

UNIVERSITY OF PENNSYLVANIA - SCHOOL OF MEDICINE
Curriculum Vitae

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The Children's Hospital of Philadelphia
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34th Street and Civic Center Boulevard
Philadelphia, Pennsylvania 19104

Education: 1964-1968 B.A. Harvard University (Magna cum laude)
1968-1972 M.D. University of Pennsylvania School of Medicine

Postgraduate Training and Fellowship Appointments:

1972-1973 Internship, The Children's Hospital of Philadelphia
1973-1974 Assistant Chief Resident,
The Children's Hospital of Philadelphia
1974-1975 Chief Resident, The Children's Hospital of Philadelphia
1975-1978 Hematology-Oncology Fellowship,
The Children's Hospital of Philadelphia

Faculty Appointments:

1977-1978 Clinical Associate in Pediatrics,
University of Pennsylvania School of Medicine
1978-1984 Assistant Professor of Pediatrics,
University of Pennsylvania School of Medicine
1984-1990 Associate Professor of Pediatrics,
University of Pennsylvania School of Medicine
1990- Professor of Pediatrics, Clinician-Educator Track
University of Pennsylvania School of Medicine
2001-2013 Chairman, Department of Pediatrics
University of Pennsylvania School of Medicine

Hospital and Administrative Appointments:

1978-1982 Assistant Physician, The Children's Hospital of Philadelphia
1982-1985 Associate Physician, The Children's Hospital of Philadelphia
1985- Senior Physician, The Children's Hospital of Philadelphia
1978-1988 Director, Hematology and Coagulation Laboratories,
The Children's Hospital of Philadelphia
1986-1990 Director of Clinical Activities, Division of Hematology,
The Children's Hospital of Philadelphia
1990-1992 Acting Chief, Division of Hematology, The Children's
Hospital of Philadelphia
1992-2001 Chief, Division of Hematology, The Children's Hospital
of Philadelphia

1994-	Senior Member, Joseph Stokes, Jr. Research Institute, The Children's Hospital of Philadelphia
2001-2013	Physician-in-Chief, The Children's Hospital of Philadelphia Leonard & Madlyn Abramson Endowed Chair in Pediatrics, The Children's Hospital of Philadelphia
2013-	Medical Advisor to the Chief Executive Officer The Children's Hospital of Philadelphia

Specialty Certification:

1973	National Board of Medical Examiners
1977	American Board of Pediatrics: Pediatrics
1978	American Board of Pediatrics: Hematology-Oncology
1993	Recertification – American Board of Pediatrics: Hematology-Oncology
2003	Recertification – American Board of Pediatrics: Hematology-Oncology
2013	Recertification – American Board of Pediatrics: General Pediatrics

Licensure:

Pennsylvania

Awards, Honors and Membership in Honorary Societies:

1968	Phi Beta Kappa
1971	Alpha Omega Alpha
1985	Christian R. and Mary F. Lindback Award for Distinguished Teaching
1985	Residents' Annual Teaching Award
1998	Dripps Memorial Award for Excellence in Graduate Medical Education
2004	American Academy of Pediatrics, Pennsylvania Chapter, Pediatrician of the Year Award

Memberships in Professional and Scientific Societies:

National Societies:

- American Pediatric Society
- Society for Pediatric Research
- American Society of Hematology
- American Academy of Pediatrics
- American Society of Pediatric Hematology-Oncology
(President, 1997-1999)
(Board of Trustees, 1995-2001)
- Cooley's Anemia Foundation
(Chair, Medical Advisory Board, 1992-2003)
- Thalassaemia International Federation Scientific Advisory Board
- American Board of Pediatrics
Subboard of Pediatric Hematology-Oncology
Member, 2001 – 2006

(Chair, 2004-2006)

Board of Directors, 2006-2012

(Chair, 2010-2011)

Finance Committee, 2007-2011

(Chair, 2008-2009)

