

7. Literaturverzeichnis

1. Augsburger, J.J., Shields, J.A. and Goldberg, R.E.: Classification and management of hereditary retinal angiomas. *Int Ophthalmol*, 1981. 4: p. 93-106.
2. Webster, A.R., Maher, E.R., Bird, A.C., Gregor, Z.J., Moor, A.T.: A clinical and molecular genetic analysis of solitary ocular angioma. *Ophthalmology*, 1999. 106(3): p. 623-9.
3. Neumann, H.P.H. and Wiestler, O.D.: Clustering of features and genetics of von Hippel-Lindau syndrome. *Lancet*, 1991. 338(8761): p. 258.
4. Latif, F., Tory, K., Gnarra, J. et al.: Identification of the von Hippel-Lindau disease tumor suppressor gene. *Science*, 1993. 260(5112): p. 1317-20.
5. Seizinger, B.R., Rouleau, G.A., Ozelius, L.J. et al.: Von Hippel-Lindau disease maps to the region of chromosome 3 associated with renal cell carcinoma. *Nature*, 1988. 332(6161): p. 268-9.
6. Maher, E.R., Yates, J.R., Harries, R. et al.: Clinical features and natural history of von Hippel-Lindau disease. *Q J Med*, 1990. 77(283): p. 1151-63.
7. van der Hoeve, J.: The Doyne Memorial Lectures: Eye symptoms in phakomatoses. *Tr Ophth Soc UK*, 1932. 52: p. 380.
8. Maher, E.R., Iselius, L., Yates, J.R. et al.: Von Hippel-Lindau disease: a genetic study. *J Med Genet*, 1991. 28(7): p. 443-7.
9. Horton, W.A., Wong, V. and Eldridge, R.: Von Hippel-Lindau disease: clinical and pathological manifestations in nine families with 50 affected members. *Arch Intern Med*, 1976. 136(7): p. 769-77.

10. Lamiell, J.M., F.G. Salazar, and Y.E. Hsia: Von Hippel-Lindau disease affecting 43 members of a single kindred. *Medicine Baltimore*, 1989. 68(1): p. 1-29.
11. Lommatsch, A. and A. Wessing: Angiomatosis retinae. *Ophthalmologe*, 1996. 93: p. 158-162.
12. Streiff, E.B.: Un nouveau cas, le premier, de maladie de von Hippel-Lindau. *Ophthalmologica*, 1951. 122: p. 367.
13. Panas, F. and Remy, A.: Decollement kystique de la retine. *Anatomie Pathologique de l’Oeil*. 1879, Paris: Delahaye & Cie. p.88-93.
14. Galezowski, J.: Traite Iconographique D` ophthalmoscopic. 1886, Paris: Bailliere et Fils., p. 33.
15. Jackson, H.: A series of cases illustrativ of cerebral pathology. Cases of intracranial tumor. *Med. Times & Gaz.*, 1872. 2. p. 541-42.
16. Wood, D.J.: Retinal detachment with unusual dilatation of retinal vessels and other changes. *Trans Ophth Soc UK*, 1892. 12. p.143.
17. Collins, E.T.: Intra-Ocular Growths. Two cases, brother and sister, with peculiar vascular new growth, probably primarily retinal, affecting both eyes. *Trans Ophth Soc UK*, 1894. 14: p. 141.
18. von Hippel, E.: Vorstellung eines Patienten mit einer sehr ungewöhnlichen Netzhaut. *Ber Deutsch Ophthal Ges*, 1895. 24: p. 269.
19. von Hippel, E.: Über eine sehr seltene Erkrankung der Netzhaut. *Ber Deutsch Ophthal Ges*, 1903. 31: p. 199.
20. von Hippel, E.: Über eine sehr seltene Erkrankung der Netzhaut. v. Graefes Arch f Opth, 1904. 59: p. 83-106.

