

Publication List

h-Index: 26

A) Original papers (IF 2015, Times Cited ISI Web of Knowledge 12/2016)

Number of original papers: 62

|   | <u>Cited times</u> | <u>Impact factor</u> |
|---|--------------------|----------------------|
| 1. Wrona D, Siler U, <b>Reichenbach J</b> . CrispR/Cas9-generated p47phox-deficient cell line for Chronic Granulomatous Disease gene therapy vector development. <i>Scientific Reports</i> (Nature Publishing Group) 2017;7:44187   | 0                  | <b>5.228</b>         |
| 2. Vaas M, Enzmann G, Perinat T, Siler U, <b>Reichenbach J</b> , Licha K, Kipar A, Rudin M, Engelhardt B, Klohs J. Non-invasive near-infrared fluorescence imaging of the neutrophil response in a mouse model of transient cerebral ischaemia. <i>J Cereb Blood Flow Metab</i> , 2016 Oct 27.  | 0                  | <b>4.929</b>         |
| 3. Marquardt L, Lacour M, Hoernes M, Opitz L, Lecca R, Volkmer B, <b>Reichenbach J</b> , Hohl D, Ansari M, Ozsahin H, Güngör T, Pachlopnik Schmid J. Unusual dermatological presentation and immune phenotype in SCID due to an IL7R mutation: the value of whole-exome sequencing and the potential benefit of newborn screening. <i>J Eur Acad Dermatol Venereol</i> , 2016 Sep 5.  | 0                  | <b>3.029</b>         |
| 4. Siler U, Romao S, Tejera E, Pastukhov O, Kuzmenko E, Valencia RG, Spaccamela VM, Belohradsky BH, Speer O, Schmutz M, Kohne E, Hoenig M, Freihorst J, Schulz AS, <b>Reichenbach J</b> . Severe glucose-6-phosphate dehydrogenase deficiency leads to susceptibility to infection and absent NETosis. <i>J Allergy Clin Immunol</i> , 2016 Jul 22.   | 4                  | <b>12.485</b>        |
| 5. Elkaim E, Neven B, Bruneau J, Mitsui-Sekinaka K, Stanislas A, Heurtier L, Lucas CL, Matthews H, Deau MC, Sharapova S, Curtis J, <b>Reichenbach J</b> , Glastre C, Parry DA, Arumugakani G, McDermott E, Kilic SS, Yamashita M, Moshous D, Lamrini H, Otremba B, Gennery A, Coulter T, Quinti I, Stephan JL, Lougaris V, Brodzki N, Barlogis V, Asano T, Galicier L, Boutboul D, Nonoyama S, Cant A, Imai K, Picard C, Nejentsev S, Molina TJ, Lenardo M, Savic S, Cavazzana M, Fischer A, Durandy A, Kracker S. Clinical and immunologic phenotype associated with activated phosphoinositide 3-kinase $\delta$ syndrome 2: A cohort study. <i>J Allergy Clin Immunol</i> , 2016 Jul;138(1):210-218.e9   | 1                  | <b>12.485</b>        |
| 6. Toubiana J, Okada S, Hiller J, Oleastro M, Lagos Gomez M, Aldave Becerra JC, Ouachée-Chardin M, Fouyssac F, Girisha KM, Etzioni A, Van Montfrans J, Camcioglu Y, Kerns LA, Belohradsky B, Blanche S, Bousfiha A, Rodriguez-Gallego C, Meyts I, Kisand K, <b>Reichenbach J</b> , Renner ED, Rosenzweig S, Grimbacher B, van de Veerdonk FL, Traidl-Hoffmann C, Picard C, Marodi L, Morio T, Kobayashi M, Lilic D, Milner JD, Holland S, Casanova JL, Puel A; International STAT1 Gain-of-Function Study Group. Heterozygous STAT1 gain-of-function mutations underlie a broad clinical phenotype: an international survey of 234 patients from 140 kindreds. STAT1 gain of function mutations underlie an unexpectedly broad clinical phenotype: an | 2                  | <b>11.847</b>        |

