



## Curriculum Vitae

Personal information **Gitte Rønde**

Work experience

01.05.19 - Department Physician, Department of Children and Adolescents, Herlev Hospital  
01.03.18 - 30.04.19: Department Physician, Children and Adolescents Clinic, Rigshospitalet (expert training)  
01.09.17 - 28.02.18: Department Physician, Epilepsy Hospital, Filadelfia, Dianalund (expert training)  
01.03.17 - 31.08.17: Department Physician, Neurological Department, Roskilde Hospital (expert training)  
01.07.16 - 28.02.17: Department Physician, Pediatric Department, Roskilde Hospital  
01.07.14 - 30.06.16: Main training, Pediatric Department, Roskilde Hospital  
20.02.13 - 30.06.14: Main training, Pediatric Department, Hvidovre Hospital  
10.03.12 - 19.02.13: Maternity leave  
01.09.11 - 09.03.12: Main training, Pediatric Department, Hvidovre Hospital  
01.03.11 - 31.08.11: Side training, Genetic Counseling, Kennedy Center  
  
31.03.07 - 28.02.11: PhD. fellow, Center for Rett Syndrome, and substitute for chief physician at the Center for PKU, Kennedy Center (KC)  
01.02.07 - 28.02.07: Side training, Department of Anesthesiology, Hvidovre Hospital  
01.11.06 - 31.01.07: Unclassified position, Pediatric Department, Roskilde Hospital  
01.11.05 - 31.10.06: Introductory position, Pediatric Department, Roskilde Hospital  
01.03.05 - 31.10.05: Unclassified position, Pediatric Department, Roskilde Hospital  
01.09.04 - 28.02.05: Rotation training, Doctors Hanne and Peter Silbye, Ølby  
01.10.03 - 31.08.04: Maternity Leave  
01.09.03 - 30.09.03: Unclassified position, Surgical Dept., Køge Hospital  
01.03.03 - 31.08.03: Internship, Surgical Dept., Køge Hospital  
01.09.02 - 28.02.03: Internship, Medical Dept., Køge Hospital  
01.08.02 - 31.08.02: Unclassified position, Medical Dept., Køge Hospital  
01.03.01 - 30.04.01: Pre-graduate, temporary, Neurological Dept., Bispebjerg Hospital

Education and training

01.06.2022 Danish diploma training in cervical dystonia and spasticity treatment  
29.02.2020 Expert in Neuropediatrics  
31.06.2016 Pediatric specialist training  
14.09.2011 PhD training at the University of Copenhagen (KU)  
31.10.2006 Introductory training, Department of Pediatrics, Roskilde Hospital  
19.06.2002 Master of Medicine, KU  
June 1993 Student exam, Støvring Gymnasium

Additional information

Publications

Book Chapter

2015, Bone Markers in Rett Syndrome, Gitte Roende and Jens-Erik Beck Jensen, Biomarkers in Disease: Methods, Discoveries and Applications. Biomarkers in Bone Disease, edited by Vinood B. Patel and Victor R. Preedy, Meteor Springer.com. doi:10.1007/978-94-007-7745-3\_21-1

Pediatric varicella zoster virus associated arterial ischemic stroke - A post-pandemic rise in incidence in Denmark. Esserlind AL, Schmidt LS, Rønde G, Miranda M, Jensen L, Bindslev JB, Born AP, Vissing NH, Møller FT, Møllerup JE, Børresen ML. Eur J Paediatr Neurol. 2025 Nov;59:100-106

