VINCENT M. RICCARDI, MD, MBA *Curriculum Vitae*, Updated: 1 April 2015

DATE AND PLACE OF BIRTH 14 October 1940; Brooklyn, New York

OFFICE ADDRESS The Neurofibromatosis Institute
5415 Briggs Avenue
La Crescenta, CA 91214
Telephone: 818/957-3508 (Office and Cell)
FAX: 818/957-4926 (Call first)
E-mail: Riccardi@MedConsumer.com

CURRENT POSITIONS:

Director, The Neurofibromatosis Institute	1980-
Clinical Professor of Pediatrics (Genetics), UCLA	1991-
Director, Genetic Connections	1992 -
Medical Director, NF California	2012 -

MEDICAL LICENSES:

Pennsylvania	1967-1979
Massachusetts	1969-1979
Colorado	1973-1978
Wisconsin	1975-1979
Texas	1977-1995
California	1990-2013 Current & Active

SPECIAL PRESENTATION

Photons, Gravity and Untranslated Nucleic Acids Are the Origin of Life and Contribute Substantially to Genetic Disease, an invited lecture presented at the **Presidential Symposium** of the annual meeting of the **American College of Medical Genetics** in Charlotte, NC, on 28 March 2012.

CITATIONS

2,000 Notable American Men, (American Biographical Institute, 5126 Bur Oak Circle, Box 31226, Raleigh, NC 27622 (1994). "Health Industry Executives and Thinkers: Crusaders and Ethicists," page 172 in the publication, ***The Managed Care 1500***, "A Complete Guide to the Most Influential Managed Care Leaders and Organizations in the United states." ***Who's Who in the West***, 24th-32nd editions, ***Who's Who in Medicine and Healthcare***, 5th & 6th editions, ***Who's Who in America***, 59th – 63rd editions (2005-2009) and **Who's Who in Science and Engineering** (2016-2017).

DEGREES:

AB	University of California at Los Angeles	1962
MD	Georgetown University School of Medicine	1966
MBA	University of La Verne (*Summa Cum Laude*)	1993

POST GRADUATE EXPERIENCE:

Intern in Medicine, Health Center Hospitals University of Pittsburgh	1966-1967
Resident in Medicine, Health Center Hospitals University of Pittsburgh	1967-1968
Clinical and Research Fellow in Pediatrics, Genetics Unit Massachusetts General Hospital and Harvard Medical School	1968-1970
Internal Medical Ward Officer, Major, Medical Corps, USA, Ft. Devens, MA	1970-1971
Clinical and Research Fellow in Pediatrics, Genetics Unit Massachusetts General Hospital and Harvard Medical School	1972
Assistant Professor of Medicine, University of Colorado Medical Center	1973-1975

Assistant Director, Genetic Counseling Clinics and Genetics Unit, UCMC	1973-1974
Co-Founder, Organizer and Associate Director, Colorado-Wyoming Regional Genetic Counseling Program	1973-1974
Director, Colorado-Wyoming Regional Genetic Counseling Program	1974-1975
Associate Professor of Pediatrics and Medicine, The Medical College of Wisconsin	1975-1977
Director, Kleberg Cytogenetics Lab., Baylor College of Medicine	1977-1980
Associate Professor of Medicine, Baylor College of Medicine	1977-1985
Associate Professor Pediatrics, Baylor College of Medicine	1977-1985
Associate Professor Ob-Gyn, Baylor College of Medicine	1977-1990
Founder and Director, Baylor Neurofibromatosis Program	1978-1990
Adjunct Associate Professor, Graduate School of Biomedical Sciences, University of Texas Health Science Center, Houston	1979-1990
Director, Research Cytogenetics Laboratory, Baylor College of Medicine	1980-1990
Founder, The Neurofibromatosis Institute	1980-
Professor of Medicine, Baylor College of Medicine	1985-1990
Professor of Pediatrics, Baylor College of Medicine	1985-1990
Adjunct Professor, Department of Experimental Carcinogenesis, University of Texas System Cancer Center/Science Park-Research Division.	1986-1990
Visiting Scientist at the Science Park-Research Division, Department of Experimental Carcinogenesis, University of Texas System Cancer Center/M.D. Anderson Hospital and Tumor Institute.	1986-1987
Medical Director, Alfigen/The Genetics Institute	1990-1992
Founder & Director, American Medical Consumers	1991-2010
Clinician, Riccardi Neurofibromatosis/Genetics Clinic	1992-
Consultant to CIGNA Healthcare of California (Credentialing and Quality of Care)	1993 & 1995-2002

