

**Professor Dr. med. Dr. h.c. mult. Hartmut P.H. Neumann**  
**Peer Review Journal Publications**

**Publications in Peer Review Journals**

**2021**

1. Bayley JP, Bausch B, Jansen JC, Hensen EF, van der Tuin K, Corssmit EP, Devilee P, **Neumann HP**.  
*SDHB* variant type impacts phenotype and malignancy in pheochromocytoma-paraganglioma.  
**J Med Genet.** 2021 Nov 8:jmedgenet-2020-107656. doi: 10.1136/jmedgenet-2020-107656. Online ahead of print. PMID: 34750193

**2020**

1. Bancos I, Atkinson E, Eng C, Young WF Jr, **Neumann HPH**; International Pheochromocytoma and Pregnancy Study Group.  
Maternal and fetal outcomes in phaeochromocytoma and pregnancy: a multicentre retrospective cohort study and systematic review of literature.  
**Lancet Diabetes Endocrinol.** 2021 Jan;9(1):13-21. doi: 10.1016/S2213-8587(20)30363-6. Epub 2020 Nov 26. PMID: 33248478
2. Därr R, Kater J, Sekula P, Bausch B, Krauss T, Bode C, Walz G, **Neumann HP**, Zschiedrich S.  
Clinical decision making in small non-functioning VHL-related incidentalomas.  
**Endocr Connect.** 2020 Aug;9(8):834-844. doi: 10.1530/EC-20-0208. PMID: 32869749 Free PMC article.
3. Casey R, **Neumann HPH**, Maher ER.  
Genetic stratification of inherited and sporadic phaeochromocytoma and paraganglioma: implications for precision medicine.  
**Mol Genet.** 2020 Oct 20;29(R2):R128-R137. doi: 10.1093/hmg/ddaa201. PMID: 33059362 Review.
4. Bayley JP, Bausch B, Rijken JA, van Hulsteijn LT, Jansen JC, Ascher D, Pires DEV, Hes FJ, Hensen EF, Corssmit EPM, Devilee P, **Neumann HPH**.  
Variant type is associated with disease characteristics in *SDHB*, *SDHC* and *SDHD*-linked phaeochromocytoma-paraganglioma.  
**J Med Genet.** 2020 Feb;57(2):96-103. doi: 10.1136/jmedgenet-2019-106214. Epub 2019 Sep 6. PMID: 31492822
5. Klingler JH, Gläsker S, Bausch B, Urbach H, Krauss T, Jilg CA, Steiert C, Puzik A, Neumann-Haefelin E, Kotsis F, Agostini H, **Neumann HPH**, Beck J.  
Hemangioblastoma and von Hippel-Lindau disease: genetic background, spectrum of disease, and neurosurgical treatment.  
**Childs Nerv Syst.** 2020 Oct;36(10):2537-2552. doi: 10.1007/s00381-020-04712-5. Epub 2020 Jun 7. PMID: 32507909 Free PMC article.

6. Larsen LV, Mirebeau-Prunier D, Imai T, Alvarez-Escola C, Hasse-Lazar K, Censi S, Castroneves LA, Sakurai A, Kihara M, Horiuchi K, Barbu VD, Borson-Chazot F, Gimenez-Roqueplo AP, Pigny P, Pinson S, Wohllk N, Eng C, Aydogan BI, Saranath D, Dvorakova S, Castinetti F, Patocs A, Bergant D, Links TP, Peczkowska M, Hoff AO, Mian C, Dwight T, Jarzab B, **Neumann HPH**, Robledo M, Uchino S, Barlier A, Godballe C, Mathiesen JS.  
Primary hyperparathyroidism as first manifestation in multiple endocrine neoplasia type 2A: an international multicenter study.  
**Endocr Connect.** 2020 Jun;9(6):489-497. doi: 10.1530/EC-20-0163. PMID: 32375120

## 2019

7. **Neumann HP**, Young WF, Eng C.  
Pheochromocytoma and Paraganglioma.  
**N Engl J Med.** 2019 Aug 8;381(6):552-565. doi: 10.1056/NEJMra1806651.
8. **Neumann HP**, Young WF Jr, Eng C.  
Pheochromocytoma and Paraganglioma. Reply.  
**N Engl J Med.** 2019 Nov 7;381(19):1883. doi: 10.1056/NEJMc1912022.
9. Schuhmacher P, Kim E, Hahn F, Sekula P, Jilg CA, Leiber C, **Neumann HP**, Schultze-Seemann W, Walz G, Zschiedrich S.  
Growth characteristics and therapeutic decision markers in von Hippel-Lindau disease patients with renal cell carcinoma.  
**Orphanet J Rare Dis.** 2019 Oct 28;14(1):235. doi: 10.1186/s13023-019-1206-2.
10. **Neumann HP**, Schmid KW, Eng C.  
Morphology and etiology of pheochromocytoma.  
**Pathologe.** 2019 Oct 7. doi: 10.1007/s00292-019-00663-0. [Epub ahead of print] Review.
11. Bayley JP, Bausch B, Rijken JA, van Hulsteijn LT, Jansen JC, Ascher D, Pires DEV, Hes FJ, Hensen EF, Corssmit EPM, Devilee P, **Neumann HP**.  
Variant type is associated with disease characteristics in SDHB, SDHC and SDHD-linked pheochromocytoma-paraganglioma.  
**J Med Genet.** 2019 Sep 6. pii: jmedgenet-2019-106214. doi: 10.1136/jmedgenet-2019-106214. [Epub ahead of print]
12. **Neumann HP**, Tsoy U, Bancos I, Amodru V, Walz MK, Tirosh A, Kaur RJ, McKenzie T, Qi X, Bandgar T, Petrov R, Yukina MY, Roslyakova A, van der Horst-Schrivers ANA, Berends AMA, Hoff AO, Castroneves LA, Ferrara AM, Rizzati S, Mian C, Dvorakova S, Hasse-Lazar K, Kvachenyuk A, Peczkowska M, Loli P, Erenler F, Krauss T, Almeida MQ, Liu L, Zhu F, Recasens M, Wohllk N, Corssmit EPM, Shafiqullina Z, Calissendorff J, Grozinsky-Glasberg S, Kunavisarut T, Schalin-Jääntti C, Castinetti F, Vlcek P, Beltsevich D, Egorov VI,

Schiavi F, Links TP, Lechan RM, Bausch B, Young WF Jr, Eng C; International Bilateral-Pheochromocytoma-Registry Group.

Comparison of Pheochromocytoma-Specific Morbidity and Mortality Among Adults With Bilateral Pheochromocytomas Undergoing Total Adrenalectomy vs Cortical-Sparing Adrenalectomy.

**JAMA Netw Open.** 2019 Aug 2;2(8):e198898. doi: 10.1001/jamanetworkopen.2019.8898.

13. Castinetti F, Waguespack SG, Machens A, Uchino S, Hasse-Lazar K, Sanso G, Else T, Dvorakova S, Qi XP, Elisei R, Maia AL, Glod J, Lourenço DM Jr, Valdes N, Mathiesen J, N, Bandgar TR, Drui D, Korbonits M, Druce MR, Brain C, Kurzawinski T, Patocs A, Bugalho MJ, Lacroix A, Caron P, Fainstein-Day P, Borson Chazot F, Klein M, Links TP, Letizia C, Fugazzola L, Chabre O, Canu L, Cohen R, Tabarin A, Spehar Uroic A, Maiter D, Laboureau S, Mian C, Peczkowska M, Sebag F, Brue T, Mirebeau-Prunier D, Leclerc L, Bausch B, Berdelou A, Sukurai A, Vlcek P, Krajewska J, Barontini M, Vaz Ferreira Vargas C, Valerio L, Ceolin L, Akshintala S, Hoff A, Godballe C, Jarzab B, Jimenez C, Eng C, Imai T, Schlumberger M, Grubbs E, Dralle H, **Neumann HP**, Baudin E. Natural history, treatment, and long-term follow up of patients with multiple endocrine neoplasia type 2B: an international, multicentre, retrospective study. **Lancet Diabetes Endocrinol.** 2019 Mar;7(3):213-220. doi: 10.1016/S2213-8587(18)30336-X. Epub 2019 Jan 16.  
Erratum in: *Lancet Diabetes Endocrinol.* 2019 Mar;7(3):e3. PMID:30660595

14. von Dobschütz E, **Neumann HPH**. [Genetics of pheochromocytoma and the relevance in surgery]. **Chirurg.** 2019 Jan;90(1):15-22. doi: 10.1007/s00104-018-0741-z. Review. German. PMID:30306232

## 2018

15. **Neumann HPH**, Young W, Krauss T, Bayley JP, Schiavi F, Opocher G, Boedeker C, Tirosh A, Castinetti F, Ruf J, Beltsevich D, Walz MK, Groeben H, Von Dobschuetz E, Gimm O, Wohllk N, Pfeifer M, Lourenço DM, Peczkowska M, Patocs A, Ngeow J, Makay O, Shah N, Tischler AS, Leijon H, Pennelli G, Villar Gómez de Las Heras K, Links TP, Bausch B, Eng C. Genetics Informs Precision Practice in the Diagnosis and Management of Pheochromocytoma. **Endocr Relat Cancer.** 2018 May 24. pii: ERC-18-0085. doi: 10.1530/ERC-18-0085. [Epub ahead of print] PMID: 29794110
16. Martinez MF, Mazzuocolo LD, Oddo EM, Iscoff PV, Muchnik C, **Neumann HPH**, Martin RS, Fraga AR, Azurmendi PJ. Co-Inheritance of Autosomal Dominant Polycystic Kidney Disease and Naevoid Basal Cell Carcinoma Syndrome: Effects on Renal Progression. **Nephron.** 2018;140(4):282-288. doi: 10.1159/000490771. Epub 2018 Oct 26. PMID:30368514

17. Krauss T, Ferrara AM, Links TP, Wellner U, Bancos I, Kvachenyuk A, Villar Gómez de Las Heras K, Yukina M, Petrov R, Bullivant G, von Duecker L, Jadhav SS, Ploeckinger U, Welin S, Schalin-Jantti C, Gimm O, Pfeifer M, Ngeow J, Hasse-Lazar K, Sanso G, Qi XP, Ugurlu U, Diaz RE, Wohlk N, Peczkowska M, Aberle J, Lourenço DM Jr, Pereira MA, Fragoso MCBV, Hoff AO, Almeida MQ, Violante AHD, Quidute ARP, Zhang Z, Recasens M, Robles Diaz L, Kunavisarut T, Wannachalee T, Sirinvaravong S, Jonasch E, Grozinsky-Glasberg S, Fraenkel M, Beltsevich D, Egorov VI, Bausch D, Schott M, Tiling N, Pennelli G, Zschiedrich S, Därr R, Ruf J, Denecke T, Link KH, Zovato S, von Dobschuetz E, Yaremchuk S, Amthauer H, Makay O, Patocs A, Walz MK, Huber TB, Seufert J, Hellman P, Kim RH, Kuchinskaya E, Schiavi F, Malinoc A, Reisch N, Jarzab B, Barontini M, Januszewicz A, Shah N, Young W, Opocher G, Eng C, **Neumann HPH**, Bausch B.  
Preventive medicine for von Hippel-Lindau disease-associated pancreatic neuroendocrine tumors.  
**Endocr Relat Cancer**. 2018 May 10. pii: ERC-18-0100. doi: 10.1530/ERC-18-0100. [Epub ahead of print] PMID: 29748190
18. Walz MK, Iova LD, Deimel J, **Neumann HPH**, Bausch B, Zschiedrich S, Groeben H, Alesina PF.  
Minimally Invasive Surgery (MIS) in Children and Adolescents with Pheochromocytomas and Retroperitoneal Paragangliomas: Experiences in 42 Patients.  
**World J Surg**. 2018 Apr;42(4):1024-1030. doi: 10.1007/s00268-018-4488-y. PMID:29392429

## 2017

19. Bausch B, Tischler AS, Schmid KW, Leijon H, Eng C, **Neumann HPH**.  
Max Schottelius: Pioneer in Pheochromocytoma.  
**J Endocr Soc**. 2017 Jul 1;1(7):957-964. doi: 10.1210/js.2017-00208. eCollection 2017 Jul 1. Review. PMID:29264546
20. Thomaschewski M, Neeff H, Keck T, **Neumann HPH**, Strate T, von Dobschuetz E.  
Is there any role for minimally invasive surgery in NET?  
**Rev Endocr Metab Disord**. 2017 Dec;18(4):443-457. doi: 10.1007/s11154-017-9436-x. Review. PMID:29127554
21. Castinetti F, Maia AL, Peczkowska M, Barontini M, Hasse-Lazar K, Links TP, Toledo RA, Dvorakova S, Mian C, Bugalho MJ, Zovato S, Alevizaki M, Kvachenyuk A, Bausch B, Loli P, Bergmann SR, Patocs A, Pfeifer M, Costa JB, von Dobschuetz E, Letizia C, Valk G, Barczynski M, Czetwertynska M, Plukker JTM, Sartorato P, Zelinka T, Vlcek P, Yaremchuk S, Weryha G, Canu L, Wohlk N, Sebagg F, Walz MK, Eng C, **Neumann HPH**.

The penetrance of MEN2 pheochromocytoma is not only determined by *RET* mutations.

**Endocr Relat Cancer.** 2017 Aug;24(8):L63-L67. doi: 10.1530/ERC-17-0189.

Epub 2017 Jun 25. PMID: 28649091

22. Bausch B, Schiavi F, Ni Y, Welander J, Patocs A, Ngeow J, Wellner U, Malinoc A, Taschin E, Barbon G, Lanza V, Söderkvist P, Stenman A, Larsson C, Svahn F, Chen JL, Marquard J, Fraenkel M, Walter MA, Peczkowska M, Prejbisz A, Jarzab B, Hasse-Lazar K, Petersenn S, Moeller LC, Meyer A, Reisch N, Trupka A, Brase C, Galiano M, Preuss SF, Kwok P, Lendvai N, Berisha G, Makay Ö, Boedeker CC, Weryha G, Racz K, Januszewicz A, Walz MK, Gimm O, Opocher G, Eng C, **Neumann HPH**; European-American-Asian Pheochromocytoma-Paraganglioma Registry Study Group. Clinical Characterization of the Pheochromocytoma and Paraganglioma Susceptibility Genes *SDHA*, *TMEM127*, *MAX*, and *SDHAF2* for Gene-Informed Prevention. **JAMA Oncol.** 2017 Sep 1;3(9):1204-1212. doi: 10.1001/jamaoncol.2017.0223.
23. Groeben H, Nottebaum BJ, Alesina PF, Traut A, **Neumann HP**, Walz MK. Perioperative  $\alpha$ -receptor blockade in phaeochromocytoma surgery: an observational case series. **Br J Anaesth.** 2017 Feb;118(2):182-189

## 2016

24. Michałowska I, Ćwikła JB, Michalski W, Wyrwicz LS, Prejbisz A, Szperl M, Nieć D, **Neumann HP**, Januszewicz A, Pęczkowska M. Growth rate of paragangliomas related to germline mutations of the *SDHx* genes. **Endocr Pract.** 2016 Dec 14. [Epub ahead of print]
25. Hoekstra AS, van den Ende B, Julià XP, van Breemen L, Scheurwater K, Tops CM, Malinoc A, Devilee P, **Neumann HP**, Bayley JP. Simple and rapid characterization of novel large germline deletions in *SDHB*, *SDHC* and *SDHD*-related paraganglioma. **Clin Genet.** 2016 Aug 3. Doi: 10.1111/cge.12843. [Epub ahead of print]
26. Suárez C, Fernández-Alvarez V, **Neumann HP**, Boedeker CC, Offergeld C, Rinaldo A, Stojan P, Ferlito A. Modern trends in the management of head and neck paragangliomas. **Eur Arch Otorhinolaryngol.** 2015 Dec;272(12):3595-9. No abstract available.
27. Castinetti F, Taieb D, Henry JF, Walz M, Guerin C, Brue T, Conte-Devolx B, **Neumann HP**, Sebag F. Management of endocrine disease: Outcome of adrenal sparing surgery in heritable pheochromocytoma.

