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Dienstanschrift:

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Qualifikationen

- Promotion zum Dr. med. an der LMU München, 13.05.1993
- Habilitation: Pädiatrie an der FAU-Erlangen-Nürnberg, 06/2001
- Schwerpunktbezeichnung: Pädiatrische Rheumatologie 02/2003
- Zertifizierung als Kinder- und Jugendrheumatologe (GKJR) 02/2005
- W2-Professur Allgemeine Pädiatrie, Universität Greifswald 01.09.2008
- Chefarzt des Deutschen Zentrums für Kinder- und Jugendrheumatologie, Garmisch-Partenkirchen, seit 01.04.2009
- Weiterbildungsbefugnis Kinderheilkunde, Bayerische Landesärztekammer (B-LÄK) 06/09
- Weiterbildungsbefugnis Kinderrheumatologie durch die B-LÄK 09/09
- Apl-Professur: Kinderrheumatologie, Ludwig-Maximilians Universität München 05/11

Werdegang

1985 - 1991 Medizinstudium an der Ludwig-Maximilians-Universität München
1992 - 1993 Kinderpoliklinik & Immungenetik, Ludwig-Maximilians-Universität, München
1993 - 2001 Klinik mit Poliklinik für Kinder und Jugendliche, Universität Erlangen-Nürnberg
2001 – 2008 Universität Greifswald, Leitender Oberarzt: Abteilungen Allgemeine Pädiatrie, Neonatologie und Pädiatrische Intensivmedizin
2008 Abteilungsleiter: Allgemeine Pädiatrie, Stellvertretender Abteilungsleiter: Neonatologie & Pädiatrische Intensivmedizin, Stellvertr. geschäftsführender Direktor der Universitätskinderklinik, W2-Professor für Allgemeine Pädiatrie an der Universität
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Ausgewählte Publikationen

1. Haas J. P., Andreas A., Rutkowski B., Brunner H., Keller E., Hoza J., Havelka S., Sierp G., Albert E. D.: A Model for the Role of HLA-DQ Molecules in the Pathogenesis of Juvenile Chronic Arthritis. *Rheumatology International*, 1991;11(4-5):191-7
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3. Haas J. P., Kimura A., Andreas A., Hochberger M., Keller E., Brännler G., Bettinotti MdIP., Nevinny-Stickel C., Sierp G., Sasazuki T., Albert E. D.: Polymorphism in the upstream regulatory region of DQA1 genes and DRB1, QAP, DQA1, DQB1 haplotypes in the german population. *Human Immunol.* 1994 Jan;39(1):31-40.
4. Haas J. P., Nevinny-Stickel C., Schoenwald U., Truckenbrodt H., Suschke J., Albert E. D.: Susceptible and Protective MHC-class II Haplotypes in Early Onset Pauciarticular Juvenile Chronic Arthritis. *Human Immunol.* 1994 Nov;41(3):225-33.
5. Paul C., Haas J. P., Schoenwald U., Truckenbrodt H., Bettinotti MdIP., Bönisch J., Brännler G., Keller E., Nevinny-Stickel C., Yao Z., Albert E. D.: HLA class I/ class II interaction in early onset pauciarticular juvenile chronic arthritis. *Immunogenetics.* 1994;39(1):61-4.
6. Haas J. P., Truckenbrodt H., Paul C., Hoza J., Scholz S., Albert E. D.: Subtypes of HLA-DRB1*03, *08, *11, *12, *13 and *14 in early onset pauciarticular juvenile chronic arthritis (OA-JIA) with and without iridocyclitis. *Clin Exp Rheumatol* 1994 Sep-Oct;12Suppl 10:S7-14.
7. Haas J. P., Kimura A., Truckenbrodt H., Suschke J., Sasazuki T., Volgger A., Albert E. D.: Early Onset Pauciarticular Juvenile Chronic Arthritis is associated with a Mutation in the Y-Box of the HLA-DQA1 Promoter. *Tissue Antigens* 1995 May;45(5):317-21.
8. Leipold G., Schütz E., Haas J. P., Oellerich M.: Azathioprine-induced severe pancytopenia due to a homocygous two-point mutation of the thiopurine methyltransferase gene in a patient with juvenile HLA-B27 associated spondylarthritis. *Arthr. Rheum.* 1997 Oct;40(10):1896-8.
9. Haas J. P., Gruenke M., Frank C., Kolowos W., Dirnecker D., Leipold G., Hieronymus T., Lorenz H. M., Herrmann M.: Increased spontaneous apoptosis in double negative T cells of humans with Fas/Apo1 mutation. *Cell Death Diff.* 1998 Sep;5(9):751-7.
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15. Meyer-Bahlburg A., Haas J. P., Haase R., Eschrich U., Wawer A., Frank L., Marsch W. C., Burdach S., Horneff G.: Treatment with cyclosporin A in a patient with Omenn's syndrome. *Arch Dis Child.* 2002 Sep;87(3):231-3.
16. Fernandez S., Wassmuth R., Knerr I., Frank C., Haas J. P.: Relative quantification of HLA-DRA1 and DQA1-expression by real time RT-PCR. *Eur. J. Immunogenet.* 2003 Apr;30(2):141-8.
17. Ruf R. G., Lichtenberger A., Karle S. M., Haas J. P., Anacleto F. E., Schultheiss M., Zalewski I., Imm A., Ruf E. M., Mucha, Bagga A., Fuchshuber A., Bakkaloglu A., Hildebrandt F. and Members of the APN Study Group: Patients with Mutations in NPHS2 (Podocin) do Not Respond to standard steroid treatment of nephrotic syndrome. *J Am Soc Nephrol.* 2004 Mar;15(3):722-32.
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53. Scheuern A, Tyrrell PN, Haas JP, Hügler B. Countermeasures against methotrexate intolerance in juvenile idiopathic arthritis instituted by parents show no effect. *Rheumatology (Oxford)*. 2017 Jan 24. pii: kew507. doi: 10.1093/rheumatology/kew507. [Epub ahead of print] PubMed PMID: 28122960.
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