Curriculum Vitae

Ramelli Gian Paolo

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Neuropediatrics Unit
Department of Paediatrics
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Personal data

Date of birth	May 14th, 1958
Place of birth	Basel
Citizenship	Switzerland

Education

1964-1969	Primary school Bellinzona
1969-1974	Gymnasium Bellinzona
1974-1977	High School, Bellinzona
1977-1983	Faculty of medicine, University of Bern (Medical State Examination November 1983)

Licensure and academic degree

1983	Diplom of Physician (Swiss Confederation)
1983	Thesis, Doctor of Medicine, University of Bern “Quality of Life in young people with Spina Bifida” (Prof. Jean-Claude Vuille) University of Bern, Switzerland
1993	Certification as Pediatrician (FMH - Title, Swiss Medical Society)
1993	Bobath certificate in Neurology developments
1995	Certification as Paediatric Neurology (FMH - Title, Swiss Medical Society)
1997	Swiss Certificate in Electroencephalography
2003	Certificate of Full Registration as a Medical Practitioner with Specialist (General Medical Council; Unit Kingdom)

Habilitation:

2007	Certificate for delegated psychotherapy
2008	Associate Professor of Pediatrics (“**Venia docendi”)** (Faculty of Medicine, Basel) „Dystrobrevin isoform expression in patients with neuromuscular disease“ Prof. Dr.med. U.B. Schaad, Universitätskinderklinik beider Basel
2014	Professor of Pediatrics and Pedatric Neurology (Faculty of Medicine, Basel)

Awards:

2021	**Engagement Award at the annual Autismus Kongress Schweiz 2021**

Postgraduate Education

1.1.-31.12.1984/ 1.4.85-31.3.1986	**Internal Medicine** Ospedale S.Giovanni Bellinzona. Chief: Prof.Dr.C.Marone, Dr.V.Tatti
1.1.-31.3.1985:	**Gynaecology and obstetrics** Ospedale S.Giovanni Bellinzona. Chief: Dr. A. Gallino
1.4.86-31.3.1988:	**Paediatric** Kinderspital Lucerna. Chief: Prof.O.Toenz
1.4.-30.6.1988:	**Department Cerebral Polsy** Inselspital Chief: Dr. U.Aebi
1.7.88-30.6.1989:	**Paediatric Surgery** Kinderspital Lucerna. Chief: Prof. A.Schärli.
1.7.89-30.6.1990:	**Internal Medicine** Kantonsspital Lucerna Chiefs: Prof.Dr.B.Truniger, Prof. Dr.M.Nager
1.7.90-31.3.1991:	**Neuropaediatric** CHUV Losanna Chief: Prof.T.Deonna
1.5.-31.12.1991:	**Endocrinology and diabetic** Inselspital Berna Chief: Prof.U.Bürgi (Endocrinology paediatric: Prof.K. Zuppinger)
1.1-31.12.1992:	**Paediatric** Universitäts-Kinderklinik Berna Chief: Prof.E.Gugler
1.7.94-30.9.1995:	**Neurology** Neurologische Klinik Inselspital Berna Director: Prof. Dr.Ch.Hess
1.1.93-30.6.1994:	(senior registrar) **Neuropediatric Universitäts-Kinderklinik Zürich.** Director: Prof.A.Fanconi Head of unity of Neuropaediatric Department: Prof.E.Boltshauser
1.10.1995-31.04.1998:	(senior registrar) **Pediatric und Neuropediatric Universitäts-Kinderklinik, Inselspital Berne** Director: Prof.E Gugler Vice-director: Prof.F.Vassella.
01.05-30.09.1998	**Department of Neonatology** CHUV Lausanne Chief: Prof. A. Moessinger

From 01.10.1998 **Chief of Paediatric Department and Neuropediatrics Unit EOC** Bellinzona