**Eur J Endocrinol.** 2016 Jan;174(1):R9-18. doi: 10.1530/EJE-15-0549. Review.

## 2015

28. **Neumann HP**, de Herder W.  
Energy and metabolic alterations in predisposition to pheochromocytomas and paragangliomas: the so-called Warburg (and more) effect, 15 years on.  
**Endocr Relat Cancer.** 2015 Aug;22(4):E5-7. doi: 10.1530/ERC-15-0340. No abstract available.
  
29. Toledo RA, Maciel RM, Erlic Z, Lourenço DM Jr, Cerutti JM, Eng C, **Neumann HP**, Toledo SP.  
RET Y791F Variant Does Not Increase the Risk for Medullary Thyroid Carcinoma.  
**Thyroid.** 2015 Aug;25(8):973-4. doi: 10.1089/thy.2015.0168. No abstract available.
  
30. Gläser S, Neumann HPH, Koch CA, Vortmeyer AO.  
Von Hippel-Lindau Disease. In: De Groot LJ, Chrousos G, Dungan K, Feingold KR, Grossman A, Hershman JM, Koch C, Korbonits M, McLachlan R, New M, Purnell J, Rebar R, Singer F, Vinik A, editors.  
**Endotext** [Internet]. South Dartmouth (MA): MDText.com, Inc.; 2000-. 2015 Jul 11.
  
31. Luchetti A, Walsh D, Rodger F, Clark G, Martin T, Irving R, Sanna M, Yao M, Robledo M, **Neumann HP**, Woodward ER, Latif F, Abbs S, Martin H, Maher ER. Profiling of somatic mutations in pheochromocytoma and paraganglioma by targeted next generation sequencing analysis.  
**Int J Endocrinol.** 2015;2015:138573. doi: 10.1155/2015/138573.
  
32. Bausch B, Wellner U, Peyre M, Boedeker CC, Hes FJ, Anglani M, de Campos JM, Kanno H, Maher ER, Krauss T, Sansó G, Barontini M, Letizia C, Hader C, Schiavi F, Zanoletti E, Suárez C, Offergeld C, Malinoc A, Zschiedrich S, Gläser S, Bobin S, Sterkers O, Ba Huy PT, Giraud S, Links T, Eng C, Opocher G, Richard S, **Neumann HP**; International Endolymphatic Sac Tumor (ELST) Consortium.  
Characterization of endolymphatic sac tumors and von Hippel-Lindau disease in the International Endolymphatic Sac Tumor Registry.  
**Head Neck.** 2016 Apr;38 Suppl 1:E673-9. doi: 10.1002/hed.24067.
  
33. Michałowska I, Ćwikła JB, Pęczkowska M, Furmanek MI, Buscombe JR, Michalski W, Prejbisz A, Szperl M, Malinoc A, Moczulski D, Szutkowski Z, Kawecki A, Antoniewicz J, Pęczkowski P, Lewczuk A, Otto M, Cichocki A, Bednarek-Tupikowska G, Kabat M, Janaszek-Sitkowska H, Przybyłowska K, Janas J, **Neumann HP**, Januszewicz A.  
Usefulness of Somatostatin Receptor Scintigraphy (Tc-[HYNIC, Tyr3]-

Octreotide) and 123I-Metaiodobenzylguanidine Scintigraphy in Patients with SDHx Gene-Related Pheochromocytomas and Paragangliomas Detected by Computed Tomography.

**Neuroendocrinology**. 2015;101(4):321-30. doi: 10.1159/000381458.

34. von Dobschuetz E, Leijon H, Schalin-Jääntti C, Schiavi F, Brauckhoff M, Peczkowska M, Spiazzi G, Demattè S, Cecchini ME, Sartorato P, Krajewska J, Hasse-Lazar K, Roszkowska-Purska K, Taschin E, Malinoc A, Akslen LA, Arola J, Lange D, Fassina A, Pennelli G, Barbareschi M, Luettgies J, Prejbsiz A, Januszewicz A, Strate T, Bausch B, Castinetti F, Jarzab B, Opocher G, Eng C, **Neumann HP**.

A registry-based study of thyroid paraganglioma: histological and genetic characteristics.

**Endocr Relat Cancer**. 2015 Apr;22(2):191-204. doi: 10.1530/ERC-14-0558.

## 2014

35. Barski D, Ezziddin S, Heikaus S, **Neumann HP**.  
Diagnosis of extra-adrenal pheochromocytoma after nephrectomy.  
**Cent European J Urol**. 2014;67(2):162-6. doi: 10.5173/cej.2014.02.art9.
36. Bachurska S, Staykov D, Belovezhov V, Sasano H, Gulinac M, Stefanov C, **Neumann HP**.  
Bilateral pheochromocytoma/intra-adrenal paraganglioma in von Hippel-Lindau patient causing acute myocardial infarction.  
**Pol J Pathol**. 2014 Mar;65(1):78-82.
37. Castinetti F, Qi XP, Walz MK, Maia AL, Sanso G, Peczkowska M, Hasse-Lazar K, Links TP, Dvorakova S, Toledo RA, Mian C, Bugalho MJ, Wohlk N, Kollyukh O, Canu L, Loli P, Bergmann SR, Biarnes Costa J, Makay O, Patocs A, Pfeifer M, Shah NS, Cuny T, Brauckhoff M, Bausch B, von Dobschuetz E, Letizia C, Barczynski M, Alevizaki MK, Czetwertynska M, Ugurlu MU, Valk G, Plukker JTM, Sartorato P, Siqueira DR, Barontini M, Szperl M, Jarzab B, Verbeek HHG, Zelinka T, Vlcek P, Toledo SPA, Coutinho FL, Mannelli M, Recasens M, Demarquet L, Petramala L, Yaremchuk S, Zabolotnyi D, Schiavi F, Opocher G, Racz K, Januszewicz A, Weryha G, Henry JF, Brue T, Conte-Devolx B, Eng C, **Neumann HPH**  
Outcomes of adrenal sparing surgery or total adrenalectomy in pheochromocytoma associated with multiple endocrine neoplasia type 2: an international retrospective population based study  
**The Lancet Oncology**, accepted March 26, 2014
38. **Neumann HP**  
My life for pheochromocytoma  
**Endocr Relat Cancer** 2014 Jan 3 (Epub ahead of print)

39. Kugelberg J, Welander J, Schiavi F, Fassina A, Bäckdahl M, Larsson C, Opocher G, Söderkvist P, Dahia PL, **Neumann HP**, Gimm O  
Role of SDHAF2 and SDHD in von Hippel-Lindau associated pheochromocytomas  
**World J Surg** 2013 Dec 10 (Epub ahead of print)

## 2013

40. **Neumann HP**, Jilg C, Bacher J, Nabulsi Z, Malinoc A, Hummel B, Hoffmann MM, Ortiz-Bruechle N, Glasker S, Pisarski P, Neeff H, Krämer-Guth A, Cybulla M, Hornberger M, Wilpert J, Funk L, Baumert J, Paatz D, Baumann D, Lahl M, Felten H, Hausberg M, Zerres K, Eng C; for the Else-Kroener-Fresenius-ADPKD-Registry.  
Epidemiology of autosomal-dominant polycystic kidney disease: an in-depth clinical study for south-western Germany.  
**Nephrol Dial Transplant.** 2013 Jan 8. [Epub ahead of print]
41. Panizza E, Ercolino T, Mori L, Rapizzi E, Castellano M, Opocher G, Ferrero I, **Neumann HP**, Mannelli M, Goffrini P.  
Yeast model for evaluating the pathogenic significance of SDHB, SDHC and SDHD mutations in PHEO-PGL syndrome.  
**Hum Mol Genet.** 2013 Feb 15;22(4):804-15. doi: 10.1093/hmg/dds487. Epub 2012 Nov 21
42. Bausch B, Wellner U, Bausch D, Schiavi F, Barontini M, Sanso G, Walz MK, Peczkowska M, Weryha G, Dall'igna P, Cecchetto G, Bisogno G, Moeller L, Bockenbauer D, Patocs A, Racz K, Zabolotnyi D, Yaremchuk S, Dzivite-Krisane I, Castinetti F, Taieb D, Malinoc A, von Dobschuetz E, Roessler J, Schmid KW, Opocher G, Eng C, **Neumann HP**.  
Long term prognosis of patients with pediatric pheochromocytoma.  
**Endocr Relat Cancer.** 2013;21:17-25 Oct 29. [Epub ahead of print]
43. Därr R, Pamporaki C, Peitzsch M, Miehle K, Prejbisz A, Peczkowska M, Weismann D, Beuschlein F, Sinnott R, Bornstein SR, **Neumann HP**, Januszewicz A, Lenders J, Eisenhofer G.  
Biochemical diagnosis of phaeochromocytoma using plasma-free normetanephrine, metanephrine and methoxytyramine: importance of supine sampling under fasting conditions.  
**Clin Endocrinol (Oxf).** 2013 Sep 18. doi: 10.1111/cen.12327. [Epub ahead of print]
44. Cama A, Verginelli F, Lotti LV, Napolitano F, Morgano A, D'Orazio A, Vacca M, Perconti S, Pepe F, Romani F, Vitullo F, di Lella F, Visone R, Mannelli M, **Neumann HP**, Raiconi G, Paties C, Moschetta A, Tagliaferri R, Veronese A, Sanna M, Mariani-Costantini R.  
Integrative genetic, epigenetic and pathological analysis of paraganglioma



- reveals complex dysregulation of NOTCH signaling.  
**Acta Neuropathol.** 2013 Oct;126(4):575-94. doi: 10.1007/s00401-013-1165-y. Epub 2013 Aug 18.
45. Boedeker CC, Hensen EF, **Neumann HP**, Maier W, van Nederveen FH, Suárez C, Kunst HP, Rodrigo JP, P Takes R, Pellitteri PK, Rinaldo A, Ferlito A. Genetics of hereditary head and neck paragangliomas.  
**Head Neck.** 2013 Aug 3. doi: 10.1002/hed.23436. [Epub ahead of print]
46. Bausch B, Jilg C, Gläsker S, Vortmeyer A, Lützen N, Anton A, Eng C, **Neumann HP**. Renal cancer in von Hippel-Lindau disease and related syndromes.  
**Nat Rev Nephrol.** 2013 Sep;9(9):529-38. doi: 10.1038/nrneph.2013.144. Epub 2013 Jul 30.
47. Gläsker S, Schatlo B, Klingler JH, Braun V, Spangenberg P, Kim IS, van Velthoven V, Zentner J, **Neumann HP**. 6.Associations of Collagen Type I  $\alpha 2$  Polymorphisms with the Presence of Intracranial Aneurysms in Patients from Germany.  
**J Stroke Cerebrovasc Dis.** 2013 Jun 22. doi:pil: S1052-3057(13)00166-3. 10.1016/j.jstrokecerebrovasdis.2013.04.038. [Epub ahead of print]
48. Jilg CA, Drendel V, Bacher J, Pisarski P, Neeff H, Drognitz O, Schwardt M, Gläsker S, Malinoc A, Erlic Z, Nunez M, Weber A, Azurmendi P, Schultze-Seemann W, Werner M, **Neumann HP**. Autosomal dominant polycystic kidney disease: prevalence of renal neoplasias in surgical kidney specimens.  
**Nephron Clin Pract.** 2013;123(1-2):13-21. doi: 10.1159/000351049. Epub 2013 Jun 4.
49. Pęczkowska M, Kowalska A, Sygut J, Waligórski D, Malinoc A, Janaszek-Sitkowska H, Prejbisz A, Januszewicz A, **Neumann HP**. Testing new susceptibility genes in the cohort of apparently sporadic pheochromocytoma/paraganglioma patients with clinical characteristics of hereditary syndromes.  
**Clin Endocrinol (Oxf).** 2013 Apr 1. doi: 10.1111/cen.12218. [Epub ahead of print]
50. Klingler JH, Krüger MT, Lemke JR, Jilg C, Van Velthoven V, Zentner J, **Neumann HP**, Gläsker S. Sequence variations in the von Hippel-Lindau tumor suppressor gene in patients with intracranial aneurysms.  
**J Stroke Cerebrovasc Dis.** 2013 May;22(4):437-43. doi: 10.1016/j.jstrokecerebrovasdis.2013.01.016. Epub 2013 Feb 20.
51. Suárez C, Rodrigo JP, Mendenhall WM, Hamoir M, Silver CE, Grégoire V,

Strojan P, **Neumann HP**, Obholzer R, Offergeld C, Langendijk JA, Rinaldo A, Ferlito A.

Carotid body paragangliomas: a systematic study on management with surgery and radiotherapy.

**Eur Arch Otorhinolaryngol.** 2013 Feb 19. [Epub ahead of print]

52. Gläsker S, Krüger MT, Klingler JH, Wlodarski M, Klompen J, Schatlo B, Hippchen B, **Neumann HP**, Van Velthoven V.

Hemangioblastomas and neurogenic polyglobulia.

**Neurosurgery.** 2013 Jun;72(6):930-5; discussion 935. doi: 10.1227/NEU.0b013e31828ba793.

## 2012

53. Schiavi F, Demattè S, Cecchini ME, Taschin E, Bobisse S, Del Piano A, Donner D, Barbareschi M, Manera V, Zovato S, Erlic Z, Savvoukidis T, Barollo S, Grego F, Trabalzini F, Amistà P, Grandi C, Branz F, Marroni F, **Neumann HP**, Opocher G

The Endemic Paraganglioma Syndrome Type 1: Origin, Spread, and Clinical Expression.