21. von Hippel, E.: Die anatomische Grundlage der von mir beschriebenen sehr seltenen Erkrankung der Netzhaut. Albrecht von Graefes Arch f Ophth, 1911. 79: p. 350-377.
22. Brandt, R.: Zur Frage der Angiomatosis retinae. von Graefes Arch. Ophthal., 1921. 106: p. 127.
23. Lindau, A.: Angiomatosis retinae (v. Hippelsche Krankheit). Acta path et microbiol Scand, 1926. Suppl 1: p. 1-129.
24. Lindau, A.: Zur Frage der Angiomatosis retinae und ihrer Hirnkomplikationen. Acta Ophth, 1927. 4: p. 193.
25. Lindau, A.: Capillary angiomatosis of the central nervous system. Acta genet, 1957. 7: p. 338.
26. Melmon, K.L. and Rosen, S.W.: Lindau's disease. Am J Med, 1964. 36: p. 595-617.
27. Green, J.S., Bowmer, M.I., and Johnson, G.J.: Von Hippel-Lindau disease in a Newfoundland kindred. Can Med Assoc J, 1986. 134(2): p. 133-8.
28. Neumann, H.P.H.: Basic criteria for clinical diagnosis and genetic counselling in von Hippel-Lindau syndrome. Vasa, 1987. 16(3): p. 220-6.
29. Hubschmann, O.R., Vijayanathan, T. and Countee, R.W.: Von Hippel-Lindau Disease with multiple manifestations: diagnosis and management. Neurosurgery, 1981. 8(1): p. 92-5.
30. Maher, E.R., Webster, A.R. and Moore, A.T.: Clinical features and molecular genetics of Von Hippel-Lindau disease. Ophthalmic Genet, 1995. 16(3): p. 79-84.
31. Ridley, M., Green, J. and Johnson, G.: Retinal angiomatosis: the ocular manifestations of von Hippel-Lindau disease. Can J Ophthalmol, 1986. 21(7): p. 276-83.

32. Moore, A.T., Maher, E.R., Rosen, P., Gregor, Z., Bird, A.C.: Ophthalmological screening for von Hippel-Lindau disease. *Eye*, 1991. 5 p. 723-8.
33. Jesberg, D.O., William, H.S and William, F.H.: Incipient Lesions of von Hippel-Lindau Disease. *Arch Ophthal*, 1968. 80: p. 632-40.
34. Webster, A.R., Maher, E.R. and Moore, A.T.: Clinical characteristics of ocular angiomas in von Hippel-Lindau disease and correlation with germline mutation. *Arch Ophthalmol*, 1999. 117: p. 371-378.
35. Knudson, A.G.J.: Mutation and Cancer: statistical study of retinoblastoma. *Proc Natl Acad Sci USA*, 1971. 68: p. 820-823.
36. Bottini, F., Canevini, M.P., Canger, R., Orzalesi, N.: Twin vessels in familial retinal cavernous hemangioma. *Am J Ophthalmol*, 1990. 109(3): p. 285-9 .
37. de Jong, P.T., Verkaart, R.J., van de Vooren, M.J., Majoor Krakauer, D.F., Wiegel, A.R.: Twin vessels in von Hippel-Lindau disease. *Am J Ophthalmol*, 1988. 105(2): p. 165-9.
38. Chang, J.H., Spraul, C.W., Lann, M.L., Drack, A., Grossniklaus, H.E.: The two-stage mutation model in retinal hemangioblastoma. *Ophthalmic Genet*, 1998. 19: p. 123-30.
39. Benson, M., Mody, C., Rennie, I., Talbot, J.: Haemangioma of the optic disc. *Graefe's Arch Clin Exp Ophthalmol*, 1990. 228: p. 332-334.
40. Imes, R.K., Monteiro, M.L.R., Hoyt, W.F.: Incipient Hemangioblastoma of the optic disk. *American Journal of ophthalmology*, 1984. 98(1): p. 116.
41. In, S., Miyagi, J., Kojho, N., Kuramoto, S., Uehara, M.: Intraorbital optic nerve hemangioblastoma with von Hippel-Lindau disease. Case report. *J Neurosurg*, 1982. 56(3): p. 426-9.

