



POSTER PRESENTATION

Open Access

Severe erythrodermic psoriasis and arthritis as clinical presentation of a CARD14-mediated psoriasis (CAMPS)

S Signa^{1*}, M Rusmini², E Campione³, I Gueli¹, A Grossi², A Omenetti¹, L Bianchi³, A Martini¹, I Ceccherini², M Gattorno¹

From 8th International Congress of Familial Mediterranean Fever and Systemic Autoinflammatory Diseases Dresden, Germany. 30 September - 3 October 2015

Introduction

Autosomal dominant gain of function mutations in caspase recruitment domain family member 14 (CARD14) were found to cause plaque psoriasis in two families and severe generalized pustular psoriasis as a monogenic form of childhood (CARD14-mediated psoriasis, CAMPS) [1]. CARD14 mutations have also been implicated in plaque-type psoriasis and pityriasis rubra pilaris [2].

Objectives

Describing a family with an unusual clinical phenotype characterized by some members with childhood-onset erythrodermic psoriasis first localized and then diffuse over all the skin surface; in some family members is also reported psoriatic arthritis.

Patients and methods

We assessed for the first time the family in december 2013, because of their skin lesions and family recurrence for erythrodermic psoriasis associated to arthritis in some cases. There are three pairs of twins, five of them presenting psoriasis and two of them presenting psoriatic arthritis. The children presented poor clinical response to topic and systemic therapy with antihistamine, steroid, retinoids, cyclosporine and etanercept. After exclusion of the most common genes associated to autoinflammatory diseases (IL36RN, IL1RN, MVK, TNFRSF1A, NLRP3, NLRP12, MEFV, IL1RN, NOD2, PSMB8, PSTPIP1, LPIN2) we approached the new gene search by subjecting to Whole Exome Sequencing (WES) analysis five members of the family. Samples were processed in outsourcing

and raw data were transferred to our lab for the bioinformatic analysis.

Results

Among variants shared by the four affected individuals and not present in the unaffected subject, a missense mutation of the CARD14 gene resulted worth of further investigation. In particular, it was the case of an exon4 heterozygous nucleotide change, c.446T>G, leading to the missense amino acid substitution p.L149R. The presence of this variant was validated by Sanger sequencing not only in the affected members undergone WES, but also assessed in all the rest of the available family members. This allowed us to confirm the expected segregation of the CARD14 mutation with the disease phenotype.

Conclusions

CARD14 gain of function mutations can give rise to unusual clinical phenotype like diffuse erythrodermic psoriasis and psoriatic arthritis. WES appears to be a powerful, suitable and proper approach to identify new SAID genes, thus also disclosing new molecular pathogenic mechanisms.

Authors' details

¹IRCCS G. Gaslini, U.O. Pediatria II, Genoa, Italy. ²IRCCS G. Gaslini, U.O. Genetica Medica, Genoa, Italy. ³Tor Vergata University of Rome, Department of Dermatology, Rome, Italy.

Published: 28 September 2015

References

1. Jordan CT, Cao L, Roberson ED, Pierson KC, Yang CF, Joyce CE, et al: **PSORS2 is due to mutations in CARD14.** *American journal of human genetics* 2012, **90**(5):784-95, Epub 2012/04/24.

¹IRCCS G. Gaslini, U.O. Pediatria II, Genoa, Italy
Full list of author information is available at the end of the article

- Fuchs-Telem D, Sarig O, van Steensel MA, Isakov O, Israeli S, Nousbeck J, *et al*: **Familial pityriasis rubra pilaris is caused by mutations in CARD14.** *American journal of human genetics*.n 2012, **91**(1):163-70, Epub 2012/06/19.

doi:10.1186/1546-0096-13-S1-P57

Cite this article as: Signa *et al*: **Severe erythrodermic psoriasis and arthritis as clinical presentation of a CARD14-mediated psoriasis (CAMPS).** *Pediatric Rheumatology* 2015 **13**(Suppl 1):P57.

**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

