

Curriculum Vitae Professor Marcus A. Mall, M.D., FERS

Professor and Director, Department of Pediatric Pulmonology,
Immunology and Critical Care Medicine

Medical Director, CharitéCenter 17 for Obstetrics and Gynecology,
and Child and Adolescent Medicine with Perinatal Center and
Human Genetics

Charité - Universitätsmedizin Berlin



Scientific Vita

- Since 2020 Medical Director, CharitéCenter 17 for Obstetrics and Gynecology, and Child and Adolescent Medicine with Perinatal Center and Human Genetics
- Since 2018 W3-Professor of Pediatric Pulmonology and Immunology and Director of the Department of Pediatric Pulmonology, Immunology and Critical Care Medicine & Cystic Fibrosis Center, Charité - Universitätsmedizin Berlin
- Since 2018 BIH Professor, Berlin Institute of Health (BIH)
- 2012 - 2018 Director of the Translational Lung Research Center (TLRC) and Department of Translational Pulmonology, University of Heidelberg
- 2009 - 2018 W3-Professor and Head of the Division of Pediatric Pulmonology & Allergy and Cystic Fibrosis Center, Department of Pediatrics, University of Heidelberg
- 2009 - 2012 W3-Heisenberg Professor of Translational Pediatric Pulmonology
- 2007 - 2018 Attending Physician and Head of the Cystic Fibrosis Center, Department of Pediatrics, University of Heidelberg
- 2007 Habilitation and venia legendi, University of Heidelberg, Germany
- 2006 - 2018 Group leader at the Molecular Medicine Partnership Unit (MMPU) of the University of Heidelberg and the European Molecular Biology Laboratory (EMBL)
- 2006 - 2018 Faculty Member of the Hartmut Hoffmann-Berling International Graduate School of Molecular & Cellular Biology Heidelberg (HBIGS)
- 2005-2009 Group leader of EU-funded Independent Junior Research Group "Cystic Fibrosis / Chronic Airway Disease" (Marie Curie Excellence Team)
- 2004 - 2006 Fellow in Pediatric Pulmonology, Department of Pediatrics, University of Heidelberg
- 2003 - 2004 Assistant Professor, CF/Pulmonary Research and Treatment Center, School of Medicine, University of North Carolina at Chapel Hill, Chapel Hill, NC, USA
- 2000 - 2003 Postdoc and Research Associate, CF/Pulmonary Research and Treatment Center, School of Medicine, University of North Carolina at Chapel Hill, Chapel Hill, NC, USA
- 1997 - 2000 Resident, University Children's Hospital, University of Freiburg

- 1997 Dissertation (M.D.), University Freiburg
 1990 – 1997 Medical School, University of Freiburg, Germany and University College
 London School of Medicine, UK

Clinical Qualifications

- 2017 Licensed for Allergology
 2011 Licensed for Infectious Diseases
 2008 Licensed for Pediatric Pulmonology
 2006 Licensed for Pediatrics and Adolescent Medicine

Other Professional Activities

- 2018-2020 Vice President of the European Cystic Fibrosis Society (ECFS)
 2017 - 2018 Chair and Co-Chair of the ECFS Basic Science Conference
 Since 2017 Member of the Program Planning Committee of the North American Cystic
 Fibrosis Conference (NACFC)
 2016 Conference President of the 19th German Cystic Fibrosis Conference,
 Würzburg, Germany
 Since 2015 Member of the Scientific Advisory Board for the SickKids-Cystic Fibrosis
 Canada Program for Individualized CF Therapy (CFIT), Toronto, Canada
 2014-2020 Member of the Board of the European Cystic Fibrosis Society (ECFS)
 2011 Conference Vice-President, 34th ECFS Conference, Hamburg, Germany
 2009 - 2012 Member of the ECFS Conference Scientific Steering Committee
 2006 - 2012 Elected Member (2006-2009) and Chairman (2010-2012) of the Research
 Council of the Germany Cystic Fibrosis Association

