

## *Curriculum vitae*

**Updated on:** January 26, 2022  
**Name:** Ramakrishna Prasad Koduru  
**Office Address:** 2330 Inwood Rd, Suite EB3.220H  
**Work Phone:** 214-648-0977  
**Work E-Mail:** Prasad.Koduru@utsouthwestern.edu  
**Work Fax:** 214-648-0976  
**Place of Birth:** Mudinepalli, India

### **Education**

Year	Degree (Honors)	Field of Study (Thesis advisor for PhDs)	Institution
1972	B.Sc	Biology	Andhra University, India
1974	M.Sc	Botany (Genetics main)	Andhra University, India
1979	Ph.D	Cytogenetics (Botany)	Andhra University, India
2000	MHA	Health Care Administration	Hofstra University, Hempstead, NY

### **Postdoctoral Training** [*Include residency/fellowship*]

Year(s)	Titles	Specialty/Discipline (Lab PI for postdoc research)	Institution
1979-1981	Postdoctoral	Cytogenetics (MK Rao)	Andhra University, India
1981-1982	Research Associate	Cytogenetics (MK Rao)	Andhra University, India
1982-1983	Sr. Research Associate	Cytogenetics (Independent)	Andhra University, India
1984-1988	Res Associate Fellow	Cancer Genetics (RSK Chaganti)	MSKCC, New York

### **Current Licensure and Certification**

Licensure: New York State Dept. of Health, CQ KODUP1: Cytogenetics, Genetic Testing, Molecular Oncology

Board and Other Certification: American Board of Medical Genetics – Clinical Cytogenetics, 1987  
National provider identification number: 1508145830

### **Honors and Awards**

Year	Name of Honor/Award	Awarding Organization
1974-1977	Junior Res Fellow	CSIR, New Delhi, India
1977-1979	Senior Res Fellow	CSIR, New Delhi, India

**Faculty Academic Appointments**

Year(s)	Academic Title	Department	Academic Institution
1990-1998	Asst. Professor	Pathology	Cornell Univ Medical College, NY
1998-2009	Associate Professor	Pathology	NYU Medical Center, NY
2009-Current	Professor	Pathology	UT Southwestern Medical Center

**Appointments at Hospitals/Affiliated Institutions**

<u>Past</u>			
Year(s)	Position Title	Department/Division	Institution
1988-1996	Director	Cancer Genetics Lab - Pathology	NSUH, Manhasset, NY
1996-2004	Director	Cell Genetics - Pathology	NSUH, Manhasset, NY
2004-2009	Chief	Div. Cell Genetics, Pathology	NS-LIJ Health System, NY
<u>Current</u>			
Year(s)	Position Title	Department/Division	Institution
2009-current	Med. Director	Genetics Diag. Labs, Pathology Cytogenetics and Molecular Genetics	UT Southwestern Medical Cntr

**Other Professional Positions** [*Industry, foundation, private practice*]

Year(s)	Position Title	Institution
2000-2009	Consulting Geneticist, Pathology	WUH, Mineola, NY
1998-2006	Consulting Cytogeneticist, Pathology	NYU Medical Center, NY

**Major Administrative/Leadership Positions** [*Do not include Professional Society positions*]

Year(s)	Position Title	Institution
2004-2009	Chief, Division of Cellular Genetics	North Shore LIJ Health System Labs, NY
2000-2009	Chief Consulting Geneticist	Winthrop Univ Hospital, Mineola, NY
2009-	Medical Director, Genetics Diagnostics Labs	UTSWMC, Dallas

**Committee Service** (*Member, unless noted otherwise*) [*Do not include Professional Society positions*]

Year(s)	Name of Committee	Institution/Organization
<u>UTSW</u>		
2010-Current	Residents Education Committee	UTSWMC, Pathology
2012-Current	Fellowship Committee	UTSWMC, Pathology
<u>Hospital</u>		
2016-Current	CLS Continuous education committee	

<u>State/Regional:</u> None		
<u>National/International:</u>		
1996-Current	Inspection team member	CAP
2009-Current	Team leader	
2014-Current	International inspector	CAP

**Professional Societies** [List all society committees, leadership, and course leadership roles here]

Dates	Society Name, member
1985-2009	American Society of Human Genetics
1996-2009	American Society of Hematology
2000-Current	Association for Molecular Pathology
2012-Current	American College of Medical Genetics
	Committees
	Fellowships
1997-Current	Founding Fellow American College of Medical Genetics
2000-2004	American College of Health Care Executives

**Grant Review Activities**

Year(s)	Name of Review Committee	Organization
1990-2006	Institutional Research Grant Review	NSUH, NY

**Editorial Activities**

Year(s)	Journal Name
<u>Editor/Associate Editor:</u> None	
<u>Editorial Board:</u> None	
<u>Ad Hoc Reviewer</u>	
2009-Current	Cancer Genetics
2012-Current	Leukemia and Lymphoma
2006-2008	American Journal of Surgical Pathology
2017-current	International Journal of Cancer Research and Molecular Mechanisms
2017-current	The Application of Clinical Genetics

**Grant Support**

<u>Present</u>	<b>Grantor:</b>
----------------	-----------------

	<i>Title of Project:</i>
	<i>Role (Principal Investigator, Co-Investigator):</i>
	<i>Annual amount and date (direct costs only):</i>
	<i>Total amount of award (if multi-year) and dates (direct costs only):</i>

<u>Past</u>	<i>Grantor:</i>
	<i>Title of Project:</i>
	<i>Role (Principal Investigator, Co-Investigator):</i>
	<i>Annual amount and date (direct costs only):</i>
	<i>Total amount of award (if multi-year) and dates (direct costs only):</i>

**Clinical Trials Activities**

<u>Present</u>	<i>Grantor:</i>
	<i>Title of Project:</i>
	<i>Role (Principal Investigator, Co-Investigator):</i>

<u>Past</u>	<i>Grantor:</i>
	<i>Title of Project:</i>
	<i>Role (Principal Investigator, Co-Investigator):</i>

**Teaching Activities**

Year(s)	Activity
	<u>Medical and graduate school didactic and small group teaching</u>
	Dissertation committees
	<u>Qualifying examination committees</u>
	<u>Committees concerned with medical and graduate student education</u>

<u>Graduate student rotations</u>	
<u>Medical student rotations</u>	
<u>Graduate student trainees</u>	
<u>Postgraduate medical education (graduate &amp; continuing medical education)</u>	
<u>Postdoctoral trainees</u>	

**Invited Lectures** [*Since last promotion/appointment*]

Year(s)	Title	Location
<u>International:</u> None		
<u>National:</u>		
9/2010	Fluorescence in situ hybridization in clinical diagnosis	Pathology, Medical City Hospital, Dallas
4/2013	Cancer biomarkers and personalized cancer care	Pathology, Univ. of Baltimore
10/2013	Unique Genomic Difference Between Diffuse Large Cell lymphoma and Burkitt Lymphoma	Pathology, West Virginia School of Medicine
3/2014	Genetic subtypes in non-Hodgkin lymphoma	Pathology, Winthrop Univ Hospital
4/2017	Genetic tests in personalized medical care	Pathology, Winthrop Univ Hospital
<u>Regional/Local</u>		
1/2011	Genetics in the management of hematopoietic tumors	Hematology/Oncology, UTSWMC
5/2012	Genetics in the care of Neonates	Neonatal-perinatal Medicine, UTSWMC
8/2012	Cancer Genetics	Hematology/Oncology, UTSWMC
7/2012	Prenatal Genetic Diagnosis	OB-GYN, UTSWMC
1/2013	Principles of ALK rearrangements in non-small cell lung cancer	Medical Oncology, UTSWMC
5/2017	Polyploidy and pregnancy loss	OB-GYN, UTSWMC
7/2017	Cancer cytogenetics	Hematology/Oncology, UTSWMC

9/2017	How do we use data from NGS for precision medicine	NW-LIJ Health System, Manhasset, NY
9/2017	New technologies in precision medicine	NYU-Winthrop Hospital, Mineola, NY

**Technological and Other Scientific Innovations:** None

Innovation
Patent, if any, pending or awarded /If described in print/on web, provide citation

**Service to the Community**

Year(s)	Role	Organization or institution
May include a brief, one-sentence description of each role if needed (optional)		

2010-2012	HOA Board Director	Chateaus of Coppell
2014-2016	HOA Board Director	Chateaus of Coppell
2010-2014	Member	Hari Hara Peetham

**Bibliography**

**Peer-Reviewed Publications** (List in chronological order with complete pagination. Authors should be listed in the same order as they appear in the published article. Do not include abstracts or submitted works.)

Original Research Articles

1. Koduru, P.R.K., Krishna Rao, M. 1978. Chromosome pairing and desynapsis in spontaneous autopolyploids of *Pennisetum typhoides*. Cytologia 43: 445-452.
2. Krishna Rao, M., Koduru, P.R.K. 1978. Asynapsis and spontaneous centromeric breakage in an inbred line of *Pennisetum typhoides* (L.) Leeke. Proc. Ind. Acad. Sci. B. 87: 29-35.
3. Krishna Rao, M., Koduru, P.R.K. 1978. Cytogenetics of a factor for formation and male sterility in *Pennisetum americanum*. Theor. Appl. Genet. 53: 1-7.
4. Krishna Rao, M., Koduru, P.R.K. 1978. Inheritance of genetic male sterility in *Pennisetum americanum* (L.) Leeke. Euphytica 27: 777-785.
5. Krishna Rao, M., Koduru, P.R.K. 1979. Bi-parental plastid inheritance in *Pennisetum americanum*. J. Heredity 69: 327-330.
6. Koduru, P.R.K., Krishna Rao, M. 1979. Inheritance of anthocyanin and way stem in *Pennisetum*. J. Cytol. Genet. 14: 18-21.
7. Krishna Rao, M., Koduru, P.R.K. 1979. Genetics of five hairy phenotypes and a linkage group of *Pennisetum americanum*. Euphytica 28: 445-451.
8. Lakshmi, K.V., Murthy, T.G.K., Koduru, P.R.K. 1979. Cytogenetic behaviour and estimation of phosphate and potassium content in desynaptic pearl millet. Theor. Appl. Genet. 55: 189-190.
9. Koduru, P.R.K. 1979. Metaphase I centromere co-orientation in interchange heterozygotes of pearl millet. Genet Res. (Camb.) 34: 69-74.