EXTRAMURAL ACTIVITIES:

Editorial Board, ***American Journal of Medical Genetics***	1980-1993
Chairman, Cytogenetics Committee, **Pediatric Oncology Group**	1980-1983
Chairman, Medical Advisory Board, **Texas NF Foundation**	1981-1990
Member, Board of Directors, **Texas Prader-Willi Association**	1982-1988
Chairman, Information and Education Committee, American Society of Human Genetics	1983-1988
Liaison to the **American Association for the Advancement of Science** from the American Society of Human Genetics	1985-1991
Editorial Board, ***Cancer Genetics and Cytogenetics***	1986-1987
Founder, Editor-in-Chief, *Neurofibromatosis*	1986-1990
Co-Chair, Clinical Care Advisory Board, **National NF Foundation**	1986-1988
Professional Advisory Board, **International Rett Syndrome Assoc.**	1986-1988
Member, Cancer Biology-Immunology Contracts Review Committee, **National Cancer Institute** (Chairman from 1988)	1986-1989
Member, **National Institutes of Health** Reviewers Review (NRR)	1989-1993
Contribution to preparation of report for the Congress of the United States (**Office of Technology Assessment**); Genetic Monitoring and Screening in the Workplace. (OTABA-455)	1989-1990
Editor, Neurogenetics Section, ***Neurosurgery***	1991-2002
Member, Quality Control Committee, **Pacific Southwest Regional Genetics Network**	1992-1993
Organizer/Facilitator, Dept. of Defense NF Research Program	1993, 1995

Member, Information and Education Committee, **American Society of Human Genetics**	2001-2004
U.S. Department of Defense Congressionally Directed Research Program, Neurofibromatosis Research Program, Programmatic Review and Vision Setting panel	2009-
Member, External Advisory Committee, **Neurofibroma Treatment Acceleration Program** (**NTAP**), Johns Hopkins University Medical School	2012-
Member, Consumer Advisory Board of **FAIR Health**, a national medical consumer advocacy organization	2012-

AWARDS:

Certification, **American Board of Internal Medicine**	1974
Fellowship, **American College of Physicians**	1975
Certification, **American Board of Medical Genetics**, Clinical Genetics and Clinical Cytogenetics	1982
Fellowship, American Association for the Advancement of Science	1992
Fellowship, **American College of Medical Genetics**	1993
Recipient of the **Friedrich von Recklinghausen Award** of the Children's Tumor Foundation	2008
Special Recognition Award, **NF Network**; Las Vegas, NV	2013
Special Recognition Award, **NF Association of Brazil**	2014

BOOKS:

1. Riccardi VM: ***The Genetic Approach to Human Disease*.** New York: Oxford University Press, 273 pg, 1977.

2. Riccardi VM & Mulvihill JJ (eds.): ***Neurofibromatosis (Von Recklinghausen Disease). Genetics, Cell Biology and Biochemistry.*** *Advances in Neurology,* Vol. 29. New York: Raven Press, 282 pg, 1981.

3. Riccardi VM & Kurtz SM: ***Communication and Counseling in Health Care*.** Springfield, Illinois: Charles C. Thomas, 248 pg, 1983.

4. Riccardi VM & Valenta SH: ***Neurofibromatosis: A Primer for Patients and Families*.** Houston: V.M. Riccardi, 16 pg, 1983.

5. Riccardi VM & Eichner JE: ***Neurofibromatosis: Phenotype, Natural History and Pathogenesis*.** Baltimore: Johns Hopkins University Press, 352 pg, 1986.