42. Kerr, D.J., Scheithauer, B.W., Miller, G.M., Ebersold, M.J., McPhee, T.J.: Hemangioblastoma of the optic nerve: case report. *Neurosurgery*, 1995. 36(3): p. 573-80.
43. Wittebol-Post, D., Hes F.J. and Lips, C.J.: The eye in von Hippel-Lindau disease. Long-term follow-up of screening and treatment: recommendations. *J Intern Med*, 1998. 243(6): p. 555-61.
44. Kreusel, K.M., Bornfeld, N., Lommatsch, A., Wessing, A., Foerster, M.H.: Ruthenium-106 brachytherapy for peripheral retinal capillary hemangioma. *Ophthalmology*, 1998. 105(8): p. 1386-92.
45. Whitson, J.T., Welch, R.B. , Green, W.R.: Von Hippel-Lindau disease: case report of a patient with spontaneous regression of a retinal angioma. *Retina*, 1986. 6(4): p. 253-9.
46. Schmidt, D., Natt, E. and Neumann, H.P.H.: Longterm results of Laser treatment for retinal angiomas in von Hippel-Lindau Disease. *EurJ Med Res* 2000, 5(2), p. 47-58.
47. Grossniklaus, H.E., Thomas, J.W., Vigneswaran, N., Jarret, W.H.: Retinal hemangioblastoma. A histologic, immunohistochemical and ultrastructural evaluation. *Ophthalmology*, 1992. 99(1): p. 140-5.
48. Wizigmann Voos, S., Breier, G., Risau, W. Plate, K.H.: Up-regulation of vascular endothelial growth factor and its receptors in von Hippel-Lindau disease - associated and sporadic hemangioblastomas. *Cancer Res*, 1995. 55(6): p. 1358-64.
49. Henkind, P. and Benjamin, J.V.: Vascular anomalies and neoplasms of the optic nerve head. *Transact Ophthalmol Soc UK*, 1976. 96: p. 418-423.
50. Naumann, G.: *Pathologie des Auges*. Springer-Verlag Berlin Heidelberg New York, 1980, p. 648

51. Hardwig, P. and Robertson, D.M.: Von Hippel-Lindau disease: a familial, often lethal, multi-system phakomatosis. *Ophthalmology*, 1984. 91(3): p. 263-70.
52. Maher, E.R. and Kaelin, W.G. Jr.: Von Hippel-Lindau disease. *Medicine (Baltimore)*, 1997. 76(6): p. 381-91.
53. Neumann, H.P.H., Eggert, H.R., Weigel, K. et al.: Hemangioblastomas of the central nervous system. A 10-year study with special reference to von Hippel-Lindau syndrome. *J Neurosurg*, 1989. 70(1): p. 24-30.
54. Neumann, H.P.H., Eggert, H.R., Scheremet, R. et al.: Central nervous system lesions in von Hippel-Lindau syndrome. *J Neurol Neurosurg Psychiatry*, 1992. 55(10): p. 898-901.
55. Filling Katz, M.R., Choyke, P.L., Patronas, N.J. et al.: Radiologic screening for von Hippel-Lindau disease: the role of Gd-DTPA enhanced MR imaging of the CNS. *J Comput Assist Tomogr*, 1989. 13(5): p. 743-55.
56. Richards, F.M., Payne, S.J., Zbar, B. et al.: Molecular genetic analysis of von Hippel-Lindau disease. *J Intern Med*, 1998. 243(6): p. 527-33.
57. Nerad, J.A., Kersten, R.C. and Anderson, R.L.: Hemangioblastoma of the optic nerve. Report of a case and review of literature. *Ophthalmology*, 1988. 95(3): p. 398-402.
58. Ginzburg, B.M., Montanera, W.J., Tyndel, F.J. et al.: Diagnosis of von Hippel-Lindau disease in a patient with blindness resulting from bilateral optic nerve hemangioblastomas. *AJR Am J Roentgenol*, 1992. 159(2): p. 403-5.
59. Glenn, G.M., Choyke, P., Zbar, P., Linehan, W.M.: Von Hippel-Lindau disease: Clinical review and molecular genetics. *Probl Urol* 1990; 4: p. 312-330.