10. Koduru, P.R.K., Krishna Rao, M. 1980. Spontaneous chlorophyll mutants of *Pennisetum americanum*: Genetics and chlorophyll quantities. Theor. Appl. Genet. 56: 137-143.
11. Koduru, P.R.K. 1980. Chromosome pairing and the meiotic behavior of univalents in synaptic mutants of pearl millet, *Pennisetum americanum* (L.) Leeke, Graminae. Genetica 54: 191-197.
12. Koduru, P.R.K., Murthy, T.G.K., Lakshmi, K.V. 1980. Sectional translocation monosomy in a plant of pearl millet, *Pennisetum americanum* (L.) Leeke. Chromosoma (Berl.) 78: 365-370.
13. Koduru, P.R.K., Lakshmi, K.V., Murthy, T.G.K. Chromosome behavior in trisomic plants of pearl millet, *Pennisetum americanum* (L.) Leeke. Beitr. Biol. Pflanzen. 55: 289-297.
14. Lakshmi, K.V., Koduru, P.R.K., Murthy, T.G.K., Rao, M.K. 1982. The effect of trisomy on the meiotic behavior of interchange complexes in pearl millet, *Pennisetum americanum* (L.) Leeke. Theor. Appl. Genet. 61: 333-335.
15. Koduru, P.R.K., Murthy, T.G.K., Lakshmi, K.V., Krishna Rao, M. 1982. Analysis of chromosome pairing and breakage in pearl millet. Genet. Res. (Camb.) 40: 165-174.
16. Koduru, P.R.K., Grace, R.J., Krishna Rao, M. 1982. Genetic and pigment analysis of a yellow virescent mutant of pearl millet, *Pennisetum americanum* (L.) Leeke. Beitr. Biol. Pflanzen. 57: 431-438.
17. Krishna Rao, M., Aswani Kumari, K., Koduru, P.R.K. 1983. Rate of early seed development in two heterotic hybrids and their inbred lines of pearl millet, *Pennisetum americanum* (L.) Leeke. Zeitschrift Pflanzenzuchtg 91: 25-35.
18. Krishna Rao, M., Subba Rao, B., Koduru, P.R.K. 1984. Genetic analysis of an induced double mutant system in pearl millet, *Pennisetum americanum* (L.) Leeke. Biol. Zbl. 103: 295-306.
19. Koduru, P.R.K. Krishna Rao, M. 1984. Cytogenetics of a semi-dwarf mutant in pearl millet, *Pennisetum americanum* (L.) Leeke. Can. J. Genet. Cytol. 26: 272-278.
20. Krishna Rao, M., Aswani Kumari, K., Uma Devi, K., Koduru, P.R.K. 1994. Some aspects of cell development in the young seed of pearl millet, *Pennisetum americanum* (L.) Leeke. Proc. Natl. Sem. Bot. Res., p83-89, Chandigarh.
21. Koduru, P.R.K., Krishna Rao, M., Lakshmi, K.V., Sybenga, J. 1984. Anther development and the orientation of an interchange quadrivalent in pearl millet. Chromosoma (Berl.) 90: 89-93.
22. Koduru, P.R.K. 1984. Centromere orientation, reorientation, and segregation in an interchange quadrivalent during anther development in pearl millet. Chromosoma (Berl.) 90: 94-102
23. Koduru, P.R.K. 1984. Metaphase I orientation of chain forming interchange quadrivalents - a theoretical consideration. Genetics 108: 707-718.
24. Koduru, P.R.K. Grace, R.J., Krishna Rao, M. 1985. Karyotype, heterochromatin content and meiotic features of *Poecilocera picta* (Orthoptera, Acrididae). Genetica 67: 31-37.
25. Burns J.P., Koduru, P.R.K. Alonso, M.L., Chaganti, R.S.K. 1986. Analysis of meiotic segregation in a man heterozygous for two reciprocal translocations using hamster in-vitro fertilization system. Am. J. Hum. Genet. 38: 954-964.
26. Chaganti, R.S.K., Balazs, I. Jhanwar, S.C., Murthy, T.G.K., Koduru, P.R.K., Grzrschik, K.-H., Stavnenzer, E. 1986. C-ski, the cellular homologue of the transforming gene of SKV avian retrovirus maps to human chromosome band 1q22-24, a site of specific translocations in tumors. Cytogenet. Cell Genet. 43: 181-197.
27. Doucette, L.A., Koduru, P.R.K., Long, L., Filippa, D. A., Chaganti, R.S.K. 1986. Molecular detection of the 14;18 translocation in non-Hodgkin's lymphoma. Am. J. Hum. Genet. 38: 197.
28. Koduru, P.R.K., Filippa, D. A., Richardson, M.E., Jhanwar, S.C., Chaganti, S.C., Koziner, B., Clarkson, B.D., Lieberman, P.H., Chaganti, R.S.K. 1987. Cytogenetic and histologic correlation in malignant lymphoma. Blood 69: 97-102.
29. Alonso, M.L., Richardson, M.E., Metroka, C.E., Mouradian, J.A., Koduru, P.R.K., Filippa, D.A., Chaganti, R.S.K. 1987. Chromosome abnormalities in AIDS-related lymphadenopathy. Blood 69: 855-858.
30. Richardson, M.E., Quanquang, C., Filippa, D.A., Offit, K.O., Hampton, A., Koduru, P.R.K., Jhanwar, S.C., Lieberman, P.H., Clarkson, B.D., Chaganti, R.S.K. 1987. Intermediate to high grade histology lymphomas carrying t(14;18) is associated with additional nonrandom chromosome changes. Blood 70: 444-447.
31. Chaganti, R.S.K., Koduru, P.R.K. 1987. Patterns of chromosome breakage in non-Hodgkin's lymphoma: significance of gene alteration in tumorigenesis. Cytogenet. Cell Genet. 45: 93-98.

32. Carroll, P., Morse, M.J., Koduru, P.R.K., Chaganti, R.S.K. 1988. Testicular germ cell tumors in a patient with Klinefelter syndrome. Urology 31: 72-74.
33. Koduru, P.R.K., Chaganti, R.S.K. 1988. Congenital chromosome breakage clusters within giemsa light bands and identifies sites of chromosome instability. Cytogenet. Cell Genet. 49: 269-274.
34. Koduru, P.R.K., Chaganti, R.S.K. 1989. Meiotic chromosome segregation in human t(11;22)(q23;q11) carriers: a theoretical consideration. Genome 32: 24-29.
35. Offit, K., Richardson, M.E., Quanaquang, C., Hampton, A., Koduru, P.R.K., Jhanwar, S.C., Filippa, D.A., Lieberman, P.H., Clarkson, B., Chaganti, R.S.K. 1989. Non-random chromosome aberrations are associated with sites of involvement in non-Hodgkin's lymphoma. Cancer Genet. Cytogenet. 37: 85-93.
36. Offit, K., Koduru, P.R.K., Hollis, R., Filippa, D.A., Jhanwar, S.C., Clarkson, B., Chaganti, R.S.K. 1989. 18q21 rearrangement in diffuse large cell lymphoma: incidence and clinical significance. Br. J. Haematol. 72:178-193.
37. Koduru, P.R.K., DiCostanzo, D., Jhanwar, S.C. 1989. Nonrandom cytogenetic changes characterize Merkel cell carcinoma. Dis. Mar. 7: 153-161.
38. Koduru, P.R.K., Offit, K., Jhanwar, S.C. 1989. Molecular analysis of structural chromosome changes affecting chromosome band 11q23. Dis. Mar. 7: 145-152.
39. Koduru, P.R.K., Offit, K., Filippa, D.A. 1989. Molecular analysis of breaks in BCL-1 proto-oncogene in B-cell lymphomas with abnormalities of 11q13. Oncogene 4: 929-934.
40. Koduru, P.R.K., Offit, K., Filippa, D.A., Lieberman, P.H., Jhanwar, S.C. 1989. Cytogenetic and molecular genetic analysis of abnormal cells in Hodgkin's disease. Cancer Genet. Cytogenet. 43: 109-119.
41. Sun, T., Susin, M., Desner, M., Pergolezzi, R., Cuomo, J., Koduru, P. 1990. The clonal origin of two cell populations in Richter's syndrome. Hum. Pathol. 21: 722-728.
42. Chaganti, R.S.K., Koduru, P.R.K., Chakraborty, R., Jones, W.B. 1990. Genetic origin of trophoblastic carcinoma. Cancer Res. 50: 6330-6333.
43. Koduru, P.R.K. 1991. Molecular structure of double reciprocal translocations: significance in B-cell lymphomagenesis. Oncogene 6: 145-148.
44. Koduru, P.R.K., Goh, J.C., Allen, S., Karp, L., Jasti, H., DeMarco, L., Lichtman, S.M. 1991. Different patterns of chromosome and molecular breakage in classic Ph1 chronic myelogenous leukemias (CMLs) and variant Ph1 CMLs. Hematol. Pathol. 5: 57-66.
45. Sun, T., Cohen, N.S., Marino, J., Koduru, P., Cuomo, J., Henshall, J. 1991. CD3+, CD4-, CD8- large granular T-cell lympho-proliferative disorder. Am. J. Hematol. 37: 173-178.
46. Foti, A., Ahuja, H.G., Allen, S., Koduru, P., Schuster, M.W., Schulman, P., Bar-Eli, M., Cline, M.J. 1991. Correlation between molecular and clinical events in the evolution of CML to blast crisis. Blood 77: 2441-2444.
47. Sun, T., Susin, M., Koduru, P., Coffey Jr. E.L., Weiss, R., Dittmar, K., Goh, J., Brody, J. 1991. Extranodal blast crisis in chronic myelogenous leukemia. Demonstration of T-cell lineage and Philadelphia chromosome in a paraspinal tumor. Cancer 68: 605-610.
48. Koduru, P.R.K., Goh, J.C., Broome, J.D. 1992. Novel restriction fragment length polymorphism in the cellular oncogene SEA. Hematol. Pathol. 6: 71-78.
49. Sun, T., Schulman, P., Koltz, J., Susin, M., Brody, J., Koduru, P., Muse, W., Hombal, S., Teichberg, S., Broome, J. 1992. A study of lymphoma of large granular lymphocytes with modern modalities: Report of two cases and review of literature. Am. J. Hematol. 40: 135-145.
50. Sun, T., Brody, J., Koduru, P., Vinciguerra, V., Weiselberg, L., Marino, J., Chaudhri, F., Pappas, J. Erickson, R. 1992. Study of the major phenotype of the large granular T-cell lympho-proliferative disorder. Am. J. Clin. Pathol. 98: 516-521.
51. Sun, T., Susin, M., Koduru, P., Dittmar, K., Yamnopoulos, K., Mahapatro, D., Rogers, C. 1992. Phenotyping and genotyping of composite lymphoma with KI-1 component. Hematol. Pathol. 6: 179-192.
52. Lichtman, S., Brody, J., Kaplan, M., Susin, M., Koduru, P., Goh, J.C. 1993. Hodgkin's disease and non-Hodgkin's lymphoma in an HIV positive patient. Leukemia Lymphoma 9: 393-398.
53. Koduru, P., Susin, M., Schulman, P., Cattell, D., Goh, J., Karp, L., Broome, J.D. 1993. Phenotypic and genotypic characterization of Hodgkin's disease. Am. J. Hematol. 44: 117-124.



