

JUDITH R. BAKER, DPH, MHSA

University of California Los Angeles, Department of Pediatric Hematology/Oncology
10833 Le Conte Avenue, Room A2-410 MDCC Los Angeles, CA 90095-1752
Tel: 310 794 6264 email: judithbaker@mednet.ucla.edu

EDUCATION

BSc Michigan State University, Magna Cum Laude, Bachelor of Science, Criminal Justice
MHSA University of Michigan, School of Public Health
Master of Health Services Administration, Department of Health Management and Policy
DrPH University of California Los Angeles, School of Public Health
Doctor of Public Health, Department of Health Policy and Management

PROFESSIONAL EXPERIENCE

1984-1986 Program Specialist, Maternity and Infant Care, Philadelphia Public Health Department, Pennsylvania
1986-1990 Executive Director, Low Birth Weight Long-term Outcome Study, Children's Hospital of Philadelphia
1990-1995 Program Coordinator, Hemophilia AIDS Help and Prevention, Children's Hospital of Los Angeles
1995-2006 Regional Coordinator, Federal Hemophilia Treatment Centers/Region IX, Children's Hospital of Los Angeles
2006-2012 Administrative Director, Federal Hemophilia Treatment Centers/Region IX, UCLA, Los Angeles, CA
2012 – Director of Public Health, Center for Inherited Blood Disorders (CIBD), Orange, CA.
Administrative Director, HRSA Western States Region Hemophilia Network, CIBD
Multi-site CDC Surveillance Manager, Federal Hemophilia Treatment Centers/Region IX, CIBD
2014 - Adjunct Assistant Professor, UCLA David Geffen School of Medicine, Department of Pediatrics, Division of Hematology/Oncology
Director of Policy and Partnerships, Pacific Sickle Cell Regional Collaborative, CIBD

PROFESSIONAL ACTIVITIES: COMMITTEE SERVICE (Federal)

US Food and Drug Administration

Special Government Employee, Center for Biologics Evaluation and Research 2008 -
Member, US Food and Drug Administration, Blood Products Advisory Committee 2005 – 2008, 2017- 2020
Temporary Voting Member, Transmissible Spongiform Encephalopathy Advisory Committee, 2007, 2011
Temporary Voting Member, Blood Products Advisory Committee, 2010, 2011, 2015 - 2017

US Centers for Disease Control and Prevention

Member, Female Bleeding Disorders Work Group, CDC Division of Blood Disorders, 2004 – 2011
Planning Committee, 1st and 2nd National Conferences on Blood Disorders and Public Health, 2010 and 2012
Co-Chair, Hemophilia Socio-economic Health Status Outcomes Work Group, CDC Division of Blood Disorders, 2007 – 2011
Member, Administrative Committee, CDC Surveillance to Reduce Complications of Bleeding and Clotting Disorders 2012 -
Advisor to CDC National Center for Disabilities, Birth Defects and Division of Blood Disorders on Sickle Cell Partnerships, 2017
Member, 2nd National Hemophilia Inhibitor Summit, January 30, 2017.
Member, Project Review Committee – CDC Blood Disorders Surveillance/American Thrombosis & Hemostasis Network, 2017 -

US Health Resources and Services Administration

Member: National Hemophilia Program Coordinating Center -- Advisory Committee 2012 - 2015; Needs Assessment Work Group, 2012 –; Regional Leadership Committee 2012 –; Minority Patient Needs Committee 2014 -

PROFESSIONAL ACTIVITIES: COMMITTEE SERVICE (National)

National Hemophilia Foundation

Member, Women's Bleeding Disorder Task Force, 1997 – 2001; Member, Women's Institute, 2009 – 2011
Planning Committee: National Conferences on Recognition and Management of Women's Bleeding Disorders, 1997
Member, Emergency Factor Work Group, 2001
Member, Blood Safety Working Group, 2002 – 2003

PROFESSIONAL ACTIVITIES: COMMITTEE SERVICE (Continued)

National Hemophilia Foundation (Continued)

Member, National Policy Summit, facilitated by The Lewin Group, 2012
Member, Regional Coordinator Work Group, 2017-
Member, Planning Committee – Familias de Sangre – 1st California Hispanic Family Education Conference, 2017

University of California Los Angeles

Quality Improvement Committee, Division of Pediatric Hematology/Oncology 2012 - 2013
Children's Discovery and Innovation Institute, Fellows Abstract Reviewer, 2015 -
Scholarship Oversight Committee:
Heather Vickers, MD, Fellow, Division of Pediatric Hematology/Oncology 2015 - 2017
Elizabeth Van Dyne, MD Fellow, Division of Pediatric Hematology/Oncology 2015 - 2016
Gavin Roach, MD, Chief Fellow, Division of Pediatric Hematology/Oncology 2012 - 2015
Pediatric Quality Council, Department of Pediatrics, 2013 -

University of California San Francisco

Mentor – Pediatric Hematology/Oncology Fellow, Tiffany Lin, MD, 2014 -

Children's Hospital Los Angeles

Member, Spiritual Care Advisory Board, 2002 – 2006
Member, Accreditation Committee, Clinical Pastoral Education, 2004 – 2006
Member, Clinical Pastoral Internship Advisory Committee, 2004 – 2006

Hemophilia Council of California

Co-Chair, Statewide Home Care/Specialty Pharmacy Forum 2004 –
Advisor, Managed Health Care Work Group 2007 –
Advisor, California Policy Summit, 2011 -

DES (Diethylstilbestrol) Action, Chair, Ann Arbor, Michigan Chapter 1981 – 1983

City of Philadelphia, Mayor's Expert Panel on Infant Mortality, Staff 1984 – 1985

Maternity Care Coalition of Philadelphia, Member, 1987 – 1989

California Department of Health Services, Member, HIV Community Planning Work Group, 1995 – 1997

University of Michigan, School of Public Health, Mentor to Master Degree Students 2004 – 2005

PROFESSIONAL ACTIVITIES: TEACHING

Practices of Evaluation: Theory and Methodology (Health Policy and Management 422), UCLA School of Public Health
Teaching Assistant to Elizabeth Yano, PhD., Executive Master of Public Health program, Spring 2013

PROFESSIONAL ACTIVITIES: PROFESSIONAL ASSOCIATIONS AND SCHOLARLY SOCIETIES

Academy Health – Member, 2014-

American Public Health Association

Member, Disability Section 1993 –
Abstract Reviewer, Community Health Planning Section, 2003 - 2004

American Thrombosis and Hemostasis Network

Member, Community Relations and Communications Committee, 2007 – 2012
Representative to Healthy People 2020 Working Group on Blood Disorders and Blood Safety, 2009

Federal Hemophilia Treatment Centers/Western States Region IX

Chair, Annual Regional clinical education conference, locations throughout CA, HI, Nevada, 1990 –
Chair, HIV Risk Reduction Retreats for Region IX Hemophilia patients, 16 throughout CA, 1990 – 1996
Chair, Statewide Women's Bleeding Disorders Conference for Consumers, Berkeley & Los Angeles, CA 1996, 1998

PROFESSIONAL ACTIVITIES: PROFESSIONAL ASSOCIATIONS AND SCHOLARLY SOCIETIES (Continued)

Federal Hemophilia Treatment Centers/Western States Region IX (Continued)

Chair, Reducing Hemophilia Morbidity through a State/Federal Partnership, for State of California Children's Medical Service Branch, Oakland and Los Angeles, CA, 1999, 2001

Co-Chair, 30 CACTI: Celebrating Access to Comprehensive Treatment Innovations, Federal Hemophilia Treatment Centers/Regions VIII, IX and X Annual Conference, Scottsdale, AZ, 2009

Co-Chair, Biennial Advance Practice Hemophilia Education Seminar for U.S. Hemophilia Regions VIII, IX and X 1994 - 2006

Co-Chair, Excessive Bleeding in Females, Seminar for Hemophilia Association of San Diego Counties, San Diego, CA, 2002

Chair, Basics of Bleeding Disorder Care and Services, various locations in CA, 2004, 2006, 2008, 2010 -

Chair, 340B Discount Drug Program Technical Assistance Seminars, Burbank, CA, 2003 - 2008

Co-Chair, Best Practices Summit for Hemophilia Center Nurses, Social Workers, Physical Therapists, San Francisco, CA 2008

Co-Chair, Program Integrity Seminar for 340B Discount Drug Programs, Los Angeles, CA, 2012 -

Co-Chair, *Improving Outcomes via Care Coordination Best Practices: Hemophilia Members in Medi-Cal Managed Health Plans*, Statewide Medical Directors Meeting with California Department of Healthcare Services, Sacramento, January 2014

US Hemophilia Treatment Center Network - Hemophilia and Thrombosis Data Set

Co-Chair, Dataset Revision Committee, 2014

Co-Chair, Steering Committee, 2016 -

US Hemophilia Treatment Center Network – National Patient Satisfaction Survey Workgroup

Co-Chair – Steering Committee, 2014 inaugural survey

Co-Chair – Dissemination Work Group

Co-Chair -- Steering Committee, 2018 survey

Hemophilia Utilization Group Study

Founding Member, 1992 -

External Advisory Committee 2005 – 2012

Steering Committee, 2013 -

Hemostasis and Thrombosis Research Network Member, 2010 -

World Federation of Hemophilia

Member, 1997 -

Co-Chair, Pacific Rim Hemophilia Workshop, 1997, Honolulu, Hawaii: 300 clinicians from 30 nations

PROFESSIONAL ACTIVITIES: EDITORIAL SERVICES

Reviewer, American Journal of Preventive Medicine, 2009

Reviewer, BMC Health Services Research, 2015 -

Reviewer, Journal of Pediatric Nursing, 2016 -

PROFESSIONAL ACTIVITIES: CONSULTING ACTIVITIES

3D Communications. FDA Advisory Committee Preparation, 2010, 2014

Biogen-Idec: NCQA Specialty Care Medical Home Advisory Committee, 2015 -

Sickle Cell Disease Foundation of California: Advisor to the CEO/President on HRSA Newborn Screening Grant, 2015 -

HONORS AND SPECIAL AWARDS

Meritorious Service Award, National Hemophilia Foundation, 2005

Distinguished Service Award, Federal Food and Drug Administration, 2008

Best Abstract Award, National Hemophilia Foundation Convention, 2009

Best Poster Award, Hemostasis and Thrombosis Research Society Conference, 2009

Best Poster Award, 1st and 2nd National Conferences on Blood Disorders and Public Health, 2010 and 2012

Ruth Roemer Scholarship, UCLA School of Public Health, 2011 - 2012

Child and Family Health Leadership Trainee, UCLA School of Public Health, 2011 - 2013

HONORS AND SPECIAL AWARDS *(Continued)*

University Fellowship, UCLA Graduate Division, 2012 – 2013

Delta Omega Honorary Society in Public Health, 2013 -

Best Poster Award, International Society for Pharmacoeconomics and Outcomes Research Conference, 2013

Nominee – CDC Charles C. Shepard Science Award for original research published in peer review journal, 2017 for Mazepa et al, *Blood* 2016.