60. Neumann, H.P.H., Berger, D.P., Siegmund, G. et al.: Pheochromocytomas, multiple endocrine neoplasia type 2, and von Hippel-Lindau disease [see comments] [published erratum appears in N Engl J Med 1994 Dec 1;331(22):1535]. *N Engl J Med*, 1993. 329(21): p. 1531-8.
61. Fill, W.L., J.M. Lamiell, and N.O. Polk: The radiographic manifestations of von Hippel-Lindau disease. *Radiology*, 1979. 133(2): p. 289-95.
62. Neumann, H.P.H. and B. Zbar: Renal cysts, renal cancer and von Hippel-Lindau disease. *Kidney Int*, 1997. 51(1): p. 16-26.
63. Neumann, H.P.H., Bender, B.U., Berger, D.P. et al.: Prevalence, morphology and biology of renal cell carcinoma in Von Hippel-Lindau disease compared to sporadic renal cell carcinoma. *J Urol*, 1998. 160: p. 1248-1254.
64. Neumann, H.P.H., Dinkel, E., Brambs, H. et al.: Pancreatic lesions in the Von Hippel-Lindau syndrome. *Gastroenterol*, 1991. 101: p. 465-471.
65. Choyke, P.L., Filling-Katz, M.R., Shawker, T.H. et al.: Von Hippel-Lindau disease: radiologic screening for visceral manifestations. *Radiology*, 1990. 174(3 Pt 1): p. 815-20.
66. Neumann, H.P.H.: The von Hippel-Lindau syndrome. *Dtsch Med Wochenschr*, 1991. 116(1): p. 28-34.
67. McGrath, F.P., Gibney, R.G., Morris, D.C. Owen, D.A., Erb, S.R.: Case report: multiple hepatic and pulmonary haemangioblastomas--a new manifestation of von Hippel-Lindau disease. *Clin Radiol*, 1992. 45(1): p. 37-9.
68. Jennings, A.M., Smith, C., Cole, D.R. et al.: Von Hippel-Lindau disease in a large British family: clinicopathological features and recommendations for screening and follow-up. *Q J Med*, 1988. 66(251): p. 233-49.

69. Blamires, T.L., Friedmann, I. and Moffat, D.A.: Von Hippel-Lindau disease associated with an invasive choroid plexus tumour presenting as a middle ear mass. *J Laryngol Otol*, 1992. 106(5): p. 429-35.
70. Kempermann, G., Neumann, H.P.H., Scheremet, R. et al.: Deafness due to bilateral endolymphatic sac tumours in a case of von Hippel-Lindau syndrome. *J Neurol Neurosurg Psychiatry*, 1996. 61(3): p. 318-20.
71. Moller, H.U.: Familial angiomas retinae and cerebelli-Lindau's disease. *Arch Ophth*, 1929. 7: p. 244.
72. Richards, F.M., Webster, A.R., McMahon, R., Woodward, E.R., Rose, S., Maher, E.R.: Molecular analysis of de novo germline mutations in the von Hippel-Lindau disease gene. *Hum Mol Genet*, 1995. 4(11): p. 2139-43.
73. Neumann, H.P.H., Kreusel, K.M., Apel, T.W., Munk, R.D., Schmidt, D., Bornfeld, N., Foerster, M.H.: Gefäßtumoren der Netzhaut. *Onkologe*, 1999. 5: p. 805-812.
74. Pearson, P.L.: The genetic analysis of cancer. *Journal of internal medicine*, 1998. 243: p. 413-417.
75. Gnarra, J.R., Tory, K., Wenig, Y. et al.: Mutation of the VHL tumor suppressor gene in renal carcinoma. *Nat Genet*, 1994. 7: p. 85-90.
76. Thrash Bingham, C.A., Greenberg, R.E., Howard, S. et al.: Comprehensive allelotyping of human renal cell carcinomas using microsatellite DNA probes. *Proc Natl Acad Sci USA*, 1995. 92: p. 2854-2858.
77. Vargas, M.P., Zhunag, Z., Wang, C. et al.: Loss of heterozygosity on the short arm of chromosomes 1 and 3 in sporadic pheochromocytoma and extra-adrenal paraganglioma. *Hum Pathol* 28, 1997. 28: p. 411-5.