**J Clin Endocrinol Metab.** 2012 2012 Apr;97(4):E637-41. doi: 10.1210/jc.2011-2597. Epub 2012 Mar 28

54. Jilg CA, **Neumann HP**, Gläsker S, Schäfer O, Leiber C, Ardelt PU, Schwardt M, Schultze-Seemann W.

Nephron sparing surgery in von Hippel-Lindau associated renal cell carcinoma; clinicopathological long-term follow-up.

**Fam Cancer** 2012;11:387-94. doi: 10.1007/s10689-012-9525-7.

55. **Neumann HP**, Bacher J, Nabulsi Z, Ortiz Brüchle N, Hoffmann MM, Schaeffner E, Nürnberger J, Cybulla M, Wilpert J, Riegler P, Corradini R, Kraemer-Guth A, Azurmendi P, Nunez M, Gläsker S, Zerres K, Jilg C.

Adult patients with sporadic polycystic kidney disease: the importance of screening for mutations in the PKD1 and PKD2 genes.

**Int Urol Nephrol** 2012;44:1753-62. doi: 10.1007/s11255-012-0125-0. Epub 2012 Feb 25.

56. Malinoc A, Sullivan M, Wiech T, Schmid KW, Jilg C, Straeter J, Deger S, Hoffmann MM, Bosse A, Rasp G, Eng C, **Neumann HP**.

Biallelic inactivation of the SDHC gene in renal carcinoma associated with paraganglioma syndrome type 3.

**Endocr Relat Cancer** 2012 May 3;19(3):283-90. doi: 10.1530/ERC-11-0324. Print 2012 Jun.

57. Hentschel M, Rottenburger C, **Neumann HP**, Brink I.

Is there an optimal scan time for 6-F-18-Fluoro-L-DOPA PET in

Paragangliomas?

**Clin Nucl Med** 2012 Feb;37(2):e24-9. doi: 10.1097/RLU.0b013e318238f550.

58. Bausch B, Malinoc A, Maruschke L, Offergeld C, Gläsker S, Rischke HC, Brauckhoff MM, Boedeker C, **Neumann HPH**.  
Genetik der Phäochromozytome.  
**Der Chirurg** 2012 Jun;83(6):511-8. doi: 10.1007/s00104-011-2191-8.
59. Offergeld C, Brase C, Yaremchuk S, Mader I, Rischke HC, Gläsker S, Schmid KW, Wiech T, Preuss SF, Suárez C, Kopec T, Patocs A, Wohlk N, Malekpour M, Boedeker CC, **Neumann HP**.  
Head and neck paragangliomas: clinical and molecular genetic classification.  
**Clinics** (Sao Paulo). 2012;67 Suppl 1:19-28.
60. Neeff HP, Pisarski P; Tittelbach-Helmrich D; Karajanev K, **Neumann HP**, Hopt UT, Drognitz O.  
One hundred consecutive kidney transplantations with simultaneous ipsilateral nephrectomy in patients with autosomal dominant polycystic kidney disease  
**Nephrol Dial Transpl** 2012 Oct 4. [Epub ahead of print]
61. Rischke HC, Benz MR, Wild D, Mix M, Dumont RA, Campbell D, Seufert J, Wiech T, Rössler J, Weber WA, **Neumann HP**  
Correlation of the genotype of paragangliomas and pheochromocytomas with their metabolic phenotype on <sup>18</sup>F-DOPA PET  
**J Nuclear Medicine** 2012 Sep;53(9):1352-8. doi:10.2967/jnumed.111.101303.  
Epub 2012 Jul 26.
62. Taïeb D, Sebag F, Sarde E, Berdah S, Doddoli C, Palazzo FF, Barlier A, **Neumann HP**, Mundler O  
First report of Harlequin Syndrome as the Presenting Feature of Carney Triad – a Diagnostic and Imaging challenge'  
**J Clin Oncol** 2012 Jul 10;30(20):e168-71. doi: 10.1200/JCO.2011.41.0159.  
Epub 2012 Jun 11.
63. **Neumann HP**, Bacher J, Nabulsi Z, Malinoc A, Ivanovas V, Hoffmann MM, Schaeffner E, Nürnberger J, Cybulla M, Wilpert J, Riegler P, Corradini R, Kraemer-Guth A, Azurmendi P, Nunez M, Gläsker S, Jilg C.  
Adult Patients with Sporadic Polycystic Kidney Disease – The Importance of Screening for Mutations in the *PKD1* and *PKD2* Genes  
**Int Urol Nephrol** 2012;44:1753-62. doi: 10.1007/s11255-012-0125-0. Epub 2012 Feb 25.
64. Schirpenbach C, Hoppert T, Aleksic I, **Neumann HP**, Hahner S, Fassnacht M, Allolio B.  
**Internist** (Berl). 2012;53:1119-24. doi: 10.1007/s00108-012-3109-3. German.

## 2011

65. Frank-Raue K, Rybicki LA, Erlic Z, Schweizer H, Winter A, Milos I, Toledo SP, Toledo RA, Tavares MR, Alevizaki M, Mian C, Siggelkow H, Hübner M, Wohlk N, Opocher G, Dvořáková S, Bendlova B, Czetwertynska M, Skasko E, Barontini M, Sanso G, Vorländer C, Maia AL, Patocs A, Links TP, de Groot JW, Kerstens MN, Valk GD, Miehle K, Musholt TJ, Biarnes J, Damjanovic S, Muresan M, Wüster C, Fassnacht M, Peczkowska M, Fauth C, Golcher H, Walter MA, Pichl J, Raue F, Eng C, **Neumann HP**; International RET Exon 10 Consortium.  
Risk profiles and penetrance estimations in multiple endocrine neoplasia type 2A caused by germline RET mutations located in exon 10.  
**Hum Mutat.** 2011;32:51-8.
66. Gläsker S, Shah MJ, Hippchen B, **Neumann HP**, Van Velthoven V.  
Doppler-Sonographically Guided Resection of CNS Hemangioblastomas.  
**Neurosurgery.** 2011 Feb 22. [Epub ahead of print]
67. Maher ER, **Neumann HP**, Richard S.  
von Hippel-Lindau disease: a clinical and scientific review.  
**Eur J Hum Genet.** 2011;19:617-23. Epub 2011 Mar 9.
68. Gonc N, Engiz O, **Neumann HP**, Demirbilek H, Ozon A, Alikasifoglu A, Kandemir N.  
Two pediatric patients with Von Hippel-Lindau disease type 2b: from patient to screening, from screening to patient.  
**J Pediatr Endocrinol Metab.** 2011;24(1-2):109-12.
69. Bennett KL, Campbell R, Ganapathi S, Zhou M, Rini B, Ganapathi R, **Neumann HP**, Eng C.  
Germline and somatic DNA methylation and epigenetic regulation of KILLIN in renal cell carcinoma.  
**Genes Chromosomes Cancer.** 2011 Aug;50(8):654-61
70. **Neumann HP**, Sullivan M, Winter A, Malinoc A, Hoffmann MM, Boedeker CC, Bertz H, Walz MK, Moeller LC, Schmid KW, Eng C.  
Germline mutations of the TMEM127 gene in patients with paraganglioma of head and neck and extraadrenal abdominal sites.  
**J Clin Endocrinol Metab.** 2011 Aug;96(8):E1279-82.
71. **Sullivan M**, Rybicki LA, Winter A, Hoffmann MM, Reiermann S, Linke H, Arbeiter K, Patzer L, Budde K, Hoppe B, Zeier M, Lhotta K, Bock A, Wiech T, Gaspert A, Fehr T, Woznowski M, Berisha G, Malinoc A, Goek ON, Eng C, **Neumann HP**.  
Age-related penetrance of hereditary atypical hemolytic uremic syndrome.  
**Ann Hum Genet.** 2011 Nov;75(6):639-47.

72. Alesina PF, Hinrichs J, Meier B, Schmid KW, **Neumann HP**, Walz MK. Minimally invasive cortical-sparing surgery for bilateral pheochromocytomas. **Langenbecks Arch Surg**. 2011 Sep 21. [Epub ahead of print]
73. Dudziak K, Rettig I, Adam P, Horger M, **Neumann HP**, Müssig K. Rare cause of insufficient metabolic control of diabetes mellitus - Case 10/2011. **Dtsch Med Wochenschr**. 2011 Oct;136(43):2196.
74. Poeppel TD, Yuece A, Boy C, Metz KA, Kaminsky E, **Neumann HP**, Rosenbaum SJ, Mann K, Moeller LC. Novel SDHD Gene Mutation (H102R) in a Patient With Metastatic Cervical Paraganglioma Effectively Treated by Peptide Receptor Radionuclide Therapy. **J Clin Oncol** 2011 Nov 20;29(33):e812-5
75. Jilg CA, **Neumann HP**, Gläsker S, Schäfer O, Ardelt PU, Schwardt M, Schultze-Seemann W. Growth Kinetics in Von Hippel-Lindau-Associated Renal Cell Carcinoma. **Urol Int**. 2011 Dec 9. [Epub ahead of print]

## 2010

76. Bhakdi SC, Ottinger A, Somsri S, Sratogno P, Pannadaporn P, Chimma P, Malasit P, Pattanapanyasat K, **Neumann HPH**. Optimized high gradient magnetic separation for isolation of Plasmodium-infected red blood cells. **Malaria J**. 2010 Feb 2;9(1):38. [Epub ahead of print]
77. Sullivan M, Erlic Z, Hoffmann MM, Arbeiter K, Patzer L, Budde K, Hoppe B, Zeier M, Lhotta K, Rybicki LA, Bock A, Berisha G, **Neumann HPH**. Epidemiological approach to identifying genetic predispositions for atypical hemolytic uremic syndrome. **Ann Hum Genet**. 2010 Jan;74(1):17-26.
78. Otto EA, Hurd TW, Airik R, Chaki M, Zhou W, Stoetzel C, Patil SB, Levy S, Ghosh AK, Murga-Zamalloa CA, van Reeuwijk J, Letteboer SJ, Sang L, Giles RH, Liu Q, Coene KL, Estrada-Cuzcano A, Collin RW, McLaughlin HM, Held S, Kasanuki JM, Ramaswami G, Conte J, Lopez I, Washburn J, Macdonald J, Hu J, Yamashita Y, Maher ER, Guay-Woodford LM, **Neumann HP**, Obermüller N, Koenekoop RK, Bergmann C, Bei X, Lewis RA, Katsanis N, Lopes V, Williams DS, Lyons RH, Dang CV, Brito DA, Dias MB, Zhang X, Cavalcoli JD, Nürnberg G, Nürnberg P, Pierce EA, Jackson PK, Antignac C, Saunier S, Roepman R, Dollfus H, Khanna H, Hildebrandt F. Candidate exome capture identifies mutation of SDCCAG8 as the cause of a retinal-renal ciliopathy. **Nat Genet**. 2010 Sep 12.

79. Erlic Z, Hoffmann MM, Sullivan M, Franke G, Peczkowska M, Harsch I, Schott M, Gabbert HE, Valimäki M, Preuss SF, Hasse-Lazar K, Waligorski D, Robledo M, Januszewicz A, Eng C, **Neumann HP**.  
Pathogenicity of DNA variants and double mutations in multiple endocrine neoplasia type 2 and von Hippel-Lindau syndrome.  
**J Clin Endocrinol Metab**. 2010 Jan;95(1):308-13.
80. Erlic Z, Ploeckinger U, Cascón A, Hoffmann MM, von Duecker L, Winter A, Kammel G, Bacher J, Sullivan M, Isermann B, Fischer L, Raffel A, Knoefel WT, Schott M, Baumann T, Schaefer O, Keck T, Baum RP, Milos I, Muresan M, Peczkowska M, Januszewicz A, Cupisti K, Tönjes A, Fasshauer M, Langrehr J, von Wussow P, Agaimy A, Schlimok G, Lamberts R, Wiech T, Schmid KW, Weber A, Nunez M, Robledo M, Eng C, **Neumann HP**  
Systematic comparison of sporadic and syndromic pancreatic islet cell tumors.  
**Endocr Relat Cancer**. 2010 Jul 26. [Epub ahead of print]
81. Wohllk N, Schweizer H, Erlic Z, Schmid KW, Walz MK, Raue F, **Neumann HP**.  
Multiple endocrine neoplasia type 2.  
**Best Pract Res Clin Endocrinol Metab**. 2010 Jun;24(3):371-87
82. Papewalis C, Kouatchoua C, Ehlers M, Jacobs B, Porwol D, Schinner S, Willenberg HS, Anlauf M, Raffel A, Eisenhofer G, **Neumann HP**, Bornstein SR, Scherbaum WA, Schott M.  
Chromogranin A as potential target for immunotherapy of malignant pheochromocytoma.  
**Mol Cell Endocrinol**. 2010 Jun 22. [Epub ahead of print]
83. von Duecker L, Walz MK, Voss C, Arnold G, Eng C, **Neumann HP**.  
Laparoscopic Organ-Sparing Resection of Von Hippel-Lindau Disease-Associated Pancreatic Neuroendocrine Tumors.  
**World J Surg**. 2010 Dec 23. [Epub ahead of print]
84. Schweizer H, Boehm J, Winterer JT, Wild D, **Neumann HP**, Wiech T, Stubanus M, Kuehn EW.  
Pheochromocytoma and thrombotic microangiopathy: favourable outcome despite advanced renal failure.  
**J Clin Pathol**. 2010 Aug;63(8):754-6. No abstract available
85. Gläsker S, Klingler JH, Müller K, Würtenberger C, Hader C, Zentner J, **Neumann HP**, Velthoven VV.  
Essentials and Pitfalls in the Treatment of CNS Hemangioblastomas and von Hippel-Lindau Disease.  
**Cen Eur Neurosurg**. 2010 Mar 12

