5 Δ 32 resulting from deletion of 32 base pairs in this gene leads to production of a non-functional isoform of the receptor, and has been implicated in a variety of autoimmune diseases.

Methods

To investigate the role of this polymorphism in the pathogenesis of preeclampsia, we evaluated the frequency of polymorphic variant CCR5 Δ 32 among women with and without PE. In order to do this, 155 pregnant women with PE and 144 pregnant women without PE were genotyped (both groups with similar age).

Results and conclusions

The presence of the mutant delta 32 was associated to protection against the development of PE, i.e. carriers of the Δ 32 allele had a lower chance of developing preeclampsia (OR 0.342, CI 95%, $p = 0.047$). We suggest that

cells involved in the modulation of the immune response during pregnancy which express this protein cannot migrate properly to certain sites of inflammation and this could alter the individual's inflammatory profile. Such an imbalance would make less likely the establishment of a pathological inflammatory profile characteristic of preeclampsia. Our study suggests the CCR5 Δ 32 as an independent factor in the susceptibility to PE together with primary hypertension, which was also associated to PE risk in our cohort (OR 8.696, CI = 95%, $p < 0.001$).

Published: 1 October 2014

Reference

1. Gurdol F, Yurdum LM, Ozturk U, Isbilen E, Cakmakoglu B: Association of the CC chemokine receptor 5 (CCR5) polymorphisms with preeclampsia in Turkish women. *Arch Gynecol Obstet* 2012, **286**(1):51-4.

doi:10.1186/1753-6561-8-S4-P70

Cite this article as: Paula Telini et al.: Association between the CCR5 Δ 32 polymorphism and preeclampsia. *BMC Proceedings* 2014 **8**(Suppl 4):P70.

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



Universidade Federal do Rio Grande do Sul, Porto Alegre, Brazil



© 2014 Paula Telini et al.; licensee BioMed Central Ltd. This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited. The Creative Commons Public Domain Dedication waiver (<http://creativecommons.org/publicdomain/zero/1.0/>) applies to the data made available in this article, unless otherwise stated.