6. Biscardi M (pseudonym): ***Where Reality Begins: Light and Color, Truth and Love*.** A volume of poetry – La Crescenta, CA: Rich Heart Press, 33 pg, 1991

7. Riccardi VM: ***Neurofibromatosis: A Primer for Patients and Families*.** Pasadena: V.M. Riccardi, 16 pg, 1991.

8. Riccardi VM: ***Neurofibromatosis: Phenotype, Natural History and Pathogenesis*.** Second Edition. Baltimore**:** Johns Hopkins University Press, 498 pg, 1992.

9. Friedman JM, Gutmann DH, MacCollin M & Riccardi VM: ***Neurofibromatosis: Phenotype, Natural History and Pathogenesis*.** Third Edition. Baltimore: Johns Hopkins University Press, 1999.

10. Riccardi VM: ***Matching Gravity: Life in Terms of Sameness, Space and Time.*** (In preparation)

11. Riccardi VM: ***Venus Complete: Amrita and the Digital Woman.*** (In preparation)

PROFESSIONAL JOURNAL ARTICLES:

1. Riccardi VM: **Chromatin states in cytodifferentiation.** *Georgetown Med Bull* 20:30-38, 1966.

2. Riccardi VM, et al.: **Absent patellae, mild retardation, skeletal and genitourinary anomalies and C-group autosomal mosaicism.** *J Pediatr* 77:220-221, 1971.

3. Hayek A, Riccardi VM, et al.: **49,XXXXY chromosomal anomaly in a newborn.** *J Med Genet* 8:220-221, 1971.

4. Riccardi VM & Littlefield JW: **Adaptive increase in phosphoribosyltransferase activity in Lesch-Nyhan fibroblasts exposed to aminopterin.** *Exptl Cell Res* 72:417-422, 1972.

5. Riccardi VM & Holmes LB: **Brachydactyly, Type E: Hereditary shortening of digits, metacarpals and long bones.** *J Pediatr* 84:251-254, 1974.

6. Atkins L, Holmes LB & Riccardi VM: **Trisomy 8.** *J Pediatr* 84:302-303, 1974.

7. Riccardi VM & Robinson A: **Regional genetic counseling in local health care.** *Rocky Mtn Med J* 71:686-688, 1974.

8. Riccardi VM, et al.: **Hereditary pancreatitis: Nonspecificity of aminoaciduria and diagnosis of occult disease.** *Arch Int Med* 135:822-825, 1975.

9. Riccardi VM & Robinson A: **Preventive medicine through genetic counseling: A regional program.** *Prev Med* 4:126-134, 1975.

10. Wenger DA & Riccardi VM: **Possible misdiagnosis of Krabbe disease.** *J Pediatr* 88:76-79, 1976.

11. Riccardi VM: **Health care and disease prevention through genetic counseling: A regional approach.** *Am J Publ Health* 66:268-272, 1976.

12. Riccardi VM: **Trisomy 8 mosaicism in the skin of a patient with leukemia.** *Birth Defects Orig Art Ser* 12(1):187, 1976.

13. Larson SA, Yeatman GW & Riccardi VM: **Deletion of 11q: Report of 2 cases and a review.** *Birth Defects Orig Art Ser* 12(5):125-130, 1976.

14. Yeatman GW & Riccardi VM: **Partial trisomy of chromosome 14: (+14q-).** *Birth Defects Orig Art Ser* 12(5):119-124, 1976.

15. Frank J & Riccardi VM: **The 11q- syndrome.** *Hum Genet* 35:241-246, 1977.

16. Riccardi VM & Kleiner B: **Neurofibromatosis - A neoplastic birth defect with two age peaks of severity.** *Birth Defects Orig Art Ser* 13(3c):131-138, 1977.

17. Riccardi VM: **Trisomy 8: An international study of 70 patients.** *Birth Defects Orig Art Ser* 13(3c):171-184, 1977.

18. Sujansky E, Riccardi VM & Matthews AL: **The familial occurrence of the Poland syndrome.** *Birth Defects Orig Art Ser* 13(3C):117- 121, 1977.

19. Riccardi VM & Grum CM: **The prune belly anomaly: Heterogeneity and X-linkage mimicry.** *J Med Genet* 14:266-270, 1977.