Credentials Committee, 2013-

Dragonfly Forest, Inc. Board

Member, 2006 -2010

Association of Medical School Pediatric Department Chairs (AMSPDC)

(President Elect, 2009-2011)

(President, 2011-2013)

(Immediate Past-President, 2013-2014)

Local Societies

Philadelphia Pediatric Society

Philadelphia Hematology Society

(Secretary-Treasurer, 1979-1986)

Editorial Positions:

Editorial Board, American Journal of Pediatric Hematology/Oncology (1998-2008)

Editorial Board, Haematologica (1997 – 2001)

Reviewer for Pediatrics, Haematologica, Blood, American Journal of Hematology, American Journal of Pediatric Hematology/Oncology, Journal of Pediatrics, New England Journal of Medicine, Pediatric Blood & Cancer

Academic Committees at the University of Pennsylvania and Affiliated Hospitals:

1982-1992	Pediatrics Residency Advisory Committee (Chairman, 1986-1992)
1985-2000	Teaching Awards Committee
1989-1992	Subcommittee for Evaluation
1995-1996	Integrative Neuroscience Course Review
2012-2013	Review of Department of Medicine
2014-	Residency Mentoring Committee (CHOP)
2014-	Intern Selection Committee (CHOP)
2014-	Conflict of Interest Committee, Chair (CHOP)
2015-	Hematology-Oncology Fellowship Committee (CHOP)
2015-	Committee on Academic Freedom and Responsibility

Major Teaching and Clinical Responsibilities at the University of Pennsylvania and Affiliated Hospitals:

1. Medicine 100 (formerly ID 103), Director, Pediatric Physical Diagnosis Course for first-year medical students, 1981-1986
2. Pediatrics 200, 1 month each year, 1979-
3. Pediatrics 300, 1 month each year, 1979-
4. Pediatrics 324, elective in Pediatric Hematology, 1979-present, course director, 1985-2001
5. Hematology attending rounds, 2-3 months each year (CHOP), 1978 -
6. Hematology Clinic, 1 day each week (CHOP), 1978 -

Lectures by Invitation (past 3 years):

- February, 2013 “Hemoglobinopathies and Iron Overload: Scientific Advances and Ethical Setbacks.” Grand Rounds, Cincinnati Children’s Hospital Medical Center, Cincinnati, Ohio.
- March, 2014 “When the Solution is the Problem: Transfusion-Derived Iron Overload.” Global Iron Summit, Athens, Greece
- January, 2015 “Medical Education in the United States” and “Teaching and Supervising Research” Conference on Education, Bangkok, Thailand.
- March, 2015 “How Much of a Threat is Iron Overload in Sickle Cell Disease?” Global Iron Summit, Berlin, Germany
- March, 2016 “Thalassemia Patients: Are We Transfusing Them Effectively?” Global Iron Summit, Madrid, Spain

Organizing Roles in Scientific Meetings:

- May, 1997 Seventh Cooley’s Anemia Symposium, Chairperson, New York Academy of Sciences, Cambridge, Massachusetts
- March, 2001 11th International Conference on Oral Chelation in the Treatment of Thalassaemia and Other Diseases, Advisory Board, Catania, Italy
- March, 2005 Cooley’s Anemia Eighth Symposium (sponsored by the New York Academy of Sciences and the Cooley’s Anemia Foundation) Conference Organizer, Lake Buena Vista, Florida
- October, 2010 Child Health Research Center (CHRC) 20th Annual Retreat, Philadelphia, PA. (co-organizer)
- March, 2012 The 2nd National Conference on Blood Disorders in Public Health, Atlanta, Georgia. (Planning Committee)
- March, 2015 Global Iron Summit, Berlin Germany. (Scientific Committee)

Bibliography:

Research Publications, peer reviewed (print or other media):

Cohen A, Schwartz E. Iron chelation therapy with deferoxamine in Cooley's anemia. J Pediatr 92:643-647, 1978.