Project Coordination and Membership in Collaborative Research Projects

- Since 2021 Coordinator of the Berlin site of the German Center for Child and Youth
 Health (DZKJ)
 Since 2021 Deputy Speaker of the SFB 1449 „Dynamic Hydrogels at Biointerfaces“
 Since 2019 Coordinator and Speaker of the Innovationsfonds project “conneCT CF:
 Coaching and Telemonitoring for Patients with Cystic Fibrosis”
 2018 - 2022 Principal Investigator of the SFB-TR 84 “Innate Immunity of the Lung:
 Mechanisms of Pathogen Attack and Host Defence in Pneumonia“
 2013-2016 Principal Investigator of the EU collaborative project CFMATTERS -
 Cystic Fibrosis Microbiome-determined Antimicrobial Therapy Trial
 in Exacerbations: Results Stratified (7. Framework Programme)
 Since 2012 Director, Disease Area Cystic Fibrosis, German Center for Lung Research
 (DZL)
 2011 – 2018 Founding Director and Member of the Executive Board, German Centre for
 Lung Research (DZL)

2009 – 2015 Principal Investigator of the BMBF collaborative project CARPuD:
Network Cellular Approaches for Rare Pulmonary Diseases

Honors & Awards

2020 Adalbert-Czerny-Prize, German Society of Pediatrics and Adolescent Medicine (DGKJ)
2020 ERS Excellence Award for Research in Cystic Fibrosis
2018 Einstein-Professorship, Einstein Foundation Berlin
2017 Fellow of ERS (FERS)
2009 Heisenberg-Professorship for Translational Pediatric Pulmonology, German Research Foundation (DFG)
2009 Research Award, German Society of Pulmonology (DGP)
2005 Marie Curie Excellence Grant (EU 6. Framework Programme)
2005 Johannes Wenner Prize, German Society of Pediatric Pulmonology (GPP)
2005 Professor David Shmeling Research Award, German Society of Pediatric Gastroenterology and Nutrition (GPGE)
2000 -2002 Research Fellowship, German Research Foundation (DFG)
1998 Albrecht Fleckenstein Young Investigator Award, University of Freiburg
1997 Adolf Windorfer Prize for Cystic Fibrosis Research, German CF Association
1994 - 1997 Scholarship of the German Student Scholarship Foundation (Studienstiftung des deutschen Volkes)
1993 - 1994 Scholarship of the German Academic Exchange Service (DAAD)

Memberships in Professional Societies

Since 2019 Member of the Fleischner Society
Since 2010 European Respiratory Society (ERS)
Since 2010 German Society of Pulmonology (DGP)
Since 2006 American Thoracic Society (ATS)
Since 2004 German Cystic Fibrosis Association (Mukoviszidose e.V.)
Since 1999 European Cystic Fibrosis Society (ECFS)
Since 1998 German Physiological Society (DPG)
Since 1998 German Society of Pediatrics and Adolescent Medicine (DGKJ)
Since 1998 German Society of Pediatric Pulmonology (GPP)

Publications

Original Articles

Rohrich M, Leitz D, Glatting FM, Wefers AK, Weinheimer O, Flechsig P, Kahn N, **Mall MA**, Giesel FL, Kratochwil C, Huber PE, von Deimling A, Heussel CP, Kauczor HU, Kreuter M, Haberkorn UA. Fibroblast Activation Protein specific PET/CT imaging in fibrotic interstitial lung diseases and lung cancer: a translational exploratory study. *J Nucl Med* 2022; 63:127-133.