20. Riccardi VM, et al.: **The FG syndrome: Further characterization by follow-up, report of a third family and a sporadic case.** *Am J Med Genet* 1:47-58, 1977.

21. Riccardi VM & Bergmann CA: **Anencephaly with incomplete twinning.** *Teratology* 16:137-140, 1977.

22. Francke U, George DL, Brown MG & Riccardi VM: **Gene dose effect: Intraband mapping of the LDH-A locus using cells from four individuals with different interstitial deletions of 11p.** *Cytogenet Cell Genet* 19:197-207, 1977.

23. Riccardi VM, et al.: **Chromosomal imbalance in the aniridia Wilms tumor association: 11p interstitial deletion.** *Pediatrics* 61:604-610, 1978.

24. Riccardi VM: **Unilateral gluteal hypoplasia and brachysyndactly: Lower extremity counterpart of the Poland anomaly.** *Pediatrics* 61:653-654, 1978.

25. Riccardi VM & Marcus ES: **Congenital hydrocephalus and cerebellar agenesis.** *Clin Genet* 13:443-447, 1978.

26. Riccardi VM, et al.: **Genetic counseling in hospital care.** *Am J Publ Health* 68:652-655, 1978.

27. Riccardi VM, Humbert JR, Peakman D: **Acute leukemia associated with trisomy 8 mosaicism and a familial translocation 46,XY,t(7;20) (p13;p12).** *Am J Med Genet* 2:15-21, 1978.

28. Michels VV & Riccardi VM: **Twin recurrence and amniocentesis: Male and MZ heritability factors.** *Birth Defects Orig Art Ser* 14(6C):201-211, 1978.

29. Riccardi VM, et al.: **The Turner syndrome and the Y- chromosome: Mechanisms of diminished Y-determined maleness.** *Birth Defects Orig Art Ser* 14(6c):123-132, 1978.

30. Riccardi VM: **A geneticist's approach to deafness.** *Volta Rev* 81:9- 4, 1979.

31. Riccardi VM, et al.: **Partial triplication and deletion of 13q: Study of a family presenting with bilateral retinoblastoma.** *Clin Genet* 15:332-345, 1979.

32. Riccardi VM & Forgason J: **Chromosome 8 abnormalities as components of neoplastic and hematologic disorders.** *Clin Genet* 15:317-326, 1979.

33. Hittner HM, Riccardi VM & Francke U: **Aniridia caused by a heritable chromosome 11 deletion.** *Ophthalmology* 86:1173-1183, 1979.

34. Francke U, Holmes LB, Atkins L & Riccardi VM: **Aniridia-Wilms tumor association: Evidence for specific deletion of 11p13.** *Cytogenet Cell Genet* 24:185-192, 1979.

35. Riccardi VM: **Cell-cell interaction as an epigenetic determinant in the expression of mutant neural crest cells.** *Birth Defects Orig Art Ser* 15(8):89-98, 1979.

36. Riccardi VM, et al.: **Neurofibromatosis: Variable expression is not intrinsic to the mutant gene.** *Birth Defects Orig Art Ser* 15(5b):283-289, 1979.

37. Kurtz SM & Riccardi VM: **Nonverbal communication as an element of genetic counseling.** *Birth Defects Orig Art Ser* 15(5c):245-256, 1979.

38. Marcus ES, Fuller B & Riccardi VM: **Triplication of chromosome arm 20p due to inherited translocation and secondary nondisjunction.** *Am J Med Genet* 4:47-50, 1979.

39. Marcus ES, Holcombe JH, Tulchinsky D, Rich RR & Riccardi VM: **Prenatal diagnosis of congenital adrenal hyperplasia.** *Am J Med Genet* 4:201-204, 1979.

40. Riccardi VM & Holmquist GP: **De novo 13q paracentric inversion in a boy with cleft palate and mental retardation.** *Hum Genet* 52:211- 215, 1979.

41. Carter DM, Goldsmith LA, Epstein WL, Alper JC, Elsas LF III, Mahoney MJ, Minor RR & Riccardi VM: **Analysis of research needs and priorities in dermatology. VI. Birth defects and genetic disorders.** *J Invest Dermatol* 73:460-472, 1979.