From 01.07.2017-30.6.2023: **President of the Swiss Society of Paediatrics**

Research Training

01.04.2003-31.03.2004 Research Fellow, Neuromuscular Unit, Department of Pediatrics and Neonatal Medicine,
Imperial College, Faculty of Medicine, Hammersmith Hospital Campus, Cane Road, London W12 ONN
Chief: Prof. F Muntoni

Research Project

1. Creation of a database of mentally retardet padiatric patients assisted by the Neuropediatrics Unit of the San Giovanni Hospital in Bellinzona
   Project supported by the Advisory Board for Scientific Research Ente Ospedaliero Cantonale (ABREOC), 2009.
   Leader investigator

2. "How to learn social cognition? A Study on occupational therapy for young children with autistic spectrum disorders"
   Project supported by Swiss National Science Foundation (13DPD3_132427/1), 2010
   Co-investigator

3. Creation of a register of pediatric patients with cerebral palsy for the Canton Ticino
   Project supported by the Advisory Board for Scientific Research Ente Ospedaliero Cantonale (ABREOC), 2016
   Leader investigator

4. Creation of a database of pediatric pantients with autism spectrum disorder in the south part of Switzerland
   Project supported by the Advisory Board for Scientific Research Ente Ospedaliero Cantonale (ABREOC), 2017
   Leader investigator

5. AutoPlay: smart games for evolutionary screening in children under 14 months
   Co-funding: Advisory Board for scientific Research, Ente Ospedaliero Cantonale (ABREOC) and University of Applied Sciences and Arts of Southern Switzerland (SUPSI), 2019
   Leader investigator

Teaching activities:

Postgraduate teaching in Pediatric Neurology for medical students, Faculty of Medicine, University of Basel.

Postgraduate teaching in Pediatric Neurology for medical students, Faculty of Medicine, Università della Svizzera Italiana.

Society Membership:

- Swiss Society of Paediatrics
- Swiss Society of Neuropaediatrics
- Swiss Society of Electrophysiology
- Swiss Society for Neuroscience
- European Paediatric Neurology Society (EPNS)
- Société Européenne de Neurologie Pédiatrique (SENP)
- European Academy of Childhood Disability
- Gesellschaft für Neuropädiatrie

Activity as Reviewer

- Pediatrics
- Acta Paediatrica
- Journal of Pediatrics
- Journal of Pediatric
- Pediatric Neurology
- Neuropediatrics
- Epilepsia
- Clinical Medicine
- Tribuna Medica Ticinese

Scientific publications (peer reviewed)