Achievement Awardee and Brilliance Award Nominee, Health Category - National Association of Counties, *A Novel Approach to Sickle Cell Disease Closes the Healthcare Gap for Adults in South Los Angeles*, 2017.

RESEARCH, SURVEILLANCE, CLINICAL CARE, AND EDUCATION GRANTS RECEIVED

1. University of California San Francisco, University-Wide AIDS Research Project. *Condom Attitudes and Behaviors of HIV+ Hispanic Hemophiliacs*. Purpose: to characterize the condom attitudes and behaviors of California Hispanics with hemophilia who were co-infected with HIV from contaminated blood products. \$29,000 in 1994. Co-Principal Investigator.
2. Centers for Disease Control and Prevention. *HIV/Hemophilia Conference Series*. Purpose: to reduce HIV transmission from Californians and Nevadans with inherited bleeding disorders to their sexual partners through 16 educational retreats that fostered the development of peer networks. \$63,000 total: \$36,000 in 1993, \$9,000 in 1994, and \$18,000 in 1996. Principal Investigator.
3. HRSA Maternal and Child Health Bureau and Centers for Disease Control and Prevention. *Region IX Hemophilia Program*. Purpose: to enhance health outcomes and reduce HIV transmission among residents of California, Hawaii, and Nevada who have genetic bleeding disorders, through providing regionally organized comprehensive diagnosis, prevention, and treatment services. \$3,018,715 total: \$996,000 in 1994, \$1,006,836 in 1995, and \$1,015,879 in 1996. Co-Principal Investigator.
4. Centers for Disease Control and Prevention. *Tri-Regional Hemophilia Education Conferences*. Purpose: to increase the clinical skills and reduce professional isolation among Hemophilia Treatment Center Nurses, Social Workers and Physical Therapists in US Region's VIII, IX, and X. \$30,207 in 1996 and \$29,836 in 1998. Principal Investigator.
5. HRSA Maternal and Child Health Bureau. *Region IX Hemophilia Program*. Purpose: to foster health among underserved residents of California, Hawaii, and Nevada and the US Pacific who have genetic bleeding disorders through expanding regionally organized comprehensive diagnosis, prevention, and treatment services. \$2,360,949 total: \$750,283 in 1997, \$820,383 in 1998, and \$790,283 in 1999. Co-Principal Investigator.
6. Centers for Disease Control and Prevention. *Preventing Hemophilia Complications in Region IX*. Purpose: to characterize viral and musculoskeletal complications among residents of California, Hawaii, Nevada and Guam inherited bleeding disorders through nationwide surveillance. \$3,583,317 total: \$706,520 in 1997, \$657,520 in 1998, \$673,754 in 1999, \$787,300 in 2000, and \$758,223 in 2001. Co-Principal Investigator.
7. HRSA Maternal and Child Health Bureau. *Region IX Hemophilia Program*. Purpose: to promote good health outcomes among residents of California, Hawaii, and Nevada and the US Pacific who have genetic bleeding disorders through expanding regionally organized comprehensive diagnosis, prevention, and treatment services. \$2,310,810 total: \$779,799 in 2000, \$765,728 in 2001, and \$765,283 in 2002. Co-Principal Investigator.
8. Centers for Disease Control and Prevention. *Preventing Central Venous Access Devices Complications in Hemophilia*. Purpose: to determine predictors of CVAD infections in California Hemophilia Treatment Center patients and thereby reduce incidence. \$482,479 in 2000 – 2001. Co-Principal Investigator.
9. HRSA Maternal and Child Health Bureau. *Region IX Comprehensive Hemophilia Treatment Center Program*. Purpose: to strengthen health outcomes among residents of California, Hawaii, and Nevada and the US Pacific with genetic bleeding disorders through expanding regionally organized comprehensive clinical services \$2,354,998 total: \$795,400 in 2003, \$799,799 in 2004 and \$799,799 in 2005. Co-Principal Investigator.
10. Giannini Family Foundation. *Advance Practice Seminar*. Purpose: to increase the clinical skills and reduce professional isolation among California Hemophilia Treatment Center Nurses, Social Workers and Physical Therapists. \$125,000 total: \$30,000 in 2003 and \$45,000 in 2004, \$50,000 in 2006. Principal Investigator.
11. Centers for Disease Control and Prevention. *Preventing Hemophilia Complications in Region IX*. Purpose: to characterize viral and musculoskeletal complications among residents of California, Hawaii, Nevada and Guam inherited bleeding disorders through conducting nationwide surveillance. \$3,791,115 total: \$758,223 annually 2002 - 2006. Co-Principal Investigator.

RESEARCH, SURVEILLANCE, CLINICAL CARE, AND EDUCATION GRANTS RECEIVED (Cont.)

12. A.P. Giannini Foundation. *Regional Technology Enhancement & 340B Seminar*. Purpose: to increase the quality and comparability of clinical and health outcomes data among Hemophilia Treatment Centers in California, Hawaii, Nevada and Guam by providing training in a new software program, and to give legal and business technical assistance to California Hemophilia Treatment Centers to start Federal 340B drug discount programs. \$90,000 total: \$40,000 in 2005 and \$50,000 in 2007. Principal Investigator.
13. ZLB Foundation. *Translating Hemophilia and Von Willebrand Disease patient education materials for low literacy Spanish, Chinese, Vietnamese and Korean readers*. Purpose: to increase health literacy among under-served populations who have inherited bleeding disorders. \$35,152 in 2005. Co-Principal Investigator.
14. Centers for Disease Control and Prevention. *Diamond Blackfan Anemia Registry*. Purpose: to characterize demographic and clinical characteristics of Californians with DBA. \$60,000 in 2006. Co-Principal Investigator.
15. Centers for Disease Control and Prevention. *Healthy Joint Range of Motion Study*. Purpose: to determine the normal joint range among healthy US adults. \$8,649 in 2006. Co-Principal Investigator.
16. ZLB Behring Foundation. *Nevada Native American Hemophilia Education/Family Support*. Purpose: to reduce health disparities by supporting clinical care and patient/family education for residents of the Duck Valley Native Reserve in Owyhee, NV. \$70,197 in 2006. Co-Principal Investigator.
17. HRSA Maternal and Child Health Bureau *Comprehensive Hemophilia Care at Region IX Hemophilia Treatment Centers*. Purpose: to continue to improve health outcomes among residents of California, Hawaii, and Nevada and the US Pacific who have genetic bleeding disorders through expanding regionally organized comprehensive diagnosis, prevention, and treatment services \$2,339,556 total: \$779,852 annually 2006 – 2008. Co-Principal Investigator.
18. NovoNordisk Foundation. *Clinical Education Conference*. Purpose: to improve diagnostic and clinical skills among 14 Hemophilia Treatment Center core teams through advance practice clinical education conference. \$72,486 in 2007. Principal Investigator.
19. Centers for Disease Control and Prevention. *Preventing Bleeding Disorder Complications through Hemophilia Treatment Centers in Region IX*. Purpose: to characterize natural history of viral and musculoskeletal complications among residents of California, Hawaii, Nevada and Guam inherited bleeding disorders through conducting nationwide surveillance. \$4 million total: \$800,000 annually 2007 – 2012. Co-Principal Investigator.
20. A.P. Giannini Foundation. *340B Seminar, Best Practices Summit*. Purpose: to provide technical assistance to California Hemophilia Treatment Centers on operating 340B discount drug programs and expedite the spread of best clinical care and health promotion practices \$50,000 in 2008. Principal Investigator.
21. AP Giannini Foundation, Baxter, Bayer, CSL Behring, Grifols, Hemophilia Alliance Foundation, NovoNordisk, Nu Factor, Red Chip, Wyeth. *Hemophilia Treatment Center Conference for U.S. Regions VIII, IX, X*. Purpose: to celebrate the 30th anniversary of the US Hemophilia Treatment Center Network and strengthen clinical skills. \$200,000 in 2009. Principal Investigator.
22. HRSA Maternal and Child Health Bureau *Comprehensive Hemophilia Care at Region IX Hemophilia Treatment Centers*. Purpose: to strengthen health outcomes among residents of California, Hawaii, and Nevada and the US Pacific who have genetic bleeding disorders through expanding regionally organized comprehensive diagnosis, prevention, and treatment services \$2,100,000 total: \$700,000 annually 2009 – 2012. Co-Principal Investigator.
23. Novo Nordisk Foundation, AP Giannini Foundation, Bayer HealthCare. *Federal Hemophilia Treatment Centers/Region IX Clinical Education Conference*. Purpose: increase advance practice clinical skills of Hemophilia Treatment Center core teams. \$61,000 total in 2010: \$32,000 Novo Nordisk, \$20,000 AP Giannini Foundation, \$9,000 Bayer. Principal Investigator.
24. AP Giannini, Baxter, Bayer, Biogen Idec, CSL Behring, Grifols, Hemophilia Alliance, Novo Nordisk, Octapharma, Pfizer, Red Chip. *Federal Hemophilia Treatment Centers/Region IX Clinical Education Conference*. Purpose: increase clinical skills of Region IX Hemophilia Treatment Center teams through advance practice conference. \$115,000 in 2011, \$125,000 in 2012, \$145,000 in 2013, \$162,000 in 2014, \$183,000 in 2015, \$190,000 in 2016 and \$200,000 in 2017. Principal Investigator.
25. HRSA Maternal and Child Health Bureau *Western States Regional Hemophilia Network* Purpose: to strengthen health outcomes among residents of California, Hawaii, and Nevada and the US Pacific who have genetic bleeding disorders through expanding regionally organized comprehensive diagnosis, prevention, and treatment, and collaboration with National Coordinating Center \$2,500,000 total: \$500,000 annually 2012 – 2017. Co-Principal Investigator/Public Health Director.