42. Hittner HM, Riccardi VM, et al.: **Two-step mutation theory for retinoblastoma: Ultrastructural support.** *Documenta Ophthalmol* 48:345-362, 1980.

43. Ferrell RE, Chakravarti A, Hittner HM & Riccardi VM: **Autosomal dominant aniridia: Probable linkage to acid phosphatase-1 on chromosome 2.** *Proc Natl Acad Sci (USA)* 77:1580-1582, 1980.

44. Hittner HM, Riccardi VM, et al.: **Variable expressivity in autosomal dominant aniridia by clinical, electrophysiologic, and angiographic criteria.** *Am J Ophthalmol* 89:531-539, 1980.

45. Ledbetter DH, Riccardi VM, Au W, Wilson DH & Holmquist GP**: Ring chromosome 15. Phenotype, secondary aneuploidy, Ag-NOR analysis and associated chromosome breakage.** *Cytogenet Cell Genet* 27:111- 112, 1980.

46. Riccardi VM & Maragos VA: **The pathophysiology of neurofibromatosis. I. Resistance in vitro to 3-nitrotyrosine.** *In Vitro* 16:706-714, 1980.

47. Riccardi VM: **The pathophysiology of neurofibromatosis. IV. Dermatologic insights into heterogeneity and pathogenesis.** *J Am Acad Dermatol* 3:157-166, 1980.

48. Pang S, Levine LS, Cederqvist LL, Fuentes M, Riccardi VM, et al.: **Amniotic fluid concentrations of delta-5 and delta-4 steroids for fetuses with congenital adrenal hyperplasia due to 21-hydroxylase deficiency and for anencephalic fetuses.** *J Clin Endocrinol Metab* 51:223-229, 1980.

49. Riccardi VM, et al.: **The aniridia-Wilms tumor association: The critical role of chromosome band 11p13.** *Cancer Genet Cytogenet* 2:131-137, 1980.

50. Strobel RJ, Riccardi VM, et al.: **Duplication 11p11.3-11p4.1 due to meiotic crossing over.** *Am J Med Genet* 7:15-20, 1980.

51. Hittner HM, Riccardi VM, et al.: **Genetic heterogeneity of aniridia: Negative linkage data.** *Metab Pediatr Ophthalmol* 4:179-182, 1980.

52. Ledbetter DH, Riccardi VM, et al.: **Deletion of chromosome 15 as a cause of the Prader-Willi syndrome.** *N Engl J Med* 304:325-329, 1981.

53. Cleaver JE, Greene AE, Coriell L & Riccardi VM: **An autosomal dominant inheritance for multiple sunlight-induced malignancy in a patient without abnormalities in DNA repair or replication.** *Cytogenet Cell Genet* 29:122-124, 1981.

54. Lewis RA & Riccardi VM: **Von Recklinghausen neurofibromatosis: I. Prevalence of iris hamartomata.** *Ophthalmology* 88:348-354, 1981.

84. Darby JK, Kidd JR, Pakstis AJ, Sparkes RS, Cann HM, Ferrell RE, Gerhard DG, Riccardi VM, et al.: **Linkage relationships of the gene for the beta-subunit of nerve growth factor (*NGFB*) with other chromosome 1 marker loci.** *Cytogenet Cell Genet* 39:158-160, 1985.

56. Strong LC, Riccardi VM, et al.: **Familial retinoblastoma and chromosome 13 deletion transmitted via an insertional translocation.** *Science* 213:1501-1503, 1981.

57. Riccardi VM: **Cutaneous manifestations of neurofibromatosis. Cellular interaction, pigmentation and mast cells.** *Birth Defects Orig Art Ser* 17(2):129-145, 1981.

58. Ferrell RE & Riccardi VM: **Catalase levels in patients with aniridia and/or Wilms tumor: Utility and limitations.** Cytogenet Cell Genet 31:120-123, 1981.