2009

86. Erlic Z, **Neumann HPH**.  
Diagnosing patients with hereditary paraganglial tumours.  
**Lancet Oncol.** 2009 Aug;10(8):741.
87. Mehta A, Beck M, Elliott P, Giugliani R, Linhart A, Sunder-Plassmann G, Schiffmann R, Barbey F, Ries M, Clarke JT; Fabry Outcome Survey investigators (with **Neumann HP**).  
Enzyme replacement therapy with agalsidase alfa in patients with Fabry's disease: an analysis of registry data.  
**Lancet.** 2009 Dec 12;374(9706):1986-96. doi: 10.1016/S0140-6736(09)61493-8. PMID: 19959221
88. Franke G, Bausch B, Hoffmann MM, Cybulla M, Wilhelm C, Kohlhase J, Scherer G, **Neumann HPH**.  
Alu-Alu recombination underlies the vast majority of large VHL germline deletions: Molecular characterization and genotype-phenotype correlations in VHL patients.  
**Hum Mutat.** 2009 May;30(5):776-86.
89. Franke G, Scherer G, **Neumann HPH**.  
Response to: Extremely low risk of pheochromocytomas in complete VHL gene deletion cases  
**Hum Mutat.** 2009 Jul 20 Letter
90. **Neumann HPH**, Erlic Z, Boedeker CC, Rybicki LA, Robledo M, Hermsen M, Schiavi F, Falcioni M, Kwok P, Bauters C, Lampe K, Fischer M, Edelman E, Benn DE, Robinson BG, Wiegand S, Rasp G, Stuck BA, Hoffmann MM, Sullivan M, Sevilla MA, Weiss MM, Peczkowska M, Kubaszek A, Pigny P, Ward RL, Learoyd D, Croxson M, Zabolotny D, Yaremchuk S, Draf W, Muresan M, Lorenz RR, Knipping S, Strohm M, Dyckhoff G, Matthias C, Reisch N, Preuss SF, Esser D, Walter MA, Kaftan H, Stöver T, Fottner C, Gorgulla H, Malekpour M, Zarandy MM, Schipper J, Brase C, Glien A, Kühnemund M, Koscielny S, Schwerdtfeger P, Välimäki M, Szyfter W, Finckh U, Zerres K, Cascon A, Opocher G, Ridder GJ, Januszewicz A, Suarez C, Eng C.  
Clinical predictors for germline mutations in head and neck paraganglioma patients: cost reduction strategy in genetic diagnostic process as fall-out  
**Cancer Res.** 2009 Apr 15;69(8):3650-6.
91. Erlic Z, Rybicki L, Peczkowska M, Golcher H, Kann PH, Brauckhoff M, Müssig K, Muresan M, Schäffler A, Reisch N, Schott M, Fassnacht M, Opocher G, Klose S, Fottner C, Forrer F, Plöckinger U, Petersenn S, Zabolotny D, Kollukch O, Yaremchuk S, Januszewicz A, Walz MK, Eng C, **Neumann HPH** for the European-American Pheochromocytoma Study Group  
Clinical Predictors and Algorithm for the Genetic Diagnosis of Pheochromocytoma Patients

**Clin Cancer Res.** 2009 Oct 15;15(20):6378-85. Epub 2009 Oct 13

92. Boedeker CC, Erlic Z, Richard S, Kontny U, Gimenez-Roqueplo AP, Cascon A, Robledo M, de Campos JM, van Nederveen FH, de Krijger RR, Burnichon N, Gaal J, Walter MA, Reschke K, Wiech T, Weber J, Rückauer K, Plouin PF, Darrouzet V, Giraud S, Eng C, **Neumann HPH**.  
Head and neck paragangliomas in von Hippel-Lindau disease and multiple endocrine neoplasia type 2  
**J Clin Endocrinol Metab.** 2009 Jun;94(6):1938-44.
93. Erlic Z, Hoffmann MM, Sullivan M, Franke G, Peczkowska M, Harsch I, Schott M, Gabbert HE, Valimäki M, Preuss SF, Hasse-Lazar K, Waligorski D, Robledo M, Januszewicz A, Eng C, **Neumann HPH**  
Pathogenicity of DNA Variants and Double Mutations in Multiple Endocrine Neoplasia Type 2 and Von Hippel-Lindau Syndrome.  
**J Clin Endocrinol Metab.** 2009 Nov 11, Epub ahead to print.
94. **Neumann HPH**, Eng C.  
The approach to the patient with paraganglioma.  
**J Clin Endocrinol Metab.** 2009 Aug;94(8):2677-83.
95. Erlic Z, **Neumann HPH**.  
When should genetic testing be obtained in a patient with pheochromocytoma or paraganglioma?  
**Clin Endocrinol (Oxf).** 2009 Mar;70(3):354-7.
96. Gkaliagkousi E, Erlic Z, Petidis K, Semertzidis P, Doumas M, Zamboulis C, **Neumann HPH**, Douma S.  
Neurofibromatosis type 1: should we screen for other genetic syndromes? A case report of co-existence with multiple endocrine neoplasia 2A  
**Eur J Clin Invest** Epub 2009 Jun 25.
97. Boedeker CC, **Neumann HPH**, Offergeld C, Maier W, Falcioni M, Berlis A, Schipper J.  
Clinical features of paraganglioma syndromes. **Skull Base.** 2009 Jan;19(1):17-25.
98. Schipper J, Spetzger U, Tatagiba M, Rosahl S, **Neumann HPH**, Boedeker CC, Maier W.  
Juxtacondylar approach in temporal paraganglioma surgery: when and why?  
**Skull Base.** 2009 Jan;19(1):43-7.
99. Reisch N, Walz MK, Erlic Z, **Neumann HPH**.  
Pheochromocytoma – still a challenge.  
**Internist (Berl).** 2009 Jan;50(1):27-35.



100. Kreusel KM, Krause L, Graul-Neumann L, Bechrakis NE, **Neumann HPH**, Foerster MH.  
Family screening in patients with retinal angiomatosis  
**Klin Monbl Augenheilkd.** 2009 Nov;226(11):939-43
101. Gaal J, van Nederveen FH, Erlic Z, Korpershoek E, Oldenburg R, Boedeker CC, Kontny U, **Neumann HPH**, Dinjens WN, de Krijger RR.  
Parasympathetic Paragangliomas are Part of the von Hippel-Lindau Syndrome  
**J Clin Endocrinol Metab** 2009;94: 4367–4371
102. Romaker D, Puetz M, Teschner S, Donauer J, Geyer M, Gerke P, Rumberger B, Dworniczak B, Pennekamp P, Buchholz B, **Neumann HPH**, Kumar R, Gloy J, Eckardt KU, Walz G.  
Increased expression of secreted frizzled-related protein 4 in polycystic kidneys.  
**J Am Soc Nephrol** 2009 Jan;20(1):48-56.

## 2008

103. **Neumann HPH**, Erlic Z  
Maternal Transmission of Symptomatic Disease with SDHD Mutation: Fact or Fiction?  
**J Clin Endocrinol Metab.** 2008;93:1573-5
104. Peczkowska M, Cascon A, Prejbisz A, Kubaszek A, Cwikła BJ, Furmanek M, Erlic Z, Eng C, Januszewicz A, **Neumann HPH**  
Extra-adrenal and adrenal pheochromocytomas associated with a germline SDHC mutation.  
**Nat Clin Pract Endocrinol Metab** 2008;4:111-5
105. Cascon A, Landa Í, López-Jiménez E, Díez-Hernández, A, Buchta M, Montero-Conde C, Leskelä S, Leandro-García LJ, Letón R, Rodríguez-Antona C, Eng C, **Neumann HPH**, Robledo M  
Molecular characterisation of a common SDHB deletion in paraganglioma patients.  
**J Med Genet.** 2008;45;233-238
106. Margetts CD, Morris M, Astuti D, Gentle DC, Cascon A, McRonald FE, Catchpoole D, Robledo M, **Neumann HPH**, Latif F, Maher ER  
Evaluation of a functional epigenetic approach to identify promoter region methylation in phaeochromocytoma and neuroblastoma.  
**Endocr Relat Cancer.** 2008 Sep;15(3):777-786. Epub 2008 May 22
107. Peczkowska M, Erlic Z, Hoffmann MM, Furmanek M, Cwikła J, Kubaszek A, Prejbisz A, Szutkowski Z, Kawecki A, Chojnowski K, Lewczuk A, Litwin M, Szyfter W, Walter M, Sullivan M, Eng C, Januszewicz A, **Neumann HPH**

Impact of Screening Kindreds for SDHD p.Cys11X as a Common Mutation Associated with Paraganglioma Syndrome Type 1

**J Clin Endocrinol Metab.** 2008 Dec;93(12):4818-25. Epub 2008 Sep 30

108. Romaker D, Puetz M, Teschner S, Donauer J, Geyer M, Gerke P, Rumberger B, Dworniczak B, Pennekamp P, Buchholz B, **Neumann HPH**, Kumar R, Gloy J, Eckardt KU, Walz G  
Increased Expression of Secreted Frizzled-Related Protein 4 in Polycystic Kidneys.  
**J Am Soc Nephrol.** 2009 Jan;20(1):48-56. Epub 2008 Oct 22
109. Hoffmann B, Beck M, Rolfs A, **Neumann HPH**  
Fabry disease - complex clinical picture, simple diagnosis procedure, causal treatment.  
**Dtsch Med Wochenschr.** 2008 Sep;133 (39):1965-72; quiz 1973-4
110. Milos IN, Frank-Raue K, Wohllk N, Maia AL, Pusiol E, Patocs A, Robledo M, Biarnes J, Barontini M, Links TP, de Groot JW, Dvorakova S, Peczkowska M, Rybicki LA, Sullivan M, Raue F, Zosin I, Eng C, **Neumann HPH**  
Age-related neoplastic risk profiles and penetrance estimations in multiple endocrine neoplasia type 2A caused by germ line RET Cys634Trp (TGC>TGG) mutation.  
**Endocr Relat Cancer** 2008 Dec;15(4):1035-1041. Epub 2008 Sep 15.
111. Erlic Z, **Neumann HPH**  
Clinical question: When should genetic testing be obtained in a patient with pheochromocytoma or paraganglioma?  
**Clin Endocrinol (Oxf).** 2008 epub ahead
112. Bockenbauer D, Rees L, **Neumann HPH**, Foo Y  
A sporadic case of paraganglioma undetected by urine metabolite screening  
**Pediatr Nephrol** 2008 Oct;23(10):1889-91. Epub 2008 May 6
113. Cotesta D, Erlic Z, Petramala L, Verrienti A, Cavallaro G, Giustini S, Divona L, Polistena A, Ciardi A, D'Erasmo E, De Toma G, Calvieri S, **Neumann HPH**, Filetti S, Letizia C  
Coincidence of neurofibromatosis type 1 and multiple endocrine neoplasia type 2 (MEN 2)  
**The Endocrinologist** 2008;18(6):277-281

## 2007

114. **Neumann HPH**, Vortmeyer A, Schmidt D, Werner M, Erlic Z, Cascon A, Bausch B, Januszewicz A, Eng C.  
Evidence of MEN-2 in the original description of classic pheochromocytoma.  
**The New England Journal of Medicine.** 2007;357:1311-5

115. Boedeker CC, **Neumann HPH**, Maier W, Bausch B, Schipper J, Ridder GJ. Malignant head and neck paragangliomas in SDHB mutation carriers. **Otolaryngol Head Neck Surg.** 2007;137:126-19
116. Cybulla M, Walter K, **Neumann HPH**, Widmer U, Schärer M, Sunder-Plassmann G, Jansen T, Rolfs A, Beck M  
Morbus Fabry: Demographische Übersicht aus dem deutschsprachigen Raum seit Einführung der Enzyersatztherapie (EET)  
**Dtsch Med Wschr** 2007;132:1505-9
117. Bausch B, Borozdin W, Mautner VF, Hoffmann MM, Boehm D, Robledo M, Cascon A, Harenberg T, Schiavi F, Pawlu C, Peczkowska M, Letizia C, Calvieri S, Arnaldi G, Klingenberg-Noftz RD, Reisch N, Fassina A, Brunaud L, Walter MA, Mannelli M, MacGregor G, Palazzo FF, Barontini M, Walz MK, Kremens B, Brabant G, Pfäffle R, Koschker AC, Lohofner F, Mohaupt M, Gimm O, Jarzab B, McWhinney SR, Opocher G, Januszewicz A, Kohlhase J, Eng C, **Neumann HPH**; European-American Pheochromocytoma Registry Study Group. Germline NF1 mutational spectra and loss-of-heterozygosity analyses in patients with pheochromocytoma and neurofibromatosis type 1. **J Clin Endocrinol Metab.** 2007 Jul;92(7):2784-92.
118. Cybulla M, **Neumann HPH**  
Morbus Fabry – Eine interdisziplinäre Herausforderung  
**Dtsch Med Wschr** 2007;133:71-7
119. Cascon A, Landa I, Lopez-Jimenez E, Díez-Hernández A, Buchta M, Montero-Conde C, Leskelä S, Leandro-García LJ, Letón R, Rodríguez-Antona C, Eng C, **Neumann HPH**, Robledo M  
Molecular characterization of a common SDHB deletion in paraganglioma patients.  
**J Med Genet** 2007[Epub ahead of print].
120. Boehm D, Bacher J, **Neumann HPH**  
Gross genomic rearrangement involving the TSC2-PKD1 contiguous deletion syndrome: characterization of the deletion event by quantitative polymerase chain reaction.  
**Am J Kidney Dis** 2007: 49:e11-21
121. Bhakdi SC, Sratongno P, Chimma P, Rungruang T, Chuncharunee A, **Neumann HPH**, Malasit P, Pattanapanyasat K.  
Re-evaluating acridine orange for rapid flow cytometric enumeration of parasitemia in malaria-infected rodents.  
**Cytometry A.** 2007;71:662-7.
122. Bender BU, Quaschnig T, **Neumann HPH**, Schmidt D, Kraemer-Guth A.

A novel frameshift mutation of the lecithin:cholesterol acyltransferase (LCAT) gene associated with renal failure in familial LCAT deficiency.