1. Tatti V, **Ramelli GP**. Pneumopathie interstitielle et amiodarone. Schweiz Med Wochenschr 1986; 116: 377-382.
2. Tönz O, **Ramelli GP**, Gebbers JO. Hepatitis-B-assoziierte membranöse Glomerulonephritis: Heilung nach medikamentös induzierter Reaktivierung der Hepatitis. Helv Paediatr Acta 1989;43:539-547.
3. **Ramelli GP**, Marone C, Truniger B. Akutes Nierenversagen bei infektiöser Mononucleose. Schweiz Med Wochenschr 1990; 120: 1590-1594.
4. **Ramelli GP**, Marone C, Gebbers JO. Sicca-Komplex bei Amyloidose. Schweiz Med Wochenschr 1990; 120: 995-958.
5. **Ramelli GP**, Gebbers JO, Tönz O. Chronische, letal verlaufende Zytomegalievirus-Erkrankung mit Hypereosinophilie und multiplem Organbefall nach neonataler Infektion. Schweiz Med Wochenschr 1990; 120: 632-640.
6. **Ramelli GP**, Tönz O, Zimmermann A, Lentze MJ. Crohn disease with sclerosing cholangitis and liver cirrhosis in adolescence. Eur J Pediatr 1991; 150: 557-559.
7. Mersin SS, **Ramelli GP**, Laux-End R, Bianchetti MG. Urinary chloride excretion distinguishes between renal and extrarenal metabolic alkalosis. Eur J Pediatr 1995; 154: 979-982.
8. Pedrozzi NE, **Ramelli GP**, Tomasetti R, Nobile-Buetti L, Bianchetti MG. Rhabdomyolysis and anesthesia: a report of two cases and review of the literature. Pediatr Neurol 1996; 15: 254-257.
9. Schneider JF, Baumgartner R, Ammann R, **Ramelli GP**, Remonda L. Unilaterales Moyamoya-Syndrom. Transkranielle Farbduplexsonographie, MRT und Angiographie. Röfo Fortschr Geb Röntgenstr Neuen Bildgeb Verfahr 1997; 167: 322-324.
10. **Ramelli GP**, Sturzenegger M, Bianchetti MG, Vassella F. Thymectomy in children with generalized myasthenia gravis. Neuropediatrics 1997; 28: 292.
11. Lövblad K, **Ramelli GP**, Remonda L, Nirkko AC, Ozdoba C, Schroth G. Retardation of myelination due to dietary vitamin B12 deficiency: cranial MRI findings. Pediatr Radiol 1997; 27: 155-158.
12. **Ramelli GP**, Bianchetti MG. Dapsone in cutaneous Henoch-Schönlein syndrome - worth a trial. Acta Paediatr 1997;86:337.
13. Lövblad K, Kelkar P, Ozdoba C, **Ramelli GP**, Remonda L, Schroth G. Pure methotrexate encephalopathy presenting with seizures: CT and MRI features. Pediatr Radiol 1998;28:86-91.
14. Ozdoba C, **Ramelli GP**, Schroth G. MRI in a patient with congenital agammaglobulinaemia. Neuroradiology 1998;40:516-518.
15. **Ramelli GP**, Sturzenegger M, Donati F, Karbowski K. EEG findings during basilar migraine attacks in children. Electroencephalogr Clin Neurophysiol 1998;107:374-378.
16. **Ramelli GP**, von der Weid N, Stanga Z, Mullis PE, Buergi U. Suprasellar germinomas in childhood and adolescence: diagnostic pitfalls. J Pediatr Endocrinol Metab 1998;11:693-697.
17. **Ramelli GP**, Donati F, Moser H, Vassella F. Concomitance of childhood absence and Rolandic epilepsy. Clin Electroencephalogr 1998;29:177-180.
18. **Ramelli GP**, Donati F, Bianchetti MG, Vassella F. Apnoeic attacks as an isolated manifestation of epileptic seizures in infants. Eur J Paediatr Neurol 1998;2:187-191.
19. **Ramelli GP**, Donati F, Kollar M, Remonda L, Vassella F. Rotatory seizures in a patient with tuberous sclerosis. Epileptic Disord 1999;1:233-235.
20. Vella S, **Ramelli GP**, Schroth G, Bianchetti MG. Circulating antineutrophil autoantibodies in a child with isolated central nervous system vasculitis. Neuropediatrics 1999;30:268-269.
21. **Ramelli GP**, Nagy L, Tuncdogan E, Mathis J. Ganglion cyst of the peroneal nerve: a differential diagnosis of peroneal nerve entrapment neuropathy. Eur Neurol 1999;41:56-58.
22. Villiger RM, von Vigier RO, **Ramelli GP**, Hassink RI, Bianchetti MG. Precipitants in 42 cases of erythema multiforme. Eur J Pediatr 1999;158:929-932.
23. Morger ID, Truttmann AC, von Vigier RO, Bettinelli A, **Ramelli GP**, Bianchetti MG. Plasma ionized magnesium in tubular disorders with and without total hypomagnesemia. Pediatr Nephrol 1999;13:50-53.
24. Donati F, **Ramelli GP**. Rotatory seizures are not so rare as described. Epileptic Disord 2000;2:177.
25. **Ramelli GP**, Remonda L, Lövblad KO, Hirsiger H, Moser H. Abnormal myelination in a patient with deletion 14q11.2q13.1. Pediatr Neurol 2000;23:170-172.

