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To cite this version:

HAL Id: inserm-00914630
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Submitted on 5 Dec 2013

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ORAL PRESENTATION

PReS-FINAL-2173: Protein kinase C delta deficiency is a new cause of monogenic SLE

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From 20th Pediatric Rheumatology European Society (PReS) Congress Ljubljana, Slovenia. 25-29 September 2013

Introduction
Systemic lupus erythematosus (SLE) is a prototype autoimmune disease. Infectious triggers, genetic background, immunological abnormalities and environmental factors are all supposed to interact in disease development. Rare causes of monogenic SLE have been described, (e.g. complement deficiencies, interferonopathies and FasL deficiency) providing unique insights into fundamental mechanisms of immune tolerance.

Objectives
Our objective was to identify the cause of an autosomal recessive form of SLE in an inbred family with three affected siblings.

Methods
We investigated three siblings and used next generation sequencing to identify mutations in the disease-associated gene. We performed extensive biochemical, immunological and functional assays to assess the impact of the identified mutations on B cell biology.

Results
Genetic mapping and targeted exome sequencing led to the identification of a homozygous mutation in PRKCD, encoding protein kinase C delta (PKCδ). Mutation of PRKCD resulted in reduced expression and activity of encoded protein PKCδ. In mouse, PKCδ plays a crucial role in the deletion of autoreactive B cells. As for mice deficient in PKCδ, we demonstrated that B cells display a resistance to calcium-dependent apoptosis and a higher proliferation rate associated with an increase of immature B cells in affected patients, and a developmental shift toward an immature phenotype of naïve B cells.

Conclusion
Our findings indicate that PKCδ is crucial in regulating B cell tolerance and preventing self-reactivity in humans.

Disclosure of interest
None declared.

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Published: 5 December 2013

doi:10.1186/1546-0096-11-S2-O8
Cite this article as: Belot et al. PReS-FINAL-2173: Protein kinase C delta deficiency is a new cause of monogenic SLE. Pediatric Rheumatology 2013