

CURRICULUM VITAE: DANIEL J. WALLACE, M.D., F.A.C.P., M.A.C.R.

Up to date as of July 1, 2018

Personal:

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Education:

University of Southern California, 2/67-6/70, BA Medicine, 1971.
University of Southern California, 9/70-6/74, M.D, 1974.

Postgraduate Training:

7/74-6/75 Medical Intern, Rhode Island (Brown University) Hospital, Providence, RI.
7/75-6/77 Medical Resident, Cedars-Sinai Medical Center, Los Angeles, CA.
7/77-6/79 Rheumatology Fellow, UCLA School of Medicine, Los Angeles, CA.

Medical Boards and Licensure:

Diplomate, National Board of Medical Examiners, 1975.
Board Certified, American Board of Internal Medicine, 1978.
Board Certified, Rheumatology subspecialty, 1982. California License: #G-30533.

Present Appointments:

Medical Director, Wallace Rheumatic Study Center
Attune Health Affiliate, Beverly Hills, CA 90211
Attending Physician, Cedars-Sinai Medical Center, Los Angeles, 1979-
Clinical Professor of Medicine, David Geffen School of Medicine at UCLA, 1995-
Professor of Medicine, Cedars-Sinai Medical Center, 2012-
Expert Reviewer, Medical Board of California, 2007-
Associate Director, Rheumatology Fellowship Program, Cedars-Sinai Medical Center, 2010-
Board of Governors, Cedars-Sinai Medical Center, 2016-
Member, Medical Policy Committee, United Rheumatology, 2017-

Honorary Appointments:

Fellow, American College of Physicians (FACP)
Fellow, American College of Rheumatology (FACR) 1979-2015
Master, American College of Rheumatology, 2015-
Member, Royal College of Physicians (London)
Organizing Committee: Lupus 2014, Quebec, Quebec, Canada

Hospital Appointments:

Cedars Sinai: **Clinical Chief of Rheumatology, 1991-1996**
Chairman, Medical Peer Review 1985-1987, Member 1982-1989
Intern Selection Committee, 1979-1993
Medical Advisory Committee, 1985-1988, 1991-1996
Performance Improvement Committee, 1991-1996
Hospital Peer Review Committee, 1986-1989, 1992-1994
Division of Rheumatology Reappointment/Peer Review, 1988-present
Pharmacy gamma globulin committee, 2002-present
Chairman of Department of Medicine Search Committee 2011-2012
Clinical Academic Promotions and Appointments Council 2011-2014
Division of Rheumatology Executive Committee 2011-

Chief Rheumatology Consultant, City of Hope Medical Center, Duarte, CA 1980-1988

Attending Physician, UCLA Center for the Health Sciences, 1981-

Prior Academic Appointments:

