Date : 23 November 2018
Place : Bambino Gesù Hospital, Rome
Organizers : Department of Paediatric Dermatology, Ospedale Pediatrico Bambino Gesù, with the support of the ERN-SKIN training group and FRT-Fondation René Touraine.

PROGRAMME

9h00 – 10h30 AUTOIMMUNE BULLOUS DISEASES AND SEVERE CUTANEOUS DRUG REACTIONS

Co-chairs : Branka Marinovic, Pascal Joly

9H00-9H30 : CLINICAL CASES

- 9h00 – Paradoxical exacerbation of Bullous Pemphigoid following Rituximab treatment, *Roberto Maglie*
- 9h05 – Paraneoplastic pemphigus in a girl with post-transplantation lymphoproliferative disorder, *Marieke Bolling*
- 9h10 – EBA in 4-year-old girl, *Ivana Martinac*
- 9h15 – Diagnostic Conundrums in Bullous Diseases, *Vivien Hébert*
- 9h20 – IgA pemphigus associated with ulcerative colitis and review of the literature, *Maria Polina Konstantinou*
- 9h25 – Severe drug induced reactions Epidermal necrolysis can be non-toxic, *Sophie Lalevée*

9H30 – DISCUSSION AND COMMENTS
Branka Marinovic, Pascal Joly

10H00 – UPDATE : WHAT'S NEW IN THE TREATMENT OF AIBD ?
Saskia Oro

SESSION 2 : 11H00 – 12H30 EPIDERMOLYSIS BULLOSA

Co-chairs : Johann Bauer, Marcel Jonkman

1H00-11H30 : CLINICAL CASES

- 11h00 – Facing the challenges of managing Laryngo-onchyo-cutaneous syndrome, *Cassandra McDonald*
- 11h05 – Mild clinical phenotype in infants unravels EB genotype, *Irina Condrat*

- 11h10 – Recessive Dystrophic EB inversa : marked mucosal phenotype with limited cutaneous involvement, *Sorcha O'Sullivan*
- 11h15 – Cardiomyopathy in epidermolysis bullosa simplex patients with KLHL24 mutations : an intriguing association, *Vamsi Yenamandra*
- 11h20 – Recessive forms of dystrophic Epidermolysis Bullosa and anemia : a complex therapeutic challenge, *Lilia Bekel*
- 11h25 – JEB gen sev – unexpected natural histories, *Christine Prodinger*

11H30 – DISCUSSION AND COMMENTS
Johann Bauer, Marcel Jonkman

12H00 – UPDATE ON EB TYPES AND THERAPIES
Christina Has

12H30-13H30 : LIGHT LUNCH BUFFET

SESSION 3 : 13H30 – 15H00 ICHTHYOSIS & PALMOPLANTAR KERATODERMA

Co-chairs : May El Hachem, Kathrin Giehl

13H30-15H30 : CLINICAL CASES

- 13h30 – Clinical case on the Neonatal management of a collodion baby, *Aurore Brun*
- 13h35 – Clinical case on the use of tazarotene, *Luana Niculescu*
- 13h40 -Clinical case on itch in ichthyoses, *Ifigenia Spanoudi*
- 13h45 – Clinical case on Netherton syndrome, *Nathalia Bellon*
- 13h50 – Clinical case on extending the spectrum of autosomal epidermolytic ichthyosis, *Leonie Frommherz*
- 13h55 – Clinical case on the treatment of Palmoplantar Keratoderma, *Bjorn Thomas*

14H00 – DISCUSSION AND COMMENTS
Kathrin Giehl

14H30 – MANAGEMENT OF CONGENITAL ICHTHYOSIS IN 2018
Juliette Mazereeuw-Hautier

SESSION 4 : 15H00 – 16H30 ECTODERMAL DYSPLASIAS INCLUDING INCONTINENTIA PIGMENTI AND P63-ASSOCIATED DISORDERS

Co-chairs : Christine Bodemer, Angus Clarke

15H00-15H50 : CLINICAL CASES

- 15h00 – A woman with severe autosomal-dominant hypohidrotic ectodermal dysplasia, *Monika Ettinger*

- 15h05 – Ectodermal dysplasia with extreme palmoplantar hyperkeratosis in a young man, *Julia Hiernickel*
- 15h10 – Management of tooth abnormalities in a patient with X-linked hypohidrotic ectodermal dysplasia, Ajit Tanday
- 15h15 – Dermatological findings in patients with p63 deficiency, *Alexia Maillard*
- 15h20 – Sweating ability of patients with EEC and AEC syndrome, *Paul Ferstl*
- 15h25 – Ocular care in patients with incontinentia pigmenti, *Sarah Michel*

15H30 – DISCUSSION AND COMMENTS↵
Holm Schneider

16H00 – UPDATE ON ED TYPES AND THERAPIES↵
Smail Hadj-Rabia

16H30-17H00 : DISCUSSION AND CONCLUSIONS OF THE TRAINING AND ANNOUNCEMENT OF THE NEXT ONE↵
Christine Bodemer, Maya El-Hachem