26. **Ramelli GP**, Vella S, Lovblad K, Remonda L, Vassella F. Swelling of the third nerve in a child with transient oculomotor paresis: a possible cause of ophthalmoplegic migraine. Neuropediatrics 2000;31:145-147.
27. **Ramelli GP**, von der Weid N, Remonda L, Mariani L, Weis J. Pleomorphic xanthoastrocytoma derived from glioneuronal malformation in a child with intractable epilepsy. J Child Neurol 2000;15:270-272.
28. Stoffel PB, Sauvain MJ, von Vigier RO, Beretta-Piccoli BC, **Ramelli GP**, Bianchetti MG. Non-infectious causes of uveitis in 70 Swiss children. Acta Paediatr 2000;89:955-958.
29. **Ramelli GP**, Slongo T, Weis J, Tschäppler H, Vassella F. Pseudoarthrose bei Neurofibromatose Typ 1. Klin Pädiatr 2000;212:26-30.
30. von Vigier RO, Mozzettini S, Truttmann AC, Meregalli P, **Ramelli GP**, Bianchetti MG. Cough is common in children prescribed converting enzyme inhibitors. Nephron 2000;84:98.
31. **Ramelli GP**, Slongo T, Tschäppeler H, Weis J. Congenital pseudarthrosis of the ulna and radius in two cases of neurofibromatosis type 1. Pediatr Surg Int 2001;17:239-341.
32. **Ramelli GP**, Wyttenbach R, von der Weid N, Ozdoba C. Anterior spinal artery syndrome in an adolescent with protein S deficiency. J Child Neurol 2001;16:134-135.
33. Lin S, **Ramelli GP** Moser H, Gallati S, Burgunder JM. A novel insert mutation in gamma-sarcoglycan gene leads to severe childhood autosomal recessive muscular dystrophy. J Neurol 2002;249:1608-1611.
34. Alkan M, **Ramelli GP**, Hirsiger H, Keser I, Remonda L, Bühler EM, Moser H. Presumptive monosomy 21 with neuronal migration disorder re-diagnosed as de novo unbalanced translocation t(18p;21q) by fluorescence in situ hybridisation. Genet Couns 2002;13:151-156.
35. Simonetti GD, Dumont-Dos Santos K, Pachlopnik JM, **Ramelli GP**, Bianchetti MG. Hemolytic uremic syndrome linked with infectious mononucleosis. Pediatr Nephrol 2003;18:1193-1194.
36. Widmer S, Ghisla R, **Ramelli GP**, Taminelli F, Widmer B, Caoduro L, Gallino A. Tele-echocardiography in paediatrics. Eur J Pediatr 2003;162:271-275.
37. Donati-Genet PC, **Ramelli GP**, Bianchetti MG. A newborn infant of a diabetic mother with refractory hypocalcaemic convulsions. Eur J Pediatr 2004;163:759-760.
38. Stauffer CM, van der Weg B, Donadini R, **Ramelli GP**, Marchand S, Bianchetti MG. Family history and behavioral abnormalities in girls with recurrent urinary tract infections: a controlled study. J Urol 2004;171:1663-1665.
39. **Ramelli GP**, Simonetti GD, Gorgievski-Hrisoho M, Aebi C, Bianchetti MG. Outbreak of coxsackie B5 virus meningitis in a scout camp. Pediatr Infect Dis J 2004;23:86-87.
40. Tosi F, Bianda NDF, Truttmann AC, Crosazzo L, Bianchetti MG, Bettinelli A, **Ramelli GP.** Normal plasma total magnesium in Gitelman's syndrome. Am J Med 2004;116:573-574.
41. **Ramelli GP**, Aebi C, Remonda L, Lovblad KO. Basal ganglia infarction complicating basilar meningitis: visualization by serial magnetic resonance imaging. Eur Neurol 2005;54:227-229.
42. **Ramelli GP**, Hammer J. Swiss physicians' practices of long-term mechanical ventilatory support of patients with Duchenne Muscular Dystrophy. Swiss Med Wkly 2005;135:599-604.
43. **Ramelli GP**, Sozzo AB, Vella S, Bianchetti MG. Benign neonatal sleep myoclonus: an under-recognized, non-epileptic condition. Acta Paediatr 2005;94:962-963.
44. Steinlin M, Pfister I, Pavlovic J, Everts R, Boltshauser E, Capone Mori A, Gubser Mercati D, Hanggeli CA, Keller E, Luetschg J, Marcoz J, **Ramelli GP**, Roulet Perez E, Schmitt-Mechelke T, Weissert M; The Swiss Societies of Paediatric Neurology and Neonatology. The first three years of the Swiss Neuropaediatric Stroke Registry (SNPSR): a population-based study of incidence, symptoms and risk factors. Neuropediatrics 2005;36:90-97.
45. Schlapbach LJ, Sozzo A, **Ramelli GP**, Bianchetti MG. Severe normotensive metabolic alkalosis in a 2-month-old boy with hyperekplexia. J Inherit Metab Dis 2006;29:238-239.
46. **Ramelli GP**, Joncourt F, Gasperini M, Tonin P, Burgunder J-M. Rather mild phenotype in a patient with homozygous null mutations in the a-sarcoglycan gene. Swiss Med Wkly 2006;136:96-97.
47. **Ramelli GP,** Gallati S, Weis J, Krähenbühl S, Burgunder JM. Point Mutation $tRNA^{Ser(UCN)}$ in a Child with Hearing Loss and Myoclonus Epilepsy. J Child Neurol 2006;21:253-5.