RESEARCH, SURVEILLANCE, CLINICAL CARE, AND EDUCATION GRANTS RECEIVED (Cont.)

26. HRSA Maternal and Child Health Bureau *Pacific Regional Sickle Cell Disease Demonstration Program*. Purpose: To improve health outcomes for people with SCD by increasing the number of providers treating with disease-modifying therapies and the numbers of patients obtaining care from providers with greater knowledge of how to treat SCD in eight western states. \$2,550,000 total: \$850,000 annually 2014 – 2017. Public Policy Director.
27. HRSA Maternal and Child Health Bureau *Western States Regional Hemophilia Network* Purpose: to strengthen health outcomes among residents of California, Hawaii, and Nevada and the US Pacific who have genetic bleeding disorders through expanding regionally organized comprehensive diagnosis, prevention, and treatment, and collaboration with National Coordinating Center \$2,500,000 total: \$500,000 annually 2017 – 2022. Co-Principal Investigator/Public Health Director.
28. HRSA Maternal and Child Health Bureau *Pacific Regional Sickle Cell Disease Demonstration Program*. Purpose: To improve health outcomes for people with SCD by increased clinical capacity and conducting quality improvement that increase access to evidence based care in thirteen western states. \$1,800,000 total: \$850,000 annually 2017 – 2022. Public Policy Director.

LECTURES AND PRESENTATIONS: *International*

1. Sexual Practices among Latino Men with Hemophilia. With V Hoffman, BV Marin, **JR Baker**, et al. Poster at Xth International Conference on AIDS. Yokohama, Japan, 1994
2. Hemophilia Case finding in the Pacific Basin, **JR Baker**, JM Hall, and JP Iser, Presentation at the XXII International Congress of the World Federation of Hemophilia, Dublin, Ireland, 1996
3. 25 Million Women Worldwide have von Willebrand Disease: How do we reach them? Paper, R and **JR Baker**, Presentation at the XXIII International Congress of the World Federation of Hemophilia, The Hague, Netherlands, 1998
4. U.S. Outreach Efforts to Women with Bleeding Disorders, Presentation at the XXIV International Congress of the World Federation of Hemophilia, Montreal, Canada, 1998
5. Advance Practice Education Seminar for Hemophilia Treatment Center Nurses, Social Workers and Physical Therapists in Three Western United States Regions. Riske B, **Baker JR**, Ingram-Rich R. Presentation at the XXVII International Congress of the World Federation of Hemophilia, Vancouver, Canada, 2006
6. Involvement in Thrombophilia care among Hemophilia Treatment Center Nurses in the Western U.S. **JR Baker**, B Riske, R Ingram-Rich. Poster presented at the XXVII International Congress, World Federation of Hemophilia, Vancouver, Canada, 2006
7. Hemophilia care in Guam: Public/Private partnership improves access to care and health, **JR Baker**, RV Zabala. Poster presented at the XXVIII International Congress of the World Federation of Hemophilia, Istanbul, Turkey, 2008
8. Factors associated with lower health related quality of life (HRQOL) in adults with factor VIII deficiency – the Hemophilia Utilization Group Study V (HUGS-V). Wu J, Globe D, Gwady-Sridhar F, Riske B, Ullman M, Huszti H, Koerper M, **Baker JR**, Johnson K, Poster presented at the 13th Annual International Meeting of the International Society for Pharmacoeconomics and Outcomes Research, Toronto, Canada, 2008
9. Healthcare utilization and factor cost in hemophilia. Zhou ZY, Globe D, Ullman M, Baker JR, Koerper M, Gwady-Sridhar F, Wu J, Forsberg A, Shapiro A, Trawinski B, Duncan N, Johnson KA, Poster at the 14th Annual Meeting of the International Society for Pharmacoeconomics and Outcomes Research, Orlando, Florida, 2009.
10. Health care utilization and cost in persons with factor viii deficiency: results of the HUGS VA Study, Zhou ZY, Globe D, Ullman M, **Baker JR**, Koerper M, Gwady-Sridhar F, Wu J, Forsberg A, Shapiro A, Trawinski B, Duncan N, Johnson KA, Poster at the 4th Asia-Pacific meeting of the International Society for Pharmacoeconomics and Outcomes Research, Phuket, Thailand, 2010
11. Suicide Among US Males with Hemophilia, 1998-2007. Gomperts E, Holtz J, **Baker JR**, Geraghty S, Hudson M, Karp S, Osip J, Presley R. Poster at the XXIV International Congress of the World Federation of Hemophilia, Buenos Aires, Argentina, 2010
12. A national surveillance of females with bleeding disorders receiving care at the United States Hemophilia Treatment Center Network: Preliminary report of the pilot study of the Female Universal Data Collection, Centers for Disease Control and Prevention. Kulkarni R, Kouides PA, Byams VR, **Baker JR** on behalf of the CDC Female UDC Working Group. Invited presentation at the XXIV International Congress of the World Federation of Hemophilia, Buenos Aires, Argentina. 2010

LECTURES AND PRESENTATIONS: *International* (Cont.)

13. The Hemophilia Utilization Group Study (HUGS-VB): Health-related quality of life in Hemophilia B. Poon JL, Lou M, Doctor J, Zhou ZY, Gwadry-Sridhar F, **Baker JR**, Ullman M, Koerper M, Johnson K. Poster at the 16th Annual Meeting of the International Society for Pharmacoeconomics and Outcomes Research, Baltimore, Maryland. 2011
14. Characterizing direct costs associated with Hemophilia A: a resource based cost analysis Zhou ZY, **Baker J**, Globe D, Ullman M, Koerper M, Gwadry-Sridhar F, Wu J, Forsberg A, Shapiro A, Trawinski B, Duncan N, Johnson KA. Poster at the 16th Annual Meeting of the International Society for Pharmacoeconomics and Outcomes Research. Baltimore, Maryland. 2011
15. Characterizing excessive school and work absenteeism in Hemophilia A Poon JL, Lou M, Zhou ZY, Riske B, Koerper M, **Baker JR**, Johnson KA. Poster at the XXV International Congress of the World Federation of Hemophilia, Paris, France, 2012
16. Indirect costs among persons with Hemophilia B. Lou M, Poon JL, Koerper M, Riske B, **Baker JR**, Kulkarni R, Cockrell E, Konkle B, Johnson KA. Poster at the XXV International Congress of the World Federation of Hemophilia, Paris, France, 2012
17. A national infrastructure for rare blood disorders: An evaluation of staffing, training, and services in the US federally supported hemophilia treatment centres. Forsberg A, Cutter S, Riske B, **Baker, JR**, Wicklund B, Voutsis M, Drake J. Poster at the XXV International Congress of the World Federation of Hemophilia, Paris, France, 2012
18. Cost to society due to underemployment in persons with Hemophilia A and B – Hemophilia Utilization Group Study V. Niu X, Poon JL, Riske B, **Baker JR**, Ullman M, Gwadry-Sridhar F, Lou M, Nichol MB. Poster at the International Society of Pharmacology Outcomes Research, New Orleans, LA, May 2013.
19. Longitudinal changes in health-related quality of life for chronic diseases: an example from the Hemophilia Utilization Group Study VA. Poon JL, Doctor J, Gwadry-Sridhar F, Ullman M, Riske B, **Baker JR**, Niu X, Lou M, Nichol MB. Poster at the International Society of Pharmacology Outcomes Research, New Orleans, LA, May 2013.
20. Association between health status of persons with hemophilia B and consumption of factor IX concentrate – Hemophilia Utilization Group Study Part Vb (HUGS Vb). Niu X, Riske B, **Baker JR**, Lou M, Ullman M, Nichol MB. Oral presentation at the XXVI International Congress:World Federation of Hemophilia, Melbourne, Australia, May 2014
21. Measuring work or school absence associated with administration of factor concentrate in children with hemophilia in the United States – Hemophilia Utilization Group Study Part V (HUGS V). Niu X, Lou M, **Baker JR**, Poon JL, Konkle B, Ullman M, Riske B, Hord J, Kulkarni R, Curtis R, Koerper M, Wu, J, Nichol MB. Poster at the XXVI International Congress of the World Federation of Hemophilia, Melbourne, Australia, May 2014
22. Association between Health Status and factor consumption among persons with Hemophilia B using propensity score matching. Niu X, Poon JL, Kulkarni R, Lou M, **Baker JR**, M, Riske B, Ullman M, Curtis R, Nichol MB. Poster at the International Society of Pharmacology Outcomes Research, Montreal, QC, Canada, June 2014.
23. Risk for Obstetric Bleeding in Von Willebrand Disease (VWD): Analysis of a US National Cohort. Roach G, **Baker JR**, Elashoff D, Oakley M. Kohn D. Poster at the International Society of Thrombosis and Hemostasis, Toronto, Canada, June 2015.
24. Clustering Health Status among Adults with Hemophilia Using SF-12 Health Survey. Niu X, **Baker JR**, Riske B, Ullman MM, Wu J, Lou M, Nichol MB. Poster at the International Society of Pharmacology Outcomes Research, Milan, Italy, November 2015.
25. Characteristics Associated with Annual Bleeding Frequency Among Hemophilia Patients in the United States. Chen CX, Ullman M, Hord J, Kulkarni R, Konkle BA, **Baker JR**, Riske B, Koerper M, Lou M, Wu J, Nichol MB. Poster at the International Society of Pharmacology Outcomes Research, Milan, Italy, November 2015.
26. Comparison of Health Utilities in Persons with Hemophilia B. Lou M, Wu J, Gwadry F, Wasserman J, **Baker JR**, Hord J, Konkle B, Doctor J, Nichol, Riske B. Poster at the World Federation of Haemophilia International Congress Orlando FL, July 2016.
27. Gap Analysis – Hemophilia Patient Perspectives on Information and Service Needs in a Remote US Pacific Island. Lin T, **Baker JR**. Poster at the World Federation of Haemophilia International Congress Orlando FL, July 2016.
28. Comparison of clinical characteristics and health care utilization among individuals with hemophilia A and B in the Hemophilia Utilization Group Studies (HUGS) cohorts. Lou M, Koerper, M, Ullman M, **Baker JR**, Riske B, Curtis R, Wu J, Nichol M. Oral Presentation at World Federation of Haemophilia International Congress Orlando FL, July 2016
29. U.S. Hemophilia Physician Prescribing Practices: Then and Now. Riske B, Koerper M, Curtis C, **Baker JR**, Ullman M. Poster at the World Federation of Haemophilia International Congress Orlando FL, July 2016.