Cohen A, Schwartz E. Excretion of iron in response to deferoxamine in sickle cell anemia. J Pediatr 92:659-662, 1978.

Cohen A, Schwartz E. Iron chelation therapy in sickle cell anemia. Am J Hematol 7:69-76, 1979.

- Cohen A, Markenson AJ, Schwartz E. Transfusion requirements and splenectomy in Cooley's anemia. *J Pediatr* 97:100-102, 1980.
- Cohen A. Effect of time on hematologic values in prediluted capillary and venous blood. *Am J Clin Pathol* 74:306-307, 1980.
- Cohen A, Schwartz E. Decreasing iron stores during intensive chelation therapy. *Ann NY Acad Sci* 344:405-408, 1980.
- Greenberg J, Curtis-Cohen M, Gill FM, Cohen A. Prolonged reticulocytopenia in autoimmune hemolytic anemia of childhood. *J Pediatr* 97:784-786, 1980.
- Poncz M, Colman N, Herbert V, Schwartz E, Cohen A. Therapeutic management of congenital folate malabsorption. *J Pediatr* 98:76-79, 1981.
- Cohen A, Trotzky M, Pincus D. Reassessment of the microcytic anemia of lead poisoning. *Pediatrics* 67:904-906, 1981.
- Cohen A, Cohen I, Schwartz E. Scurvy and altered iron stores in thalassemia major. *N Engl J Med* 304:158-160, 1981.
- Cohen A, Martin M, Schwartz E. Response to long-term deferoxamine therapy in thalassemia. *J Pediatr* 99:689-694, 1981.
- Cohen A. Chelation therapy in the treatment of iron overload. *Drug Ther* 6:47-58, 1981.
- Graziano JH, Piomelli S, Seaman C, Wang T, Cohen A, Kelleher JF, Schwartz E. A simple technique for preparation of young red cells for transfusion from ordinary blood units. *Blood* 59:865-868, 1982.
- Cohen A, Martin M, Schwartz E. Depletion of excessive liver iron stores with desferrioxamine. *Br J Haematol* 58:369-373, 1984.
- Cohen A, Schmidt JM, Martin M, Barnsley W, Schwartz E. Clinical trial of young red cell transfusions. *J Pediatr* 104:865-868, 1984.
- Cohen A, Mizanin J, Schwartz E. Treatment of iron overload in Cooley's anemia. *Ann NY Acad Sci* 445:274-281, 1985.
- Nelson L, Elfman J, Cohen A. Management of Ludwig's angina in a patient with severe hemophilia A with factor VIII inhibitors: Report of a case. *J Dent Child* 52:306-311, 1985.
- Poncz M, Greenberg J, Gill FM, Cohen A. Hematologic changes during acute chest syndrome in sickle cell disease. *Am J Pediatr Hematol Oncol* 7:96-99, 1985.
- Hoyt R, Scarpa N, Wilmott RW, Cohen A, Schwartz E. Pulmonary function abnormalities in homozygous beta thalassemia. *J Pediatr* 109:452-455, 1986.
- Cohen A, Seidl-Friedman J. HemoCue system for hemoglobin measurement: evaluation in anemic and non-anemic children. *Am J Clin Pathol* 90:302-305, 1988.
- Schwartz E, Cohen A, Surrey S. Overview of the beta thalassemias: Clinical and genetic aspects. *Hemoglobin* 12:551-564, 1988.