48. Pavlovic J, Kaufmann F, Boltshauser E, Capone Mori A, Gubser Mercati D, Haenggeli CA, Keller E, Lutschg J, Marcoz JP, **Ramelli GP**, Roulet Perez E, Schmitt-Mechelke T, Weissert M, Steinlin M. Neuropsychological problems after paediatric stroke: two year follow-up of Swiss children. Neuropediatrics 2006;37:13-19.
49. **Ramelli GP,** Joncourt F, Lütschg J, Weis J, Tolnay M, Burgunder JM. Becker muscular dystrophy with marked divergence between clinical and molecular genetic findings: case series. Swiss Med Wkly 2006;136:189-193.
50. **Ramelli GP**, Zanda N, Wyttenbach M, Bronz L, Schnider A. The prognosis of agenesis of the corpus callosum might mostly be favorable. Swiss Med Wkly 2006; 136: 404-405.
51. **Ramelli GP**, Aloysius A, King C, Davies T, Muntoni F. Gastrostomy placement in paediatric neuromuscular patients: indications and outcome. Dev Med Child Neurol 2007; 49: 367-371.
52. **Ramelli GP**, Boscherini D, Kehrli P, Rilliet B. Spontaneous Spinal Epidural Hematomas in Children: Can we prevent a negative prognosis? Reflections on 2 Cases. J Child Neurol 2008; 23: 564-567.
53. **Ramelli GP**, Sillaci C, Ferrarrini A, Cattaneo C, Visconti P, Pescia G. Microduplication 22q11.1 in a child with autism spectrum disorder: clinical and genetic study. Dev Med Child Neurol 2008; 50: 953-955.
54. Wohlrab G, Leiba H, Kästle R, **Ramelli GP**, Schmitt-Mechelke T, Schmitt B, Landau L. Vigabatrin therapy in infantile spasms: Solving one problem and inducing another. Epilepsia 2009; 50: 2008-2009.
55. Garzoni L, Vanoni F, Rizzi M, Simonetti G, Goeggel-Simonetti B, Bianchetti MG, **Ramelli GP**. Nervous system dysfunction in Henoch-Schönlein syndrome: systematic review of the literature. Rheumatology 2009; 48: 1524-1529.
56. Ferrarini A, **Ramelli GP**, Bianchetti MG, Hedman J, Sharathkumar A, Shapiro A, Bourland C. Index of Suspicion. Pediatr Rev 2009; 30: 479-485.
57. Belotti E, Taddeo I, Ragazzi M, Pifferini R, Simonetti G, Bianchetti MG, **Ramelli GP.** Chronic impact of topiramate on acid-base balance and potassium in childhood. Eur J Paediatr Neurol 2010; 14: 445-448.
58. **Ramelli GP**, Zanda N, Bianchetti MG, Leoni F. Age-dependent presentation in children with attention deficit hyperactivity disorder. World J Pediatr 2010; 6: 90.
59. Maurer VO, Rizzi M, Bianchetti MG, **Ramelli GP.** Benign Neonatal Sleep Myoclonus: A Review of the Literature. Pediatrics 2010; 125; e919-e924.