Bi-allelic pathogenic variants in TRMT1 disrupt tRNA modification and induce a neurodevelopmental disorder. Efthymiou S, Leo CP, Deng C, Lin SJ, Maroofian R, Lin R, Karagoz I, Zhang K, Kaiyrzhanov R, Scardamaglia A, Owrang D, Turchetti V, Jahnke F, Huang K, Petree C, Derrick AV, Rees MI, Alvi JR, Sultan T, Li C, Jacquemont ML, Tran-Mau-Them F, Valenzuela-Palafoll M, Sidlow R, Yoon G, Morrow MM, Carere DA, O'Connor M, Fleischer J, Gerkes EH, Phornphutkul C, Isidor B, Rivier-Ringenbach C, Philippe C, Kurul SH, Soydemir D, Kara B, Sunnetci-Akkoyunlu D, Bothe V, Platzter K, Wiczorek D, Koch-Hogrebe M, Rahner N, Thuresson AC, Matsson H, Frykholm C, Bozdoğan ST, Bisgin A, Chatron N, Lesca G, Cabet S, Tümer Z, Hjortshøj TD, Rønde G, Marquardt T, Reunert J, Afzal E, Zamani M, Azizimalamiri R, Galehdari H, Nourbakhsh P, Chamanrou N, Chung SK, Suri M, Benke PJ, Zaki MS, Gleeson JG, Calame DG, Pehlivan D, Yilmaz HI, Gezdirici A, Rad A, Abumansour IS, Oprea G, Bereketoglu MB, Banneau G, Julia S, Zeighami J, Ashoori S, Shariati G, Sedaghat A, Sabri A, Hamid M, Parvas S, Tajudin TA, Abdullah U, Baig SM, Chung WK, Glazunova OO, Sabine S, Cheema HA, Zifarelli G, Bauer P, Sidpra J, Mankad K, Vona B, Fry AE, Varshney GK, Houlden H, Fu D. Am J Hum Genet. 2025 May 1;112(5):1117-1138. doi: 10.1016/j.ajhg.2025.03.015. Epub 2025 Apr 16

A Novel UPF1 Variant Associated With a Rare UPF1-Related Neurodevelopmental Disorder. Tümer Z, Dalsberg J, Rønde G, Sørensen JK, Østergaard E. Clin Genet. 2025 Feb 24;108(3):313-7

Seizure and movement disorder in CACNA1E developmental and epileptic encephalopathy: Two sides of the same coin or same side of two different coins? Di Micco V, Affronte L, Khinchi MS, Rønde G, Miranda MJ, Hammer TB, Specchio N, Beniczky S, Olofsson K, Møller RS, Gardella E. Epileptic Disord. 2024 Aug;26(4):520-526

Clinical features and genotype-phenotype correlations in epilepsy patients with de novo DYNC1H1 Variants. Cuccurullo C, Cerulli Irelli E, Ugga L, Riva A, D'Amico A, Cabet S, Lesca G, Bilo L, Zara F, Ilescu C, Barca D, Fung F, Helbig K, Ortiz-Gonzalez X, Schelhaas HJ, Willemsen MH, van der Linden I, Canafoglia L, Courage C, Gommarsaschi S, Gonzalez-Alegre P, Bardakjian T, Syrbe S, Schuler E, Lemke JR, Vari S, Roende G, Bak M, Huq M, Powis Z,

Johannesen KM, Hammer TB, Møller RS, Rabin R, Pappas J, Zupanc ML, Zadeh N, Cohen J, Naidu S, Krey I, Saneto R, Thies J, Licchetta L, Tinuper P, Bisulli F, Minardi R, Bayat A, Villeneuve N, Molinari F, Salimi Dafsari H, Møller B, Le Roux M, Houdayer C, Vecchi M, Mammi I, Fiorini E, Proietti J, Ferri S, Cantalupo G, Battaglia DI, Gambardella ML, Contaldo I, Brogna C, Trivisano M, De Dominicis A, Bova SM, Gardella E, Striano P, Coppola A. *Epilepsia*. 2024 Jul 2. doi: 10.1111/epi.18054. Online ahead of print.

Cerebral Palsy - Early Diagnosis and Intervention Trial: protocol for the prospective multicentre CP-EDIT study with focus on diagnosis, prognostic factors, and intervention. Hoei-Hansen CE, Weber L, Johansen M, Fabricius R, Hansen JK, Viuff AF, Rønde G, Hahn GH, Østergaard E, Duno M, Larsen VA, Madsen CG, Røhder K, Elvrum AG, Laugesen B, Ganz M, Madsen KS, Willerslev-Olsen M, Debes NM, Christensen J, Christensen R, Rackauskaite G. *BMC Pediatr*. 2023 Oct 30;23(1):544