- international survey of 274 patients. *Blood*, 2016 Jun 23;127(25):3154-64.
7. Kuehn HS\*, Boisson B\*, Cunningham-Rundles C, **Reichenbach J**, Stray-Pedersen A, Gelfand EW, Maffucci P, Pierce KR, Abbott JK, Voelkerding KV, South ST, Augustine NH, Bush JS, Dolen WK, Wray BB, Itan Y, Cobat A, Sorte HS, Ganesan S, Prader S, Martins TB, Lawrence MG, Orange JS, Calvo KR, Niemela JE, Casanova JL, Fleisher TA, Hill HR, Kumánovics A, Conley ME\*, Rosenzweig SD\*. Loss of B Cells in Patients with Heterozygous Mutations in IKAROS. *N Engl J Med* 2016 Mar 17;374(11):1032-43. \* *Equal contribution* 5 **59.558**
  8. Weisser M, Demel UM, Stein S, Chen-Wichmann L, Touzot F, Santilli G, Sujer S, Brendel C, Siler U, Cavazzana M, Thrasher AJ, **Reichenbach J**, Essers MA, Schwäble J, Grez M. Hyperinflammation in Chronic Granulomatous Disease leads to impairment of hematopoietic stem cell functions. *J Allergy Clin Immunol* 2016, 138(1):219-228.e9. 0 **12.485**
  9. Romao S, Tejera Puente E, Nytko KJ, Siler U, Münz C, **Reichenbach J**. Defective nuclear entry of hydrolases prevents NETosis in Chronic Granulomatous Disease. *J Allergy Clin Immunol* 2015; 136(6):1703-1706.e5 0 **12.485**
  10. Harbort CJ, Soeiro-Pereira PV, von Bernuth H, Kaindl AM, Costa-Carvalho BT, Condino-Neto A, **Reichenbach J**, Roesler J, Zychlinsky A, Amulic B. Neutrophil oxidative burst activates ATM to regulate cytokine production and apoptosis. *Blood* 2015; 126(26):2842-51 3 **11.847**
  11. Dreyer AK, Hoffmann D, Lachmann N, Ackermann M, Steinemann D, Timm B, Siler U, **Reichenbach J**, Grez M, Moritz T, Schambach A, Cathomen T. TALEN-mediated functional correction of X-linked chronic Granulomatous disease in patient-derived induced pluripotent stem cells. *Biomaterials* 2015; 69:191-200. 9 **8.387**
  12. Marschall K, Hoernes M, Bitzenhofer-Grüber M, Jandus P, Duppenhaler A, Wuillemin WA, Rischewski J, Boyman O, Heiningner U, Hauser T, Steiner U, Posfay-Barbe K, Seebach J, Recher M, Hess C, Helbling A, **Reichenbach J**. The Swiss National Registry for Primary Immunodeficiencies: Report on the first 6 years' activity 2008-2014. *Clin Exp Immunol* 2015;182: 45-50. 1 **3.148**
  13. Siler U\*, Paruzynski A\*, Holtgreve-Grez H, Kuzmenko E, Koehl U, Renner ED, Alhan C, van de Loosdrecht AA, Schwäble J, Pfluger T, Tchinda J, Schmugge M, Jauch A, Naundorf S, Kühlcke K, Notheis G, Güngör T, v. Kalle C, Schmidt M\* *Equal contribution*, Grez M\*, Seger R\*, **Reichenbach J**\*. Sequential gene therapy and successful rescue HSCT in two children with X-CGD – importance of timing. *Curr Gene Ther* 2015;15(4):416-27 \* *Equal contribution* 0 **2.738**
  14. Bielas H, Jud A, Lips U, **Reichenbach J**, Wieser I, Landolt MA. Preliminary Evidence for a Compromised T-Cell Compartment in Maltreated Children with Depression and Posttraumatic Stress Disorder. *Neuroimmunomodulation* 2015;22(5):303-10 0 **2.361**
  15. Romao S, Gasser N, Becker AC, Guhl B, Bajagic M, Vanoaica D, Ziegler U, Roesler J, Dengjel J, **Reichenbach J**, Munz C. Autophagy proteins stabilize pathogen-containing phagosomes for prolonged MHC II antigen processing. *J Cell Biol* 2013;203(5):757-66. 34 **8.717**