60. Milani G, Ragazzi M, Simonetti GD, **Ramelli GP**, Rizzi M, Bianchetti MG; Fossali EF. Superior palatability of crusche lercanidipine compared to amlopidine among children. Br J Clin Pharmacol 2010, 69: 204-206.
61. Buerki S, Roellin K, Remonad L, Mercati DG, Jeannet PY, Keller E, Luetschg J, Menache C, **Ramelli GP**, Schmitt-Mechelke T, Weissert M, Boltshauser E, Steinlin M. Neuroimaging in childhood arterial ischaemic stroke: evaluation of imaging modalities and aetiologies. Dev Med Child Neurol 2010; 52: 1033-1037.
62. **Ramelli GP**. Dystrobrevin isoform expression in patients with neuromuscular disease. Clin Neuropathol 2010; 29: 254-261.
63. Poncini C, Willi B, Pezzoli V, Bianchetti MG, **Ramelli GP.** Rotavirus and acute cerebellar ataxia. J Paediatr Child Health 2010; 46: 448.
64. Gauthey M, Poloni CB, **Ramelli GP**, Roulet-Perez E, Korff CM. Status Epilepticus in Fragile X Syndrome. Epilepsia 2010; 51: 2470-2473.
65. **Ramelli GP**, Cortesi C, Boscherini D, Faggin R, Bianchetti MG. Age-dependent presentation of tectal plate tumors: preliminary observations. J Child Neurol 2011; 26: 377-380.
66. Faré PB, Lava SA, Simonetti GD, **Ramelli GP**, Bianchetti MG. Blue breath-holding spells caused by an alcoholic vitamin D_3 preparation in infancy. Acta Paediatr 2011; 100: e4.
67. Jacquemont S, Reymond A, Zufferey F, Harewood L, Walters RG, Kutalik Z, Martinet D, Shen Y, Valsesia A, Beckmann ND, Thorleifsson G, Belfiore M, Bouquillon S, Campion D, de Leeuw N, de Vries BB, Esko T, Fernandez BA, Fernández-Aranda F, Fernández-Real JM, Gratacòs M, Guilmatre A, Hoyer J, Jarvelin MR, Frank Kooy R, Kurg A, Le Caignec C, Männik K, Platt OS, Sanlaville D, Van Haelst MM, Villatoro Gomez S, Walha F, Wu BL, Yu Y, Aboura A, Addor MC, Alembik Y, Antonarakis SE, Arveiler B, Barth M, Bednarek N, Béna F, Bergmann S, Beri M, Bernardini L, Blaumeiser B, Bonneau D, Bottani A, Boute O, Brunner HG, Cailley D, Callier P, Chiesa J, Chrast J, Coin L, Coutton C, Cuisset JM, Cuvellier JC, David A, de Freminville B, Delobel B, Delrue MA, Demeer B, Descamps D, Didelot G, Dieterich K, Disciglio V, Doco-Fenzy M, Drunat S, Duban-Bedu B, Dubourg C, El-Sayed Moustafa JS, Elliott P, Faas BH, Faivre L, Faudet A, Fellmann F, Ferrarini A, Fisher R, Flori E, Forer L, Gaillard D, Gerard M, Gieger C, Gimelli S, Gimelli G, Grabe HJ, Guichet A, Guillin