**Clin Chem Lab Med.** 2007;45:483-6

123. **Neumann HPH**, Cybulla M, Gläsker S, Coulin C, Van Velthoven V, Berlis A, Hader C, Schäfer O, Treier M, Brink I, Schultze-Seemann W, Leiber C, Rückauer K, Junker B, Agostini FJ, Hetzel A, Boedecker CC.  
Von Hippel-Lindau Erkrankung. Interdisziplinäre Patientenversorgung  
**Ophthalmologe.** 2007;104:119-26
124. Kreusel KM, Bechrakis NE, **Neumann HPH**, Foerster MH  
Juxtapapillary capillary retinal angioma with epiretinal membrane of the macula in familial Von-Hippel-Lindau-Syndrome.  
**Ophthalmologe** 2007;317-20.
125. Kreusel KM, Bechrakis NE, **Neumann HPH**, Schmidt D, Foerster MH.  
Solitary juxtapapillary capillary retinal angioma and von Hippel-Lindau disease.  
**Can J Ophthalmol.** 2007;42:251-5.
126. Langrehr JM, Bahra M, Kristiansen G, **Neumann HPH**, Neumann LM, Plöckinger U, Lopez-Hänninen E.  
Neuroendocrine tumor of the pancreas and bilateral adrenal pheochromocytomas. A rare manifestation of von Hippel-Lindau disease in childhood.  
**J Pediatr Surg.** 2007;42:1291-4
127. Peczkowska M, Januszewicz A, Jarzab B, **Neumann HPH**, Kubaszek A, Janaszek-Sitkowitzka H, Litwin M, Antoniewicz J, Aksamit-Bialoszewska E, Roslonowska E, Prejbisz A, Januszewicz M, Michalowska I, Ciwla J, Furmanek M, Walecki J  
Pheochromocytoma in children and adolescents based on the Polish Pheochromocytoma registry  
**Ann Diagn Paed Pathol** 2007;11:15-20
128. Brink I, Hentschel M, **Neumann HPH**, Schäfer O, Moser E  
FDOPA-PET als Paradigma molekularer Bildgebung in der Onkologie  
**Der Nuklearmediziner** 2007;30:70-82

## 2006

129. Bausch B, Borozdin W, **Neumann HPH** and the European-American Pheochromocytoma Study working Group.  
Clinical and genetic characteristics of patients with neurofibromatosis type 1 and pheochromocytoma.  
**The New England Journal of Medicine** 2006;354(25): 2729-31

130. Fliegau M, Horvath J, Schnakenburg von C, Olbrich H, Müller D, Thumfart J, Schermer B, Pazour GJ, **Neumann HPH**, Zentgraf H, Benzing T and Omran H  
Nephrocystin Specifically Localizes to the Transition Zone of Renal and Respiratory Cilia and Photoreceptor Connecting Cilia.  
**The Journal of the American Society of Nephrology** 2006;17: 2424 - 2433.
131. Bausch B, Koschker AC, Fassnacht M, Stoevesandt J, Hoffmann MM, Eng C, Allolio B and **Neumann HPH**  
Comprehensive mutation scanning of NF1 in apparently sporadic cases of pheochromocytoma.  
**The Journal of Clinical Endocrinology & Metabolism** 2006;91(9): 3478-81
132. Reisch N, Peczkowska M, Januszewicz A, **Neumann HPH**  
Pheochromocytoma: Presentation, diagnosis and treatment  
**The Journal of Hypertension** 2006;24(12): 2331-2339
133. Müller AMS, Geibel A, **Neumann HPH**, Kühnemund A, Schmitt-Gräff A, Böhm J, Engelhardt M.  
Primary (AL) Amyloidosis in Plasma Cell Disorders.  
**The Oncologist** 2006;11: 824 – 830.
134. Cybulla M, Kleber M, Walter KN, Kroeber SM, **Neumann HPH**, Engelhardt M  
Is Fabry associated with leukemia?  
**The British Journal of Haematology** 2006;135: 264-275
135. Hering A, Guratowska M, Bucsky P, Claussen U, Decker J, Ernst G, Hoepfner W, Michel S, **Neumann HPH**, Parlowsky T, Loncarevic I.  
Characteristic genomic imbalances in pediatric pheochromocytoma.  
**Genes, Chromosomes & Cancer** 2006;45(6): 602-7
136. Kreusel KM, Bechrakis NE, Krause L, **Neumann HPH**, Foerster MH  
Retinal angiomas in von Hippel-Lindau disease: a longitudinal ophthalmologic study.  
**Ophthalmology** 2006;113(8): 1418-24
137. Gimenez-Roqueplo AP, Lehnert H, Mannelli M, **Neumann HPH**, Opocher G, Maher ER, Plouin PF  
Pheochromocytoma, new genes and screening strategies.  
**Clinical Endocrinology** 2006;65(6):699-705
138. Walz MK, Alesina PF, Wenger FA, Deligiannis A, Szuczik E, Petersenn S, Ommer A, Groeben H, Peitgen K, Janssen OE, Philipp T, **Neumann HPH**, Schmid KW, Mann K  
Posterior retroperitoneoscopic adrenalectomy – results of 560 procedures in 520 patients-  
**Surgery** 2006;140(6): 943-950

139. Kreusel KM, Bechrakis NE, **Neumann HPH**, Foerster MH  
Pars plana vitrectomy for juxtapapillary capillary retinal angioma.  
**American Journal of Ophthalmology** 2006;141(3): 587-9
140. Koch CA, Brouwers FM, Vortmeyer AO, Tannapfel A, Libutti SK, Zhuang Z, Pacak K, **Neumann HPH**, Paschke P  
Somatic VHL gene alterations in MEN2-associated medullary thyroid carcinoma. **BMC Cancer** 2006;6: 131
141. Bausch B, Boedeker CC, Berlis A, Brink I, Cybulla M, Walz MK, Januszewicz A, Opocher G, Eng C, **Neumann HPH**  
Genetic and Clinical Investigation of Pheochromocytoma: A 22-year experience, from Freiburg, Germany to International Effort.  
**Annals of the New York Academy of Sciences** 2006;1073: 112-121
142. Brink I, Schaefer O, Walz M, **Neumann HPH**  
Fluorine-18 DOPA PET imaging of paraganglioma syndrome.  
**Clinical Nuclear Medicine** 2006;31(1): 39-41
143. Walz MK, Alesina PF, Wenger FA, Koch JA, **Neumann HPH**, Petersenn S, Schmid KW, Mann K  
Laparoscopic and Retroperitoneoscopic Treatment of Pheochromocytomas and Retroperitoneal Paragangliomas: Results of 161 Tumors in 126 Patients.  
**World Journal of Surgery** 2006;30: 1-10
144. Fuentes C, Menendez E, Pineda J, Martinez De Esteban JP, Anda E, Goni MJ, Bausch B, **Neumann HPH**  
The malignant potential of a succinate dehydrogenase subunit B germline mutation. **Journal of Endocrinological Investigation** 2006;29(4): 350-2

## 2005

145. Schiavi F, Boedeker CC, Bausch B, Peczkowska M, Gomez CF, Strassburg T, Pawlu C, Buchta M, Salzmann M, Hoffmann MM, Berlis A, Brink I, Cybulla M, Muresan M, Walter MA, Forrer F, Valimaki M, Kawecki A, Szutkowski Z, Schipper J, Walz MK, Pigny P, Bauters C, Willet-Brozick JE, Baysal BE, Januszewicz A, Eng C, Opocher G, **Neumann HPH** for the European-American Paraganglioma Study Group.  
Predictors and prevalence of paraganglioma syndrome associated with mutations of the *SDHC* gene.  
**JAMA** 2005;294(16):2057-63
146. Margetts CD, Astuti D, Gentle DC, Cooper WN, Cascon A, Catchpoole D, Robledo M, **Neumann HPH**, Latif F, Maher ER.  
Epigenetic analysis of HIC1, CASP8, FLIP, TSP1, DCR1, DCR2, DR4, DR5,

- KvDMR1, H19 and preferential 11p15.5 maternal-allele loss in von Hippel-Lindau and sporadic pheochromocytomas.  
**Endocrine-Related Cancer** 2005;12(1):161-72
147. Cybulla M, Schaefer E, Wendt S, Ling H, Krober SM, Hovelborn U, Schandelmaier S, Rohrbach R, **Neumann HPH**  
Chronic renal failure and proteinuria in adulthood: Fabry disease predominantly affecting the kidneys.  
**American Journal of Kidney Diseases** 2005;45(5):e82-9
148. Walz MK, Petersenn S, Koch JA, Mann K, **Neumann HPH**, Schmid KW.  
Endoscopic treatment of large primary adrenal tumours.  
**British Journal of Surgery** 2005;92(6):719-23)
149. Nambirajan T, Leeb K, **Neumann HPH**, Graubner UB, Janetschek G.  
Laparoscopic adrenal surgery for recurrent tumours in patients with hereditary pheochromocytoma.  
**European Urology** 2005;47(5):622-6
150. Boedeker CC, **Neumann HPH**, Ridder GJ, Maier W, Schipper J  
Paragangliomas in patients with mutations of the SDHD gene.  
**Otolaryngology - Head & Neck Surgery** 2005;132(3):467-70
151. Joerger M, Koeberle D, **Neumann HPH**, Gillessen S  
Von Hippel-Lindau disease--a rare disease important to recognize.  
**Onkologie** 2005;28(3):159-63
152. Schmidt D, **Neumann HPH**  
Spontaneous Regression of Retinal Angiomatous Lesions in v. Hippel-Lindau Disease (VHL).  
**European Journal of Medical Research** 2005;10(12):532-4 (
153. Pawlu C, Bausch B, Reisch N, **Neumann HPH**  
Genetic testing for pheochromocytoma-associated syndromes.  
**Annales d Endocrinologie** 2005;66(3):178-85
154. **Neumann HPH**, Cybulla M, Shibata H, Oya M, Naruse M, Higashihara E, Terachi T, Ling H, Takami H, Shuin T, Murai M.  
New genetic causes of pheochromocytoma: current concepts and the clinical relevance.  
**Keio Journal of Medicine** 2005;54(1):15-21
155. Pawlu C, Bausch B, **Neumann HPH**  
Mutations of the SDHB and SDHD genes.  
**Familial Cancer** 2005;4(1):49-54

## 2004

156. Leverkus M, **Neumann HPH**, Bröcker EB, Hamm H, Mayer J  
Soft Nodules at the Tip of the Tongue of a 26-year-old Man: Mucosal Neuromas in Multiple Endocrine Neoplasia (MEN) type 2 B  
**Arch Dermatol** 2004;139:1647-1652
157. Astuti D, Morris M, Krona C, Abel F, Gentle D, Martinsson T, Kogner P, **Neumann HP**, Voutilainen R, Eng C, Rustin P, Latif F, Maher ER.  
Investigation of the role of SDHB inactivation in sporadic pheochromocytoma and neuroblastoma.  
**Br J Cancer**. 2004 Nov 15;91(10):1835-41.
158. **Neumann HPH**, Pawlu C, Peczkowska M, Bausch B, McWhinney SR, Muresan M, Buchta M, Franke G, Klisch J, Bley TA, Hoegerle S, Boedeker CC, Opocher G, Schipper J, Januszewicz A, Eng C for the European-American Paraganglioma Study Group.  
Distinct clinical features of paraganglioma syndromes associated with SDHB and SDHD gene mutations.  
**JAMA** 2004;292:943-51 Erratum in: **JAMA**. 2004;13;292:1686
159. McWhinney SR, Buchta M, Vanharanta S, Pezkowska M, Morrison CD, Virta SK, Lehtonen R, Järvinen H, Juhola M, Mecklin JP, Herva R, Nupponen NN, Januszewicz A, Aaltonen LA, **Neumann HPH**, Eng C  
Early onset renal cell carcinoma as novel extra-paraganglial component of SDHB-associated hereditary paraganglioma  
**Am J Hum Genet** 2004;74:153-159
160. Hoefele J, Otto E, Felten H, Kühn K, Bley TA, Zäuner I, Hildebrandt F, **Neumann HPH**  
Clinical and Histological Presentation of Three Siblings with Mutations in the NPHP4 Gene  
**Am J Kidney Dis** 2004;43:358-364
161. Birkenfeld A, Bergmann M, Bräsen JH, Luft FC, **Neumann HPH**  
A Paraganglioma Parasitizing the Left Circumflex Coronary Artery  
**Am J Med** 2004;116:787-788
162. Walz MK, Peitgen K, Diesing D, Petersen S, Janssen OE, Philipp T, Metz KA, Mann K, Schmid KW, **Neumann HPH**  
Partial versus Total Adrenalectomy by the Posterior Retroperitoneoscopic Approach - Early and Long-term Results of 325 Consecutive Procedures in Primary Adrenal Neoplasias  
**World J Surg** 2004;28:1323-9
163. Astuti D, Morris M, Krona C, Abel F, Gentle D, Martinsson T, Kogner P,



- Neumann HPH**, Voutilainen R, Eng C, Farida L, Maher ER  
Epigenetic Inactivation of SDHB by Promoter Region Hypermethylation in Pheochromocytoma and Neuroblastoma  
**Brit J Cancer** 2004;15:1835-41
164. Eisenhofer G, Bornstein SR, Brouwers FM, Cheung NKV, Dahia PL, de Krijger RR, Giordano TJ, Greene LA, Goldstein DS, Lehnert H, Manger WM, Maris JM, **Neumann HPH**, Pacak K, Shulkin BL, Smith DI, Tischler AS, Young WF  
Malignant pheochromocytoma: Current status and Initiatives for Future Progress  
**Endocrine Related Cancer** 2004;11:423-36
165. Allibhai Z, Rodrigues G, Brecevic E, **Neumann HPH**, Winqvist E.  
Malignant pheochromocytoma associated with germline mutation of the SDHB gene.  
**J Urol** 2004;172:1409-10
166. Vanharanta S, Buchta M, McWhinney SR, Virta SK, Peczkowska M, Morrison CD, Lehtonen R, Januszewicz A, Jarvinen H, Juhola M, Mecklin JP, Pukkala E, Herva R, Kiuru M, Nupponen NN, Aaltonen LA, **Neumann HPH**, Eng C.  
Early-onset renal cell carcinoma as a novel extraparaganglial component of SDHB-associated heritable.  
**Am J Hum Genet.** 2004;74:153-9.
167. Beck O, Fassbender WJ, Beyer P, Kriener S, **Neumann HPH**, Klingebiel T, Lehrnbecher T  
Pheochromocytoma in Childhood - Implication for Further Diagnostic Procedures  
**Acta Paediatr** 2004;93:1630-1634
168. Gimm O, Koch CA, Januszewicz A, Opocher G, **Neumann HPH**  
The genetic basis of pheochromocytoma.  
**Horm Res** 2004;31:45-60
169. Ling H, Cybulla M, Schaefer O, Arnold C, Schories M, **Neumann HPH**  
When to look for Von Hippel-Lindau disease in gastroenteropancreatic neuroendocrine tumors?  
**Neuroendocrinology** 2004;80 Suppl 1:39-46

## 2003

170. Hoegerle S, Ghanem N, Althoefler C, Schipper J, Brink I, Moser E, **Neumann HPH**  
18F DOPA positron emission tomography for detection of glomus tumors: comparison to MRI  
**Eur J Nucl Med** 2003;30:689-694

