Substantiating atypical phenotypes of epidermolysis bullosa
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Kasia, Groningen 2016
A novel PLEC isoform rescues the phenotype in epidermolysis bullosa simplex

KB Gostyńska, H Lemmink, J Bremer, HH Pas, M Nijenhuis, PC van den Akker, RJ Sinke, MF Jonkman, AM Pasmooij
Excerpts resubmitted to the Journal of Investigative Dermatology

Carriers with functional null mutations in LAMA3 have localized enamel abnormalities due to haploinsufficiency

KB Gostyńska, WY Yuen, AM Pasmooij, C Stellingsma, HH Pas, H Lemmink, MF Jonkman
Accepted for publication in the European Journal of Human Genetics.

In-frame exon skipping in KRT5 due to novel intronic deletion causes epidermolysis bullosa simplex, generalized severe

KB Gostyńska, J Bremer, KK van Dijk-Bos, RJ Sinke, AM Pasmooij, MF Jonkman
Accepted for publication in Acta Dermato-Venereologica

Epidermolysis Bullosa Pruriginosa Excoriée: A Deceptive Pruritic Variant in Two Female Patients.

KB Gostyńska, MF Jonkman
Acta Derm Venereol. 2016 Jan 20;96(1):140-141

Mutation in exon 1a of PLEC, leading to disruption of plectin isoform 1a, causes autosomal-recessive skin-only epidermolysis bullosa simplex

KB Gostyńska, M Nijenhuis, H Lemmink, HH Pas, AM Pasmooij, K Kernland Lang, MJ Castañón, G Wiche, MF Jonkman

Epidermolysis bulosa pruriginosa excoriée

KB Gostyńska, MF Jonkman
Nederlands Tijdschrift voor Dermatologie en Venereologie, Volume 23, September 2013

Residents’corner July 2012. sQUIZ your knowledge! Diagnosis: Laptop induced erythema ab igne.

K Karolak, MF Jonkman

Coexistence of Klippel-Trenaunay-Weber syndrome with Sturge-Weber syndrome
A Terlikowska-Brzósko, K Karolak, M Sińczak, B Kwiek
Postępy Dermatologii i Alergologii, XXV, 2008
Katarzyna (Kasia) Gostyńska (née Karolak) was born on December 2nd 1985 in Warsaw, Poland. In her youth, she emigrated with her parents and two older brothers to Canada where she lived until completing high school at Handsworth Secondary School in North Vancouver, B.C.

At 17, she returned to her native Poland to study medicine at the II Medical Faculty of the Medical University of Warsaw. She obtained her MD title in 2009 and worked as a medical intern in Międzyleski Szpital Specjalistyczny in Warsaw, Poland until 2010. She moved to Groningen, the Netherlands in 2011 and began her residency in Dermatology and Venereology in July, 2011. In July 2012 she briefly stepped out of clinical work to pursue research on the works presented in this thesis under the supervision of Prof. dr. Marcel Jonkman, Prof. dr. Richard Sinke, Dr. Marjon Pasmooij. Concurrently she was and continues to be the coordinating investigator of a clinical trial on hematopoietic stem cell therapy for children affected with severe epidermolysis bullosa in collaboration with the Wilhelmina Children's Hospital of the University of Utrecht, Utrecht the Netherlands.

In July 2014, she resumed her clinical residency. Her interests in dermatology include pediatric dermatology and genodermatoses.

Kasia currently lives in Groningen with her husband Antoni and their daughter Wiktoria.