**Prof. Dr. med. Janine Reichenbach**

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| 16. | Brendel C, Hanseler W, Wohlgensinger V, Bianchi M, Tokmak S, Chen-Wichmann L, Kouzmenko E, Cesarovic N, Nicholls F, <b>Reichenbach J</b> , Seger R, Grez M, Siler U. Human miR223 Promoter as Novel Myelospesific Promoter for CGD Gene Therapy. <i>Human Gene Ther</i> 2013; 24(3):151-9.   | 7   | 4.062  |
| 17. | Bielas H, Jud A, Lips U, <b>Reichenbach J</b> , Landolt MA. Increased number of activated T cells in lymphocyte subsets of maltreated children: Data from a pilot study. <i>J Psychosom Res</i> 2012;73(4):313-8.  | 3   | 2.840  |
| 18. | Patel K, Akhter J, Kobrynski L, Gathman B, Davis O, Sullivan KE; International DiGeorge Syndrome Immunodeficiency Consortium ( <b>Reichenbach J</b> as ESID Member). Immunoglobulin deficiencies: the B-lymphocyte side of DiGeorge Syndrome. <i>J Pediatr</i> 2012;161(5):950-3.  | 24  | 4.122  |
| 19. | Cliffe ST, Bloch DB, Suryani S, Kamsteeg EJ, Avery DT, Palendira U, Church J, Wainstein B, Trizzino A, Lefranc G, Akatcharian C, Megarbané A, Gilissen C, Moshous D, <b>Reichenbach J</b> , Misbah S, Salzer U, Abinun M, Ong P, Stepensky P, Ruga E, Ziegler JB, Wong M, Tangye SG, Lindeman R, Buckley MF and Roscioli T. Clinical, Molecular and Cellular Immunology Findings in SP110-associated Venooclusive Disease with Immunodeficiency Syndrome. <i>J Allergy Clin Immunol</i> 2012;130(3):735-742.e6.  | 12  | 12.485 |
| 20. | Jiang Y, Cowley SA, Siler U, Melguzo D, Tilgner K, Browne K, deWilton A, Pryzborski S, Saretzki G, Seger RA, James WS, <b>Reichenbach J</b> , Lako M, Armstrong L. Derivation and functional analysis of patient specific induced pluripotent stem cells as an in vitro model of Chronic Granulomatous Disease. <i>Stem Cells</i> 2012; 30:599-611.  | 29  | 5.902  |
| 21. | Meyer-Bahlburg A, Renner ED, Rylaarsdam S, <b>Reichenbach J</b> , Schimke LF, Marks A, Tcheurekdjian H, Hostoffer R, Brahmandam A, Torgerson TR, Belohradsky BH, Rawlings DJ, Ochs HD. Heterozygous STAT3 mutations in hyper-IgE syndrome result in altered B cell maturation. <i>J Allergy Clin Immunol</i> 2012; 129(2), 559–562.e2  | 11  | 12.485 |
| 22. | Liu L, Okada S, Kong XF, Kreins AY, Cypowyj S, Abhyankar A, Toubiana J, Itan Y, Audry M, Nitschke P, Masson C, Toth B, Flatot J, Migaud M, Chrabieh M, Kochetkov T, Bolze A, Borghesi A, Toulon A, Hiller J, Eyerichs S, Eyerichs K, Gulácsy V, Chernyshova L, Chernyshov V, Bondarenko A, Cortés Grimaldo RM, Blancas Galicia L, Madrigal Beas IM, Roesler J, Magdorf K, Engelhardt D, Thumerelle C, Burgel PR, Hoernes M, Drexel B, Seger R, Kusuma T, Jansson AF, Sawalle-Belohradsky J, Belohradsky B, Jouanguy E, Bustamante J, Bué M, Bodemer C, Lortholary O, Fischer A, Blanche S, Al-Muhsen S, <b>Reichenbach J</b> , Kobayashi M, Torres Lozano C, Sebnem Kilic S, Oleastro M, Etzioni A, Traidl-Hoffmann C, Renner ED, Abel L, Picard C, Maródi L, Boisson-Dupuis S, Puel A and Casanova JL. Gain-of-function human STAT1 mutations impair IL-17 immunity and underlie chronic mucocutaneous candidiasis. <i>J Exp Med</i> 2011;208(8):1635-48. | 238 | 11.240 |
| 23. | Nakagawa N, Imai K, Kanegane H, Sato H, Yamada M, Kondo K, Okada S, Kobayashi M, Takada H, Mitsui N, Oshima K, Ohara O, Suri D, Rawat A, Singh S, Pan-Hammarström Q, Hammarström L, <b>Reichenbach J</b> , Seger R, Ariga T, Hara T, Miyawaki T, Nonoyama S. Newborn screening of B cell   | 27  | 12.485 |

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25. Bianchi M, Niemiec MJ, Siler U, Urban CF\*, **Reichenbach J\***. Restoration of anti-Aspergillus defense by NETs in human CGD is Calprotectin-dependent. *J Allergy Clin Immunol* 2011;127(5):1243-1252. \* *Equal contribution* 90 **12.485**
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**B) Other publications**

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