171. Walz MK, **Neumann HPH**, Peitgen K, Petersenn S, Janssen OE, Mann K  
Endoscopic Treatment of Recurrent Pheochromocytomas and  
Retroperitoneal Paragangliomas  
**World J Surg** 2003;35:93-96
172. Manuelian T, Hellwege J, Seppo Meri S, Jessica Caprioli J, Marina Noris M,  
**Neumann HPH**, Remuzzi G, Zipfel PF  
Factor H Gene Mutations in Atypical Hemolytic Uremic Syndrome Affect  
Protein Stability and Function - Single Amino Acids Mutations of Factor H in  
Hemolytic Uremic Syndrome Affect Binding to C3b, to Heparin and Surface  
Attachment  
**J Clin Invest** 2003;111:1181-1190
173. Arnold S, Strecker R, Scheffler K, Spreer J, Schipper J, **Neumann HPH**, Klisch  
J  
Dynamic Contrast Enhancement of Paragangliomas of the Head and Neck:  
Evaluation with Time-Resolved 2D MR Projection Angiography  
**Eur Radiol** 2003;13:1608-1611
174. Kreusel KM, Bechrakis NE, **Neumann HPH**, Foerster MH  
A sporadic case of von Hippel-Lindau disease with a secondary maculopathy  
as the presenting sign  
**Acta Ophthalm Scand** 2003;81:309-310
175. **Neumann HPH**, Salzmann M, Bohnert-Iwan B, Mannuelian T, Skerka C, Lenk  
D, Bender BU, Cybulla M, Riegler P, Königsrainer A, Neyer U, Bock A, Widmer  
U, Male DA, Franke G, Zipfel PF  
Hemolytic Uremic Syndrome and Mutations of the Factor H Gene. A Registry –  
based Study of German Speaking Countries  
**J Med Genet** 2003;40:676-681
176. Berlis, Schumacher M, Spreer J, **Neumann HPH**, van Velthoven V  
Subarachnoid haemorrhage due to cervical spinal cord haemangioblastomas in  
a patient with von Hippel-Lindau disease  
**Acta Neurochir** 2003;145:1009-1013
177. McWhinney SR, Boru G, Binkley PK, Januszewicz A, **Neumann HPH**, Eng C  
Intronic Single Nucleotide Polymorphisms in the RET Proto-oncogene are  
Associated with a Subset of Apparently Sporadic Pheochromocytoma and May  
Modulate Age of Onset  
**J Clin Endocrinol Metab** 2003;88:4911-4916
178. Wolf MTF, Mucha BE, Attanasio M, Zalewski I, Karle SM, **Neumann HPH**,  
Rahmen N, Bader B, Baldamus CA, Otto E, Witzgall R, Fuchshuber A,  
Hildebrandt F

Mutations of the Uromodulin gene in MCKD type 2 patients cluster in exon 4 which encodes three EGF-like domains  
**Kidney International** 2003;64:1580-1587

Velthoven van V, Reinacher PC, Klisch J, **Neumann HPH**, Gläsker S  
 Treatment of intramedullary hemangioblastomas, with special attention to von Hippel-Lindau disease  
**Neurosurgery** 2003;53:1306-1314

## 2002

179. Högerle S, Nitzsche E, Althöfer C, Ghanem N, Manz T, Brink I, Reincke M, Moser E, **Neumann HPH**  
<sup>18</sup>Fluoro-DOPA whole-body positron emission tomography for detection of pheochromocytomas: initial results.  
**Radiology** 2002;22:507-512

180. Reichardt P, Apel TW, Domula M, Tröbs RB, Krause I, Bierbach U, **Neumann HPH**, Kiess W  
 Recurrent polytopic chromaffin paragangliomas in a 9 year old boy due to a novel germline mutation in the von Hippel-Lindau gene  
**Journal of Pediatric Hematology/Oncology** 2002;24:145-148

181. **Neumann HPH**, Högerle S, Manz T, Brenner K, Iliopoulos O  
 How Many Pathways to Pheochromocytoma?  
**Seminars in Nephrology**, 2002;22:89-99

182. Lui WO, Chen JD, Gläsker S, Bender BU, Larrson C, **Neumann HPH**, Teh BT  
 VHL-related pheochromocytoma: Selection of chromosome 11p Loss  
**Oncogene** 2002;21:1117-1122

183. Zipfel PF, **Neumann HPH**  
 Komplement Faktor H Mutation führen zur Ausbildung der atypischen Form des Hämolytischuräemischen Syndroms  
**Nieren und Hochdruckkrankheiten** 2002;31:172-179

184. **Neumann HPH**, Bausch B, McWhinney SR, Bender BU, Gimm O, Franke G, Schipper J, Klisch J, Althoefer C, Zerres K, Januszewicz A, Smith WM, Munk R, Manz T, Glaesker S, Apel TW, Treier M, Reineke M, Walz MK, Hoang-Vu C, Brauckhoff M, Klein-Franke A, Klose P, Schmidt H, Maier-Woelfle M, Peçzkowska M, Szmigielski C, Eng C for the Freiburg–Warsaw–Columbus Pheochromocytoma Study Group  
 Germ-Line Mutations in Nonsyndromic Pheochromocytoma  
**The New England Journal of Medicine**, 2002;346:1459-1466

185. **Neumann HPH**, Schipper J, Eng C

Germ-Line Mutations in Nonsyndromic Pheochromocytoma - Correspondence  
**The New England Journal of Medicine**, 2002;347:854-855

186. Peczkowska M, Gessek J, Januszewicz A, **Neumann HPH**, Janaszek-Sitkowska H, Kabat M, Januszenwicz M, Skierski J, Prejbisz A, Ciesla W, Szostek M  
 Pheochromocytoma of the bladder  
**Blood Pressure**, 2002;11:101-105
187. Walz MK, Peitgen K, **Neumann HPH**, Philipp T, Mann K  
 Endoscopic treatment of solitary, bilateral, multiple and recurrent pheochromocytomas and paragangliomas  
**World Journal of Surgery**, 2002;26:1005-1012
188. **Neumann HPH**  
 Imaging vs biochemical testing for pheochromocytoma – letter / correspondence  
**The Journal of the American Medical Association (JAMA)** 2002;288:314-315
189. **Neumann HPH**, Högerle S, Manz T, Brenner K, Iliopoulos O  
 How Many Pathways to Pheochromocytoma?  
**Seminars in Nephrology**, 2002;22:89-99
190. Zipfel PF, **Neumann HPH**  
 Komplement Faktor H Mutationen führen zur Ausbildung der atypischen Form des hämolytisch-urämischen Syndroms  
**Nieren- und Hochdruckerkrankungen**, 2002;31:172-179
- 2001**
191. Janetschek G, **Neumann HPH**  
 Laparoscopic surgery for Pheochromocytoma  
**Urologic Clinics of North America** 2001;28:97-105
192. **Neumann HPH**, Schipper J, Eng C  
 Mutations in *SDHD*, a mitochondrial complex II gene, in pheochromocytomas  
**Cancer Research Alert** 2001;10:107-109
193. **Neumann HPH**, Riegler P, Huber W, Corradini R, Sessa A, Fontana D, Wetterauer U, Janetschek G  
 The challenge of kidney lesions in Von Hippel-Lindau disease  
**Contributions to Nephrology** 2001;136:193-207
194. Zipfel PF, Skerka C, Caprioli J, Manuelian T, **Neumann HPH**, Noris M, Remuzzi G

- Complement factor H and hemolytic uremic syndrome  
**International Immunopharmacology** 2001;1:461-468
195. **Neumann HPH**, Hildebrandt F, Weber M, Zerres K  
Hereditäre Nephropathie (Editorial, Gastedition: Neumann HPH)  
**Nieren- und Hochdruckkrankheiten** 2001;30:241
196. **Neumann HPH**, Schulenburg S, Apel TW  
Familiäre Nierentumoren im Erwachsenenalter  
**Nieren- und Hochdruckkrankheiten** 2001;30:267-277
197. Abel K-B, Apel TW, Beck M, **Neumann HPH**  
Morbus Fabry  
**Nieren- und Hochdruckkrankheiten** 2001;30:261-266
198. Zipfel PF, Skerka C, Munk RD, **Neumann HPH**  
Immunregulator Faktor H und Hämolytisch-Urämisches Syndrom  
**Nieren- und Hochdruckkrankheiten** 2001;7:291-297
199. Gläsker S, Bender BU, Apel TW, v. Velthoven V, Mulligan LM, Zentner J,  
**Neumann HPH**  
Reconsidering of biallelic inactivation of the VHL tumour suppressor gene in  
haemangioblastomas of the central nervous system  
**Journal of Neurology, Neurosurgery & Psychiatry** 2001;70:644-648
200. Chatha RK, Johnson AM, Rothberg PG, Townsend RR, **Neumann HPH**, Gabow  
PA  
Von Hippel – Lindau disease masquerading as autosomal dominant polycystic  
kidney disease  
**American Journal of Kidney Diseases** 2001;37:852-858
201. Manz T, Grotz W, Orszagh M, Volk B, Kirste G, **Neumann HPH**  
A patient with neurological deficits and seizures after renal transplantation  
**Nephrology Dialysis Transplantation**, 2001;16:631-633
202. Frenzel S, Dörr HG, Apel TW, Heidemann PH, Zerres K, **Neumann HPH**  
Pheochromocytoma associated with a de novo VHL mutation as form fruste of  
the Von Hippel-Lindau disease  
**European Journal of Pediatrics** 2001;160:421-424
203. Bender BU, Eng C, Olschewski M, Berger DP, Laubenberger J, Althöfer C,  
Kirste G, Orzagh M, van Velthoven V, Mioszcza H, Schmidt D, **Neumann HPH**  
No increased mortality from germline *VHL* c.505 T>C mutation: the necessity of  
*VHL* mutation-specific assessment  
**Journal of Medical Genetics**, 2001;38:508-514

204. **Neumann HPH**, Reincke M, Eng C  
Genetic Testing in Young Patients with Apparently Isolated Pheochromocytoma  
Letter to the editor:Case 13-2001  
**The New England Journal of Medicine**, 2001;345:547
205. Astuti D, Agathangleo A, Honorio S, Dallol A, Martinsson T, Kogner P, Cummins C, **Neumann HPH**, Voutilainen R, Dahia P, Maher ER, Latif F  
RASSF1A promoter region CpG island hypermethylation in pheochromocytomas and neuroblastoma tumors  
**Oncogene** 2001;20:7573-7577

## 2000

206. Riegler P, Huber W, Corradini R, **Neumann HPH**, Gläsker S, Sessa A  
Von Hippel-Lindau disease: the role of gene analysis in affected families. A case report  
**Nephron** 2000;84:95-97
207. Schmidt D, Natt E, **Neumann HPH**  
Long-term results of Laser treatment for retinal angiomatosis in Von Hippel-Lindau disease  
**European Journal of Medical Research** 2000;5:47-58
208. Rouers JFM, Apel T, **Neumann HPH**, v. Arnim U, Lips CJM, Höppener JWM  
Internally shortened menin protein as a consequence of alternative RNA splicing due to a germline deletion in the multiple endocrine neoplasia type 1 gene  
**International Journal of Molecular Medicine** 2000;5:611-614
209. Kreusel K-M, Bechrakis NE, Heinichen T, Neumann L, **Neumann HPH**, Foerster MH  
Retinal angiomatosis and von Hippel-Lindau disease  
**Clinical Investigation** 2000;238:916-921
210. Januszewicz A, **Neumann HPH**, Lon I, Szmigielski C, Symnuides B, Kabat M, Apel TW, Wocial B, Lapinski M, Januszewicz W  
Incidence and clinical relevance of RET proto-oncogene germline mutations in pheochromocytoma patients  
**Journal of Hypertension** 2000;18:1019-1023
211. Radmayr, **Neumann HPH**, Bartsch G, Janetschek G  
Laparoscopic Partial Adrenalectomy for Bilateral Pheochromocytomas in a Boy with von Hippel Lindau disease  
**European Urology** 2000; 38: 344-8
212. Gimm O, Armanios M, Dziema H, **Neumann HPH**, Eng C

Somatic and occult germline mutations in SDHD, a mitochondrial complex II gene, in non-familial pheochromocytoma  
**Cancer Research** 2000;60:6822-6825

213. Bender BU, Gutsche M, Gläsker S, Müller B, Kirste G, **Neumann HPH**  
 Genetic alterations in von Hippel-Lindau syndrome associated and sporadic pheochromocytomas  
**Journal of Clinical Endocrinology & Metabolism** 2000;85:4568-4574
214. Zerres K, Eggermann T, Hildebrandt F, Konrad M, Fuchshuber A, **Neumann HPH**, Zimmerhackl B, Rudnik-Schöneborn S  
 Erbliche Nierenkrankheiten – eine Übersicht  
**Medizinische Genetik** 2000; 12:163-169
215. **Neumann HPH**, Munk RD, Manuelian T, Zipfel PF  
 Familiäres hämolytisch-urämisches Syndrom  
**Medizinische Genetik** 2000; 12:203-206
216. Schulenburg S, Apel TW, **Neumann HPH**  
 Familiäre Nierentumoren im Erwachsenenalter  
**Medizinische Genetik** 2000; 12:239-245
- 1999**
217. **Neumann HPH**, Bender BU, Reincke M, Eggstein S, Laubenberger J, Kirste G  
 Adrenal sparing surgery for pheochromocytoma  
**British Journal of Surgery** 1999;84:94-97
218. **Neumann HPH**, Hildebrandt F  
 Genetic mechanisms of renal disease (Congress report)  
**Kidney and Blood Pressure Research** 1999;22:172-174
219. **Neumann HPH**  
 The spectrum of renal cysts in adulthood  
**Nephrology Dialysis Transplantation** 1999;14:2234-2244
220. Hemberger, Himmelbauer H, **Neumann HPH**, Plate KH, Schwarzkopf G, Fundele R  
 Expression of the Von Hippel-Lindau binding protein-1 (*Vbp1*) in fetal and adult mouse tissues  
**Human Molecular Genetics** 1999;8:229-236
221. **Neumann HPH**, Krumme B, van Velthoven V, Orszagh M, Zerres K  
 Multiple intracranial aneurysms in a patient with autosomal-recessive polycystic kidney disease

**Nephrology Dialysis Transplantation** 1999;14:936-939

222. Kreusel KM, Bornfeld N, Bender BU, Neumann L, Foerster MH, **Neumann HPH**  
Kapilläres retinales Angiom. Klinische und molekulargenetische Untersuchungen  
**Der Ophthalmologe** 1999;96:71-76
223. Frenzel S, **Neumann HPH**, Hümmer PH, Dörr HG  
Biadrenales Phäochromozytom bei einem Jungen  
**Monatsschrift für Kinderheilkunde** 1999;147:477-479
224. **Neumann HPH**, Reincke M, Bender BU, Elsner R, Janetschek G  
Preserved adrenocortical function after laparoscopic bilateral adrenal sparing  
surgery for hereditary pheochromocytoma  
**The Journal of Clinical Endocrinology & Metabolism** 1999;84:2608-2610
225. Schmidt L, Junker K, Nakaigawa N, Kinjerski T, Weirich G, Miller M, Lubensky I,  
**Neumann HPH**, Brauch H, Decker J, Bocke C, Brown JA, Jenkins R, Richard S,  
Bergerheim U, Gerrard B, Dean M, Linehan WM, Zbar B  
Novel mutation of the MET protooncogene in papillary renal carcinomas  
**Oncogene** 1999;18:2343-2350
226. Gläsker S, Bender BU, Apel TW, Natt E, van Velthoven V, Scheremet R, Zentner  
J, **Neumann HPH**  
The impact of molecular genetic analysis of the VHL gene in patients with  
haemangioblastomas of the central nervous system  
**The Journal of Neurology, Neurosurgery and Psychiatry** 1999;67:758-762