- Michelson A, Cohen A. Incidence and treatment of fractures in thalassemia. *J Ortho Trauma* 2:29-32, 1988.
- Cohen A, Gayer R, Mizanin J. Long-term effect of splenectomy on transfusion requirements in thalassemia major. *Am J Hematol* 30:254-256, 1989.
- Manno CS, Cohen AR. Splenectomy in spherocytosis: Is it worth the risk? *Am J Pediatr Hematol/Oncol* 11:300-303, 1989.
- Cohen AR, Mizanin J, Schwartz E. Rapid removal of excessive iron with daily, high-dose intravenous chelation therapy. *J Pediatr* 115:151-155, 1989.
- Goedert JJ, Kessler CM, Aledort LM, Biggar RJ, Andes WA, White GC, Drummond JE, Vaidya K, Mann DL, Eyster ME, Ragni MV, Lederman MM, Cohen AR, Bray GL, Rosenberg PS, Friedman RM, Hilgartner MW, Blattner WA, Kroner B, Gail MH. Rates, markers and cofactors of human immunodeficiency virus type I infection and AIDS in subjects with hemophilia. *N Engl J Med* 321:1141-1148, 1989.
- Spear ML, Spear M, Cohen AR, Pereira GR. Effect of fat infusions on platelet concentration in premature infants. *J Parenter Enteral Nutr* 14(2):165-168, 1990.
- Cohen A, Martin M, Mizanin J, Konkle DF, Schwartz E. Vision and hearing during deferoxamine therapy. *J Pediatr* 117:326-30, 1990.
- Cohen A, Martin M, Ohene-Frempong, K. Increased blood requirements during long-term transfusion therapy in sickle cell disease. *J Pediatr* 118:405-407, 1991.
- Young TL, Schaffer DB, Cohen AR. Infantile glaucoma associated with the Diamond-Blackfan syndrome. *J Pediatr Ophthal Strab* 29:55-58, 1992.
- Rabkin CS, Hilgartner MW, Cohen AR, Goedert JJ. Incidence of lymphomas and other cancers in HIV-infected and HIV-uninfected patients with hemophilia. *JAMA* 267:1090-94, 1992.
- Burk CD, Miller L, Handler SD, Cohen AR. Pre-operative history and coagulation screening in children undergoing tonsillectomy. *Pediatrics* 89:691-695, 1992.
- Cohen AR, Martin MB, Silber JH, Kim HC, Ohene-Frempong K, Schwartz E. A modified transfusion program for prevention of stroke in sickle cell disease. *Blood* 79:1657-1661, 1992.
- Rackoff WR, Ohene-Frempong K, Month S, Scott JP, Neahring B, Cohen AR. Neurologic events after partial exchange transfusion for priapism in sickle cell disease. *J Pediatr* 120:882-885, 1992.
- Brittenham GM, Cohen AR, McLaren, CE, Martin MB, Griffith PM, Nienhuis AW, Young NS, Allen CJ, Farrell DE, Harris JW. Hepatic iron stores and plasma ferritin concentration in patients with sickle cell anemia and thalassemia major. *Am J Hematol* 42:81-85, 1993.
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- Kim HC, Dugan NP, Silber JH, Martin MB, Schwartz E, Ohene-Frempong K, Cohen AR. Erythrocytapheresis therapy for reduction of iron overload in chronically transfused patients with sickle cell disease. *Blood* 83:1136-1142, 1994.
- Goedert JJ, Cohen AR, Kessler CM, Eichinger S, Seremetis SV, Rabkin CS, Yellin FJ, Rosenberg PS, Aledort LM. Risks of immune deficiency, AIDS and death by purity of factor VIII concentrate. *Lancet* 344:791-792, 1994.
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- Kattamis AC, Shankar S, Cohen AR. Neurologic complications of treatment of childhood acute immune thrombocytopenic purpura with intravenous immunoglobulin G. *J Pediatr* 130:281-283, 1997.
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- Cohen AR, Galanello R, Piga A, DiPalma A, Vullo C, Tricta F. Safety profile of the oral iron chelator deferiprone: A multi-center study. *Br J Haematol* 108:305-312, 2000.
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- CM, Cohen AR, DiMichele D, Hilgartner MW, Aledort LM, Kroner BL, Rosenberg PS, Hatzakis A. End-stage liver disease in persons with hemophilia and transfusion-associated infections. *Blood* 100:1584-1589, 2002.
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- Hoffbrand AV, Cohen A, Hershko C. Role of deferiprone in chelation therapy for transfusional iron overload: *Blood* 102:17-24, 2003.
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- Cohen AR, Galanello R, Piga A, DeSanctis V, Tricta F. Safety and effectiveness of long-term therapy with the oral iron chelator deferiprone. *Blood* 102:1583-1587, 2003.
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- Chu Z, Cohen AR, Muthupillai R, Chung T, Wang ZJ. MRI measurement of hepatic magnetic susceptibility — phantom validation. *Magnetic Resonance in Medicine* 52:1318-1327, 2004.
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- Cappellini MD, Bejaoui M, Agaoglu L, Canatan D, Capra M, Cohen A, Drelichman G, Economou M, Fattoum S, Kattamis A, Kilinc Y, Perrotta S, Pga A, Porter JB, Griffel L, Dong V, Clark J, Aydinok Y. Iron chelation with deferasirox in adult and pediatric patients with thalassemia major: efficacy and safety during 5 years' follow-up. *Blood* 118:884-893, 2011.
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Neufeld EJ; Thalassemia Clinical Research Network. Chelation use and iron burden in North America and British thalassemia patients: a report from the Thalassemia Longitudinal Cohort. *Blood* 119:2746-2753, 2012.