Georg P, Astaburuaga-Garcia R, Bonaguro L, Brumhard S, Michalick L, Lippert LJ, Kostevc T, Gabel C, Schneider M, Streitz M, Demichev V, Gemund I, Barone M, Tober-Lau P, Helbig ET, Hillus D, Petrov L, Stein J, Dey HP, Paclik D, Iwert C, Mulleder M, Aulakh SK, Djudjaj S, Bulow RD, Mei HE, Schulz AR, Thiel A, Hippenstiel S, Saliba AE, Eils R, Lehmann I, **Mall MA**, Stricker S, Rohmel J, Corman VM, Beule D, Wyler E, Landthaler M, Obermayer B, von Stillfried S, Boor P, Demir M, Wesselmann H, Suttorp N, Uhrig A, Muller-Redetzky H, Nattermann J, Kuebler WM, Meisel C, Ralser M, Schultze JL, Aschenbrenner AC, Thibeault C, Kurth F, Sander LE, Bluthgen N, Sawitzki B, Group P-C-S. Complement activation induces excessive T cell cytotoxicity in severe COVID-19. *Cell* 2021. [Epub ahead of print].

Ruf S, Hommes F, van Loon W, Seybold J, Kurth T, **Mall MA**, Mockenhaupt FP, Theuring S. A Retrospective Outbreak Investigation of a COVID-19 Case Cluster in a Berlin Kindergarten, November 2020. *Int J Environ Res Public Health* 2021;19:36.

Graeber SY, Vitzthum C, Pallenberg ST, Naehrlich L, Stahl M, Rohrbach A, Drescher M, Minso R, Ringshausen FC, Rueckes-Nilges C, Klajda J, Berges J, Yu Y, Scheuermann H, Hirtz S, Sommerburg O, Dittrich AM, Tummeler B, **Mall MA**. Effects of Elexacaftor/Tezacaftor/Ivacaftor Therapy on CFTR Function in Patients with Cystic Fibrosis and One or Two F508del Alleles. *Am J Respir Crit Care Med* 2021. [Epub ahead of print].

Metzger MI, Graeber SY, Stahl M, Sommerburg O, **Mall MA**, Dalpke AH, Boutin S. A Volatile and Dynamic Longitudinal Microbiome Is Associated With Less Reduction in Lung Function in Adolescents With Cystic Fibrosis. *Front Cell Infect Microbiol* 2021;11:763121.

Sutharsan S, McKone EF, Downey DG, Duckers J, MacGregor G, Tullis E, Van Braeckel E, Wainwright CE, Watson D, Ahluwalia N, Bruinsma BG, Harris C, Lam AP, Lou Y, Moskowitz SM, Tian S, Yuan J, Waltz D, **Mall MA**, group VXs. Efficacy and safety of elexacaftor plus tezacaftor plus ivacaftor versus tezacaftor plus ivacaftor in people with cystic fibrosis homozygous for F508del-CFTR: a 24-week, multicentre, randomised, double-blind, active-controlled, phase 3b trial. *Lancet Respir Med* 2021. [Epub ahead of print].

Barry PJ, **Mall MA**, Polineni D, Group VXS. Triple Therapy for Cystic Fibrosis Phe508del-Gating and -Residual Function Genotypes. Reply. *N Engl J Med* 2021;385:2208.

Hey J, Paulsen M, Toth R, Weichenhan D, Butz S, Schatterny J, Liebers R, Lutsik P, Plass C*, **Mall MA***. Epigenetic reprogramming of airway macrophages promotes polarization and inflammation in muco-obstructive lung disease. *Nat Commun* 2021;12:6520. (*equal contribution)

van Loon W, Hommes F, Theuring S, von der Haar A, Korner J, Schmidt M, von Kalle C, **Mall MA**, Seybold J, Kurth T, Mockenhaupt FP. Renewed absence of SARS-CoV-2 infections in the day care context in Berlin, January 2021. *Clin Infect Dis* 2021;73:1944-1945.