**1998**

227. Kempermann G, **Neumann HPH**  
Endolymphatic sac tumors  
**Histopathology** 1998;33:2-10
228. Bender BU, **Neumann HPH**  
Molekularpathologie des Klarzellkarzinoms der Niere  
**Spektrum der Nephrologie** 1998;11:3-10
229. **Neumann HPH**, Schwarzkopf G, Henske EP  
Renal angiomyolipomas, cysts, and cancer in tuberous sclerosis complex  
**Seminars of Pediatric Neurology** 1998;5:269-275
230. Goldfarb DA, **Neumann HPH**, Penn I, Novick AC  
Results of renal transplantation in patients with renal cell carcinoma in Von  
Hippel-Lindau disease  
**Transplantation** 1998;64:1726-1729



231. **Neumann HPH** and Bender BU  
Genotype-phenotype correlations in Von Hippel-Lindau disease  
**The Journal of Internal Medicine** 1998;243:541-545
232. Janetschek G, Finkenstedt G, Gasser R, Waibel UG, Peschel R, Bartsch G, **Neumann HPH**  
Laparoscopic surgery for pheochromocytoma: adrenalectomy, partial resection, excision of paragangliomas  
**The Journal of Urology** 1998;160:330-334
233. **Neumann HPH**, Bender BU, Berger DP, Laubenberger J, Schultze-Seemann W, Wetterauer U, Ferstl FJ, Herbst EW, Schwarzkopf G, Hes FJ, Lips CJM, Lamiell JM, Masek O, Riegler P, Glavac D, Brauch H  
Prevalence, morphology and biology of renal cell carcinoma in Von Hippel-Lindau disease compared to sporadic renal cell carcinoma  
**The Journal of Urology** 1998;160:1248-1254
234. Henske EP, Ao X, Short P, Greenberg R, **Neumann HPH**, Kwiatkowski DJ, Russo I  
Frequent progesterone receptor immunoreactivity in tuberous sclerosis-associated renal angiomyolipomas  
**Modern Pathology** 1998;11:665-668
235. Bender BU, Altehöfer C, Hasse H, **Neumann HPH**  
The hypertensive lady with elevated plasma norepinephrine concentration and no demonstrable tumour - the search for pheochromocytoma  
**Nephrology Dialysis Transplantation** 1998;13:1295-1296
236. Watnick TJ, Gandolph MA, Weber H, **Neumann HPH**, Germino GG  
Gene conversion is a likely cause of mutation in PKD1  
**Human Molecular Genetics** 1998;7:1239-1243
237. Schmidt L, Junker K, Weirich G, Glenn G, Choyke P, Lubensky I, Z. Zhuang, Jeffers M, Vande Woude G, **Neumann HPH**, Walther M, Lineham WM, Zbar B  
Two North American families with hereditary papillary renal carcinoma and identical novel mutations in the MET proto-oncogene  
**Cancer Research** 1998;58:1719-1722
238. Martin P, Heiskari N, Zhou J, Leinonen A, Tumelis T, Hertz JM, Barker D, Gregory M, Atkin C, **Neumann HPH**, Springate J, Shows T, Petterson E, Tryggvason K  
High mutation detection rate in the COL4A5 collagen gene in suspected Alport syndrome using PCR and direct DNA sequencing  
**The Journal of the American Society of Nephrology** 1998;9:2291-2301

1997

239. Kopf D, Steinert H, Bockisch A, Beyer J, Hensen J, **Neumann HPH**, Lehnert H  
Octreotide scintigraphy does not predict catecholamine response to octreotide in malignant pheochromocytoma  
**Clinical Endocrinology** 1997;46:39-44
240. Hildebrandt F, Strahm B, Nothwang H-G, Gretz N, Schnieders B, Singh-Sawhney I, Kutt R, Vollmer M, Brandis M, and members of the APN Study Group (incl. **Hartmut P. H. Neumann**)  
Molecular genetic identification of families with juvenile nephronophthisis type 1: Rate of progression to renal failure  
**Kidney International** 1997;51:261-269
241. **Neumann HPH**, Zäuner I, Strahm B, Bender BU, Schollmeyer P, Blum U, Rohrbach R, Hildebrandt F  
Late occurrence of cysts in autosomal dominant medullary cystic kidney disease  
**Nephrology Dialysis Transplantation** 1997;12:1242-1246
242. Bender BU, Wetterauer U, Schollmeyer P, **Neumann HPH**  
An incidental finding - bilateral multifocal renal oncocytoma  
**Nephrology Dialysis Transplantation** 1997;12:1034-1036
243. Bender BU, Altehöfer C, Januszewicz A, Gärtner R, Schmidt H, Hoffmann MM, Heidemann PH, **Neumann HPH**  
Functioning thoracic paraganglioma - association with Von Hippel-Lindau syndrome  
**The Journal of Clinical Endocrinology & Metabolism** 1997;82:3356-3360
244. Marsh DJ, Zheng Z, Arnold A, Andrew SC, Learoyd D, Frilling A, Komminoth P, **Neumann HPH**, Ponder BAJ, Rollins BJ, Shapiro GI, Robinson BG, Mulligan LM and Eng C  
Mutation analysis of glial cell line-derived neurotrophic factor (GDNF), a ligand for the RET/co-receptor complex, in MEN 2 and sporadic neuroendocrine tumors  
**The Journal of Clinical Endocrinology & Metabolism** 1997;82:3025-3028
245. Watnick TJ, Piontek KB, Cordal TM, Weber H, Gandolph MA, Quian F, Lens XM, **Neumann HPH**, Germino GG  
An unusual pattern of mutation in the duplicated portion of *PKD1* is revealed by use of a novel strategy for mutation detection  
**Human Molecular Genetics** 1997;6:1473-1481
246. **Neumann HPH** and Zbar B  
Renal cysts, renal cancer and Von Hippel-Lindau disease  
**Kidney International**, 1997;51:16-26

1996

247. **Neumann HPH**, Bender B, Zäuner I, Berger DP, Eng C, Brauch H, Zbar B  
Monogenetic hypertension and pheochromocytoma  
**American Journal of Kidney Diseases** 1996;28:229-233
248. Kempermann G, **Neumann HPH**, Scheremet R, Volk B, Mann W, Gilsbach J, Laszig R  
Deafness due to bilateral endolymphatic sac tumor in a case of Von Hippel-Lindau syndrome  
**The Journal of Neurology, Neurosurgery and Psychiatry** 1996;61:318-320
249. Glavac D, **Neumann HPH**, Wittke C, Jaenig H, Rödl H, Masek O, Streicher T, Pausch F, Engelhardt D, Plate K, Höfler H, Chen F, Zbar B, Brauch H  
Mutations in the *VHL* tumor suppressor gene and associated lesions in families with Von Hippel-Lindau disease from Central Europe  
**Human Genetics** 1996;98:271-280
250. **Neumann HPH**, Eng C, Mulligan LM  
Von Hippel-Lindau disease and pheochromocytoma (letter)  
**Journal of the American Medical Association (JAMA)** 1996;275:839-840
251. Heiskari N, Zhang X, Zhou J, Leinonen A, Barker D, Gregory M, Atkin C, Netzer K-O, Weber M, Reeders S, Grönhagen-Riska C, **Neumann HPH**, Trembath R, Tryggvason K  
Identification of 17 mutations in 10 exons in the Col4A5 collagen gene, but no mutation s found in 4 exons in Col4A6: a study of 250 patients with Alport syndrome  
**The Journal of the American Society of Nephrology (JASN)** 1996;7:702-709
252. Anding K, Köhler G, Böhm N, Petersen KG, Schollmeyer P, **Neumann HPH**  
Primäre pigmentierte noduläre adrenocorticale Dysplasie, eine seltene Ursache des Cushing Syndroms  
**Deutsche Medizinische Wochenschrift** 1996;121:1321-1324
253. Zbar B, Kishida T, Chen F, Maher ER, Richards FM, Crossey PA, Webster A, Affara NA, Ferguson-Smith MA, Brauch H, **Neumann HPH**, Tisherman S, Mulvihill JJ, Gross D, Shuin T, Whaley J, Seizinger B, Kley N, Olschwang S, Boisson C, Richard S, Lips CJM, Linehan WM, Lerman M  
Germline mutations in the Von Hippel-Lindau disease (VHL) gene in families from North America, Europe, and Japan  
**Human Mutation** 1996;8:8348-8357

## 1995

254. **Neumann HPH**, Brügggen V, Berger DP, Herbst EW, Blum U, Morgenroth A, Schollmeyer P, Wetterauer U

- Tuberous Sclerosis Complex with Endstage Renal Failure  
**Nephrology Dialysis Transplantation** 1995;10:349-353
255. Eng C, Mulligan LM, Smith DP, Healey CS, Frilling A, Raue F, **Neumann HPH**, Pfragner R, Behmel A, Lorenzo MJ, Stonehouse TJ, Ponder MA, Ponder BAJ  
Mutation of the *RET* proto-oncogene in sporadic medullary thyroid carcinoma  
**Genes, Chromosomes and Cancer** 1995;12:209-212
256. Schmidt D, **Neumann HPH**  
Retinal vascular hamartoma in Von Hippel-Lindau disease  
**Archives of Ophthalmology** 1995;113:1163-1167
257. Henske EP, **Neumann HPH**, Scheithauer BW, Herbst EW, Kwiatkowski DJ  
Loss of heterozygosity in the TSC2 region of chromosome 16p13 in sporadic angioliomas  
**Genes Chromosomes and Cancer** 1995;13:295-298
258. Eng C, Mulligan LM, Smith DP, Healey CS, Frilling A, Raue F, **Neumann HPH**, Ponder MA, Ponder BAJ  
Low frequency of germline mutations in the *RET* proto-oncogene in patients with apparently sporadic medullary thyroid carcinoma  
**Clinical Endocrinology** 1995;43:123-127
259. **Neumann HPH**, Eng C, Mulligan LM, Glavac D, Zäuner I, Ponder BAJ, Crossey PA, Maher PR, Brauch H  
Consequences of direct genetic testing for germline mutations in the clinical management of families with multiple endocrine neoplasia type 2  
**The Journal of the American Medical Association (JAMA)** 1995;274:1149-1151
260. **Neumann HPH**, Brauch H  
Molekulare Diagnose des Von Hippel-Lindau Syndroms (letter)  
**Deutsche Medizinische Wochenschrift** 1995;120:1416
261. Brauch H, Kishida T, Glavac D, Chen F, Pausch F, Höfler H, Latif F, Lerman MI, Zbar B, **Neumann HPH**  
Von Hippel-Lindau (VHL) disease with pheochromocytoma in the Black Forest region of Germany: evidence for a founder effect  
**Human Genetics** 1995;95:551-556
262. **Neumann HPH**  
Von Hippel-Lindau syndrome - a rare syndrome as the clue for the molecular basis for common renal disorders (Editorial)  
**Nephrology Dialysis Transplantation** 1995;10:1498-1499
263. **Neumann HPH**, Lips CJM, Hsia YE, Zbar B

Inherited neoplastic syndromes involving the nervous system: Von Hippel-Lindau syndrome

**Brain Pathology** 1995;5:181-193

#### 1994

264. **Neumann HPH**

Von Hippel-Lindau Syndrome (teaching point)

**Nephrology Dialysis and Transplantation** 1994;9:313-315

265. **Neumann HPH**

Postoperative flank pains after surgery of pheochromocytoma (teaching point)

**Nephrology Dialysis and Transplantation** 1994;9:721-722

266. **Neumann HPH**

Rupturgefahr bei Nierenzysten (Korrespondenz)

**Deutsche Medizinische Wochenschrift** 1994;119:1486

267. **Neumann HPH**

Genetics of Hypertension: The Pheochromocytoma Model (invited lecture)

**Clinical Investigator** 1994;72:729-730

268. **Neumann HPH** and Wiestler OD

Von Hippel-Lindau disease: A syndrome providing insights into growth control and tumorigenesis (Congress report)

**Nephrology Dialysis Transplantation** 1994;9:1832-1833

269. Mulligan LM, Eng C, Healey CS, Clayton D, Kwok JBJ, Gardner E, Ponder MA, Frilling A, Jackson CE, Lehnert H, **Neumann HPH**, Thibodeau SN, Ponder BAJ  
Specific Mutations of the RET proto-oncogene are related to disease phenotype in MEN 2A and FMTC

**Nature Genetics** 1994;6:70-74

270. Bathmann J, **Neumann HPH**, Sigmund G, Moser E

False positive diagnosis of a pheochromocytoma with I-123 Metaiodobenzyl-guanidine

**Clinical Nuclear Medicine** 1994;19:221-223

271. **Neumann HPH**

Pheochromocytomas, multiple endocrine neoplasia type 2, and Von Hippel-Lindau syndrome. Correspondence

**The New England Journal of Medicine** 1994;330:1091-1092

272. Zerres K, Mücher G, Bachner L, Deschennes G, Eggermann T, Kääriäinen H, Knapp M, Lennert T, Misselwitz J, von Mühlendahl KE, **Neumann HPH**, Pirson Y, Rudnik-Schöneborn S, Steinbicker V, Wirth B & Schärer K

Mapping of the Gene for Autosomal Recessive Polycystic Kidney Disease (ARPKD) to Chromosome 6p21-cen  
**Nature Genetics** 1994;7:429-432

273. **Neumann HPH**  
 Tuberos Sclerosis  
**The New England Journal of Medicine** (letter) 1994;331:813-814
274. Whaley JM, Naglich J, Gelbert L, Hsia YE, Lamiell JM, Green JS, Collins D, **Neumann HPH**, Laidlaw J, Li FP, KLein-Szanto AJP, Seizinger BR, Kley N  
 Germline mutations in the von Von Hippel-Lindau tumor suppressor gene are similar to somatic Von Hippel-Lindau aberrations in sporadic renal cell carcinoma  
**The American Journal of Human Genetics** 1994;55:1092-1102
275. Decker H-J, Klauck SM, Lawrence JB, McNeil J, Smith D, Gemmill RM, Sandberg AA, **Neumann HPH**, Simon B, Green J, Seizinger BR  
 Cytogenetic and fluorescence in-situ hybridization studies on sporadic and hereditary tumors associated with Von Hippel-Lindau syndrome  
**Cancer Genetics and Cytogenetics** 1994;77:1-13