- Marx A, Shaw K, Taylor A, Cohen A. Department of Pediatrics Chair's Initiatives: Promoting innovation and improvement through collaboration, accountability, and faculty development. *J Pediatr* 161:377-378, 2012.
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- Aygun B, Mortier NA, Kesler K, Lockhart A, Schultz WH, Cohen AR, Alvarez O, Rogers ZR, Kwiatkowski JL, Miller ST, Sylvestre P, Iyer R, Lane PA, Ware RE; Stroke With Transfusions Changing to Hydroxyurea (SWITCH) Trial Investigators. Therapeutic phlebotomy is safe in children with sickle cell anaemia and can be effective treatment for transfusional iron overload. *Br J Haematol* 169:262-266, 2015.
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- Ware RE, Davis BR, Schultz WH, Brown, RC, Aygun B, Sarnaik SA, Odame I, Fuh B, George A, Owen W, Luchtman-Jones L, Rogers ZR, Hilliard L, Gauger C, Piccone CM, Lee MT, Kwiatkowski JL, Jackson S, Miller ST, Roberts CW, Heeney MM, Kalfa TA, Nelson SC, Imran H, Nottage KA, Alvarez OA, Rhodes M, Thompson AA, Rothman J, Helton KJ, Roberts D, Coleman J, Bonner MJ, Kutlar A, Patel N, Wood JC, Piller L, Wei P, Luden J, Mortier NA, Stuber S, Luban NLC, Cohen AR, Pressel SL, and Adams RJ. TCD with transfusions changing to hydroxyurea (TWITCH): hydroxyurea therapy as an alternative to transfusions for primary stroke prevention in children with sickle cell anemia. *Lancet* 387:661-670, 2015.
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transfused children with sickle cell anemia: baseline results from the TWITCH trial. *Br J Haematol* 172: 122-130, 2016.

Submitted manuscripts:

Taher AT, Saliba AN, Kuo KH, Giardina PJ, Cohen AR, Neufeld EJ, Aydinok Y, Kwiatkowski JL, Chapin JC, Kramer W, Jeglinski BI, Pietropaulo K, Berk G, Viprakasit V. A phase 1b study evaluating the safety and pharmacokinetics of SP-4210, a novel iron chelator for transfusional iron overload. Submitted, June 2017.

Vichinsky E, Cohen A, Thompson AA, Giardina PJ, Lal A, Paley C, Cheng WY, McCormick N, Sasane M, Qiu Y, Kwiatkowski JL: Epidemiologic and clinical characteristics of thalassemia intermedia in the United States. Submitted, June 2017.