54. Koduru, P.R.K., Lichtman, S.M., Smilari, T.F., Sun, T., Hall, W, Chiorazzi, N. Goh, J., Karp, L., Hassimoto, S., Broome, J.D. 1993. Serial phenotypic and genotypic studies in Richter's syndrome: Demonstration of lymphomagenesis through de-differentiation in chronic lymphocytic leukemia cells. Br. J. Haematol. 85: 613-616.
55. Koduru, P.R.K., Lichtman, S.M., Broome, J.D., Goh, J.C., Pergolizzi, R., Schuster, M., Schulman, P. 1993. Molecular analysis of a variant t(9;22;11)(q34;q11;q13) reveals the translocation of the 3'-part of *bcr* to 11q23. Oncogene 8: 3239-3247.
56. Sun, T., Brody, J., Susin, M., Marino, J., Teichberg, S., Koduru, P., Hall, W.W., Urmacher, C., Hajdu, S.H. 1993. Aggressive natural killer cell lymphoma/leukemia. A recently recognized clinicopathologic entity. Am. J. Surg. Pathol. 17: 1289-1299.
57. Morzek, K. Arthur, D.C., Karakousis, C.P., Koduru, P.R.K., Le Beau, M.L., et al., 1995. Der(16)t(1;16) is a nonrandom secondary chromosome aberration in many types of human neoplasia, including myxoid liposarcoma, rhabdomyosarcoma and Philadelphia chromosome-positive acute lymphoblastic leukemia. Int. J. Oncol. 6: 531-538.
58. Koduru, P.R.K., Susin, M., Kolitz, J., Soni, M., Teichberg, T., Siques, M.J., Sun, T., Amorosi, E., Budman, D.R. 1995. Morphological, ultrastructural, and genetic characterization of an unusual T-cell lymphoma in a patient with sinus histiocytosis with massive lymphadenopathy. Am. J. Hematol. 48: 192-200.
59. Gong, J.Z., Zhou, H., Hu, Z., Schulman, P., Vinciguerra, V., Broome, J.D., Koduru, P.R.K. 1995. Absence of Somatic changes in p21 gene in non-Hodgkin's lymphoma and chronic myelogenous leukemia. Hematol. Pathol. 9: 171-177.
60. Koduru, P.R.K., Zariwala, M. Soni, M., Gong, J.Z., Xiong, Y., Broome, J.D. 1995. Deletion of CDK4 inhibitor genes p15 and p16 in non-Hodgkin's lymphoma. Blood 86: 2900-2905.
61. Brody, J., Allen, S., Schulman, P., Sun, T., Chan, W.C., Friedman, H.D., Teichberg, S., Koduru, P.R.K., Cone, R.W., Loughran, Jr, T.P. 1995. Acute agranular CD4-positive natural killer cell leukemia. Cancer 75: 2464-2483.
62. Soni, M., Brody, J., Allen, S., Schulman, P., Kolitz, J., Rai, K., Broome, J.D., Koduru, P.R.K. 1996. Clinical and morphological features of cases of trisomy 13 in acute non-lymphocytic leukemia. Leukemia 10: 619-
63. Sun, T., Dittmar, K., Koduru, P., Susin, M., Teichberg, S., Brody, J. 1996. Relationship between hairy cell leukemia variant and splenic lymphoma with villous lymphocytes: Presentation of a new concept. Am. J. Hematol. 51: 282-288.
64. Koduru, P., Raju, K., Vadmal, V., Meneges, G., Susin, M., Kolitz, J., Broome, J. 1997. Correlation between mutation in P53, p-53 expression, cytogenetic, histologic type and duration of survival in patients with B-cell non-Hodgkin's lymphoma. Blood 90: 4078-4091.
65. Morzek, K., Heinonen, K., Lawrence, D., Carroll, A.,J., Koduru, P.R.K., Rao, K.W., Strout, M.P., Mutchson, R.E., Moore, J.O., Mayer, R.J., Schiffer, C.A., Bloomfield, C.D. 1997. Adult patients with de novo acute myeloid leukemia and t(9;11)(p22;q23) have a superior outcome to patients with other translocations involving band 11q23. A Cancer and Leukemia Group B study. Blood 90: 4532-4538.
66. Sun, T., Susin, M., Tomao, F.T., Brody, J., Koduru, P., Hajdu, S.H. 1997. Histiocyte-rich B-cell lymphoma. Hum. Pathol. 28: 1321-1324.
67. Vadmal MS, Smilari TF, Brody JP, Koduru P, Hajdu SI. 1998. Cytodiagnosis of primary effusion lymphoma. A case report. Acta Cytol., 42: 374-376.
68. Sun T, Susin M, Brody J, Tack K, Marsh J, Teichberg S, Koduru P. Schwartz, P. 1998. T-cell lymphom associated with natural killer-like T-cell reaction. Am J Hematol., 57: 331-337.
69. Quain L., Gong, J.Z., Liu, J., Broome, J.D., Koduru, P. 1999. Cyclin-D2 promoter disrupted by t(12;22) (p13;q11) during transformation of chronic lymphocytic leukemia (CLL) to non-Hodgkin's lymphoma. Br. J. Haematol. 106: 477-485.
70. Mrozek K, Prior TW, Edwards C, Marcucci G, Carroll AJ, Snyder PJ, Koduru PRK, Theil KS, Pattenati MJ, Archer KJ, Caligiuri MA, Vardiman JW, Kolitz JE, Larson RA, Bloomfield CD. Comparison of cytogenetic and molecular genetic detection of t(8;21) and inv(16) in a prospective series of adults with de novo acute myeloid leukemia: A cancer and leukemia group B study. J Clin Oncol 19: 2482-2492, 2001.
71. Peng H, Shen N, Qian L, Koduru P, Goodwin LO, Iss J-P, Broome JD. Hypermethylation of CpG islands in Mouse asparagine synthetase gene: Relationship to asparaginase sensitivity in lymphoma cells. Partial

- methylation in aging normal cells. *Br J Cancer*, 85: 930-935, 2001.
72. Demiroglu A, Steer JE, Heath C, Taylor K, Bentley M, Allen SL, Koduru P, Brody JP, Hawson G, Rodwell R, Doody M-L, Carnicero F, Reiter A, Goldman JM, Melo JV, Cross NCP. The t(8;22) in chronic myeloid leukemia fuses BCR to FGFR1: transforming activity and specific inhibition of FGFR1 fusion proteins. *Blood*, 98: 3778-3783, 2001.
  73. Byrd JC, Mrozek K, Dodge RK, Carroll AJ, Edwards CG, Arthur DC, Pettenati MJ, Patil SR, Rao KW, Watson MS, Koduru PR, Moore JO, Stone RM, Mayer RJ, Feldman EJ, Davery FR, Schiffer CA, Larson RA, Bloomfield CD; Cancer and Leukemia Group B (CALGB 8461). Pretreatment cytogenetics abnormalities are predictive of induction success, cumulative incidence of relapse, and overall survival in adult patients with de novo acute myeloid leukemia: results from Cancer and Leukemia Group B (CALGB 8461). *Blood* 100: 4325-4336. 2002.
  74. Byrd, JC, Mrojek, K, Carroll AJ, Pettenati MJ, Arthur DC, Koduru P, Bloomfield CD. Repetitive cycles of high-dose cytarabine benefits patients with acute myeloid leukemia and inv(16)(p13q22): Results from Cancer and Leukemia Group B (CALGB 8461). 2004. *J Clin Oncol*. 22: 1087-1094, 2004.
  75. Marcucci G, Mrozek K, Ruppert AS, Archer KJ, Pettenati MJ, Heerema NA, Carroll AJ, Koduru PRK, Kolitz JE, Sterline LJ, Edwards CG, Anastasi J, Larson RA, Bloomfield CD. Abnormal cytogenetics at date of morphologic complete remission predicts short overall and disease-free survival, and higher relapse rate in adult acute myeloid leukemia: results from Cancer and Leukemia Group B study 8461. *J. Clin. Oncol*. 22: 2410-2418, 2004.
  76. Farag SS, Ruppert AS, Mrozek K, Mayer RJ, Stone RM, Carroll AJ, Powell BL, Moore JO, Pettenati MJ, Koduru PRK, Stamberg J, Baer MR, Block, AMW, Vardiman JW, Kolitz JE, Schiffer CA, Larson RA, Bloomfield CD. Outcome of induction and postremission therapy in younger adults with acute myeloid leukemia with normal karyotype: A cancer and Leukemia Group B study. *J Clin. Oncol*. 23: 482-493, 2005
  77. Messmer BT, Messmer D, Allen SL, Kolitz J, Kuldaker P, Cesar D, Murphy EJ, Koduru P, Ferrarini M, Zupo S, Cutrona G, Damle RN, Wasil, T, Rai KR, Hellerstein MK, Chiorazzi N. In vivo measurements document the dynamic cellular kinetics of chronic lymphocytic leukemia B cells. *J. Clin. Invest*. 115: 755-764, 2005.
  78. Grand FH, Koduru P, Cross NCP, Allen SL. NUP98-LEGDF fusion and t(9;11) in transformed chronic myeloid leukemia. *Leuk. Research* 29: 1469-1472, 2005.
  79. Farag SS, Archer KJ, Mrozek K, Ruppert AS, Carroll AJ, Vardiman JW, Pettenati MJ, Baer MR, Qumsiyeh MB, Koduru PR, Ning Y, Mayer RJ, Stone RM, Larson RA, Bloomfield CD. Pretreatment cytogenetics add to other prognostic factors predicting complete remission and long-term outcome in patients 60 years of age and older with acute myeloid leukemia: results from Cancer and Leukemia Group B 8461. *Blood* 108: 73-73, 2006.
  80. Li L, Gupta S, Bashir T, Koduru PR, Brody J, Allen SL. Serial cytogenetic alterations resulting in transformation of a low-grade follicular lymphoma to Burkitt lymphoma. *Cancer Genet Cytogenet*. 170: 140-146, 2006.
  81. Langer C, Radmacher MD, Ruppert AS, Whitman SP, Paschka P, Mrózek K, Baldus CD, Vukosavljevic T, Liu CG, Ross ME, Powell BL, de la Chapelle A, Kolitz JE, Larson RA, Marcucci G, Bloomfield CD; Cancer and Leukemia Group B (CALGB). High BAALC expression associates with other molecular prognostic markers, poor outcome, and a distinct gene-expression signature in cytogenetically normal patients younger than 60 years with acute myeloid leukemia: a Cancer and Leukemia Group B (CALGB) study. *Blood* 111: 5371-5379, 2008.
  82. Mrózek K, Carroll AJ, Maharry K, Rao KW, Patil SR, Pettenati MJ, Watson MS, Arthur DC, Tantravahi R, Heerema NA, Koduru PR, Block AW, Qumsiyeh MB, Edwards CG, Sterling LJ, Holland KB, Bloomfield CD. Central review of cytogenetics is necessary for cooperative group correlative and clinical studies of adult acute leukemia: the Cancer and Leukemia Group B experience. *Int J Oncol*. 33: 239-244, 2008.
  83. Heerema NA, Byrd JC, Dal Cin PS, Dell' Aquila ML, Koduru PR, Aviram A, Smoley SA, Rassenti LZ, Greaves AW, Brown JR, Rai KR, Kipps TJ, Kay NE, Van Dyke DL; Chronic Lymphocytic Leukemia Research Consortium. Stimulation of chronic lymphocytic leukemia cells with CpG oligodeoxynucleotide gives consistent karyotypic results among laboratories: a CLL Research Consortium (CRC) Study. *Cancer Genet Cytogenet*. 203: 134-140. 2010.