### 1993

276. Raue F, Kotzerke J, Reinwein D, Schröder S, Frilling A, Deckart H, Höfer R, Ritter M, Seif F, Buhr H, Beyer J, Schober O, Becker W, **Neumann HPH**, Calvi J, Winter J, Vogt H, and the German Medullary Thyroid Carcinoma Study Group  
 Prognostic factors in medullary thyroid carcinoma: Evaluation of 741 patients from the german medullary thyroid carcinoma register  
**Clinical Investigator** 1993;71:7-12
277. **Neumann HPH**  
 Rudolf Kaltenbach - Zum 150. Geburtstag und 100. Todestag.  
**Geburtshilfe und Frauenheilkunde** 1993;53:204-211
278. **Neumann HPH**  
 Arvid Lindau zum 100. Geburtstag  
**Der Pathologe** 1993;14:178-180
279. **Neumann HPH**, Coester A, Zäuner I, Kanz L, Würtemberger G, Schollmeyer P, Blum U, Wetterauer U  
 Echinococcosis of the kidney  
**Nephrology Dialysis Transplantation** 1993;8:757-760
280. Berger DP, Zäuner I, Mihatsch MJ, Herb H-M, **Neumann HPH**  
 Ungewöhnlicher Verlauf einer Antibasalmembran-Antikörper-Nephritis  
**Medizinische Klinik** 1993;88:478-481

281. **Neumann HPH**, Berger DP, Sigmund G, Blum U, Parmer RJ, D. Schmidt, Volk B, Kirste G  
Pheochromocytomas, multiple endocrine neoplasia type 2, and Von Hippel-Lindau syndrome  
**The New England Journal of Medicine** 1993;329:1351-1358
282. **Neumann HPH**, Zäuner I  
Der Komplex der Nachsorge beim C-Zell-Karzinom  
**Deutsche Medizinische Wochenschrift** 1993;118:25-29
283. **Neumann HPH**  
Von Hippel-Lindau Syndrom - Unterschätzt und häufig verkannt  
**Deutsches Ärzteblatt** 1993;90:786-793
284. **Neumann HPH**  
Das Von Hippel-Lindau Syndrom  
**Der Pathologe** 1993;14:150-157
285. **Neumann HPH**, Kandt RS  
Klinik und Genetik der Tuberösen Sklerose  
**Deutsche Medizinische Wochenschrift** 1993;118:1577-1583
286. **Neumann HPH**  
Hereditäre Nephropathien  
**Aktuelle Nephrologie** 1993
- 1992**
287. **Neumann HPH**, Schumacher M  
Das Von Hippel-Lindau Syndrom aus neuroradiologischer Sicht  
**Klinische Neuroradiologie** 1992;2:203-209
288. **Neumann HPH**, Eggert HR, Scheremet R, Schumacher M, Mohadjer M, Wakhloo AK, Volk B, Hettmannsperger U, Riegler P, Schollmeyer P, Wiestler OD  
Lesions of the central nervous system in Von Hippel-Lindau syndrome  
**Journal of Neurology, Neurosurgery and Psychiatry** 1992;55:898-901
289. **Neumann HPH**, Hofmann V, Zäuner I, Sigmund G, Blum U, Schümichen C, Schmidt D, Kirste G  
Phäochromozytom als dominierende Manifestation des Von Hippel-Lindau Syndroms  
**Deutsche Medizinische Wochenschrift** 1992;117:1709-1716

**1991**

290. Takiyyuddin MA, Baron AD, Cervenka JH, Barbosa JA, **Neumann HPH**, Parmer JR, Sullivan PA, O'Connor DT  
Suppression of chromogranin A release from neuroendocrine sources in man: Pharmacological Studies  
**The Journal of Clinical Endocrinology and Metabolism** 1991;2:616-622
291. Takiyyuddin MA, **Neumann HPH**, Cervenka JH, Kennedy B, Dinh TQ, Ziegler MG, Baron AD, O'Connor DT  
Ultradian Variations of Chromogranin A in Humans  
**The American Journal of Physiology** 1991;261:R939-944
292. **Neumann HPH**, Jelkmann W, Eggert HR, Schollmeyer P, Wiestler OD  
Serum Erythropoietin levels in Von Hippel-Lindau syndrome  
**Journal of Neurology, Neurosurgery and Psychiatry** 1991;54:746-747
293. Seizinger BR, Smith DI, Filling-Katz MR, **Neumann HPH**, Green JS, Choyke PL, Anderson KM, Freiman RN, Klauck SM, Whaley J, Decker H-JH, Hsia YE, Collins D, Halperin J, Lamiell JM, Oostra B, Waziri MH, Gorin MB, Scherer G, Drabkin HA, Aronin N, Schinzel A, Martuza RL, Gusella JF, Haines JL  
Genetic flanking markers refine diagnostic criteria and provide insights into the genetics of Von Hippel-Lindau disease  
**Proceedings of the National Academy of Science USA** 1991;88:2864-2868
294. **Neumann HPH**, Wiestler OD  
Clustering of features of Von Hippel-Lindau syndrome: evidence for a complex genetic locus  
**The Lancet** 1991;337:1052-1054
295. **Neumann HPH**, Wiestler OD  
Clustering of features and genetics of Von Hippel-Lindau syndrome. Correspondence  
**The Lancet** 1991;338:258
296. **Neumann HPH**, Dinkel E, Brambs H, Wimmer B, Friedburg H, Sigmund G, Riegler P, Haag K, Schollmeyer P, Wiestler OD  
Pancreatic lesions in the Von Hippel-Lindau syndrome  
**Gastroenterology** 1991;101:465-471
297. Riegler P, Bonatti G, Ortore PG, Psenner K, Huber W, Königsrainer A, Margreiter R, **Neumann HPH**  
Sindrome di Von Hippel-Lindau con Tumori multipli bilaterali.  
**Giornale di Clinica Medica** 1991;72:53-59
298. Kovacs G, Emanuel A, **Neumann HPH**, Kung H  
Cytogenetics of renal cell carcinomas associated with Von Hippel-Lindau disease  
**Genes, Chromosomes and Cancer** 1991;3:256-262



299. **Neumann HPH**  
 Das Von Hippel-Lindau Syndrom  
**Deutsche Medizinische Wochenschrift** 1991;116:28-34

300. **Neumann HPH**  
 Aktuelle Diagnostik des Phäochromozytoms  
**Nieren- und Hochdruckkrankheiten** 1991;20:343-350

#### 1990

301. Telenius H, Mathew CGP, Nakamura Y, Easton DF, Clark J, **Neumann HPH**,  
 Ziegler WH, Schinzel A, Ponder BAJ  
 Application of linked DNA markers to screening families with multiple endocrine  
 neoplasia type 2a  
**The European Journal of Surgical Oncology** 1990;16:134-140

302. Hsiao RJ, **Neumann HPH**, Parmer RJ, Barbosa JA, O'Connor DT  
 Chromogranin A in familial pheochromocytoma: Diagnostic screening value,  
 prediction of tumor mass, and post-resection kinetics indicating two-compartment  
 distribution  
**The American Journal of Medicine** 1990;88:607-613

303. Takiyyuddin MA, Cervenka JH, Pandian MR, Stuenkel CA, **Neumann HPH**,  
 O'Connor DT  
 Neuroendocrine sources of Chromagranin A in normal man: Clues from selective  
 stimulation of endocrine glands  
**The Journal of Clinical Endocrinology and Metabolism** 1990;71:360-369

304. Bonatti G, Ortore P, Riegler P, Königsrainer A, **Neumann HPH**  
 La diagnostica integrata nelle localizzazioni renali della sindrome di von Hippel-  
 Lindau  
**La Radiologia Medica** (Torino) 1990;80:938-941

#### 1989

305. Kiechle-Schwarz M, **Neumann HPH**, Decker H, Dietrich C, Wullich B, Schempp  
 W  
 Cytogenetic studies on 3 pheochromocytomas derived from patients with Von  
 Hippel-Lindau syndrome  
**Human Genetics** 1989;82:127-130

306. **Neumann HPH**, Eggert H, Weigel K, Friedburg H, Wiestler O, Schollmeyer P  
 Hemangioblastomas of the central nervous system: a ten year study with special  
 reference to Von Hippel-Lindau syndrome  
**The Journal of Neurosurgery** 1989;70:24-30

307. **Neumann HPH**, Müller O, Ponder B, Mathew C, Telenius H, Schempp W, Schümichen C, Freudenberg N, Schollmeyer P  
Early diagnosis of multiple endocrine neoplasia type IIa  
**Klinische Wochenschrift** 1989;67:951-956

### 1988

308. Bender K, Bissbort S, Crone H, Senff H, Steiert A, **Neumann HPH**, Koch M, Nagel M, Wienker T  
Linkage relations of JK, CO, KEL and IgK with each other and with AHCY  
**Human Heredity** 1988;38:12-17
309. **Neumann HPH**, Schempp W, Wienker T  
High resolution chromosome banding and fragile site studies in von Hippel-Lindau syndrome  
**Cancer Genetics and Cytogenetics** 1988;31:41-46
310. **Neumann HPH**, Hofmann T, Köster W, Billmann P, Kauffmann G  
Extraction of an intracardial catheter embolus using combined radiography and transoesophageal echocardiography  
**Clinical Cardiology** 1988;11:427-429
311. Schmidt D, **Neumann HPH**, Eggert H, Friedburg H  
Neuro-Ophthalmologischer Befund bei Hämangioblastom des Kleinhirns und des Hirnstamms  
**Fortschritte der Ophthalmologie** 1988;85:427-433
312. Decker H, **Neumann HPH**, Walter T, Sandberg A  
3p Involvement in a renal cell carcinoma in Von Hippel-Lindau syndrome: region of tumor breakpoint clustering on 3p?  
**Cancer Genetics and Cytogenetics** 1988;33:59-65
313. **Neumann HPH**, Zerres K, Fischer C, Wolff G, Schäfer H, Gal A, Kröpelin T, Haag K, Schollmeyer P  
Late manifestation of autosomal-recessive polycystic kidney disease in two sisters  
**The American Journal of Nephrology** 1988;8:194-197
314. Decker H, Gemill R, **Neumann HPH**, Walter T, Sandberg A  
Loss of heterozygosity on 3p in an Von Hippel-Lindau renal cell carcinoma  
**Cancer Genetics and Cytogenetics** 1988;39:289-293

### 1987

315. Schmidt D, **Neumann HPH**

Atypische retinale Veränderungen bei Von Hippel-Lindau Syndrom  
**Fortschritte der Ophthalmologie** 1987;84:187-189

316. **Neumann HPH**  
 Basic criteria for clinical diagnosis and genetic counselling in Von Hippel-Lindau syndrome  
**VASA** 1987;16:220-226
317. **Neumann HPH**  
 Prognosis of Von Hippel-Lindau syndrome  
**VASA** 1987;16:309-311
318. Wirth B, Zerres K, Fischbach M, Claus D, **Neumann HPH**, Lennert T, Brodehl J, Neugebauer M, Müller-Wiefel D, Geisert J, Gal A  
 Autosomal recessive and dominant forms of polycystic kidney disease are not allelic  
**Human Genetics** 1987;77:221-222

#### 1986

319. **Neumann HPH**, Schollmeyer P, Schmidt D  
 Bedeutung der interdisziplinären Zusammenarbeit bei Angiomatosis retinae  
**Fortschritte der Ophthalmologie** 1986;83:230-232
320. Stahl RAK, Oberle G, **Neumann HPH**, Schollmeyer P  
 Einzelniere - Risiko oder tolerabler Organverlust?  
**Deutsche Medizinische Wochenschrift** 1986;9:350-354
321. Schmidt D, **Neumann HPH**, Witschel H  
 Mikroläsionen der Retina bei Patienten mit Von Hippel-Lindau Syndrom  
**Fortschritte der Ophthalmologie** 1986;83:233-235
322. Koup J, Keller E, **Neumann HPH**, Stöckel K  
 Ceftriaxone pharmacokinetics during peritoneal dialysis  
**European Journal of Clinical Pharmacology** 1986;30:303-307

#### 1985

323. **Neumann HPH**, Herz R, Baum C  
 Granulomatöse und eosinophile Myositis durch Onchozerca volvulus  
**Der Pathologe** 1985;6:101-107
324. **Neumann HPH**, Dietze W, Poll M, Willig F  
 Colitis cystica profunda - Beitrag zur Differentialdiagnose des Rektumkarzinoms  
**Leber Magen Darm** 1985;15:112-116

325. Schollmeyer P, **Neumann HPH**  
Die Therapie der Glomerulonephritis einschließlich des nephrotischen Syndroms  
**Therapiewoche** 1985;35:2691-2704

326. Oberle G, **Neumann HPH**, Schollmeyer P, Boesken W, Stahl RAK  
Mild proteinuria in patients with unilateral kidney  
**Klinische Wochenschrift** 1985;63:1048-1051

#### 1984

327. **Neumann HPH**, Karte H  
Riesenaneurysma der intrakraniellen Arterien bei Kindern und Jugendlichen  
**VASA** 1984;3:258-261

328. **Neumann HPH**, Kamphues R  
Neue Venenkathetermarkierungen - eine röntgenologische Ortungshilfe  
**Intensivmedizin** 1984;21:25-30

#### 1983

329. **Neumann HPH**, Karte H  
Erwiderung auf Leserbrief zu: Extremer intrauteriner mikrozephaler Minderwuchs  
mit Riesenaneurysma des Circulus arteriosus Willisii  
**Pädiatrische Praxis** 1983;28:126-128

330. **Neumann HPH**  
Morphologie und Klinik des Hämangioperizytoms - eine Analyse von 84 Fällen  
mit einem eigenen Beitrag  
**Der Pathologe** 1983;4:64-70

331. **Neumann HPH**, Löffler H, Brass H, Wegener K  
Akutes Nierenversagen und Aorteninsuffizienz bei großem nicht perforiertem  
Sinus-Valsalvae-Aneurysma  
**VASA** 1983;2:185-189

332. **Neumann HPH**, Kampschulte R  
Plötzlicher nicht-traumatischer Tod im Sport  
**Deutsche Zeitschrift für Sportmedizin** 1983;34:141-148

333. **Neumann HPH**  
Allergische und toxische Nebenwirkungen von Ethylendiamin  
**Allergologie** 1983;6:27-30

#### 1982

334. **Neumann HPH**

Akute Urtikaria bei Ethylendiamin-Theophyllin-Therapie  
**Deutsche Medizinische Wochenschrift** 1982;107:116

335. **Neumann HPH**

Extremer intrauteriner mikrozephaler Minderwuchs mit Riesenaneurysma des  
Circulus arteriosus Willisii

**Pädiatrische Praxis** 1982/83;27:21-31

**1981**

336. **Neumann HPH**

Beitrag zur Diagnostik von Katheterembolien

**Anästhesie, Intensivtherapie und Notfallmedizin** 1981;16: 279-281

337. **Neumann HPH, Kamphues R**

Vereinfachung der Ortung zentraler Venenkatheter durch neue Markierungen

**Der Radiologe** 1981;21:493-495

338. **Neumann HPH, Kühn W**

Oxyuriasis der Tuba uterina

**Zentralblatt für Gynäkologie** 1981;103:156-1564