LECTURES AND PRESENTATIONS: *International* (Cont.)

30. Hemophilia Utilization Group Studies Part VI (HUGS VI): An Adherence Study among Participants with Hemophilia. Lou M, Koerper, M, Ullman M, **Baker JR**, Riske B, Curtis R, Wu J, Nichol M. Poster at the World Federation of Haemophilia International Congress Orlando FL, July 2016

LECTURES AND PRESENTATIONS: *National*

1. Preliminary Analysis of Knowledge, Attitudes, and Behaviors among Hemophilic Youth with HIV Disease Regarding Sexuality and Condoms, **JR Baker** and K Parish. Presentation at the American Public Health Association, San Francisco, 1993
2. HIV Risk Reduction Retreats for People with Hemophilia and HIV. Presentation at the Centers for Disease Control and Prevention Division of STD/HIV Prevention Grantee Meeting, Washington, DC, 1994.
3. Acculturation and Chronic Serious Conditions: The Case of Hemophilia. **JR Baker**, V Hoffman, et al. Presentation at the American Public Health Association Convention, San Diego, CA, 1995
4. AIDS and Adolescents: Characteristics of HIV+ and HIV- Youth with Hemophilia. K Parish and **JR Baker**. Presentation at American Psychological Assn. Annual Conference, New York, NY, 1995
5. Latinos with Hemophilia: HIV Prevention and Access to Care. **JR Baker**, V Hoffman, et al Presentation at 7th National AIDS Update Conference, San Francisco, CA, 1995
6. Females Bleed, Too! Under-recognized and under-treated inherited bleeding disorders. **JR Baker** and R Paper, Poster -American Psychological Association's Women's Health Conference, Washington, DC, 1996
7. Needles in a Haystack: Hemophilia and the Pacific Basin. **JR Baker**, JM Hall, and JP Iser., Presentation at American Public Health Association Convention, New York, NY, 1996
8. Women's Bleeding Disorders: A National Call to Action. **JR Baker** and R Paper, Presentation at the National Hemophilia Foundation's National Conference on Prevention Education, Louisville, KY, 1997
9. Only Women Bleed: von Willebrand's Disease--the hidden bleeding disorder of one million American women. **JR Baker** and R Paper, Poster at the Fifth Annual Congress on Women's Health. Washington, DC, 1997
10. Rare, Chronic Conditions and Managed Care. **JR Baker** and C Kasper, Presentation at the Federal Hemophilia Program Annual Regional Director/Regional Coordinator Meeting, Houston, TX, 1997
11. Developing a Hemophilia Program in Guam. L Meadows, **JR Baker**, R Paper, Poster at the National Health Service Corps Annual Conferences in Atlanta. Tucson, AZ and San Jose, CA, 1998
12. Federal and Regional Outreach Models for Women's Bleeding Disorders, Presentation at the National Hemophilia Foundation Annual Conference, Orlando, FL, 1998
13. Initiating Hemophilia Services in the US Pacific Jurisdictions. **JR Baker**, JP Iser, and R Zabala, Presentation at the American Public Health Association Convention, Washington, DC, 1998
14. Managed Care and vWD: New Opportunities in Recognizing the Bleeding Disorder **JR Baker** and R Paper, Abstract presented at the U.S. Public Health Service's Office on Women's Health Conference on The Future of Managed Care and Women's Health: New Directions for the 21st Century. Washington, D.C. 1998
15. Structuring Regional Hemophilia Treatment Networks in Region IX. **JR Baker** and D Nugent. Presentation at the Federal Hemophilia Program Annual Regional Director/Regional Coordinator Meeting, Santa Fe, NM, 1999.
16. Actions for Women's Health Advocates. Plenary Panel. National Conference on Recognition and Management of Women's Bleeding Disorders, National Hemophilia Foundation. San Francisco, CA. 1999
17. Building Hemophilia Treatment and Advocacy Services in the U.S. Pacific Territories: a capacity building model in Guam and the Commonwealth of the Northern Mariana Islands. **JR Baker** and R Zabala, Presentation at the American Public Health Association. Boston, MA, 2000
18. Developing a kit to educate the public about von Willebrand disease in females. **JR Baker**, R Paper and K Larsen. Poster at the American Public Health Association. Boston, MA, 2000
19. Remedies to Address Aggressive Marketing. Presentation at the Tri-regional Advance Practice Seminar for Hemophilia
20. Treatment Center Nurses, San Francisco, CA, and at the CDC/MCHB Federal Hemophilia Program Annual Business Meeting, Rockville, MD, 2000
21. Expanded Roles/Responsibilities of Regional Hemophilia Directors and Coordinators Presentation at the CDC/MCHB Federal Hemophilia Program Annual Business Meeting. Atlanta, GA, 2001
22. Helping the Chapter/Staff Navigate the New Legal/Ethical Environment, Presentation at the National Hemophilia Foundation Convention/Chapter Staff Organization Seminar. Orlando, FL, 2002

LECTURES AND PRESENTATIONS: *National (Cont.)*

23. Hemophilia Morbidity in Region IX: Data from the first three years enrollment in CDC's nationwide hemophilia surveillance system, Presentation at the American Public Health Association Convention. Philadelphia, PA. 2002
24. Reducing Morbidity of Children with Special Health Care Needs Enrolled in Managed Care: A Federal, State, Community Hemophilia Partnership, Poster - American Public Health Association Convention. Philadelphia, PA. 2002
25. Menorrhagia, Post Partum Hemorrhage and Von Willebrand Disease. **JR Baker** and RB Paper. Presentation at the American College of Nurse Midwives Annual Convention. Atlanta, GA, 2002
26. Rare Chronic Disorders in Guam: Regional partnership improves hemophilia treatment, research and advocacy. CDC National Conference on Chronic Disease Prevention and Control. Atlanta, GA, 2002
27. Improving nursing competencies in rare/chronic bleeding disorders: The tri regional seminar **JR Baker**, B Riske and R Ingram Rich. Poster at the American Public Health Association Convention. San Francisco, CA, 2003
28. Improving Hemophilia Treatment Center Nurse Coordinator, Social Worker and Physical Therapist skills in the Western United States **JR Baker**, B Riske and R Ingram Rich., Poster at the National Hemophilia Foundation Annual Convention. Salt Lake City, UT, 2003
29. Pacific partnerships for rare disorders: The Region IX Hemophilia model. Presentation at the Global Public Health Conference, Honolulu, HI. 2003
30. Hemophilia morbidity in Hawaii and Guam: Data from the CDC surveillance system. Presentation at the Global Public Health Conference. Honolulu, HI, 2003
31. A model for regional systems of care to promote health and well being for persons with rare genetic disorders. **JR Baker** (Symposium Chair and Presenter), S Crudder, B Riske, A Forsberg, V Bias. Centers for Disease Control and Prevention's National Center on Birth Defects and Developmental Disabilities Conference. Washington. DC, 2004
32. Helping the female patient advocate for herself. Presentation at the National Hemophilia Foundation Conference on Women with Bleeding Disorders. Seattle, WA. 2004
33. Advocacy for the Female Patient. Presentation -- National Hemophilia Foundation Convention. San Diego, CA, 2005
34. Reducing health disparities by strengthening clinical competencies in rare/chronic bleeding disorders: the tri regional seminar. **JR Baker**, B Riske, RI Rich. Poster at the CDC's 19th National Conference on Chronic Disease Prevention and Control. Atlanta, GA, 2005
35. Tracking hemophilia joint morbidity disparities in Hawaii and Guam using CDC's surveillance system. Poster at the 21st Annual Pacific Rim Disabilities Conference. Honolulu, HI, 2005
36. Strengthening Healthy Communities for MCH Populations with Rare Disorders, Presentation at the Association of Maternal and Child Health Programs Annual Conference. Arlington, VA, 2007
37. Association of social demographics and clinical factors with barriers to hemophilia care in patients with Factor VIII deficiency-The hemophilia utilization group study-part V. B Riske, A Forsberg, ZY Zhou, F Gwady-Sridhar, M Ullman, **JR Baker**; D Globe, C Smith, J Wu, K Johnson. Poster at the American Public Health Association Convention. Washington, DC, 2007
38. Hemophilia Utilization Group Study (HUGS V): Summary of Baseline Data-Demographics, Clinic Characteristics, Barriers to Care, Arthropathy and Quality of Life. ZY Zhou, **JR Baker**, D Globe, B Riske, A Forsberg, J Wu, M Ullman, N Duncan, C Smith, F Gwady-Sridhar, R Curtis, M, Koerper, K, Johnson, Poster at National Hemophilia Foundation Convention. Orlando, FL, 2007
39. Bringing Hemophilia Treatment Center care and Chapter services to Nevada's Rural/Frontier and Native American Communities. SL Ewing, R Berkowitz, **JR Baker**. Poster at the National Hemophilia Foundation Convention. Denver, CO, 2008
40. Health Insurance Coverage for Adults and Children with Hemophilia: Hemophilia Utilization Group Study (HUGS-V: A). Y Zhou, M Ullman, M Koerper, D Globe, **JR Baker**, B Miller, B Riske, C Smith, A Forsberg, J Wu, R Curtis, K Johnson. Poster at the National Hemophilia Foundation Convention. Denver, CO. 2008
41. Adopting a Standards Based Information Infrastructure for Rare Disorders Communities. **JR Baker**, D Aschman Poster at the 136th Annual American Public Health Association Meeting & Exposition. San Diego, CA, 2008
42. Partners in Progress: How to work with your State Medicaid Program for a Win-Win-Win, **JR Baker**, P Pontrelli. Sixth Annual Conference of the Hemophilia Alliance, Inc. Las Vegas, NV, 2008
43. CDC Universal Data Collection Project Surveillance on Females with Bleeding Disorders National Hemophilia Foundation Convention. San Francisco, CA, 2009