59. Riccardi VM: **Von Recklinghausen neurofibromatosis.** *N Engl J Med* 305:1617-1628, 1981.

60. Hayman LA, Evans RA, Ferrell RE, Fahr LM, Ostrow P & Riccardi VM: **Familial cavernous angiomas: Natural history and genetic study over a five year period.** *Am J Med Genet* 11:147-160, 1982.

61. Ledbetter DH, Mascarello JT, Riccardi VM, et al.: **Chromosome 15 abnormalities and the Prader-Willi syndrome: A follow-up report of 40 cases.** *Am J Hum Genet* 34:278-285, 1982.

62. Michels VV, Medrano C, Venne VL & Riccardi VM: **Chromosome translocations in couples with multiple spontaneous abortions.** *Am J Hum Genet* 34:507-513, 1982.

63. Riccardi VM, et al.: **Wilms tumor with aniridia/iris dysplasia with apparently normal chromosomes.** *J Pediatr* 100:574-577, 1982.

64. Riccardi VM: **The multiple forms of neurofibromatosis.** *Pediatr Rev* 3:293-298, 1982.

65. Riccardi VM, Riccardi SL: **Von Recklinghausen disease: New Perspectives.** *Tex Med* 78:43-44, 1982.

66. Riccardi VM: **Neurofibromatosis: Clinical heterogeneity.** *Curr Probl Cancer* 7(2):1-34, 1982.

67. Sillence DO, Lachman RS, Jenkins T, Riccardi VM & Rimoin DL: **Spondylohumerofemoral hypoplasia (giant cell chondrodysplasia): A neonatally lethal short-limb skeletal dysplasia.** *Am J Med Genet* 13:7-14, 1982.

68. Riccardi VM: **Early manifestations of neurofibromatosis: Diagnosis and management.** *Comprehensive Therapy* 8(10):35-40, October 1982.

69. Hittner HM, King RA, Riccardi VM, et al.: **Oculocutaneous albinoidism as a manifestation of reduced neural crest derivatives in the Prader-Willi syndrome.** *Am J Ophthalmol* 94:328-337, 1982.

70. Riccardi VM: **Neurofibromatosis in children.** *Cont Ed Fam Phys* 18:565-571, 1983.

71. Horwich A, Riccardi VM & Francke U: **Aqueductal stenosis leading to hydrocephalus: An unusual manifestation of neurofibromatosis.** *Am J Med Genet* 14:577-581, 1983.

72. Cheek WR, Riccardi VM & Laurent JP: **Neurofibromatosis of childhood: Neurosurgical implications.** *Concepts Pediatr Neurosurg* 4:319-334, 1983.

73. Parke JT, Riccardi VM, et al.: **A syndrome of microcephaly and retinal pigmentary abnormalities. without mental retardation in a family with coincidental autosomal dominant hyperreflexia.** *Am J Med Genet* 17:535-594, 1984.

74. Riccardi VM, et al.: **The pathophysiology of neurofibromatosis. Angiosarcoma as a complication.** *Cancer Genet Cytogenet* 12:275-280, 1984.

75. Lewis RA, Gerson LP, Axelson KA, Riccardi VM & Whitford RP: **Von Recklinghausen neurofibromatosis. II. Incidence of optic-nerve gliomata.** Ophthalmology 91:929-935, 1984.

76. Riopelle RJ, Riccardi VM, et al.: **Serum neuronal growth factors in Von Recklinghausen neurofibromatosis.** *Ann Neurol* 16:54-59, 1984.

77. Riccardi VM, et al.: **The pathophysiology of neurofibromatosis. IX. Paternal age as a factor in the origin of new mutations.** *Am J Med Genet* 18:169-176, 1984.

78. Shapiro SD, Abramovitch K, Van Dis ML, Skoczylas LF, Langlais RP, Jorgensen RJ, Young RS & Riccardi VM: **Neurofibromatosis: Oral and radiographic manifestations.** *Oral Surg* 58:493-498, 1984.

79. Reed SD, Hall JG, Riccardi VM, et al.: **Chromosomal abnormalities associated with congenital contractures (arthrogryposis).** *Clin Genet* 27:353-372, 1985.