84. Smoley SA, Van Dyke DL, Kay NE, Heerema NA, Dell' Aquila ML, Dal Cin P, Koduru P, Aviram A, Rassenti L, Byrd JC, Rai KR, Brown JR, Greaves AW, Eckel-Passow J, Neuberg D, Kipps TJ, Dewald GW. Standardization of fluorescence in situ hybridization studies on chronic lymphocytic leukemia (CLL) blood and marrow cells by the CLL Research Consortium. *Cancer Genet Cytogenet.* 203: 141-148, 2010.
85. Tirado CA, Chen W, Huang LJ, Laborde C, Hiemenz MC, Valdez F, Ho K, Winick N, Lou Z, Koduru P. Novel JAK2 rearrangement resulting from a t(9;22)(p24;q11.2) in B-acute lymphoblastic leukemia. *Leuk Res.* 34: 1674-1676, 2010.
86. Kritharis A, Brody J, Koduru P, Teichberg S, Allen SL. Acute basophilic leukemia associated with loss of gene ETV6 and protean complications. *J Clin Oncol* 29: 623-626, 2011.
87. Chu CC, Zhang L, Dhayalan A, Agagnina BM, MagiL AR, Fraher G, Didier S, Johnson LP, Kennedy WJ, Damle RN, Yan X-J, Patten PEM, Teichberg S, Koduru P, Kolitz JE, Allen SL, Rai, KR, Chiorazzi N. Torgue teno virus 10 isolated by genome amplification technique from a patient with concomitant chronic lymphocytic leukemia and polycythemia vera. *Mol Med.* 17: 1338-1348, 2011.
88. Tirado CA, Gotway G, Torgbe E, Iyer S, Dallaire S, Appleberry T, Suterwala M, Garcia R, Valdez F, Patel S, Koduru P. Recombinant chromosome 7 in a mosaic 45,X/47,XXX patient. *Am J Med Genet A.* 158A; 206-214, 2012.
89. Walker A, Mrózek K, Kohlschmidt J, Rao KW, Pettenati MJ, Sterling LJ, Marcucci G, Carroll AJ, Bloomfield CD, Crawford J, Bigner SH, Budman DR, Koduru PR, Bloomfield CD, Heerema NA, Levine EG, Block AW, Burstein HJ, Tantravahi R, Dal Cin P, Bartlett NL, Watson MS, Crawford EC, Garcia-Heras J, Hurd DD, Pettenati MJ, Vaena DA, Patil SR, Kindler HL, Le Beau MM, Peterson BA, Arthur DC, Nattam S, Bader PI, Graziano SL, Stein CK, Parker BA, Bernstein R, Edelman MJ, Testa JR, Lister J, Diggans GR, Diasio R, Carroll AJ, Dragnev K, Wurster-Hill DH, Grunberg SM, Tang M, Roberts JD, Jackson-Cook C, Rafla S, Verma RS, Freter CE, Miles JH, Shea TC, Rao KW, Sikov W, Padre-Mendoza T, Niell HB, Tharapel SA, Ganti A, Sanger WG, Ryan C, Richkind KE, Grubbs SS, Borgaonkar DS, Van Echo D, Richkind KE. New recurrent balanced translocations in acute myeloid leukemia and myelodysplastic syndromes: cancer and leukemia group B 8461. *Genes Chromosomes Cancer.* 52(4):385-401,2013.
90. Wei Y, Zou Z, Becker N, Anderson M, Sumpter R, Xiao G, Kinch L, Koduru P, Christudass CC, Veltri RW, Grishin NV, Peyton M, Minna J, Bhagat G, Levine B. EGFR-mediated phosphorylation of Beclin-1 in autophagy suppression, tumor progression and tumor chemoresistance. *Cell,* 154: 1269-1284, 2013.
91. Kumar KR, Koduru P, Timmons C, Monaghan S, Cavalier ME, Luu HS. Myelodysplastic syndrome (MDS)-associated cytogenetic abnormalities in pediatric chronic myelogenous leukemia. *Pediatr Blood Cancer,* 2013.
92. Owusu-Brackett N, Johnson R, Schindel DT, Koduru P, Cope-Yokoyama S. A novel ALK rearrangement in an inflammatory myofibroblastic tumor in a neonate. *Cancer Genet* 206: 353-356, 2013.
93. Dang DN, Morris HD, Feusner JH, Koduru P, Wilson K, Timmons CF, Cavalier M, Luu HS. Therapy-induced secondary acute myeloid leukemia with t(11;19)(q23;p13.1) in a pediatric patient with relapsed acute promyelocytic leukemia. *J Pediatr Hematol Oncol.* 2014 Nov;36(8):e546-8. doi: 10.1097/MPH.0000000000000183.
94. Wang L, Peters JM, Fuda F, Li L, Karandikar NJ, Koduru P, Wang HY, Chen W. Acute megakaryoblastic leukemia associated with trisomy 21 demonstrates a distinct immunophenotype. *Cytometry B Clin Cytom.* 2014 Oct 31. doi: 10.1002/cyto.b.21198.
95. Kim J, Bu L, Koduru PR, Wilson KS, Fuda FS, Kumar KR, Timmons CF, Slone TL, Luu HS. Novel r(2)(p25q31) Cytogenetic Abnormality in a Pediatric Patient With Acute Leukemia of Ambiguous Lineage. *Pediatr Dev Pathol.* 18: 76-79, 2015.
96. Kasten J, Chen W, Fuda F, Koduru P, Monaghan SA. Diffuse large B-cell lymphoma with prominent cytoplasmic vacuoles involving peripheral blood, bone marrow and cerebrospinal fluid (2014).” Case Reports with Multiple Choice Questions, *BloodMed.Com.*
97. Li L, Puliappadamba VT, Chakraborty S, Rehman A, Vemireddy V, Saha D, Souza RF, Hatanpaa KJ, Koduru P, Burma S, Boothman DA, Habib AA. EGFR wild type antagonizes EGFRvIII-mediated activation of Met in glioblastoma. *Oncogene* 34: 129-134, 2015.
98. Dennis J, Parsa R, Chau D, Koduru P, Peng Y, Fang Y, MD, PhD, Sarode VR. Quantification of Human Epidermal Growth Factor Receptor 2 Immunohistochemistry using the Ventana Image Analysis System:

- Correlation with Gene Amplification by Fluorescence in Situ Hybridization: The importance of instrument validation for achieving high (>95%) concordance rate. *Am J Surg Pathol* 2015.
99. Kumar KR, Chen W, Koduru PR, Luu HS. Myeloid and lymphoid neoplasm with abnormalities of FGFR1 presenting with trilineage blasts and RUNX1 rearrangement: a case report and review of literature. *Am J Clin Pathol*. 2015 May;143(5):738-748.
  100. Witkiewicz AK, McMillan EA, Balaji U, Baek G, Lin WC, Mansour J, Mollae M, Wagner KU, Koduru P, Yopp A, Choti MA, Yeo CJ, McCue P, White MA, Knudsen ES. Whole-exome sequencing of pancreatic cancer defines genetic diversity and therapeutic targets. *Nat Commun*. 2015 Apr 9;6:6744. doi: 10.1038/ncomms7744.
  101. Hatanpaa KJ, Fuda F, Koduru P, Young K, Lega B, Chen W. Lymphomatosis Cerebri: A Diagnostic Challenge. *JAMA Neurol*. 72:1066-1067, 2015.
  102. Koduru PR, Chen W, Garcia R, Fuda F. Acquisition of a t(11;14)(q13;q32) in clonal evolution in a follicular lymphoma with a t(14;18)(q32;q21) and t(3;22)(q27;q11.2). *Cancer Genet*. 208:303-309, 2015.
  103. Doshi S, Ray D, Stein K, Zhang J, Koduru P, Fogt F, Wellmann A, Wat R, Mathews C. Economic Analysis of Alternative Strategies for Detection of ALK Rearrangements in Non Small Cell Lung Cancer. *Diagnostics* 2016, 6, 4; doi:10.3390/diagnostics6010004.
  104. Ramirez M, Rajaram S, Steininger III RJ, Osipchuk D, Roth MA, Morinishi LS, Evans L, Ji W, Hsu C-H, Thurley K, Wei S, Zhou A, Koduru PR, Posner BA, Wu LF, Altschuler SJ. Diverse drug-resistance mechanisms can emerge from drug-tolerant cancer persister cells. *Nature Communications*, 2016.
  105. Koduru P, Guruju N, Patel P, Wen J, Wilson K, Monaghan S. A unique rearrangement of *PDGFRα* and *ETV6* in a patient with acute myeloid leukemia with myelodysplasia-related changes progressed from chronic myelomonocytic leukemia. *Hematology and Leukemia* 2016, doi: 10.7243/2052-434X-4-1.
  106. Chen P, Chu A, Zia H, Koduru P, Collins R, Winick N, Fuda F, Chen W. CD25 Expression in B Lymphoblastic Leukemia/Lymphoma Predicts t(9;22)(q34;q11)/Philadelphia Chromosome Translocation (Ph) and Is Associated With Residual Disease in Ph-Negative Patients. *Am J Clin Pathol*. 2016;146:632-638.
  107. Gehlbach D, Koduru P, John G, Fuda F, Frankel AE, Chen W. Blastic plasmacytoid dendritic cell neoplasm with t(11;19)(q23;p13.3);*KMT2A(MLL)* rearranged: a diagnostic challenge. *Br J Haematol* 176:8, 2017.
  108. Koduru P, Wilson K, Wen J, Garcia R, Patel S, Monaghan SA. Cytogenetic and cytogenomic microarray characterization of chromothripsis in chromosome 8 affecting *MOZ/NCOA2 (TIF2)*, *FGFR1*, *RUNX1T1*, and *RUNX1* in a pediatric acute myeloid leukemia. *J Pediatr Hematol Oncol* 39(4):e227-e232. doi:10.1097/MPH.0000000000000770. PMID: 28085746
  109. Babu R, Van Dyke DL, Dev VG, Koduru P, Rao N, Mitter N, Liu M, Fuentes E, Fuentes S, and Papa S. Interphase Chromosome Profiling (ICP): a method for conventional banded chromosome analysis using interphase nuclei. *Arch Pathol Lab Med*. 2017, 142:213-228. Oct 5. doi: 10.5858/arpa.2016-0621-OA.
  110. Babu R, Van Dyke DL, Bhattacharya S, Dev VG, Liu M, Kwon M, Gu G, Koduru P, Rao N, Williamson C, Fuentes E, Fuentes S, Papa S, Kopuri S, Lal V. A rapid and reliable chromosome analysis method for products of conception using interphase nuclei. *Mol Genet Genomic Med* 2018; 1012, DOI: 10.1002/mgg3.381. PMID: 29573570
  111. García R, Chen W, Koduru P. Clinical impact of MYC abnormalities in plasma cell myeloma. *Cancer Genet*. 2018 Dec;228-229:115-126. doi: 10.1016/j.cancergen.2018.10.007. Epub 2018 Oct 30. PMID: 30553464.
  112. Alsuwaidan A, Pirruccello E, Jasso J, Koduru P, Garcia R, Krueger J, Doucet M, Chaudhry R, Fuda F, Chen W. Bright CD38 expression by flow cytometric analysis is a biomarker for double/triple hit lymphomas with a moderate sensitivity and high specificity. *Cytometry B Clin Cytom* 2019. PMID 30734478.
  113. Sifuentes-Domingues L, Starokadomsky P, Welch J, Gurram B, Park J, Koduru P, Burtein. Mosaic tetrasomy 9p associated with inflammatory bowel disease. *J Crohns Colitis* 2019 Apr 25. Pii:jjz079. Doi: 10.1093/ecco-jcc/jjz079. PMID 31104071
  114. Koduru P, Chen W, Haley B, Ho K, Oliver D, Wilson K. Cytogenomic characterization of double minute heterogeneity in therapy related acute myeloid leukemia. *Cancer Genet*. 8: 69-75, 2019. PMID: 31425928.

115. Webb C, Partain N, Koduru P, Hwang H, Sarode V. Secondary angiosarcoma with C-MYC amplification following prophylactic bilateral mastectomy and autologous breast reconstruction: Report of a case and review of the literature.2020 DOI: 10.1177/1066896920930100 . PMID 32552130.
116. Churchill HRO, Fuda F, Xu J, Deng M, Zhang CC, An Z, Zhang N, Chen P, Bergstrom C, Kansagra A, Collins R, John S, Koduru P, Chen W. Leukocyte Immunoglobulin-like Receptor B1 and B4 (LILRB1 and LILRB4): Highly Sensitive and Specific Markers of Acute Myeloid Leukemia with Monocytic Differentiation. Cytometry B Clin Cytom. 2020. Doi: 10.1002/cyto.b.21952. PMID 32918786
117. Montgomery-Goecker C, Koduru P, Botten G, Xu J, Ghisoli M, Goldman SC, Krueger JE, Bhushan V, Fuda F, Chen W. Mixed Phenotype Acute Leukemia, B/myeloid (Bilineal and Biphenotypic), with t(2;22)(q35;q12);EWSR1-FEV. J Ped Hematol Oncol 2020. doi: 10.1097/MPH.0000000000001934. PMID 32925408
118. Germans SK, MD, Kulak O, Prasad Koduru P, Oliver D, Gagan J, Patel P, Anderson Jr LD, Fuda FS, Chen W, Manuel Jaso JM. Lenalidomide-Associated Secondary B-Lymphoblastic Leukemia/Lymphoma—A Unique Entity A Report of Two Cases With Low-Level Bone Marrow Involvement and Review of the Literature. Am J Clin Pathol 2020;XX:1-12 DOI: 10.1093/ajcp/aqaa109. PMID 32880627
119. Li H-D, Lu C, Zhang H, Hu Q, Zhang J, Cuevas IC, Sahoo SS, Mitzi Aguilar M, Elizabeth G. Maurais EG, Zhang S, Wang X, Akbay EA, Li G-M, Li B, Koduru P, Ly P, Fu Y-X, Castrillon DH. A *Pole*<sup>P286R</sup> mouse model of endometrial cancer recapitulates high mutational burden and immunotherapy response. JCI Insight 2020. July 23;5(14):e138829.doi:10.1172/ichi.insight.138829. PMID 32699191.
120. Bishop JA, Koduru P, Veremis BM, Weinreb I, Rooper IM, Dickson BC, Demicco EG. SS18 Break-Apart fluorescence In situ hybridization is a practical and effective method for diagnosing microsecretor adenocarcinoma of salivary glands. Head and Neck Pathology, 2021. DOI: 10.1007/s12105-020-1280- PMID: 33394377.
121. Murphy KM, Carrick K, Gwin K, Rogers V, Koduru P, Ronnett BM, Castrillon DH. Rare complete hydatidiform mole with p57 expression in villous mesenchyme: Case report and review of discordant p57 expression in hydatidiform moles. Int J Gynecol Pathol 2021; doi: 10.1097/PGP.0000000000000773. PMID: 33900230.
122. Garcia R, Hussain A, Koduru P, Wilson K, Park JY, and others. Identification of **potential antiviral** compounds against SARS-CoV-2 structural and non structural protein targets: A pharmacoinformatics study of the CAS COVID-19 dataset. Computers in Biology and Medicine 133:104364. doi: 10.1016/j.combiomed.2021.104364. PMID 33895457.
123. Xu J, Shi G, Kulak O, Chen W, Koduru P, Gagan J. A CLL/SLL case with distinctive molecular and cytogenetic changes during different stage of disease progression. AMP Case Report. CAP Today, March 2021.
124. Rosado FG, Coberly J, Gupta A, John G, Naina H, Koduru P, Chen W. PD1/PD-L1 expression in plasmablastic lymphoma with clinicopathological correlation. Ann Clin Lab Sci 2021, 51:174-181. PMID: 33941556.
125. Chen P, Redd L, Schmidt Y, Koduru P, Fuda F, Montgomery-Goecker C, Kumar K, Xu-Monette Z, Young K, Collins R, Chen W. MYC protein expression does not correlate with MYC abnormalities detected by

FISH but predicts an unfavorable prognosis in de novo acute myeloid leukemia. *Leuk Res* 2021, 24: 106:106584. doi: 10.1016/j.leukres.2021.106584. PMID: 33933715.

126. Webb C, Partain N, Koduru P, Hwang H, Sarode VR. Secondary angiosarcoma with C-MYC amplification following prophylactic bilateral mastectomy and autologous breast reconstruction: Report of a case and review of the literature. *Int J Surg Pathol* 2021:2021 Apr;29(2):205-210. doi: 10.1177/1066896920930100. PMID 32552130.
127. Lin E; Koduru P; Chen M. Circulating dividing plasma cells with giant dumbbell-shaped nuclei. *eJHaem*. 2021, doi.org/10.1002/jha2.330
128. Cox A, Park C, Koduru P, Wilson K, Weinberg O, Chen W, Garcia R, Kim D Automated classification of cytogenetic abnormalities in hematolymphoid neoplasms. *Bioinformatics*.2021 Dec 7 (PMID: 34874998)
129. Willis KR, A Sathe, C Xing, P. Koduru, M Artunduaga, E Butler, J Park, R Kurmasheva, P Houghton, K Chen, D Rakheja. Extrarenal anaplastic wilms tumor: A case report with genomic analysis and tumor models. *J Ped Hem Oncol*
130. Rakheja D, Park JY, Yang MS, Koduru P, Wilson KS, Garcia R, Uddin N. Epithelioid rhabdomyosarcoma with NSD3::FOXO1 fusion resulting from t(8;13)(p11.2;q14) translocation: evidence for reconsideration of previously reported FOXO1::FGFR1 fusion in rhabdomyosarcoma. *Cancer Genetics*
131. Wick N, Hitto I, Welder D, Slone T, Koduru P, Fuda F, Rakheja D, Weinberg O. Acute myeloid leukemia with RAM immunophenotype presenting with extensive lymphadenopathy: A case report and review of the literature. *Leuk Rese Rep*
132. Panwar V, Tintle SJ, Germens S, Koduru P, Jia L. MYC amplification in epithelioid angiosarcoma of the bladder and prostate (EASBP) following prostate radiotherapy: a case report with a novel molecular alteration. *Int J Surg pathol*

#### Reviews, Chapters, Monographs and Editorials

1. Koduru, P.R.K., Krishna Rao, M. 1981. Cytogenetics of synaptic mutants of higher plants. *Theor. Appl. Genet.* 59: 197-214.
2. Koduru, P.R.K., Krishna Rao, M. Genetics of qualitative traits and linkage studies in pearl millet *Zeitschrift Pflanzenzuchtg* 90: 1-23.

