Accepted Manuscript

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PII: S0378-1119(16)30254-2

DOI: doi: 10.1016/j.gene.2016.04.006

Reference: GENE 41270

To appear in: Gene

Received date: 28 January 2016 Revised date: 11 February 2016 Accepted date: 4 April 2016

Please cite this article as: Kobbe, Robin, Kolster, Manuela, Fuchs, Sebastian, Schulze-Sturm, Ulf, Jenderny, Jutta, Kochhan, Lothar, Staab, Julia, Tolosa, Eva, Grimbacher, Bodo, Meyer, Thomas, Common variable immunodeficiency, impaired neurological development and reduced numbers of T regulatory cells in a 10-year-old boy with a STAT1 gain-of-function mutation, *Gene* (2016), doi: 10.1016/j.gene.2016.04.006

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ACCEPTED MANUSCRIPT

Common variable immunodeficiency, impaired neurological development and reduced numbers of T regulatory cells in a 10-year-old boy with a STAT1 gain-of-function mutation

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