78. Prowse, A.H., Webster, A.R. Richards, F.M. et al.: Somatic inactivation of the VHL gene in Von Hippel-Lindau disease tumors. Am J Hum Genet, 1997. 60: p. 765-71.
79. Quesnel, S., Malkin, D.: Genetic predisposition to cancer and familiar cancer syndromes. Pediatr Clin North Am 44 (1997) p. 791-808.
80. Knudson, A.G.: Genetics of human cancer. Annu Rev Genet, 1986. 29: p. 231-251.
81. Knudson, A.G. and Strong, L.C.: Mutation and Cancer: neuroblastoma and pheochromocytoma. Am J Hum Genet, 1972. 24: p. 514-532.
82. Maher, E.R., Yates, J.R. and Ferguson Smith, M.A.: Statistical analysis of the two stage mutation model in von Hippel-Lindau disease, in sporadic cerebellar hemangioblastoma and renal cell carcinoma. J Med Genet, 1990. 27: p. 331-4.
83. Neumann, H.P.H, Lips, C.J.: Von Hippel-Lindau syndrome. Brain Pathol, 1995. 5(2): p. 181-93.
84. Neumann, H.P.H. and Bender, B.U.: Genotype-phenotype correlations in von Hippel-Lindau disease. J Intern Med, 1998. 243(6): p. 541-5.
85. Zbar, B., Kishida, T., Chen, F. et al.: Germline mutations in the von Hippel-Lindau (VHL) gene in families from North America, Europe and Japan. Human Mutations, 1996. 8: p. 348-57.
86. Richards, F.M., Maher, E.R. and Latif, F.M.: Detailed genetic mapping of the von Hippel-Lindau disease tumor suppressor gene. J Med Genet, 1993. 30: p. 104-107.
87. Stolle, C., Glenn, G., Zbar, B. et al.: Improved detection of germline mutations in the von Hippel-Lindau disease tumor suppressor gene. Hum Mutat, 1998. 12(6): p. 417-23

88. American Society of Clinical Oncology: Genetic testing for cancer susceptibility. *J Clinical Oncology*, 1996. 14(5): p. 1730-1736.
89. Orita, M., Suzuki, Y., Sekiya, T., Hayashi, K.: Rapid and Sensitive Detection of Point Mutations and DNA Polymorphism. Using the Polymerase Chain Reaction. *Genomics*, 1989. 5: p. 874-879.
90. Iliopoulos, O., Kibel, A., Gray, S., Kaelin, W.G., Jr.: Tumor suppression by the human von Hippel-Lindau gene product. *Nature Med*, 1995. 1: p. 822-6.
91. Kibel, A., Iliopoulos, O., DeCaprio, J.D. Kaelin, W.G.: Binding of the von Hippel-Lindau tumor suppressor protein to elongin B and C. *Science*, 1995. 269: p. 1444-6.
92. Duan, D.R., Humphrey, J.S. Chen, D.Y. et al.: Inhibition of transcriptional elongation by the VHL tumor suppression protein. *Science*, 1995. 269: p. 1402-6.
93. Aso, T., Haque, D., Barstead, R.J., Conaway, R.C., Conaway, J.W.: Elongin (S III): a multisubunit regulator of elongation by RNA polymerase II. *Science*, 1995. 269: p. 1439-43.
94. Kishida, T., Stackhous, T.M., Chen, F., Lerman, M.I.: Cellular proteins that bind the von Hippel-Lindau disease gene product: mapping of binding domains and the effect of missense mutations. *Cancer Res*, 1995. 55: p. 4544-8.
95. Tsuchiya, H., Iseda, T. and Hino, O.: Identification of a novel protein (VBP-1) binding to the von Hippel-Lindau (VHL) tumor suppressor gene product. *Cancer Res*, 1996. 56: p. 2881-5.
96. Aso, T., Haque, D., Barstead, R.J., Conaway, R.C., Conaway, J.W.: The inducible elongin A elongation activation domain: structure, function and interaction with elongin BC complex. *EMBO J*, 1996. 15: p. 101-10.