Sommerburg O, Stahl M, Hammerling S, Gramer G, Muckenthaler MU, Okun J, Kohlmüller D, Happich M, Kulozik AE, **Mall MA***, Hoffmann GF*. Final results of the southwest German pilot study on cystic fibrosis newborn screening - Evaluation of an IRT/PAP protocol with IRT-dependent safety net: Results of the Southwest German CFNBS pilot study. *J Cyst Fibros* 2021. [Epub ahead of print]. (*equal contribution)

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Witkowski M, Tizian C, Ferreira-Gomes M, Niemeyer D, Jones TC, Heinrich F, Frischbutter S, Angermair S, Hohnstein T, Mattioli I, Nawrath P, Mc Ewen S, Zocche S, Viviano E, Heinz GA, Maurer M, Kolsch U, Chua RL, Aschman T, Meisel C, Radke J, Sawitzki B, Roehmel J, Allers K, Moos V, Schneider T, Hanitsch L, **Mall MA**, Conrad C, Radbruch H, Duerr CU, Trapani JA, Marcenaro E, Kallinich T, Corman VM, Kurth F, Sander LE, Drosten C, Treskatsch S, Durek P, Kruglov A, Radbruch A, Mashreghi MF, Diefenbach A. Untimely TGFβ responses in COVID-19 limit antiviral functions of NK cells. *Nature* 2021;600:295-301.

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Chung J, Wunnemann F, Salomon J, Boutin S, Frey DL, Albrecht T, Joachim C, Eichinger M, **Mall MA**, Wielputz MO, Sommerburg O. Increased Inflammatory Markers Detected in Nasal Lavage Correlate with Paranasal Sinus Abnormalities at MRI in Adolescent Patients with Cystic Fibrosis. *Antioxidants (Basel)* 2021;10.

van Loon W, Theuring S, Hommes F, **Mall MA**, Seybold J, Kurth T, Mockenhaupt FP. Prevalence of SARS-CoV-2 Infections Among Students, Teachers, and Household Members During Lockdown and Split Classes in Berlin, Germany. *JAMA Netw Open* 2021;4:e2127168.

Gertler M, Krause E, van Loon W, Krug N, Kausch F, Rohardt C, Rossig H, Michel J, Nitsche A, **Mall MA**, Nikolai O, Hommes F, Burock S, Lindner AK, Mockenhaupt FP, Pison U, Seybold J. Self-collected oral, nasal and saliva samples yield sensitivity comparable to professionally collected oro-nasopharyngeal swabs in SARS-CoV-2 diagnosis among symptomatic outpatients. *Int J Infect Dis* 2021;110:261-266.

Frey DL, Boutin S, Dittrich SA, Graeber SY, Stahl M, Wege S, Herth FJF, Sommerburg O, Schultz C, **Mall MA***, Dalpke AH*. Relationship between airway dysbiosis, inflammation and lung function in adults with cystic fibrosis. *J Cyst Fibros* 2021;20:754-760. (*equal contribution)

Schroder A, Lunding LP, Zissler UM, Vock C, Webering S, Ehlers JC, Orinska Z, Chaker A, Schmidt-Weber CB, Lang NJ, Schiller HB, **Mall MA**, Fehrenbach H, Dinarello CA, Wegmann M. IL-37 regulates allergic inflammation by counterbalancing pro-inflammatory IL-1 and IL-33. *Allergy* 2021. [Epub ahead of print].

Loske J, Rohmel J, Lukassen S, Stricker S, Magalhaes VG, Liebig J, Chua RL, Thurmann L, Messingschlager M, Seegebarth A, Timmermann B, Klages S, Ralser M, Sawitzki B, Sander LE, Corman VM, Conrad C, Laudi S, Binder M*, Trump S*, Eils R*, **Mall MA***, Lehmann I*. Pre-activated antiviral innate immunity in the upper airways controls early SARS-CoV-2 infection in children. *Nat Biotechnol* 2021. [Epub ahead of print]. (*equal contribution)

Goss CH, Fajac I, Jain R, Seibold W, Gupta A, Hsu MC, Sutharsan S, Davies JC, **Mall MA**. Efficacy and safety of inhaled ENaC inhibitor BI 1265162 in patients with cystic fibrosis: BALANCE-CF 1 - a randomised, Phase II study. *Eur Respir J* 2021. [Epub ahead of print].