Books/Textbooks: None

#### Case Reports

1. Jabbar S, Weina Chen W, Sara Monaghan S, Koduru, P, Kirthi R. Kumar KR. Acute Myeloid Leukemia with a novel t(7;14)(q21,q32) and poor clinical outcome. A case report. *Lab Med*. 2017; 48:376-380

#### Letters to the Editor

1.	
----	--

Proceedings of Meetings: Abstracts presented before 1998 are not listed

1. Bialer M, McLaughlin JA, Yenamandra A, Koduru P. Partial trisomy 4p and partial tetrasomy 4p, a wide spectrum of anomalies. *Am J Hum Genet*, 1998, abst# 716
2. Yenamandra A, Trinchitella L, Serotikn A, Buckwald S, Brar G, Small K, Koduru, P. Clinical and cytogenetic findings in chromosome 13 abnormalities. *Am J Hum Genet*, 1998, abst# 814
3. Koduru P, Zhou, X, Raymond D, Yenamandra A, Susin M, Broome J. Progressive translocation involving immunoglobulin (IG) heavy chain, BCL-2 and BCL-6 genes in non-Hodgkin's lymphoma (NHL). *Blood* 94s: 176-b-177b, 1999
4. Yenamandra A, Trinchetella L, Haberman G, Perrone R, McKenna C, Bialer M, Koduru P. A novel 21p+ chromosome with duplication of Down syndrome critical region in the amniotic fluid cells of a second trimester pregnancy. *Am J Hum Genet*, 65s: A363, 1999
5. Lundberg JK, Bialer M, Yenamandra A, Schneideer R, Perrone R, Zhou Z, Kahn E, Koduru P. An abnormal fetus with trisomy 20 and trisomy 9 mosaicism. *Am J Hum Genet*, 65s: A350, 1999
6. Mehta L, Cervantes C, Small K, Yenamandra A, Koduru P. Proximal deletion of chromosome 14q. *Am J Hum Genet*, 65s: A351, 1999
7. Koduru P, Schiff R. Yenamandra A, Arnold E, Bialer M. Tricuspid atresia in mosaic tetrasomy 8p. *Am J Hum Genet*, 67s: A161, 2000
8. Yenamandra A, Buckwald S, Schiff R, Cagungun N, Trinchitella L, Koduru P, Mehta L. Tetralogy of fallot and dysmorphic features associated with partial trisomy 8q/monosomy 1q. *Am J Hum Genet.*, 67s, A115, 2000
9. Mehta L, Yenamandra A, Perrone R, Koduru P. De-novo chromosomal insertion in a child with autism. *Am J Hum Genet*, 67s, A115, 2000
10. Hirawat S, Gupta R, Schulman P, Yenamandra A, Koduru P. Translocation 9;14 – a new cytogenetic abnormality in adult acute lymphoblastic leukemia. *J Am Soc Clin Oncol.*, p106, 2000
11. Steer EJ, Demiroglu A, Heath C, Pourgourides E, Taylor K, Bentley M, Allen S, Koduru P, Brody J, Hawson G, Rodwell R, Doody M-L, Carnicero F, Reiter A, Goldman J, Melo J, Cross N. The t(8;22) in chronic myeloid leukemia fuses BCR to FGFR1: transforming activity and specific inhibition of FGFR1 fusion protein. *Blood* 98s: 468b, 2001
12. Brody J, Basham K, Stanek A, Liu J, Koduru P, Cross C, Allen S. Rapid transformation to biphenotypic (B and T-cell) lymphoid blast crisis in a patient with CML associated with t(8;22)(p11;q11) and fusion of BCR to FGFR1. *Blood* 98s: 255b, 2001
13. Koduru P, Orner S, Yenamandra A, Tepperberg J, Bialer M. Dysmorphic features in two neonates with mosaic ring chromosome syndromes. *Am J Hum Genet*, 69s: A313, 2001
14. Yenamandra A, Mehta L, Hentze L, Shanmugham A, Ahmed A, Koduru P. Prenatally diagnosed complex chromosome 4 rearrangement: A counseling dilemma. *Am J Hum Genet*, 69s: A658, 2001
15. Yenamandra A, Hentze L, Krim E, Sadr I, Ahmed A, Fried W, Koduru P. Truncus arteriosus in a 45,X male fetus with SRY-autosome translocation. *Am J Hum Genet*, 71s: A557, 2001
16. Kudalkar P, Yenamandra A, Ludlum S, Koduru P, Kolitz J. Deletion of chromosome 20q in lymphoproliferative disorders. *Blood* 102s: 183, 2003
17. Mehta L, Yenamandra A, Koduru P, McLaughlin J, Schiff R, Vohra N, Rochelson B. Multiple fetal anomalies including holoprosencephaly caused by 7q monosomy/5q trisomy in the offspring of a double translocation carrier. *Am J Hum Genet*, 73s: A592, 2003
18. McKenna C, Heckman L, Schmidt R, Morett M, Yenamandra A, Koduru P, Mehta L. Phenotypic effects and pregnancy outcome of an unbalanced familial 5;21 chromosome translocation. *Am J Hum Genet*, 73s: A304, 2003
19. Koduru P, Batish SD, Dittmar S, Moore C, Mehta, L, Gupta S. Further characterization of a 46,X,abn(X)/45,X karyotype detected prenatally. *Am. Soc Hum. Genet. Abst.* P 199, 2004
20. Gupta, S, Perrone R, Gallo J, Allen SL, Koduru P. Constitutional t(8;22)(q24.1;q11.2): prevalence and clinical significance. *Am. Soc. Hum. Genet. Abst.* P 193, 2004

21. Shanmugham A, Gupta S, Koduru P, Yenamandra A, Mehta L. Familial tandem duplication of chromosome 20p. *Am. Soc. Hum. Genetic. Abst.* P 150, 2004
22. Shah HO, Koduru P, Karnik A, Miller B, Diaz-Barrios V, Zhang Y, Zhuang M, Sherman J, Lin JH. Retinoblastoma in a newborn baby with a heterozygous Rb gene deletion. *Am. Soc. Hum. Genet. Abst* P 81, 2004
23. Farag SS, Archer KJ, Mrozek K, Vardiman JW, Carroll AJ, Pettenati MJ, Powell, BL, Moore JO, Kolitz JK, Baer MR, Bigner SH, Koduru PR, Stamberg J, Mayer RJ, Stone RM, Schiffer CA, Larson RA, Bloomfield CD. Pre-Treatment cytogenetics predict complete remission and long-term outcome in patients (pts)>60 years with acute myeloid leukemia (AML): Results from cancer and leukemia group B (CALGB) 8461. *Blood* 104 S, p 164, 2004
24. Mrozek K, Carroll AJ, Maharry K, Rao KW, Patil SR, Pettenati MJ, Watson MS, Arthur DC, Tantravahi R, Heerema NA, Koduru PR, Bigner SH, Stamberg J, Block AM, Edwards CG, Sterling LJ, Bloomfield CD. Central review of cytogenetics is essential for cooperative group clinical and correlative studies of acute leukemia: The cancer and Leukemia Group B (CALGB) 8461 experience. *Blood* 104s, p307, 2004.
25. Gupta S, Paliou M, Schlosser J, Iglesias A, Koduru P. A 46,XX male with site inversus, dextracardia and Grave's disease.
26. Gupta S, Brody J, John V, Koduru P. Complex translocation (8;16;21): A new variant of t(8;21) with t(13;22) in acute myeloid leukemia. *Am. Soc Hematol Mtg. Blood* 106S; 119b, 2005.
27. Ding Y, Koduru P, Gupta S. Changing signal patterns – Fluorescence in-situ hybridization in monitoring CML. *Am. Soc. Hum. Genet. Mtg.* 2005: p77.
28. Yeboa K, Koduru P, Tambini L, Perrone R, Gupta S. Deletion 13q in infant with IUGR and dysmorphism – a case report. *Am. Soc. Hum. Genet. Mtg.* 2005: p125.
29. Koduru P, Gupta S, Perrone R, Antonelli J, Krietzler P, Fox J. Mild to moderate postnatal abnormalities in a child with del(6)(q25.1q25.2). *Am. Soc. Hum. Genet. Mtg.* 2005: p128.
30. Gupta S, Mehta L, Shanmugham A, Thomas A, Koduru P. Aplasia cutis congenital in an infant with de novo t(1;20)(p34.1;q13.1). *Am. Soc. Hum. Genetic. Mtg.:* 2005: p154.
31. Young KH, Leroy K, Moller MB. Et al. Structural profiles of p53 gene mutations predict clinical outcome in diffuse large B-cell lymphoma: An international collaborative study. *Blood* 108, p243a, 2006.
32. Smoley SA, Van Dyke D, Kary NE., et al. Validation of CLL FISH panel scoring by members of the chronic lymphocytic leukemia research consortium. *Blood* 112, p391, 2008.
33. Trevino Y, Garcia R, Koduru, P. Recurrent cytogenetic aberrations between MYC+ and MYC- groups predict MYC rearrangements in diffuse large B-cell lymphoma: A bivariate and comparison study of cytogenetic data. *AGT*, 2012
34. 34. Kumar K, Koduru P, Timmons C, Monaghan S, Cavalier ME, Luu H. Myelodysplastic-syndrome (MDS)-associated cytogenetic abnormalities in a pediatric case of chronic myelogenous leukemia. *CAP* 2012
35. Hoffines A, Karandikar N, Fuda F, Monaghan S, Koduru P, Levenson B. Greater incidence of cytogenetic aberrations in HIV-positive bob-Hodgkin lymphoma. *USCAP*, March 2012,
36. Hoffines A, Levenson B, Fuda F, Koduru P. A unique MYC rearrangement with a non-immunoglobulin partner occurring in a case of childhood acute B-cell lymphoblastic leukemia with a t(12;21) translocation. *ASCP*, Boston, 2012
37. Tricia Makin, Rolando García, Naga Guruju, Sangeeta Patel, Prasad Koduru. Recurrent Cytogenetic Aberrations in Myelodysplastic Syndrome Subtypes: A Multivariable Association Study of Cytogenetic Data. *AGT meeting*, Los Vegas, 2013
38. Prasad Koduru, Franklin Fuda. A triple-hit follicular lymphoma with t(3;22)(q27;q11),t(11;14)(q13;q32),t(14;18)(q32;q21). *AMP meeting*, Phoenix, AZ, 2013
39. Pediatric Case of Chronic Myelogenous Leukemia Presenting as B-Lymphoblastic Crisis with Monosomy 7. *CAP* 2013.
40. Yan Z, Guruju N, Garcia R, Luu H, Koduru P. Quadruplication of chromosome region 11q13 in multiple myeloma. A case report. *CAP*, 2013