LECTURES AND PRESENTATIONS: *National (Cont.)*

44. One-year health care utilization and cost in patients with factor VII deficiency: first year results of the HUGS V-A study. Z Zhou, B Riske, **J Baker**, A Forsberg, M Koerper, M Lou, A Shapiro, M Ullman, K Johnson, Poster at the National Hemophilia Foundation Annual Convention. San Francisco, CA, 2009
45. Healthcare Utilization in Persons with Severe Hemophilia A on Prophylactic vs. Episodic Replacement. Z Zhou, M Koerper, B Riske, **J Baker**, R Miller, M Ullman, A Forsberg, R Curtis, A Shapiro, K Johnson, Poster at the Hemophilia and Thrombosis Research Society Annual Meeting. Chicago, Ill, 2009
46. Rare Disorders Partnership Successes: The U.S. Hemophilia Model. **JR Baker**, D Aschman, V Bias Presentation at the Annual Genetic Alliance Conference. Washington, DC, 2009
47. Healthcare Utilization in Persons with Severe Hemophilia A on Prophylactic vs. Episodic Replacement. Z Zhou, M Koerper, B Riske, **J Baker**, R Miller, M Ullman, A Forsberg, R Curtis, A Shapiro, K Johnson, Poster at the Hemophilia and Thrombosis Research Society Annual Scientific Meeting. Chicago, Ill, 2009
48. Aligning Hemophilia Treatment Center Future Goals/Objectives with Federal Public Health Initiatives. Presentation at CDC/HRSA Regional Hemophilia Treatment Center Directors and Coordinators Meeting. New Orleans, LA, 2010
49. Burden of Illness: Bleeding Episodes and Indirect Costs among Persons with Hemophilia A: Hemophilia Utilization Group Study Part Va (HUGS Va) ZY Zhou, **J Baker**, B Riske, M Ullman, A Forsberg, C Smith, R Curtis, B Miller, M Koerper; K Johnson Presentation and poster at the National Hemophilia Foundation Annual Convention. New Orleans, LA, 2010
50. The Hemophilia Utilization Group Study (HUGS VB): Characterization of Baseline Data. M Lou, ZY Zhou, M Ullman, B Riske, JL Poon, A Forsberg, **J Baker**, F Gwady-Sridhar, R Miller, K Johnson. Poster at the National Hemophilia Foundation Annual Convention. New Orleans, LA, 2010
51. Reducing Medicaid Spending on catastrophically expensive blood disorders **JR Baker**, V Bias. Presentation at the National Conference on Blood Disorders in Public Health. Atlanta, GA, 2010
52. Trends in comprehensive care center patient populations. BK Riske, **JR Baker**. Presentation at the National Conference on Blood Disorders in Public Health. Atlanta, GA, 2010
53. Reducing Geographic and Resource Disparities for Rare Blood Disorder Populations in the U.S. Pacific PA Carhill, **JR Baker**. Presentation at the National Conference on Blood Disorders in Public Health. Atlanta, GA, 2010
54. Race and physical outcomes in males with hemophilia in the United States: 1998-2008. PE Monahan, **JR Baker**, BK Riske, JM Soucie. Presentation at the National Conference on Blood Disorders in Public Health. Atlanta, GA, 2010
55. Barriers to Care in Patients with Hemophilia A – The Hemophilia Utilization Group Study. Zhou, ZY, Riske B, Forsberg A, Gwady-Sridhar F, Ullman M, Koerper, M, **Baker J**, Globe D, Shapiro A, Wu J, Johnson, K. Presentation National Conference on Blood Disorders in Public Health. Atlanta, GA, 2010
56. Developing National Surveillance on Females with Bleeding Disorders and Baseline Characteristics of Females with Bleeding Disorders Receiving Care in the US Hemophilia Treatment Centers. V Byams, **JR Baker**, D Brown, J Gill, A Grant, A James, B Konkle, R Kulkarni, J Maahs, M Malone, S McAlister, D Nugent, C Philipp, JM Soucie, E Stang, P Kouides, Presentations at the National Conference on Blood Disorders in Public Health. Atlanta, GA, 2010
57. Sociodemographics, Clinical Characteristics and Health-Related Quality of Life among people with Hemophilia A and B. JL Poon, M Lou, ZY Zhou, R Curtis, J Doctor, F Gwady-Sridhar, B Riske, M Ullman, **J Baker**, M Koerper, A Forsberg, E Cockrell, R Kulkarni, K Johnson. Poster -- National Hemophilia Foundation Convention. Chicago, Ill, 2011
58. Hemophilia Data Set: History, Uses, Trends. HRSA National Hemophilia Program Data Meeting. Bethesda, MD, 2011
59. The Future of the Blood Disorders Workforce – Non-Physicians. Closing Plenary Panel. 2nd National Conference on Blood Disorders in Public Health. Atlanta, GA, 2012
60. Burden of Illness: Direct and Indirect Costs Among Persons with Hemophilia A. K Johnson, ZY Zhou, M Koerper, B Riske, **J Baker**, M Ullman, R Curtis, J Poon, M Lou Poster presented at 2nd National Conference on Blood Disorders in Public Health 1st prize poster awardee. Atlanta, GA, 2012
61. National Survey of State Medical Assistance Programs: Variations in Eligibility, Coverage and Hemophilia Care Standards. RE Ingram-Rich, **JR Baker**, Poster presented at 2nd National Conference on Blood Disorders in Public Health. Atlanta, GA, 2012
62. Trends in Care of Patients with Bleeding Disorders: 20 years of data B Riske, **JR Baker**. Presentation at 2nd National Conference on Blood Disorders in Public Health. Atlanta, GA, 2012
63. Healthcare Services Utilization in Persons with Hemophilia B. X Niu, J Poon, B Riske **JR Baker**, M Lou, R Kulkarni, J Hord, B Konkle, M Ullman, MB Nichols. Poster -- Hemostasis and Thrombosis Research Society, Charlotte, NC, 2013