Theuring S, Thielecke M, van Loon W, Hommes F, Hulso C, von der Haar A, Korner J, Schmidt M, Bohringer F, **Mall MA**, Rosen A, von Kalle C, Kirchberger V, Kurth T, Seybold J, Mockenhaupt FP, Group BS. SARS-CoV-2 infection and transmission in school settings during the second COVID-19 wave: a cross-sectional study, Berlin, Germany, November 2020. *Euro Surveill* 2021;26:2100184.

Terliesner N, Rosen A, Kaindl AM, Reuter U, Lippold K, **Mall MA**, von Bernuth H, Gratopp A. Maintenance of Elective Patient Care at Berlin University Children's Hospital During the COVID-19 Pandemic. *Front Pediatr* 2021;9:694963.

Barry PJ*, **Mall MA***, Alvarez A, Colombo C, de Winter-de Groot KM, Fajac I, McBennett KA, McKone EF, Ramsey BW, Sutharsan S, Taylor-Cousar JL, Tullis E, Ahluwalia N, Jun LS, Moskowitz SM, Prieto-Centurion V, Tian S, Waltz D, Xuan F, Zhang Y, Rowe SM, Polineni D, Group VXS. Triple Therapy for Cystic Fibrosis Phe508del-Gating and -Residual Function Genotypes. *N Engl J Med* 2021;385:815-825. (*equal contribution)

Meisel C, Akbil B, Meyer T, Lankes E, Corman VM, Staudacher O, Unterwalder N, Kolsch U, Drosten C, **Mall MA**, Kallinich T, Schnabel D, Goffinet C, von Bernuth H. Mild COVID-19 despite autoantibodies to type I IFNs in Autoimmune-Polyendocrine-Syndrome Type 1 (APS-1). *J Clin Invest* 2021;131:e150867.

Hagner M, Albrecht M, Guerra M, Braubach P, Halle O, Zhou-Suckow Z, Butz S, Jonigk D, Hansen G, Schultz C, Dittrich AM*, **Mall MA***. IL-17A from innate and adaptive lymphocytes contributes to inflammation and damage in cystic fibrosis lung disease. *Eur Respir J* 2021;57. (*equal contribution)

Engelmann TA, Knudsen L, Leitz DHW, Duerr J, Beers MF, **Mall MA**, Ochs M. Linking Fibrotic Remodeling and Ultrastructural Alterations of Alveolar Epithelial Cells after Deletion of *Nedd4-2*. *Int J Mol Sci* 2021;22:7607.

Zemanick ET, Taylor-Cousar JL, Davies J, Gibson RL, **Mall MA**, McKone EF, McNally P, Ramsey BW, Rayment JH, Rowe SM, Tullis E, Ahluwalia N, Chu C, Ho T, Moskowitz SM, Noel S, Tian S, Waltz D, Weinstock TG, Xuan F, Wainwright CE, McColley SA. A Phase 3 Open-Label Study of Elexacaftor/Tezacaftor/Ivacaftor in Children 6 through 11 Years of Age with Cystic Fibrosis and at Least One F508del Allele. *Am J Respir Crit Care Med* 2021;203:1522-1532.

Leitz DHW, Duerr J, Mulugeta S, Agircan AS, Zimmermann S, Kawabe H, Dalpke AH, Beers MF, **Mall MA**. Congenital Deletion of *Nedd4-2* in Lung Epithelial Cells Causes Progressive Alveolitis and Pulmonary Fibrosis in Neonatal Mice. *International Journal of Molecular Sciences* 2021;22:6146.

Graeber SY, Boutin S, Wielputz MO, Joachim C, Frey DL, Wege S, Sommerburg O, Kauczor HU, Stahl M, Dalpke AH, **Mall MA**. Effects of Lumacaftor-Ivacaftor on Lung Clearance Index, Magnetic Resonance Imaging, and Airway Microbiome in Phe508del Homozygous Patients with Cystic Fibrosis. *Ann Am Thorac Soc* 2021;18:971-980.