80. Lebo RV, Cheung MC, Bruce BD, Riccardi VM, et al.: **Mapping parathyroid hormone, beta-globin, insulin and LDH-A genes within the human chromosome 11 short arm by spot blotting sorted chromosomes.** *Hum Genet* 69:316-320, 1985.

81. Darby JK, Feder J, Selby M, Riccardi VM, et al.: **A discordant sibship analysis between beta-NGF and neurofibromatosis.** *Am J Hum Genet* 37:52-59, 1985.

82. Zoghbi HY, Percy AK, Glaze DG, Butler IJ, Riccardi VM: **Reduction of biogenic amine levels in the Rett syndrome.** *N Engl J Med* 313:921- 924, 1985.

83. Scoggin CH, Fisher JH, Shoemaker SA, Morse H, Leigh T & Riccardi VM: **The E7-associated cell surface antigen: A marker for the11p13 chromosomal deletion associated with aniridia-Wilms tumor.** *Am J Hum Genet* 37:883-889, 1985.

84. Darby JK, Kidd JR, Pakstis AJ, Sparkes RS, Cann HM, Ferrell RE, Gerhard DG, Riccardi VM, et al.: **Linkage relationships of the gene for the beta-subunit of nerve growth factor (*NGFB*) with other chromosome 1 marker loci.** *Cytogenet Cell Genet* 39:158-160, 1985.

85. Dunn BG, Ferrell RE & Riccardi VM: **A genetic linkage study in 15 families of individuals with von Recklinghausen neurofibromatosis.** *Am J Med Genet* 22:403-407, 1985.

86. Percy AK, Zoghbi H & Riccardi VM: **Rett syndrome: Initial experience with an emerging clinical entity.** *Brain Dev* 7:300-304, 1985.

87. Greenberg F, Stein F, Gresik MV, Finegold MJ, Carpenter RJ, Riccardi VM & Beaudet AL: **The Perlman familial nephroblastomatosis syndrome.** *Am J Med Genet* 24:101-110, 1986.

88. Van Eys J, Pullen J, Head D, Boyett J, Crist W, Falletta J, Humphrey GB, Jackson J, Riccardi VM & Brock B: **The French-American-British (FAB) classification of leukemia: The Pediatric Oncology Group experience with lymphocytic leukemia.** *Cancer* 57:1046-1051, 1986.

89. Riccardi VM & Elder DW: **Multiple cytogenetic aberrations in neurofibrosarcomas complicating neurofibromatosis.** *Cancer Genet Cytogenet* 23:199-209, 1986.

90. Riccardi VM: **The Rett syndrome: Genetics and the future.** *Am J Med Genet* 24 (Suppl 1):389-402, 1986.

91. Lockwood DH, Riccardi VM, et al.: **Prophase chromosome unique band sequences: Definition and utilization**. *Cytogenet Cell Genet* 42:141-153, 1986.

92. Riccardi VM: **Growth promoting factors in neurofibroma crude extracts.** *Ann NY Acad Sci* 486:206-226, 1986.

93. Darby JK, Goslin K, Riccardi VM, et al.: **Linkage analysis between the nerve growth factor gene and other chromosome 1p markers and disseminated neurofibromatosis.** *Ann NY Acad Sci* 486:304-310, 1986.

94. Riccardi VM: **Neurofibromatosis and the Albright syndrome.** *Dermatol Clin* 5:193-203, 1987.

95. Riccardi VM & Wald JS: **Discounting an adverse maternal effect on neurofibromatosis severity.** *Pediatrics* 79:386-393, 1987.

96. Gebicke-Haerter PJ, Darby JK, Shooter EM, Riccardi VM, et al.: **Apolipoprotein E synthesis in neurofibrosarcoma and schwannoma cell cultures from two individuals with neurofibromatosis.** *Exptl Neurol* 95:323-335, 1987.

97. Spira M & Riccardi VM: **Neurofibromatosis.** *Clin Plast Surg* 14:315-325, 1987

98. Schroeder WT, Chao L-Y, Dao DD, Strong LC, Pathak S, Riccardi VM, et al.: **Nonrandom loss of maternal chromosome 11 alleles in Wilms tumors.** *Am J Hum Genet* 40:413-420, 1987.