LECTURES AND PRESENTATIONS: *National (Cont.)*

64. Burden of Bleeding Episodes among Persons with Hemophilia B – Hemophilia Utilization Group Study Vb (HUGS Vb) Niu X, Poon JL, Lou M, Ullman M, Koerper M, **Baker J**, Hord J, Konkle BA, Riske B, Buranahirun C, Gwadry-Sridhar F, Curtis RG, Kulkarni R, Soni A, Nichol MB. Poster - National Hemophilia Foundation Convention. Anaheim, CA, 2013
65. Survey of Support Needs of the Hemophilia Treatment Center Staff: The National Hemophilia Program Coordinating Center. Forsberg A, Butler R, McLaughlin K, **Baker J**, Oldfield D, Aschman D. Poster presented at National Hemophilia Foundation Convention. Anaheim, CA, 2013.
66. Working together: the Hemophilia Alliance and the Regional Hemophilia programs for 340B. **Baker JR**, Riske BK, Droze K. Presentation at the Winter meeting of the Hemophilia Alliance. Phoenix, AZ, January 2014.
67. Embracing HUGS: The Hemophilia Utilization Group Study's Value to 21st Century Hemophilia Researchers - Documenting Cost, Quality, and Outcomes. **Baker JR**, Ullman M. Poster presented at Thrombosis and Hemostasis Society of North America Symposium. Chicago, IL, April 2014.
68. National patient perspectives on US federally-funded hemophilia treatment center care, services and gaps. Forsberg AF, Butler R, Cutter S, Sharathkumar A, **Baker JR**, Ahuja S, Aschman D Cheadle A. Presentation at the American Public Health Association Annual Convention, New Orleans LA, November 2014
69. Hemophilia and Thrombosis Data Set 2014 Revision: Adding Value Now and for the Future. **JR Baker**, B Riske, J Drake. HRSA National Hemophilia Program Regional Leadership Meeting. Washington, DC, April 2015
70. Inaugural National Patient Satisfaction Survey for the US Hemophilia Treatment Center Network. **JR Baker**, B Riske, K Droze. HRSA National Hemophilia Program Regional Leadership Meeting. Washington, DC, April 2015
71. National Patient Needs Assessment – US Hemophilia Treatment Center Network: Views of Minority Patients. HRSA National Hemophilia Program Regional Leadership Meeting. **JR Baker**, A Forsberg. Washington, DC, April 2015
72. Associations Between Annual Bleeding Episodes and Burden of Illness Among Persons with Hemophilia A and B in the United States. Curtis RG, Riske B, **Baker JR**, Ullman M, Niu X, Norton K, Koerper M Lou M, Nichol MB. Poster presented at National Hemophilia Foundation Convention, Dallas, TX, 2015
73. Zoris in the Sand: Hemophilia Education and Capacity Building in Saipan. T. Lin, **JR Baker**, P Carhill, J Huang. Poster presented at National Hemophilia Foundation Convention. Dallas, TX, 2015
74. Patient Satisfaction with US Hemophilia Treatment Centers 2015: National Results **JR Baker**, B Riske, K Droze, R Shearer. Poster presented at National Hemophilia Foundation Convention. Dallas, TX, 2015
75. The value of National Patient Satisfaction Data to US Hemophilia Treatment Center 340B Programs. **JR Baker**, B Riske, K Droze, R Shearer. Lecture presented at The Hemophilia Alliance Membership meeting, Denver, CO, September 20, 2015
76. Regional Hemophilia Networks: rare disorder healthcare delivery systems meeting current / future patient needs. Presentation at the HRSA/CDC Regional Hemophilia Network Leadership Meeting, Baltimore, MD, March 21, 2016.
77. Treatment Adherence Predictors: Indicators from the US Hemophilia Treatment Center Patient Satisfaction Survey 2015. **JR Baker**, KA Droze, R Shearer, B Riske. Poster presented at the 2016 Thrombosis and Hemostasis Summit of North America, Chicago, IL, April 14-16, 2016.
78. Sickle cell anemia infant screening and care initiative in Malawi: program development and local capacity building in detection and preventive services. EA Van Dyne, ES Nkosi, RB Manda Kamanga, JL Ivers, B Ritz, K Seydel, **JR Baker**, K Nielsen-Saines, and TB Moore. Poster presented at the 29th annual meeting of the American Society of Pediatric Hematology/Oncology, Minneapolis, MN, May 11-14, 2016.
79. Methods that Mobilize: Building and mobilizing networks of stakeholders in HRSA's regional Sickle Cell Treatment Demonstration Program. Inaugural SCDTDP All State Partner Webinar, Invited lecture. May 25, 2016
80. Voices of 5000+ Patients: The First National Survey on Experience of Care at US Hemophilia Treatment Centers. **JR Baker**, B Riske, R Shearer. CDC Public Health Webinar Series on Blood Disorders. September 26, 2016. <https://www.youtube.com/watch?v=4YnyHH7ANUQ>
81. Curtis R, Robertson D, **Baker JR**. Burden of Illness in Sickle Cell Disease: Capitalizing on a Successful Methodology that Moves Data to Action. Oral presentation at the Sickle Cell Disease Association of America's 44th Annual Convention Baltimore, MD, September 2016.
82. Brooks T, Brown M, Abdul-Rashid N, **Baker JR**. Building Sickle Cell Services in Nevada: Community Health Worker Impact Bridging HRSA's Sickle Cell Newborn Screenings and Treatment Demonstration Projects. Oral presentation at the Sickle Cell Disease Association of America's 44th Annual Convention Baltimore, MD, September 2016.

LECTURES AND PRESENTATIONS: *National (Cont.)*

83. Ferrerosa J, Avila M, Bryant M, **Baker JR**. Advancing Sickle Cell Care, Education and Awareness for Hispanic Communities. Oral presentation at the Sickle Cell Disease Association of America's 44th Annual Convention Baltimore, MD, September 2016.
84. Bender M, Brown M, **Baker, JR**. Sickle Cell Disease Myths, Misconceptions and *HOPE*: Mobilizing Regional Partnerships to Improve Health. Presentation at the US Conference on African Immigrant Health. New York NY, September 2016
85. **Baker JR**. Advancing sickle cell care, education and awareness in Hispanic communities. Presentation at the National Association of Hispanic Nurses-Orange County Chapter General Membership Meeting, Orange, CA, September 2016.
86. **Baker JR**, Avila M, Ferrerosa J. Advancing Sickle Cell Care for Hispanics: The Pacific Sickle Cell Regional Collaborative and the National Association of Hispanic Nurses-LA Chapter Partnership. Oral presentation at the American Public Health Association Annual Convention, Denver, CO. October 2016.
87. **Baker JR**, Rothman E, Claster S. Creating a New SCD Clinic: A Case Study. Webinar organized by The California Rare Disease Surveillance Program. January 25, 2017. "Webinar 6" [here](#)
88. **Baker JR**, Brown M. Regional Blood Disorder Collaboratives: *Synergizing Partnerships to Improve Health*. Invited presentation to the CDC – National Center for Birth Defects and Developmental Disabilities. January 31, 2017.
89. Lin T, Carhill P, Huang J, **Baker JR**. Trends in Building Bleeding Disorders Clinical Knowledge throughout an Under-served US Pacific Island: 2014 – 2016. Poster presentation at the Hemostasis and Thrombosis Research Society Annual Meeting. Scottsdale, AZ. April 2017.
90. Curtis R, Koerper M, Riske B, **Baker JR**. Trends in Hemophilia Clinician Prescribing Practices: 1999 and 2015. Poster presentation at the Hemostasis and Thrombosis Research Society Annual Meeting, April 2017, Scottsdale, AZ.
91. **Baker JR**. Sickle Cell Health Policy – Regional Rare Blood Disorder Synergies. Panelist at National Minority Quality Forum Leadership Summit on Health Disparities and Spring Health Braintrust, April 24-25, 2017, Washington, DC.
92. **Baker JR**, Ferrerosa, J, Avila M, Espinoza-Bryant, M. Mobilizing to Reduce Sickle Cell Disparities among Hispanics: The Pacific Sickle Cell Regional Collaborative and the National Association of Hispanic Nurses-LA Chapter Partnership. Lecture at the National Association of Hispanic Nurses Annual Convention, July 19, 2017, Phoenix, AZ.
93. Lou M, Ullman M, **Baker JR**, Koerper M, Riske B, Wu J, Curtis R, Nichol M. Living with hemophilia B: examining quality of life and associated characteristics in the Hemophilia Utilization Group Studies (HUGS Vb) cohort. Poster at the National Hemophilia Foundation Annual Convention, August 25, 2017. Chicago, IL.

LECTURES AND PRESENTATIONS: *Regional, State, County, University*

1. Perinatal Educational Resources for Low Literacy Populations. 50th Anniversary Conference of the International Childbirth Education Association. Philadelphia, PA, 1985
2. Improving Care for the Nation's Hemophilia Population: Initiating Programs in the Pacific. Guam Department of Public Health and Social Services. Hagatna, Guam, 1997
3. Federal Hemophilia Program Outreach to Guam...is Saipan the Next Step? Commonwealth of the Northern Mariana Islands Department of Public Health, Saipan, Commonwealth of the Northern Mariana Islands, 1998
4. Federal Hemophilia Program: Overview and Pacific Basin Initiatives. American Pacific Nurse Leadership Council's Twenty-First Annual Conference. Saipan, Commonwealth of the Northern Mariana Islands, 1999
5. Bleeding Disorders and Women. Iris Cantor-UCLA Women's Health Education Resource Center, Los Angeles, CA, 2000
6. Healthy People 2010: Implications for Hemophilia. Tri-regional Seminar for Hemophilia Centers. San Francisco, CA, 2001
7. Federal Hemophilia Treatment Center Program/Outcomes. Reducing Hemophilia Morbidity through a State/Federal Partnership Seminar. Statewide CCS Hemophilia Conference. Los Angeles, CA, 2001
8. Excessive Bleeding in Females: Could it be Von Willebrand Disease? Federal/Region IX Women's Health Advisory Council. Honolulu, HI, 2002
9. The CDC Universal Data Collection Project on Hemophilia Complications and Helping the Hemophilia Community in the new Legal/Ethical Environment. Hemophilia Foundation of Hawaii, Honolulu, HI, 2002
10. Michigan's Adolescent Transition Protocol. Region IX Hemophilia Program Annual Conference. Monterey, CA, 2002
11. Regional Centers of Excellence for Von Willebrand Disease. Excessive Bleeding in Females Seminar. San Diego, CA, 2002

LECTURES AND PRESENTATIONS: Regional, State, County, University (Cont.)