Contributions to peer-reviewed clinical research publications, participation cited but not by authorship:

Walter PB, Macklin EA, Porter J, Evans P, Kwiatkowski JL, Neufeld EJ, Coates T, Giardina PJ, Vichinsky E, Olivieri N, Alberti D, Holland J, Harmatz P; Thalassemia Clinical Research Network (Cohen AR, collaborator). Inflammation and oxidant-stress in beta-thalassemia patients treated with iron chelators deferasirox (ICL670) or deferoxamine: an ancillary study of the Novartis CICAL670A0107 trial. *Haematologica* 93:817-825, 2008.

Harmatz P, Jonas MM, Kwiatkowski JL, Wright EC, Fischer R, Vichinsky E, Giardina PJ, Neufeld EJ, Porter J, Olivieri N; Thalassemia Clinical Research Network (Cohen AR, collaborator). Safety and efficacy of pegylated interferon alpha-2a and ribavirin for the treatment of hepatitis C in patients with thalassemia. *Haematologica* 93:1247-1251, 2008.

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Morris CR, Kim HY, Trachtenberg F, Wood J, Quinn CT, Sweeters N, Kwiatkowski JL, Thompson AA, Giardina PJ, Boudreaux J, Olivieri NF, Porter JB, Neufeld EJ, Vichinsky EP; Thalassemia Clinical Research Network (Cohen AR, collaborator). Risk factors and mortality associated with an elevated tricuspid regurgitant jet velocity measured by Doppler-echocardiography in thalassemia: a Thalassemia Clinical Research Network report. *Blood* 118:3794-3802, 2011.

Walter PB, Porter J, Evans P, Kwiatkowski JL, Neufeld EJ, Coates T, Giardina PJ, Grady RW, Vichinsky E, Olivieri N, Trachtenberg F, Alberti D, Fung E, Ames B, Higa

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Abstracts: (selected from past 5 years)

Wood JC, Cohen A, Aygun B, Imran H, Luchtman-Jones L, Thompson AA, Fuh B, Davis BR, Pressel SL, Schultz WH, Ware RE. Extrahepatic iron deposition in chronically transfused children with sickle cell anemia – baseline findings from the TWiTCH trial. American Society of Hematology Annual Meeting, New Orleans, Louisiana, 2013.

Wood JC. Rogers ZR, Odame I, Kwiatkowski J, Lee M,. Owen WC, Cohen A, St. Pierre, T, Davis BR, Parker C, Schultz WH and Ware RE. Liver iron concentration by MRI in chronically transfused children with sickle cell anemia in the TWiTCH trial. American Society of Hematology Annual Meeting, New Orleans, Louisiana, 2013.

Ware WR, Davis BR, Schultz WH, Brown C, Aygun B, Sarnaik SA, Odame I, Fuh B, George A, Owen W, Luchtman-Jones L, Rogers ZR, Hilliard L, Gauger C, Piccone CM, Lee MT, Kwiatkowski J, Jackson S, Miller ST, Roberts CW, Heeney MM, Kalfa TA, Nelson SC, Imran H, Nottage KA, Alvarez OA, Rhodes M, Thompson AA, Rothman J, Helton KJ, Roberts D, Coleman J, Bonner MJ, Kutlar A, Patel N, Wood JC, Piller L, Wei P, Luden J, Mortier NA, Stuber S, Luban NLC, Cohen AR, Pressel SL, and Adams RJ. TCD with transfusions changing to hydroxyurea (TWiTCH): hydroxyurea therapy as an alternative to transfusions for primary stroke prevention in children with sickle cell anemia. American Society of Hematology Annual Meeting (plenary session), Orlando, FL 2015.

Aygun B, Mortier N, Rogers ZR, Owen W, Fuh B, George, A, Kalfa TA, Kwiatkowski J, Lee M, Imran H, Miller ST, Wood JC, Cohen AR, Pressel SL and Ware RE.. Iron unloading by therapeutic phlebotomy in previously transfused children with sickle cell anemia: the TWiTCH Experience. American Society of Hematology Annual Meeting, San Diego, CA, 2016.