97. Takahashi, A., Sasaki, H., Kim, S.J. et al.: Markedly Increased Amounts of Messenger RNA's for Vascular Growth Factor and Placenta Growth in Renal Cell Carcinoma Associated with Angiogenesis. *Cancer Research*, 1994. 54: p. 4233-4237.
98. Berse, B., Brown, L., Livingston, V., Dvorak, H., Senger, D.: Vascular permeability factor (vascular endothelial growth factor) gene is expressed differentially in normal tissues, macrophages and tumors. *Mol Biol Cell*, 1992. 3: p. 211-20.
99. Sato, K., Terada, K., Sugiyama, T. et al.: Frequent overexpression of vascular endothelial growth factor gene in human renal cell carcinoma. *Tohoku J Exp Med*, 1994. 173: p. 355-60.
100. Krieg, M., Marti, H.H. and Plate, K.H.: Coexpression of erythropoietin and vascular endothelial growth factor in nervous system tumors associated with von Hippel-Lindau tumor suppressor gene loss of function. *Blood*, 1998. 92(9): p. 3388-93.
101. Horton, J.C., Harsh, G.R., Fisher, J.W., Hoyt, W.F.: Von Hippel-Lindau disease and erythrocytosis: Radioimmunassay of erythropoietin in cyst fluid from a brainstem hemangioblastoma. *Neurology*, 1991. 41: p. 753-754.
102. Gnarra, J., Zhou, S., Merrill, M.J. et al.: Post-transcriptional regulation of vascular endothelial growth factor mRNA by the VHL tumor suppressor gene product. *Proc Natl Acad Sci USA*, 1996. 93: p. 10589-94.
103. Iliopoulos, O., Jiang, C., Levy, A.P. Kaelin, W.G., Goldberg, M.A.: Negative Regulation of Hypoxia-Inducible Genes by the von Hippel-Lindau Protein. *Proc Natl Acad Sci USA*, 96. 93: p. 10595-9.
104. Siemeister, G., Weindel, K., Mohrs, K. et al.: Reversion of deregulated expression of vascular endothelial growth factor in human renal carcinoma cells by von Hippel-Lindau tumor suppression Protein. *Cancer Res*, 1996. 56: p. 2299-301.

105. Ohh, M., Yauch, R.L., Lonergan, K.M. et al.: The von Hippel-Lindau tumor suppressor protein is required for proper assembly of an extracellular fibronectin matrix. *Mol Cell*, 1998. 1: p. 959-68.
106. Vaheri, A., Alitalo, K., Hedman, K. et al.: Fibronectin and the pericellular matrix of normal and transformed adherent cells. *Ann NY Acad Sci*, 1978. 312: p. 343-53.
107. Schmidt, D. and Neumann, H.P.H.: Atypical retinal changes in v. Hippel-Lindau syndrome. *Fortschr Ophthalmol*, 1987. 84(2): p. 187-9.
108. Weve, H.J.: On diathermie in ophthalmic practice. *Trans Ophthalmol Soc*, 1939. 59: p. 43-80.
109. Meyer-Schwickerath, G.: Light Coagulation. Trans. by Stephen M. Drance. St. Louis, Mosby, 1960, p. 75.
110. Wessing, A.: 10 Jahre Lichtkoagulation bei angiomatosis retinae. *Klin Monatsbl Augenheilk*, 1967. 150: p. 57-71.
111. Goldberg, M.F. and Duke, J.R.: Von Hippel-Lindau disease. Histopathologic findings in a treated and an untreated eye. *Am J Ophthalmol*, 1968. 66(4): p. 693-705.
112. Apple, D.J., Goldberg, M.F. and Wyhinny, G.J.: Argon laser treatment of von Hippel-Lindau retinal angiomas. II. Histopathology of treated lesions. *Arch Ophthalmol*, 1974. 92(2): p. 126-30.
113. Goldberg, M.F. and Koenig, S.: Argon laser treatment of von Hippel-Lindau retinal angiomas.. Clinical and angiographic findings. *Arch Ophthalmol*, 1974. 92(2): p. 121-5.
114. Blodi, C.F., Russel, S.R., Pulido, J.S. Folk, J.C.: Direct and feeder vessel photocoagulation of retinal angiomas with dye yellow laser. *Ophthalmology*, 1990. 97(6): p. 791-5.