O, Hartikainen AL, Heron D, Hippolyte L, Holder M, Homuth G, Isidor B, Jaillard S, Jaros Z, Jiménez-Murcia S, Joly Helas G, Jonveaux P, Kaksonen S, Keren B, Kloss-Brandstätter A, Knoers NV, Koolen DA, Kroisel PM, Kronenberg F, Labalme A, Landais E, Lapi E, Layet V, Legallic S, Leheup B, Leube B, Lewis S, Lucas J, Macdermot KD, Magnusson P, Marshall C, Mathieu-Dramard M, McCarthy MI, Meitinger T, Antonietta Mencarelli M, Merla G, Moerman A, Mooser V, Morice-Picard F, Mucciolo M, Nauck M, Coumba Ndiaye N, Nordgren A, Pasquier L, Petit F, Pfundt R, Plessis G, Rajcan-Separovic E, **Ramelli GP**, Rauch A, Ravazzolo R, Reis A, Renieri A, Richart C, Ried JS, Rieubland C, Roberts W, Roetzer KM, Rooryck C, Rossi M, Saemundsen E, Satre V, Schurmann C, Sigurdsson E, Stavropoulos DJ, Stefansson H, Tengström C, Thorsteinsdóttir U, Tinahones FJ, Touraine R, Vallée L, van Binsbergen E, Van der Aa N, Vincent-Delorme C, Visvikis-Siest S, Vollenweider P, Völzke H, Vulto-van Silfhout AT, Waeber G, Wallgren-Pettersson C, Witwicki RM, Zwolinksi S, Andrieux J, Estivill X, Gusella JF, Gustafsson O, Metspalu A, Scherer SW, Stefansson K, Blakemore AI, Beckmann JS, Froguel P. Mirror extreme BMI phenotypes associated with gene dosage at the chromosome 16p11.2 locus. Nature 2011; 478: 97-102.