12. Conflict of Interest. U.S. Region VIII and X Annual Hemophilia Treatment Center Meeting, Boulder, CO. 2003
13. Regional system of care for youth with rare blood disorders. Coordinating Services for Children with Special Health Care Needs in the Community Conference. Los Angeles, CA. 2007
14. Hemophilia Treatment Centers – Meeting the Challenges. Hemophilia Foundation of Northern California's Family Symposium. Oakland, CA, 2009
15. Hemophilia Treatment Center Update at Lucile Packard Children's Hospital's Hemophilia and Von Willebrand Disease Family Day. Palo Alto, CA, 2009
16. Hemophilia Treatment Centers- Dynamic and Growing. Hemophilia Association of San Diego Counties Family Information Day. San Diego, CA, 2009
17. Improving Health Care and Reducing Costs for Rare Expensive Genetic Blood Disorders: Policy Priorities for California's Hemophilia Treatment Centers. Hemophilia Council of California/National Hemophilia Foundation's California Policy Stakeholder Meeting. Sonoma, CA, 2011, 2012, 2013, 2014, 2015
18. Challenges of Funding Hemophilia Treatment: 50th Anniversary Celebration and Symposium, Hemophilia Treatment Center at Los Angeles Orthopaedic Hospital, April 2012, Los Angeles, CA
19. Children with Chronic Illness and Populations with Special Needs UCLA School of Public Health graduate course in Child Health, Programs and Policies, guest lecturer May 17, 2012
20. HTC/Chapter: effective collaboration. National Hemophilia Foundation's Regional Leadership Seminar. May 21, 2012, Irvine, CA
21. US Hemostasis Treatment Center Network: care, outcomes, costs. Lecture for Pediatric Hematology/Oncology Fellows. UCLA David Geffen School of Medicine, September 25, 2012
22. Inherited Bleeding Disorders: Improving Care via Medicine and Public Health Synergies. Lecture for Pediatric Residents, UCLA David Geffen School of Medicine, August 13, 2012 and November 25, 2013
23. Patient Population Trends at Western Region Hemophilia Treatment Centers and 340B Public Health Impact at Hemophilia Treatment Centers. Presentations at Western States Regional Hemophilia Network Conference, April 28 – 29, 2014
24. Sustainable capacity building strategies for improving clinical care and surveillance for rare genetic disorder populations. Lecture for the UCLA Center for World Health. May 15, 2014.
25. Guam's Public Law 32-235 – Policy Implications. Guam Hemophilia Awareness Month and Proclamation Signing. Hagathna, Guam. March 13, 2015.
26. National and Regional Trends in Patient Satisfaction at US Hemophilia Treatment Centers: Preliminary data; US Hemophilia Treatment Center Network 101 – Mission, Vision, Organizations, and Hemophilia and Thrombosis Dataset launch: Trends at Western Region Hemophilia Treatment Centers. Presentations at Western States Regional Hemophilia Network Conference, April 10-12, 2015
27. Leveraging Strategic Partnerships. Presentation at the Pacific Regional Sickle Cell Collaborative's Clinical Leadership meeting. Oakland, CA, November 5, 2015.
28. Sickle Cell Disease Provider Education and Partnerships. Presentation at the Pacific Regional Sickle Cell Collaborative's 2nd Government Partner's Meeting for Regions IX & X. San Francisco, CA, February 10, 2016
29. Leveraging Partners to Improve Health. Presentation at the National Association of Hispanic Nurses – Los Angeles Board of Directors Meeting. Covina, CA, February 24, 2016,
30. Sickle Cell Disease Myths, Misconceptions and *HOPE*: Mobilizing Regional Partnerships to Improve Health. Presentation at the Northwest Regional Conference on African Immigrant Health. Seattle, WA, March 19, 2016
31. The Pacific Sickle Cell Regional Collaborative: Mobilizing Partnerships to Improve Health. Presentation at the National Association of Hispanic Nurses – Los Angeles Annual Conference, Los Angeles, CA, April 21, 2016
32. DJ Nugent, **JR Baker**: New Staff Orientation – Hemophilia Treatment Center Clinicians, Administrators, 340B Pharmacy – Presentation at Mountain/Western States Regional Hemophilia Treatment Centers Annual Conference. Scottsdale, AZ, April 24, 2016
33. **JR Baker**, BK Riske. Quality Improvement Environmental Scan – Presentation at Mountain/Western States Regional Hemophilia Treatment Centers Annual Conference. Scottsdale, AZ, April 24, 2016
34. Methods that mobilize: advancing rare disorder care in the US Pacific. Lecture for the UCLA Center for World Health. May 16, 2016.

LECTURES AND PRESENTATIONS: Regional, State, County, University (Cont.)

35. Public Health and Rare Disorders – Eliminating the Oxymoron. Lecture for Pediatric Residents, UCLA David Geffen School of Medicine. June 6, 2016.
36. Hemophilia Treatment Centers: Evidence for Advocacy. Lecture at the Hemophilia Council of California's Bleeding Disorder Health Advocacy Summit, Manhattan Beach, CA. August 27, 2016.
37. Methods that Mobilize: Capacity building for Rare Chronic Disorders. Lecture for UCLA's Community Engagement and Research Program/Health Services Research Community Partners & Faculty Meeting. October 20, 2016.
38. Sickle Cell Disease: Building Adult services in Los Angeles County. Pediatric Grand Rounds with Ellen Rothman, MD and Susan Claster, MD at Miller Children's and Women's Hospital, Long Beach, CA. October 28, 2016.
39. Strategic Regional Sickle Cell Partnerships and Policy Update - Presentation at the Pacific Regional Sickle Cell Collaborative's Clinical Leadership meeting. Manhattan Beach, CA. November 17, 2016.
40. Strategies for Success: Building New Sickle Cell Health Services in Los Angeles. Lecture for UCLA's General Internal Medicine-Health Services Research Noon Seminar Series, Los Angeles, CA. January 13, 2017
41. Successful Researcher/Stakeholder Research Partnerships – the Pacific Regional Sickle Cell Experience. Panelist with Mary Brown, CEO of the Sickle Cell Disease Foundation of California, at the UCLA Stakeholder and Researcher Workgroup Retreat of UCLA's Community Engaged Research Program, Los Angeles, CA. January 17, 2017.
42. DJ Nugent, **JR Baker**: New Staff Orientation – Hemophilia Treatment Center Clinicians, Administrators, 340B Pharmacy; and **JR Baker** - Regional HTC Patient Population Trends and Policy Issues. Presentations at Western States Regional Hemophilia Treatment Centers Annual Conference. San Diego, CA, April 2-3, 2017.
43. Federal Blood Disorders Grants: who, what, why? Lunch and Learn at Center for Inherited Blood Disorders clinical, pharmacy and administrative staff. June, 16, 2017, Santa Ana, CA.
44. **JR Baker** and M Brown: Regional Blood Disorders Collaboratives: Synergizing Partnerships to Improve Health. Presentation at the Organizing CBO to be Effective in Meeting the Needs of Sickle Cell Patients & Family Conference, Tucson, AZ. September 23, 2017.

PUBLICATIONS/BIBLIOGRAPHY

A. RESEARCH PAPERS (PEER REVIEWED - published)

1. MC McCormick, **JR Baker**, J Brooks-Gunn, et al, Cohort Reconstruction: Which Infants can be restudied at School Age? *Pediatric and Perinatal Epidemiology*. 1991; 5: 410-412.
2. RV Zabala, **JR Baker**. Guam's Quest for Improved Hemophilia Care. *Pacific Health Dialog*. 2002; 9(2): 317-320.
3. **JR Baker**, SO Crudder, B Riske, V Bias, A Forsberg, A Model for a Regional System of Care to Promote the Health and Well-Being of People with Rare Chronic Genetic Disorders. *Am J Public Health*. 2005; 95: 1910-1916.
4. JH Drake, JM Soucie, SC Cutter, AD Forsberg, **JR Baker**, BK Riske. High School Completion Rates among Individuals with Hemophilia. *American Journal of Preventive Medicine*. 2010; 38(4S): S489-S494. doi: 10.1016/j.amepre.2009.12.024
5. **JR Baker**, BK Riske, M Voutsis, S Cutter, R Presley. Insurance, Home Therapy, and Prophylaxis in US Youth with Severe Hemophilia. *American Journal of Preventive Medicine* 2011; 41(6S4): S338-345.
6. PE Monahan, **JR Baker**, BK Riske, JM Soucie. Physical functioning in boys with hemophilia in the United States 1998 – 2008. *American Journal of Preventive Medicine*. 2011; 41(6S4): S360-368. doi: 10.1016/j.amepre.2011.09.017
7. ZY Zhou, BK Riske, A Forsberg, M Ullman, **JR Baker**, M Koerper R Curtis, M Lou, J Wu, K Johnson. Self-Reported Barriers to Hemophilia Care in People with Factor VIII Deficiency. *American Journal of Preventive Medicine*. 2011; 41(6S4): S346-S353.
8. VR Byams, PA Kouides, R Kulkarni, **JR Baker**, DL Brown, JC Gill, et al. Surveillance of Females with Inherited Bleeding Disorders in United States Hemophilia Treatment Centers. *Haemophilia*. 2011; 17(Suppl.1): S6-S13. doi: 10.1111/j.1365-2516.2011.02558.x
9. ZY Zhou, J Wu, **JR Baker**, R Curtis, A Forsberg, H Huszti, M Koerper, M Lou, R Miller, K Parish, B Riske, A Shapiro, M Ullman, K Johnson. Hemophilia Utilization Group Study - Part Va (HUGS Va): Design, Methods and Baseline Data. *Haemophilia*. 2011; 17(5): 729-736. doi: 10.1111/j.1365-2516.2011.02595.x
10. **JR Baker**, B Riske, JH Drake, AD Forsberg, R Atwood, M Voutsis, R Shearer. U.S. Hemophilia Treatment Center population trends 1990 – 2010: patient diagnoses, demographics, health services utilization. *Haemophilia*. 2013; 19: 21–26. doi: 10.1111/j.1365-2516.2012.02915.x
11. X Niu, JL Poon, B Riske, ZY Zhou, M Ullman, M Lou, **JR Baker**, M Koerper, R Curtis, M Nichol. Physical activity and health outcomes in persons with Hemophilia B. *Haemophilia* 2014: 1-8. doi: 10.1111/hae.12485
12. ZY Zhou, M Koerper, K Johnson, B Riske, **JR Baker**, R Curtis, M Ullman, R Curtis, JL Poon, M Lou, M Nichol. Burden of Illness: Direct and Indirect Costs among Persons with Hemophilia A in the United States. **J Med Econ**. June 2015, Vol. 18, No. 6 , Pages 457-465 (doi:10.3111/13696998.2015.1016228)
13. R Curtis, **JR Baker**, B Riske, M Ullman, X Niu, K Norton, M Lou, M Nichol. Young adults with hemophilia in the US: demographics, comorbidities, and health status. November 2015. *American Journal of Hematology*, 90(S2). DOI: 10.1002/ajh.24218
14. T Lin, P Carhill, J Huang, **JR Baker**. A Capacity Building Paradigm for Rare Diseases in Remote Areas: Bleeding Disorders in the Commonwealth of the Northern Mariana Islands. *American Journal of Public Health*. April 2016, Vol. 106, No. 4, pp. 658-661. doi: 10.2105/AJPH.2016.303093
15. MA Mazepa, PE Monahan, **JR Baker**, BK Riske, JM Soucie. Men with severe hemophilia in the United States: birth cohort analysis of a large national database. *Blood*. 16 JUNE 2016 x VOL. 127, No. 24., pp.3073-3081. DOI: 10.1182/blood-2015-10-675140.
16. Chen CX, **Baker, JR**, Nichol MB.: Economic Burden of Illness Among Persons with Hemophilia B from HUGS Vb: Examining the Association of Severity and Treatment Regimens with Costs and Annual Bleed Rates. *Value in Health*, published online June 17, 2017. DOI: <http://dx.doi.org/10.1016/j.jval.2017.04.017>

