

## Most important publications

Mohammadi J, C Liu, Aghamohammadi A, Bergbreiter A, Du L, Lu J, Rezaei N, Amirzargar AA, Moin M, Salzer U, Pan-Hammarström Q, Hammarström L. Novel Mutations in TAC1 (TNFRSF13B) Causing Common 5 Variable Immunodeficiency J Clin Immunol, DOI 10.1007/s10875-009-9317-5.

Salzer U\*, Bacchelli C\*, Buckridge S, Pan-Hammarström Q, Jennings S, Lougaris V, Hagena T, Birmelin J, Plebani A, Webster ADB, Peter HH, Suez D, Chapel H, Maclean-Tooke A, Spickett GP, Anover-Sombke S, Ochs HD, Urschel S, Belohradsky BH, Kumararatne DS, Lawrence TC, Holm AM, Franco JL, Schulze I, Schneider P, Hammarström L, Thrasher AJ, Gaspar HB, Grimbacher B Relevance of biallelic versus monoallelic TNFRSF13B mutations in distinguishing disease causing from disease modifying TNFRSF13B variants in common variable immunodeficiency. Blood 2009 113:1967-1976.

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Salzer U, Hagena T, Webster ADB and Grimbacher B. Sequence analysis of XIAP in male patients with common variable immunodeficiency. Int Arch Allergy Immunol 2008, 147:147-151.

Salzer U, Neumann C, Thiel J, Woellner C, Pan-Hammarstrom Q, Lougaris V, Hagena T, Jung J, Birmelin J, Du L, Metin A, Webster ADB, Plebani A, Moschese V, Hammarstrom L, Schaffer AA, Grimbacher B. Screening of functional and positional candidate genes in families with common variable immunodeficiency. BMC Immunology 2008, 9:3.

Losi CG, Salzer U, Gatta R, Lougaris V, Cattaneo G, Meini A, Soresina A, Grimbacher B, Plebani A. Mutational Analysis of Human BLYS in Patients with Common Variable Immunodeficiency. J Clin Immunol. 2006, 26:396-399.

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Warnatz K, Bossaller L\*, Salzer U\*, Skrabl-Baumgartner A, Schwinger W, van der Burg M, van Dongen JJ, Orlowska-Volk M, Knoth R, Durandy A, Draeger R, Schlesier M, Peter HH, Grimbacher B. Human ICOS-deficiency abrogates the germinal center reaction and provides a monogenic model for common variable immunodeficiency. Blood. 2006, 107:3045-3052. \*equal contribution

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