68. Martin C, von Elm E, El-Koussy M, Boltshauser E, Capone A, Fluss J, Gubser MD, Keller E, Müller A, **Ramelli GP**, Roulet-Perrez E, Schmid R, Schmitt-Mechelke T, Wehrli E, Weissert M Steinlin M. Delayed diagnosis of acute ischemic stroke in children - a registry-based study in Switzerland. Swiss Med Wkly 2011; 141: w13281.
69. Sidler JA, Filges I, Boesch N, **Ramelli GP**, Röthlisberger B, Huber AR, Tercanli S, Bronz L, Miny P, Heinimann K. TRPS1 codon 952 constitutes a mutational hot spot in trichorhinophalangeal syndrome type I and could be associated with intellectual disability. Clin. Dysmorphol 2012; 21: 87-90.
70. Salvaterra E, Giorda R, Bassi MT, Borgatti R, Knudsen LE, Martinuzzi A, Nobile M, Pozzoli U, **Ramelli GP**, Reni GL, Rivolta D, Stazi MA, Strazzer S, Thijs C, Toccaceli V, Trabacca A, Turconi AC, Zanini S, Zucca C, Bresolin N, Lenzi L, on behalf of the Pediatric Biobank ELSI Working Group. Pediatric biobanking: a pilot qualitative survey of practices, rules, and researcher opinions in ten European countries. Biopreserv Biobank 2012; 10: 29-36.
71. Lava SA, Simonetti GD, **Ramelli GP**, Tschumi S, Bianchetti MG. Symptomatic management of fever by Swiss board-certified pediatricians: results from a cros-sectional, Web-based survey. Clin Ther 2012; 34: 250-256.
72. Salmina C, Taddeo I, Falesi M, Weber P, Bianchetti MG, Ramelli GP. Paroxysmal tonic upgaze in normal children: A case series and a review of the literature. Eur J Paediatr Neurol 2012; 16: 683-687.
73. Schlapbach LJ, Adams M, Proietti E, Aebischer M, Grunt S, Borradori-Tolsa C, Bickle-Graz M, Bucher HU, Zeilinger G, Capone A, Steiner F, Schulzke S, Weber P, **Ramelli GP**, Nelle M, Steinlin M, Grunt S, Hassink R, Bär W, Keller E, Killer Ch, Fuhrer K, Tolsa JF, Bickle-Graz M, Pfister RE, Hüppi PS, Borradori-Tolsa C, Berger TM, Schmitt-Mechelke T, Pezzoli V, Ecoffey M, Mueller A, Malzacher A, Micallef JP, Schaefer Ch, von Rhein M, Arlettaz Mieth R, Bernet V, Latal B, Natalucci G. Outcome at two years of age in a Swiss national cohort of extremely preterm infants born between 2000 and 2008. BMC Pediatr 2012; 12: 198.
74. Datta AN, Oser N, **Ramelli GP**, Gobbin NZ, Lantz G, Penner IK, Weber P. BECTS evolving to Landau-Kleffner Syndrome and back by subsequent recovery: A longitudinal language reorganization case study using fMRI, source EEG, and neuropsychological testing. Epilepsy Behav 2013; 27: 107-114.
75. Lava SA, Simonetti GD, Ferrarini A, **Ramelli GP**, Bianchetti MG. Regional differences in symptomatic fever management among pediatricians in Switzerland: The results of a cross-sectional Web-based survey. Br J Clin Pharmacol 2013; 75: 236-243.
76. Antonini U, Soldini EA, D'Apuzzo V, Brunner R, **Ramelli GP**. Longitudinal neurodevelopmental evolution in children with severe non-progressive encephalopathy. Brain Dev 2013; 35: 548-554.
77. Datta AN, Oser N, Bauder F, Maier O, Martin F, **Ramelli GP**, Steinlin M, Weber P, Penner IK. Cognitive impairment and cortical reorganization in children with benign epilepsy with centrotemporal spikes. Epilepsia. 2013; 54: 487-494 .
78. Capobianco S, Lava S, Bianchetti MG, Martinet D, Belfiore M, **Ramelli GP**, Ferrarini A. Array CGH among children with mental retardation: a useful diagnostic tool for skilled geneticists. Dev Med Child Neurol 2013. Dev Med Child Neurol. 2014;56:290.
79. Dell'Orto VG, Belotti EA, Goeggel-Simonetti B, Simonetti GD, **Ramelli GP**, Bianchetti MG, Lava SAG. Metabolic disturbances and renal stone promotion on treatment with topiramate: a systematic review. Br J Clin Pharmacol. 2014;77:958-64.

80. Dell'Orto V, Bernasconi E, Pezzoli V, Queirolo S, **Ramelli GP**. Severe, life-threatening Epstein- Barr Virus encephalitis in an immunocompetent adolescent girl. J Pediatr Neurol 2014; 12: 217–219.
81. Ferrarini A, Lava SAG, Simonetti GD, **Ramelli GP**, Bianchetti MG; Swiss Italian Society of Pediatrics. Influenzavirus B-associated acute benign myalgia cruris: an outbreak report and review of the literature. Neuromuscul Disord. 2014;24:342-346.
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Supervised Doctoral Thesis

1. Vitamin D status among children and adolescents on anticonvulsant drugs in southern Switzerland. 2014
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Bellinzona, 30.11.2023 Prof. Dr. Med. Gian Paolo Ramelli