Wood JC, St Pierre T, Aygun B, Mortier N, Schultz WH, Piccone CM,, Hankins J, Rogers ZR, Owen W, Odame I, Sarnaik SA, Hilliard L, Cohen AR and Ware RE. Agreement between R2 and R2* liver iron estimates is independent of the type of

iron removal therapy: results from the TWITCH trial. American Society of Hematology Annual Meeting, San Diego, CA, 2016.

Editorials, Reviews, Chapters, including participation in committee reports (print or other media):

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- Cohen A, Schwartz E. Thalassemia, in *Brenneman's Practice of Pediatrics*, Harper and Row, Inc., Hagerstown, 1989 (revised, 1983).
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Heeney M, Cohen A, editors. Iron Disorders. Hematology-Oncology Clinics of North America, Elsevier Publishing, August 28, 2014

Grant Support:

PAST

H87-MC24055 (Kwiatkowski) HRSA Comprehensive Thalassemia Program	06/01/2012 – 05/31/2015 \$138,204	0.12 calendar
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The major goal of this project is to design and implement multi-center clinical research studies in thalassemia.

5 R01 HL095647-05 (Ware) NIH/NHLBI	08/01/2009-07/31/2015 \$80,640	0.6 calendar
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TCD with Transfusions Changing to Hydroxyurea

The major goal of this project is to compare transfusions with hydroxyurea and phlebotomy for the prevention of secondary stroke and the management of iron overload in children with sickle cell disease.

2 K12 HD043245-11 (Cohen, changed to St. Geme in 2013) 12/01/2012-11/30/2017
0.6 calendar
NIH/NICHD \$432,000
CHOP Pediatric Scholars Program

The goal of this project is to provide pediatric physician-scientists who are junior faculty members with mentorship and resources to increase their research skills.

5 T32-HD43021-09 (Cohen) 05/01/2009-04/30/2014 0.0 calendar
CHOP Institutional Training in Pediatric Research \$253,425

The goal of this project is to enhance the development of physician-scientists by providing mentorship and research training for fellows in pediatric subspecialties.

1U01 HL65232 (Cohen) 7/1/00 – 6/30/05 25%
NIH \$268,235
The Mid-Atlantic Thalassemia Consortium

The major goal of this project is to design and implement multi-center clinical research studies in thalassemia.

R01-HL61185-01 (Cohen) 8/1/98 – 7/31/03 30%
NIH \$209,189
Erythrocytapheresis to lower iron loading in thalassemia

The major goal of this project is to test the efficacy of erythrocytapheresis in lowering iron loading in patients with thalassemia.

R01-HL61182 (Wang) 9/30/98 – 7/31/01 10%
NIH \$118,272
Quantitative Assessment of Iron Overload by MRI

The major goal of this project is to develop magnetic resonance techniques to assess body iron stores.

CDC U27-CCU313193-03 (Cohen) 9/30/96 – 9/29/01 5%
Centers for Disease Control \$66,059
Prevention of Complications of Hemophilia

The major goal of this project is to develop strategies to prevent complications of hemophilia and its treatment in children with bleeding disorders, with special emphasis on inhibitors, hepatitis, and joint disease.

MCJ 422007-09 (Cohen) 10/1/96 – 5/31/01 0%
Maternal Child Health Bureau \$46,905
Regional Hemophilia Diagnostic and Treatment Center

The major goal of this project is to provide comprehensive, multidisciplinary, culturally competent family-centered care to children with bleeding disorders and their families.

P01 HL64190 (High)

12/1/99 – 11/30/04

20%

NIH/NHLBI

\$259,623

Gene Therapy for Hemophilia (PPG)

Project 4: Human application of AAV-mediated muscle-directed factor IX gene transfer. The major goal of this project is to conduct a clinical trial of gene therapy for hemophilia B.