Acute flaccid rhombencephalomyelitis with radiculitis in a child with an enterovirus A71 infection seen for the first time in Denmark: a case report. Foli-Andersen PJ, Munkholm A, Rønde G, Børresen ML, Nielsen JEK, Midgley S, Bang D. *J Med Case Rep*. 2022 Jan 25;16(1):32

Decline in gross motor skills in adult Rett syndrome; results from a Danish longitudinal study. Bisgaard AM, Wong K, Højfeldt AK, Larsen JL, Schönewolf-Greulich B, Rønde G, Downs J, Stahlhut M. *Am J Med Genet A*. 2021 Dec;185(12):3683-3693

ZMYND11 variants are a novel cause of centrotemporal and generalised epilepsies with neurodevelopmental disorder. Oates S, Absoud M, Goyal S, Bayley S, Baulcomb J, Sims A, Riddett A, Allis K, Brasch-Andersen C, Balasubramanian M, Bai R, Callewaert B, Hüffmeier U, Le Duc D, Radtke M, Korff C, Kennedy J, Low K, Møller RS, Nielsen JEK, Popp B, Quteineh L, Rønde G, Schönewolf-Greulich B, Shillington A, Taylor MR, Todd E, Torring PM, Tümer Z, Vasileiou G, Yates TM, Zweier C, Rosch R, Basson MA, Pal DK. *Clin Genet*. 2021 Oct;100(4):412-429

Nonpharmacological Interventions Addressing Pain, Sleep, and Quality of Life in Children and Adolescents with Primary Headache: A Systematic Review. Klausen SH, Rønde G, Tornøe B, Bjerregaard L. *J Pain Res*. 2019 Dec 23;12:3437-3459

Bisgaard AM, Schönewolf-Greulich B, Ravn K, Rønde G. Is it possible to diagnose Rett syndrome before classical symptoms become obvious? Review of 24 Danish cases born between 2003 and 2012. *Eur J Paediatr Neurol*. 2015 Nov;19(6):679-87

Roende G, Petersen J, Ravn K, Fuglsang K, Andersen H, Nielsen JB, Brøndum-Nielsen K, Jensen JE. Low bone turnover phenotype in Rett syndrome: results of biochemical bone marker analysis. *Pediatr Res*. 2014 Apr;75(4):551-8

Ravn K, Roende G, Duno M, Fuglsang K, Eiklid KL, Tümer Z, Nielsen JB, Skjeldal OH. Two new Rett syndrome families and review of the literature: expanding the knowledge of MECP2 frameshift mutations. *Orphanet J Rare Dis*. 2011 Aug 30;6:58

Roende G, Ravn K, Fuglsang K, Andersen H, Nielsen JB, Brøndum-Nielsen K, Jensen JE. DXA measurements in Rett syndrome reveal small bones with low bone mass. *J Bone Miner Res*. 2011 Sep;26(9):2280-6

Roende G, Ravn K, Fuglsang K, Andersen H, Vestergaard A, Brøndum-Nielsen K, Jensen JE, Nielsen JB. Patients with Rett syndrome sustain low-energy fractures. *Pediatr Res*. 2011 Apr;69(4):359-64

Rønde G, Blichfeldt S. [Children with commotio cerebri at paediatric departments in Denmark]. [Article in Danish]. *Ugeskr Laeger*. 2008 May 12;170(20):1740-3

## Projects

### Memberships

- Member of the national steering committee for CPOP 2024-
- Member of the Grant Board for Ragnhild Folkmanns Grant 2023-
- Member of the regional steering committee for CPOP, RegionH, 2022-
- CPOP responsible, Department of Children and Youth, Herlev Hospital, 2022-
- BørneNeuroCenter (BNC) professional responsible, Department of Children and Youth, Herlev Hospital, 2022-
- Chairman of DNPS, 2019-2022
- Member of the Board, DNPS, 2018-2022
- Co-organizer of NNPS meeting 2018
- Chairman of the Neurology Committee, DPS, 2020-2023
- Member of the Neurology Committee, DPS, since 2017, since 2023 as an associate member

### Other Relevant Information