41. Guruju, Garcia R, Wilson S, Monghan S, Koduru P. A Unique Rearrangement of PDGFRA and ETV6 in a Three-way Translocation t(4;12;6) in a Patient with Acute Myeloid Leukemia Progressed from Chronic Myelomonocytic Leukemia. ASHG 2013 Boston.
42. Garcia R, Guruju N, Koduru P. Unique recurrent cytogenetic aberrations distinguish between molecular subtypes of DLBCL and Burkitt lymphoma: An analysis of unsupervised clusters and logistic regression based models. 3335S, ASHG Meeting, 2014, San Diego.
43. Koduru P, Garcia N, Guruju N. Specific gene expression profiles of diffuse large B-cell lymphoma and Burkitt lymphoma identified a set of enriched genes that positively correlates with cytogenetic data. 3337T, ASHG Meeting 2014, San Diego.
44. Babu VR, Dev VG, Koduru P, Rao N, Liu M, Fuentes E, Fuentes S, Papa S, Van Dyke DL. Interphase Chromosome Profiling (ICP): Development and validation of a novel technology and its clinical applications. Cytogenet Genome Res 2014;142:226, Abstract # 22.
45. Daniel Gehlbach, Prasad Koduru, George John, Franklin Fuda, Arthur Frankel, Weina Chen. Blastic Plasmacytoid Dendritic Cell Neoplasm with t(11;19)(q23;p13.3); *MLL-ENL*, a Diagnostic Challenge. Accepted and presented in 2014 CAP meeting
46. Kumar K, Fuda, F, Garcia R, Chen, W, Koduru P. Myelodysplastic syndrome - Refractory anemia with excess blasts (RAEB) involving RUNX1 in a t(13;21)(q22;q22). Abstract. Presented at the 2015 ACMG Annual Clinical Genetics Meeting, March 25, Salt Lake City, UT.
47. Koduru, Strickland A, Garcia R, Monaghan S. Clonal Cytogenetic Abnormality in a Case of IgG4-related Lymphadenopathy. Presented at the 2015 ACMG Annual Clinical Genetics Meeting, March 25, Salt Lake City, UT.
48. Wein J, Garcia R, Wilson K, Koduru P. A clinicocytogenetic study of ocular lymphoma. Presented at the 2015 ACMG Annual Clinical Genetics Meeting, March 25, Salt Lake City, UT.
49. Babu VR, Dev VG, Koduru P, Rao N, Mitter N, Liu M, Fuentes E, Fuentes S, Papa S and Van Dyke DL. Interphase Chromosome Profiling in the Workup of Products of Conception and Hematologic Malignancies. Time to do away with Classical Karyotype? Presented at the 2015 ACMG Annual Clinical Genetics Meeting, March 25, Salt Lake City, UT.
50. Redd L, Schmidt Y, Oliver D, Koduru P, Luu H, Chen W. Detection of IDH1 (R132H) mutations in acute myeloid leukemia (AML) by immunohistochemistry. Abstract \_\_\_\_\_. 2015 USCAP Annual Meeting, March 21, Boston, MA.
51. Wachsmann M, Borja B, Knudsen E, Koduru P, Witkiewicz A. C-MYC amplification in PDA is associated with poor outcome and adenosquamous subtype. Abstract \_\_\_\_\_. 2015 USCAP Annual Meeting, March 21, Boston, MA.
52. Gehlbach D, Koduru P, Redd L, Luu HS, Monaghan SA, Fuda F, Frankel AE, Chen W. Recurrent 11q23 chromosomal abnormalities in blastic plasmacytoid dendritic cell neoplasm: a clinicopathologic study on 6 patients. Accepted to present in 2015 USCAP meeting.
53. Lucas Redd, Prasad Koduru, Yao Schmidt, Crystal Montgomery-Goecker, George John, Kirthi Kumar, Weina Chen. MYC Expression In Acute Myeloid Leukemia (AML) by Immunohistochemistry with FISH Correlation. Accepted to present in 2015 USCAP meeting.
54. Babu R, Koduru P. An interphase chromosome profiling assay to determine the need for CD 138 enrichment in the genetic workup of multiple myeloma. Blood 126: 2015.
55. Dennis J, Zia H, Chen P, Chu A, Koduru P, Luu HS, Fuda F, Chen W. Adult mixed phenotype acute leukemia: a single institution experience and recommendations on further refinement of classification criteria. 2016 USCAP Annual Meeting, March 21, 2016.
56. Chen P, Redd L, Koduru P, Dennis J, Montgomery-Goecker C, John P, Xu-Monette Z, Kumar K, Schmidt Y, Chen W. MYC protein expression does not correlate with MYC abnormalities but predicts an unfavorable prognosis in adult de novo acute myeloid leukemia. USCAP Annual Meeting, March 21, 2016.
57. Pirruccello E, Koduru P, Kruger J, Fuda F, Chen W. Distinct immunophenotypic signature of MYC-single hit lymphoma from MYC-double or triple hit lymphomas. USCAP Annual Meeting, March 21, 2016.

58. Chen P, Chu A, Zia H, Koduru P, Fuda F, Chen W. CD25 expression in B-lymphoblastic leukemia/lymphoma (B-ALL) is a biomarker for the Philadelphia chromosome (Ph) (BCR-ABL1 rearrangement) and an adverse risk factor in Ph negative B-ALL. USCAP Annual Meeting, March 21, 2016.
59. R. García, S. Patel, K. Wilson & P. Koduru., A Cytogenetic Data Analysis Reveals Unique Chromosome Abnormalities Involved in Transformation of Follicular Lymphoma (FL) to Diffuse Large B-cell lymphoma; (Abstract #161). Presented at the 2016 ACMG Annual Clinical Genetics Meeting, March 9-11, 2016, Tampa, FL.
60. Prasad Koduru, Sara Monaghan, Jaidi Wen, Rolando Garcia, Sangeeta Patel, Kathleen. Wilson. Concurrent Rearrangements in *FGFR1*, *MYST3*, *RUNX1T1* and *RUNX1* in an Infantile Acute Myeloid Leukemia. (Abstract #161). Presented at the 2016 ACMG Annual Clinical Genetics Meeting, March 9-11, 2016, Tampa, FL.
61. Prasad. K, Wen, B. Reed, P. Jimenez and V. Babu. Interphase Chromosome Profiling (ICP) Identifies Y-chromosome Origin of a Ring(X) Chromosome Called by G-band Pattern in a Patient with Turner Syndrome. (Abstract #171). Presented at the 2016 ACMG Annual Clinical Genetics Meeting, March 9-11, 2016, Tampa, FL.
62. R. Garcia and P. Koduru. A text mining and Temporal Analysis of Cytogenetic Data Correlated with Gene Expression Profiles Reveals a Dependence Network of Chromosome Aberrations Outlined by Early and Late Cytogenetic Events in Prognostic Subgroups of Diffuse Large B-cell Lymphoma; (Abstract #2855T). Presented at the 2016 ASHG Annual Meeting, October 18-22, 2016, Vancouver, BC. Canada.
63. Scheuerle AE, Koduru P, Wilson KW. Is some maternal mosaicism really an artifact of fetal microchimerism. Smith Meeting, July 2017, Smithtown, NY.
64. Garcia R, Flores M, Chen W, Koduru P. Impact of MYC abnormalities, trisomy of chromosome 8 and estimated tumor progression values in plasma cell myeloma. AMP Annual meeting, November 2017, Denver CO. JMD 19: 957
65. Babu R, Van Dyke DL, Bhattacharya S, Dev VG, Liu M, Kwon M, Gu G, Koduru P, Rao N, Williamson C, Fuentes E, Fuentes S, Papa S, Kopuri S, Lal V. A rapid and reliable chromosome analysis method using interphase nuclei from products of conception. ASHG Annual Meeting, Orlando, 2017
66. Babu R, Van Dyke D, Fuentes E, Fuentes S, Papa S, Kopuri C, Williamson C, Koduru P. ALL-ICP, a simple and comprehensive method to detect chromosome abnormalities in acute lymphoblastic leukemia. JMD, 19: 963, 2017 (AMP Meeting, November 2017).
67. Babu R, Van Dyke DL, Fuentes E, Fuentes S, Kopuri S, Williamson C and Papa S, Koduru P. A simple, fast and comprehensive method - NextGen ICP to detect chromosome abnormalities in Acute Lymphocytic Leukemia patients. AMP annual meeting, November 2017, Denver CO.
68. Garcia R, Elias B, Patel S, Koduru P. Clonal evolution pathways and genetic progression scores backtract specific genetic events and outline unique recurrent chromosome aberrations that predict cell of origin and clinical outcome in transformed follicular lymphoma. AGT Annual meeting, St Luis, June 2017.
69. Chen W, Krueger JE, Fuda F, Koduru P. Mixed phenotype acute leukemia (MPAL), B/myeloid, with t(2;22)(q34;q12);*EWSR1* rearranged. Presented at 2017 Workshop of Society for Hematopathology/European Association for Hematopathology.
70. Chen W, Collins R, Zhang CC, Wu G, Rakheja D, Monaghan S, Kumar K, Koduru P. Myelodysplastic/myeloproliferative neoplasm (MDS/MPN) with complex genetic abnormalities including t(5;12)(q31;p13);*ETV6-ACSL6*. Presented at 2017 Workshop of Society for Hematopathology/European Association for Hematopathology.
71. Garcia R, Wick N, Chen W, Koduru P. Functional Driver and Passenger Chromosomal Imbalances along with Clone Evolution Patterns Identify Cell of Origin and Predict Outcome in Diffuse Large B-cell Lymphoma. Annual meeting of American College of Medical Genetics and Genomics (ACMG). April 10-14 2018; Charlotte, NC
72. Garcia R, Wilson K, Koduru P. Cytogenomic Characterization of B-Acute Lymphoblastic Leukemia (ALL) with intrachromosomal Amplification of Chromosome 21 (iAMP21) Reveals Co-existing Cytogenetic Manifestations and Outlines Driver Alterations that Play a Role in Leukemogenesis. American Society of Human Genetics (ASHG); Oct 16-20; San Diego, CA. 2018.