Frey DL, Guerra M, **Mall MA**, Schultz C. Monitoring Neutrophil Elastase and Cathepsin G Activity in Human Sputum Samples. *J Vis Exp* 2021.

Thee S, Stahl M, Fischer R, Sutharsan S, Ballmann M, Muller A, Lorenz D, Urbanski-Rini D, Puschner F, Amelung VE, Fuchs C, **Mall MA**. A multi-centre, randomized, controlled trial on coaching and telemonitoring in patients with cystic fibrosis: connect CF. *BMC Pulm Med* 2021;21:131.

Maschmeyer P, Heinz GA, Skopnik CM, Lutter L, Mazzoni A, Heinrich F, von Stuckrad SL, Wirth LE, Tran CL, Riedel R, Lehmann K, Sakwa I, Cimaz R, Giudici F, **Mall MA**, Enghard P, Vastert B, Chang HD, Durek P, Annunziato F, van Wijk F, Radbruch A, Kallinich T, Mashreghi MF. Antigen-driven PD-1(+) TOX(+) BHLHE40(+) and PD-1(+) TOX(+) EOMES(+) T lymphocytes regulate juvenile idiopathic arthritis in situ. *Eur J Immunol* 2021;51:915-929.

Sommerburg O, Hammerling S, Schneider SP, Okun J, Langhans CD, Leutz-Schmidt P, Wielputz MO, Siems W, Graber SY, **Mall MA***, Stahl M*. CFTR Modulator Therapy with Lumacaftor/Ivacaftor Alters Plasma Concentrations of Lipid-Soluble Vitamins A and E in Patients with Cystic Fibrosis. *Antioxidants (Basel)* 2021;10:483. (*equal contribution)

Hommel F, van Loon W, Thielecke M, Abramovich I, Lieber S, Hammerich R, Gehrke-Beck S, Linzbach E, Schuster A, von dem Busche K, Theuring S, Gertler M, Martinez GE, Richter J, Bergmann C, Bolke A, Bohringer F, **Mall MA**, Rosen A, Krannich A, Keller J, Bethke N, Kurzmann M, Kurth T, Kirchberger V, Seybold J, Mockenhaupt FP, Study Group B. SARS-CoV-2 Infection, Risk Perception, Behaviour and Preventive Measures at Schools in Berlin, Germany, during the Early Post-Lockdown Phase: A Cross-Sectional Study. *Int J Environ Res Public Health* 2021;18:2739.

Ferreira-Gomes M, Kruglov A, Durek P, Heinrich F, Tizian C, Heinz GA, Pascual-Reguant A, Du W, Mothes R, Fan C, Frischbutter S, Habenicht K, Budzinski L, Ninnemann J, Jani PK, Guerra GM, Lehmann K, Matz M, Ostendorf L, Heiberger L, Chang HD, Bauherr S, Maurer M, Schonrich G, Raftery M, Kallinich T, **Mall MA**, Angermair S, Treskatsch S, Dorner T, Corman VM, Diefenbach A, Volk HD, Elezkurtaj S, Winkler TH, Dong J, Hauser AE, Radbruch H, Witkowski M, Melchers F, Radbruch A, Mashreghi MF. SARS-CoV-2 in severe COVID-19 induces a TGF-beta-dominated chronic immune response that does not target itself. *Nat Commun* 2021;12:1961.

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for the F508del mutation. *Am J Respir Crit Care Med* 2020;202:1589-1592. (*equal contribution)

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Vega G, Guequén A, Philp AR, Gianotti A, Arzola L, Villalón M, Zegarra-Moran O, Galiotta LJ, **Mall MA**, Flores CA. Lack of Kcnn4 improves mucociliary clearance in muco-obstructive lung disease. *JCI Insight* 2020;5:e140076.

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