115. Amoils, S.P.: Cryotherapy of Angiomatosis Retinae. *Arch Ophthal*, 1969. 81(May): p. 689-691.
116. Watzke, R.C.: Cryotherapy for Retinal Angiomatosis. *Arch Ophthalmol*, 1974. 92: p. 399-401.
117. Lincoff, H.J.: The cryotherapy of intraocular tumors. *Am J Ophthalmol*, 1967. 63: p. 389-399.
118. Welch, R.B.: Von Hippel-Lindau disease: the recognition and treatment of early angiomatosis retinae and the use of cryosurgery as an adjunct to therapy. *Trans Am Ophthalmol Soc*, 1970. 68: p. 367-424.
119. Lommatsch, P. and Vollmar, R.: Ein neuer Weg zur konservativen Therapie intraokularer Tumoren mit Betastrahlen (Ruthenium¹⁰⁶) unter Erhalt der Sehfähigkeit. *Klin Monatsbl Augenheilkd*, 1966. 148: p. 682-699.
120. Rohrschneider, K., Burk, R.O., Bornfeld, N., Volcker, H.E.: Capillary hemangioma of the retina. Laser scanning tomography follow-up after radiotherapy. *Fortschr Ophthalmol*, 1991. 88(6): p. 623-8.
121. Balazs, E., Berta, A., Rozsa, L., Kolozsvari, L., Rigo, G.: Hemodynamic changes after ruthenium irradiation of Hippel's angiomatosis. *Ophthalmologica*, 1990. 200(3): p. 128-32.
122. Lane, C.M., Turner, G., Gregor, Z.J., Bird, A.C.: Laser treatment of retinal angiomatosis. *Eye*, 1989. 3: p. 33.
123. Meyer-Schwickerath, G.: The preservation of vision by treatment of introcular tumors with light coagulation. *Arch Ophthalmol*, 1961. 66: p. 458-466.
124. Wessing, A. and Weiler, W.: Angiomatosis retinae-Neue Wege der Behandlung (Vortrag). 1987, 28. Tagung der österreichischen ophthalmologischen Gesellschaft: Gmunden.

125. Mulholland, D.A., Johnston, P.B. and Sharkey, J.A.: Pars plana vitrectomy in von Hippel Lindau disease. Eye, 1996. 10: p. 758-9.
126. Nicholson, D.H., Anderson, L.S. and Blodi, C.B.: Rhegmatogenous Retinal Detachment in Angiomatosis Retinae. Am J Ophthalmol, 1986. 101: p. 187-189.
127. Schwartz, P.L., Fastenberg, D.M. and Shakin, J.L.: Management of macular puckers associated with retinal angiomas. Ophthalmic Surg, 1990. 21: p. 550-6.
128. Machemer, R. and J.M. Williams, Sr.: Pathogenesis and therapy of traction detachment in various retinal vascular diseases [published erratum appears in Am J Ophthalmol 1988 Jun 15;105(6):714]. Am J Ophthalmol, 1988. 105(2): p. 170-81.
129. Karsdorp, N., Elderson, A., Wittelbol-Post, D. et al.: Von Hippel-Lindau disease: new strategies in early detection and treatment. Am J Med, 1994. 97(2): p. 158-68.
130. Glenn, G.M., Linehan, W.M., Hosoe, S. et al.: Screening for von Hippel-Lindau disease by DNA polymorphism analysis. Jama, 1992. 267(9): p. 1226-31.
131. Neumann, H.P.H.: Die Von Hippel-Lindau'sche Erkrankung: Informationsschrift für Patienten und Familien. 1996, Medizinische Universitätsklinik: Freiburg.
132. Homepage der von Hippel-Lindau Family Alliance, <http://www.vhl.org>, . 1999.
133. Salazar, F.G. and Lamiell, J.M.: Early identification of retinal angiomas in a large kindred von Hippel-Lindau disease. Am J Ophthalmol, 1980. 89: p. 540-5.

134. Bengtsson, B. and Krakau, C.E.T.: Correction of optic disc measurements on fundus photographs. *Graefe's Arch Clin Exp Ophthalmol*, 1992. 230: p. 24-28.
135. Chen, F., Kishida, T., Yao, M. et al.: Germline mutations in the von Hippel-Lindau disease tumor suppressor gene: correlations with phenotype. *Hum Mutat*, 1995. 5(1): p. 66-75.
136. Zbar, B., Brauch, H., Talmadge, C., Linehan, M.: Loss of alleles of loci on the short arm of chromosome 3 in renal cell carcinoma. *Nature*, 1987. 327: p. 721-724.
137. Miller, R.G.: The jackknife-a review. *Biometrika*, 1974. 61: p. 1-17.
138. Gläsker, S., Bender, B.U., Apel, T.W. et al.: The impact of molecular genetic analysis of the VHL gene in patients with haemangioblastoma of the central nervous system. *J Neurol Neursurg Psychiatry*, 1999. 67: p. 758-762.
139. Webster, A.R., Richards, F.M., MacRonald, F.E., Moore, A.T., Maher, E.R.: An analysis of phenotypic variation in the familial cancer syndrome von Hippel-Lindau disease: evidence for modifier effects. *Am J Hum Genet*, 1998. 63(4): p. 1025-35.