99. Riopelle RJ & Riccardi VM: **Neuronal growth factors from tumours of von Recklinghausen neurofibromatosis.** *Can J Neurol Sci* 14:141-144, 1987.

100. Riccardi VM: **Mast cell stabilization to decrease neurofibroma growth: Preliminary experience with ketotifen.** *Arch Dermatol* 123:1011-1016, 1987.

101. Riccardi VM: **Neurofibromatosis.** *Neurol Clin* 5:337-349, 1987.

102. Riccardi VM & Carey JC: **Von Recklinghausen neurofibromatosis genetic linkage studies: Clinical considerations.** *J Med Genet* 24:521-522, 1987.

103. Ferrell RE, Buetow KH, Darby JK, Eichner JE, Murray JC, Smith R, Waziri M, Huson S & Riccardi VM: **Von Recklinghausen neurofibromatosis: A linkage study of candidate and random marker genes.** *J Med Genet* 24:522-524, 1987.

104. Dao DD, Schroeder WT, Chao L-Y, Kikuchi H, Strong LC, Riccardi VM, et al.: **Genetic mechanisms of tumor-specific loss of 11p DNA sequences in Wilms tumor.** *Am J Hum Genet* 41:202-217, 1987.

105. Brown EW, Riccardi VM, et al.: **MR imaging of optic pathways in patients with neurofibromatosis.** *AJNR* 8:1031-1036, 1987.

106. Pettigrew AL, Gollin SM, Greenberg F, Riccardi VM & Ledbetter DH: **Duplication of proximal 15q as a cause of Prader-Willi syndrome.** *Am J Hum Genet* 28:791-802, 1987.

107. Wasserteil V, Bruce S & Riccardi VM: **Non-von Recklinghausen's neurofibromatosis presenting as hemifacial neurofibromas and contralateral cafe au lait spots.** *J Am Acad Dermatol* 16:1090-1096, 1987.

108. Stephens K, Riccardi VM, et al.: **Linkage studies with chromosome 17 DNA markers in 45 Neurofibromatosis-1 families.** *Genomics* 1:353-357, 1987.

109. Riccardi VM & Lewis RA: **Penetrance of von Recklinghausen neurofibromatosis: A distinction between predecessors and descendants.** *Am J Hum Genet* 42:284-289, 1988.

110. Teixeira F, Martinez-Palomo A, Riccardi VM & Fernandez-Diez J: **Vascular changes in cutaneous neurofibromas.** *Neurofibromatosis* 1:5-16, 1988.

111. Riccardi VM: **American paternal age data for selected years from 1876-1981.** *Neurofibromatosis* 1:93-99, 1988.

112. Riccardi VM & Schmickel RD: **Human genetics as a component of medical school curricula: A report to the American Society of Human Genetics.** Am J Hum Genet 42:639-643, 1988.

113. Tam AW, Darby JK & Riccardi VM: **Expression of selected growth factors and oncogenes in neurofibrosarcomas complicating von Recklinghausen disease.** *Neurofibromatosis* 1:69-84, 1988.

114. Riccardi VM: **Guidelines for organizing a comprehensive neurofibromatosis program.** *Neurofibromatosis* 1:105-119, 1988.

115. Lockwood DH, Johnston DA, Riccardi VM & Zimmerman SO: **The use of subchromosome length unique band sequences in the analysis of prophase chromosomes.** *Am J Hum Genet* 44:934-47, 1988.

116. Riccardi VM: **Neurocutaneous syndromes.** *Curr Opinion Neurol Neurosurg* 1:357-60, 1988.

117. Lacson JM, Riccardi VM & Morizot DC: **Possible genetic etiology of damselfish neurofibromatosis (DNF): Genetic differentiation of bicolor damselfish (Pomacentrus partitus) populations.** *Neurofibromatosis* 1:253-9, 1988.

118. Trevathan E, Moser HW, Opitz JM, Percy AK, Naidu S, Holm V, Boring CC, Janssen RS, Yeargin-Allsopp M, Adams MJ, Riccardi VM, et al: **Diagnostic criteria for the Rett syndrome.** *Ann Neurol* 23:425-8, 1988.

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