73. Alsuwaidan A, P. Koduru, M. Chen, J. Jasso, H. Luu, N. Sweed, R. Garcia, F. Fuda, W. Chen. Shade of Gray Between Double/Triple-hit High-Grade Large B Cell Neoplasm (DTH-HGBCL) and CD34 Negative Lymphoablasic Leukemia (B-LL): a subset of DTH-HGBCL Bridging These Entities and Biological Implication. US CAP; March 17-23 2018; Vancouver, Canada. Published in Mod Pathol.
74. Alsuwaidan A, Koduru P, Chen M, Jaso J, Luu H, Sweed N, Garcia R, Fuda F, Chen W. A combined biomarker of MYC and bright CD38 improves predictive power for identifying double/triple hit high grade B-cell lymphomas. US CAP; March 17-23 2018; Vancouver, Canada. published in Mod Pathol
75. Nathan Sweed, Rolando Garcia, Weina Chen, Prasad Koduru. MYC Translocation Partner Matters in Diffuse Large B-Cell Lymphoma and Double/Triple-Hit High-Grade B-Cell Lymphoma. US CAP; March 17-23 2018; Vancouver Canada. Published in Mod Pathol.
76. Prasad Koduru, Weina Chen, Barbara Haley, Kathleen Wilson. Cytogenomic Characterization of Double Minute Heterogeneity in a Patient with Therapy Related Acute Myeloid Leukemia. Accepted to present at 2018 annual meeting of American College of Medical Genetics and Genomics (ACMG). April 10-14 2018; Charlotte, NC.
77. Alsuwaidan A, Koduru P, Fuda F, Vusirikala M, Sadeghi N, Zhang CC, Chen W. Therapy-related acute myeloid leukemia, characterized by t(8;16)(p11;p13);*MYST3-CREBBP* and co-occurring *TET2* and *ASXL1* mutations. 11<sup>th</sup> Annual Transfusion and Laboratory Medicine Conference, 2-8-9, 2018.
78. Frame IJ, Chen W M.D, Garcia R, Koduru P, Fuda F, Rosado F, Chen M. Acute promyelocytic leukemia mimicking paroxysmal nocturnal hemoglobinuria. ICCS (International Clinical Cytometry Society) 2018 Annual Meeting, 10-1-2018.
79. Montgomery-Goecker C, Koduru P, Krueger JE, Fuda F, Chen W. A Rare Case Report: Mixed Phenotype Acute Leukemia (MPAL), B/myeloid, with t(2;22)(q34;q12); *EWSRI* rearranged. 11<sup>th</sup> Annual Transfusion and Laboratory Medicine Conference, 2-8-9, 2018.
80. Mir M, Sahoo S, Fang Y, Peng Y, Hwang H, Czapl A, Koduru P, Sarode V. Stability of HER2 Expression in Residual Breast Cancer Following Neoadjuvant Anti-HER2 Therapy. Mod Pathology, 91, 76, 2018.
81. Rahman Chaudhry, Abdullah Alsuwaidan, Prasad Koduru, Franklin Fuda, Rolando Garcia, and Weina Chen. MYC-IGH is Associated with Inferior Survival in De Novo High-Grade B-cell Lymphomas with Double-Hit (*MYC* and *BCL2* and/or *BCL6* Rearrangements), CMC 2019 meeting
82. Rahman Chaudhry, Abdullah Alsuwaidan, Elaina Pirruccello, Prasad Koduru, Franklin Fuda, Rolando Garcia, Mingyi Chen, Flavia Rosado, Jesse Jaso, Hung Luu, Hsiao-Ching Li, Prapti Patel, Madhuri Vusirikala, Navid Sadeghi, Syed Rizvi, Praveen Ramakrishnan Geethakumari, Neil B. Desai, Robert Collins Jr., and Weina Chen. *MYC* Partner Matters: *MYC-IGH* is Associated with Inferior Survival in *De Novo* High-Grade B-cell Lymphomas with *MYC* and *BCL2* and/or *BCL6* Rearrangements; Presented at 2019 USCAP
83. Rood T, Liu Y-L, Koduru P, Sahoo S, Hwang H, Peng Y, Fang Y, Sarode VR. Neo-adjuvant anti-HER2 therapy in breast cancer: Predictors of pathological complete response and survival. Presented at 2020 USCAP.
84. McCammon N, Koduru P, Sohani A, Chen W, Vos J, Chen M, Fuda F, Jaso J, Rosado F. Detection of *MYC* rearrangement is useful in distinguishing plasmablastic lymphoma from blasmablastic variant of plasma cell myeloma. Presented at 2020 USCAP.
85. Mani M, Koduru P, Gagan J, Fuda F, Xu J, Zhang Y, Botten G, Luu H, Malter J, Chen W. Clinicopathological and genetic characterization of acute myeloid leukemia with t(8;16)(p11;p13)/*KAT6A-CREBBP*: Single institution experience. Presented at 2020 USCAP.
86. Shi G, Fuda F, Garcia R, Ho K, Patel S, Malter J, Gagan J, Koduru P. Cytogenetic and molecular genetic profiling of B-lineage leukemia with *ZNF384* rearrangement identifies cryptic gene fusions and novel somatic alterations in *NF1* and epigenetic regulators. Annual ACMG meeting San Antonio, March 2020.
87. Garcia R, Timmons C, Fuda F, **Chen W**, Koduru P. Clonally Related Classical Hodgkin Lymphoma with Mantle Cell Lymphoma Presenting with an (11;14) Translocation: A Case Report with Clone Evolution and Pathway Analysis, ASHG 2020 meeting
88. Koduru P, Chen W, Garcia R, Shi G, Gagan J. Novel Fusion of *PVT1 - RCOR1* in B-cell prolymphocytic leukemia (BCPCLL) producing false FISH fusion of *MYC - IGH* with an atypical pattern. AMP November, 2020

89. García R, Hussain A, Shi G, Chen W, Koduru P. An artificial intelligence system applied to recurrent cytogenetic aberrations and derived genetic scores reveals high fidelity in predicting *MYC* rearrangements in diffuse large B-cell lymphoma. AMP November, 2020
90. Koduru RP, Wilson K, Garcia R, Weinberg O, Patel P, Kumar K, Chen W. Heterogeneity in cytological presentation of gene amplification and concurrent amplification of *MYC* and *PVT1* in therapy related acute myeloid leukemia (AML). ASHGG Annual Meeting, October 2021.
91. Chen D, Chen W, Fuda F, Koduru P, Oliver D, Ramakrishnan P, Weinberg O. Chronic lymphocytic leukemia/small lymphocytic lymphoma transformation into diffuse large B-cell lymphoma with complexed chromosomal abnormalities. Society for Hematopathology/European Association for Haematopathology, Meeting, Italy, September 2021.
92. Chen D, Fuda F, Rosado F, Koduru P, Chen W. Emergence of CD19(-) B-lymphoblastic leukemia in CD19-targeting immunotherapy and LILRB1 as a novel diagnostic B-cell marker. 100<sup>th</sup> Annual Meeting of the TSP, 2021 (Awarded Investigative Posters: 3rd Place)
93. Chen D, Fuda F, Rosado F, John S, Koduru P, Chen W. LILRB1 as a novel diagnostic B-cell marker in B-lymphoblastic leukemia in the era of CART-19 therapy. 14th Annual Transfusion and Laboratory Medicine Conference, 2021
94. Chen D, Fuda F, Rosado F, Saumell S, Samuel J, Koduru P, Chen W. Clinicopathologic features of relapsed CD19(-) B-ALL in CD19-targeted immunotherapy: insights into mechanism of relapse and LILRB1 as a novel B-cell marker for CD19(-) B lymphoblasts. USCAP, Los Angeles, March 2022.
95. Kirtek T, Chen W, Laczko D, Bagg A, Koduru P, Foucar K, Rogers HJ, Tam W, Orazi A, His E, Hasserjian R, Arber DA, Nicholas MM, Weinberg OK. Acute Leukemias with Complex Karyotype Show a Similarly Poor Outcome Irrespective of Mixed, Myeloid and Lymphoblastic Immunophenotype: A Study from the Bone Marrow Pathology Group. USCAP Meeting, Los Angeles, March, 2022
96. Prasad Koduru, Franklin Fuda, Mingyi Chen, Samuel John, Gurbakhash Kaur, Weina Chen, Jeffrey SoRelle, Jeffrey Gagan, Rolando Garcia. Discordant fluorescence in situ hybridization and RNASeq results in the identification of fusion partners in recurring translocations in hematological malignancies. ACMG Annual Meeting, Nashville, March 2022
97. Monique Morrison, Sangeeta Patel, Sou Saukam, Alycia Willard, Maria Grace Santiago, Diana Martinez, Valerie Miller, Micah Jacobs, Angela E. Scheuerle, and Prasad Koduru. Isodicentric(Y)(p11.2) mosaicism in newborn with 46,XX cells. ACMG Annual Meeting, Nashville, March 2022

### Clinical Practice Guidelines

1.	
----	--

### **Non-peer reviewed scientific or medical publications/materials in print or other media**

1.	
----	--