B. RESEARCH PAPERS – PEER REVIEWED (IN PRESS) - NONE

PUBLICATIONS/BIBLIOGRAPHY (Cont.)

C. RESEARCH PAPERS – PEER REVIEWED (SUBMITTED)

1. CX Chen, M Ullman, **JR Baker**, BK Riske, R Curtis, MB Nichol. The impact of adherence to prophylactic clotting factor replacement therapy on bleeding episodes among patients with hemophilia. Submitted to *Value in Health*, August 2017.
2. J Wu, M Lou, F Gwady-Sridhar, J Wasserman, B Riske, **J Baker**, J Hord, B Konkle, R Kulkarni, M Ullman, J Doctor, M Nichol. Comparison of Health Utilities in Persons with Hemophilia B. Submitted to *Value in Health*, August 2017.

D. RESEARCH PAPERS NON-PEER REVIEWED

1. Women's Bleeding Disorders U.S. DHHS HRSA *Women's Health Data Book*. P. 31. 2009
<http://mchb.hrsa.gov/publications/pdfs/womenhealth2009.pdf>
2. Von Willebrand Disease in *Our Bodies, Ourselves 40th Anniversary Edition*, Boston Women's Health Book Collective, New York: Simon & Schuster. 2011

LETTERS TO THE EDITOR

1. DJ Nugent, **JR Baker**. Rebuttal to Research Letter published *Outpatient pharmacy expenditures for children with serious chronic illness in California, 2010-2012*, published in the July 28, 2015 issue of the Journal of the American Medical Association. Submitted August 2015

REVIEWS - NONE

PAPERS IN PREPARATION (RESEARCH COMPLETED) - NONE

ABSTRACTS –

1. R Curtis, M Koerper, BK Riske, **JR Baker**, Physician Prescribing Practices Trends in Hemophilia: 1999 and 2015. Accepted as Poster Presentation - American Society of Hematology Annual Convention, December 2017.
2. S Claster, E Rothman, M Brown, JR Baker, DJ Nugent. Improving Access to Care for Adults with Sickle Cell Disease in Los Angeles County. Accepted as Poster Presentation - American Society of Hematology Annual Convention, December 2017.

PUBLICATIONS – Opinion Leader, Manuals, Newspapers, Trade journals, Unpublished Reports, Videos, Radio

1. *Recommendations for the Reduction of Infant Mortality in Philadelphia*, Report of the Mayor's Expert Panel. Philadelphia Department of Public Health, 1985.
2. JR Baker and R Zabala, Guam's New Hemophilia Programs: Models of Collaboration, *Hemophilia World*, World Federation of Hemophilia, 1999: 6(1) 12 - 13.
3. JR Baker and R Paper, *Women Can Have Bleeding Disorders*, Education Kit for the National Hemophilia Foundation's Project Red Flag. 1999. Over 250 disseminated 10 countries.
4. Strengthening the State/Federal Partnership: the California Model, *HemAware*, Journal of the National Hemophilia Foundation, Oct/Nov 1999.
5. Brochures: Federal Hemophilia Treatment Centers/Region IX and Hemophilia Programs in Guam. 1999.
6. Excessive Bleeding: Could it be Von Willebrand Disease? *Women's Rx*, Iris Cantor-UCLA Women's Health Education and Resource Center, lecture and video. April 2002.
7. Understanding, Implementing and Operating an Outpatient Factor Distribution Program under Section 340B of the Public Health Service Act – Manual. 2003.
8. The Federal Hemophilia Treatment Center Program: Manual for new clinicians. 2003.
9. Editor, Chaplain Internship Program Manual for Students, ACPE Accreditation Handbook. Children's Hospital Los Angeles. 2005.

PUBLICATIONS – Opinion Leader, Manuals, Newspapers, Trade journals, Unpublished Reports

10. Laying the groundwork for a career in MCH and Fostering professional development, Maternal and Child Health Leadership Skills Development Series, video publication of the Women's and Children's Health Policy Center, Johns Hopkins Bloomberg School of Public Health. 2008. <http://www.jhsph.edu/wchpc/MCHLDS/index.html>
11. Administrative Director Strengthens Region IX Hemophilia Treatment Centers in *National Hemophilia Foundation's HemeAware Online Magazine* July 2011, highlighted in *UCLA Health Sciences Media Report* July 22 – 28, 2011;
12. Safe Blood: UCLA Staffer Plays a National Role in Safeguarding the Blood Supply on *UCLA website homepage* week of 2/1/11. UCLA Employee Plays National Role in Ensuring Blood Product Safety in *UCLA Health System Employee News* February 2011; She Safeguards Blood Supply for US Consumers in *UCLA Today*, 1/20/2011;
13. Helping Hemophiliacs is in Judith Baker's Blood in *Los Angeles Daily Breeze*, 2/9/11. Blood Expert Profiled for Daily Breeze in *UCLA Health Sciences & Media Report*, February 4-11, 2011.
14. Contributor, *Addressing the needs of members with hemophilia in Medicaid Managed Care: Issues and implications for health plans*, Medicaid Health Plan of America Center for Best Practices, July 2013.
15. UCLA hemophilia advocate featured guest at proclamation signing in Guam. *UCLA Newsroom Faculty Bulletin Board*, 3/19/2015.
16. 5000 Patients Rate Satisfaction with US Hemophilia Treatment Centers. www.htcsurvey.com Article distributed throughout US Hemophilia Treatment Center Network, National Hemophilia Foundation, Hemophilia Federation of America. June 2016.
17. Sickle Cell Disease-Pathway to a Cure. UCLA Mattel Children's Hospital Pediatric Hematology/Oncology Newsletter, authors Moore T, Roach G and Baker, JR. Vol. V, Issue 1, December 2016.
18. Patient Satisfaction with U.S. Hemophilia Treatment Centers: What Did We Learn? August 2017
19. Voices of Females with von Willebrand Disease: Patient Satisfaction with U.S. Hemophilia Treatment Centers, August 2017
20. A 'potentially powerful model' for treating sickle cell disease: KPCC Radio 89.3 FM, 9.25.2017
<http://www.scpr.org/news/2017/09/25/75900/a-potentially-powerful-model-for-treating-sickle-c/>

FILMS PRODUCED

1. Producer, 7-minute documentary on Renee Paper, RN, CCRN, Founding Director, Hemophilia Foundation of Nevada, and advocate for women's bleeding disorders, 2008
2. Executive Producer, 34-minute documentary *Blood Roots: Hemophilia Treatment Center Pioneers in Comprehensive Health Care (2010)*. <http://youtu.be/H01d0cfzuFA> Trailer <http://vimeo.com/37995804> premiered at National Hemophilia Foundation Annual Convention 2009, San Francisco, CA. International debut at World Federation of Hemophilia Congress, Buenos Aires, Argentina (2010). Accepted at Film Festival of American Public Health Association's 2011 Convention, Washington, DC; Presented at National Hemophilia Foundation's 2011 Convention, Chicago, IL; American Thrombosis and Hemostasis Network's 2011 Data Summit, Chicago, IL; and the 2nd National Conference on Blood Disorders in Public Health, Atlanta, GA 2012. Uploaded to these organizations' websites in 2015: The Centers for Disease Control and Prevention, The Hemophilia Alliance, The Hemophilia Federation of America, and countless Hemophilia Treatment Centers in the US.
3. Executive Producer, *Hemophilia Treatment Centers: improving care, reducing costs*. <https://www.youtube.com/watch?v=rw6sD2ynE1U> 7-minute video for insurers and policy makers, 2011. Presented at The Hemophilia Alliance Meeting January 2011 Las Vegas, NV; the 2nd National Conference on Blood Disorders in Public Health, Atlanta, GA 2012; distributed nationwide.
4. Executive Producer, 30 second PSA (English/Spanish) 2011. Signs/symptoms of bleeding disorders, provides website of US Hemophilia Treatment Centers, telephone number of National Hemophilia Foundation. Distributed nation-wide.