Associate Clinical Professor, UCLA School of Medicine 1988-1995

Assistant Clinical Professor, UCLA School of Medicine, 1981-1988

Assistant Clinical Professor, USC School of Medicine 1979-1981

Books

1. Wallace, DJ and Dubois EL, eds., *Dubois' Lupus Erythematosus, 3rd Edition*, Lea and Febiger, Philadelphia, PA, 1987, 775 pp
2. Wallace, DJ and Hahn, BH, eds, *Dubois' Lupus Erythematosus, 4th Edition*, Lea and Febiger, Philadelphia, PA, 1993, 953 pp
3. Wallace DJ, *The Lupus Book*, Oxford University Press, NY/London, 1995, 258 pp
4. Wallace DJ, Hahn BH, Eds., *Dubois' Lupus Erythematosus, 5th Edition*, Williams and Wilkins, Baltimore, MD, 1997, 1289 pp
5. Wallace DJ, Wallace JB, *Making sense of fibromyalgia*, Oxford University Press, NY/London, 1999, 242pp
6. Wallace DJ, *The Lupus Book, New and Revised*, Oxford University Press, NY/ London, 2000, 271 pp
7. Lane NE, Wallace DJ, *All About Osteoarthritis*, Oxford University Press, NY/London, 2001, 258 pp.
8. Wallace DJ, Hahn BH, Eds, *Dubois' Lupus Erythematosus, 6th Edition*, Lippincott Williams & Wilkins, Philadelphia, PA, 2002, 1348 pp
9. Wallace DJ, Wallace JB, *All About Fibromyalgia*, Oxford University Press, NY/London, 2002, 258 pp
10. Wallace DJ, Wallace JB, *Fibromyalgia: An essential guide for patients and their families*, Oxford University Press, NY/London, 2003, 196 pp.
11. Wallace DJ, Editor, *The New Sjogren's Syndrome Handbook*, Oxford University Press, NY/London, Revised and Expanded 3rd Edition, 2005, 265 pp.
12. Wallace DJ, Clauw DJ, *Fibromyalgia and other central pain syndromes*, Lippincott Williams & Wilkins, Philadelphia, PA, 421 pages, 2005. 10.
13. Wallace DJ, *The Lupus Book: A Guide for Patients and their Families*, Oxford U Press, NY/London, 2005, 291 pp.
14. Wallace DJ, Hahn BH Eds, *Dubois' Lupus Erythematosus, 7th Edition*, Lippincott Williams & Wilkins, 2007, Philadelphia, PA, 1441 pp.
15. Wallace DJ, *Lupus: The Essential Clinician's Guide*, Oxford American Rheumatology Library, Oxford University Press, New York, 2008.
16. Clauw DJ, Wallace DJ, *Fibromyalgia: The Essential Clinician's Guide*, Oxford American Rheumatology Library, Oxford University Press, New York, 2009.
17. Wallace DJ, Editor, *The Sjogren's Book: Sjogren Syndrome Foundation*, Fourth Edition, Oxford University Press, New York 2012, 395 pages.
18. *Rheumatoid Arthritis*, Michael Weisman, Oxford American Rheumatology Library, Daniel J Wallace, Series Editor, 90 pages, 2011.
19. Wallace DJ, Hahn BH, *Dubois' Lupus Erythematosus and Related Syndromes, 8th edition*, Elsevier Saunders, Philadelphia, PA, 694 pages, 2013
20. Wallace DJ, *The Lupus Book: A Guide for patients and their families*, Oxford U Press, NY/Oxford, 2013.
21. Wallace DJ, *Lupus: The Essential Clinician's Guide*, Oxford American Rheumatology Library, Oxford U Press, Oxford/New York, 2014. 115 pp.
22. Wallace DJ, Wallace JB, *Making Sense of Fibromyalgia*, Oxford U Press, NY/Oxford, 2nd Edition, 2014, 257 pp.
23. Wallace DJ, Hahn BH, *Dubois' Lupus Erythematosus and Related Syndromes, 9th edition*, Elsevier, Philadelphia, PA 2018 (in press)

Book Chapters/Supplements

1. Plasma-lymphocytapheresis for the treatment of rheumatoid arthritis, D. Goldfinger, D. Wallace, J. Klinenberg, in *Plasma Exchange Therapy*, Ed. by H. Borberg, R Reuther, pp. 215-219, GeorgThiemeVerlag, Stuttgart, 1981.
2. The current status of therapeutic apheresis in the management of rheumatoid arthritis, DJ Wallace, D. Goldfinger, J. Klinenberg, in *Therapeutic Apheresis and Plasma Perfusion*, Ed. by J Tyndall, AR Liss, New York, pp. 27-32, 1982. (Also published as Prog. Clin. Biol. Res. 106:27-32 1982).
3. A controlled study on the use of plasmapheresis in steroid/immunosuppressive resistant systemic lupus erythematosus with nephrotic syndrome, DJ Wallace, D Goldfinger, S. Nichols, D Goodman, M Fichman, M Stewart, JR Klinenberg, in *Proceedings of the First International Congress of the World Apheresis Association*, Ed. by T Uda, Y Shlokawa, N Inoue, ISAO Press, pp. 91-102, 1987.
4. New and experimental therapeutic modalities, JR Klinenberg and DJ Wallace, in *Diagnosis and Management of Rheumatic Diseases*, W. Katz, ed, Lippincott, Philadelphia, pp 855-858, 1988.

5. Wallace D, The Treatment of Mild Lupus, in *Proceedings of the 2nd Intl Conference on S.L.E., Professional Postgraduate Services*, Singapore, 1989, pp. 126-129.
6. Wallace DJ, Peter JB, Bowman RL, Wormsley SB, Silverman S, Fibromyalgia, cytokines, fatigue syndromes, and immune regulation, in *Advances in Pain Research and Therapy: Myofascial Pain and Fibromyalgia, Volume 17*, JR Friction and EA Awad, Eds, Raven Press, NY, 1990, pp. 277-287.
7. Wallace D, Metzger AL, The antiphospholipid syndrome, in *A Decade of Lupus*, H Aladjem editor, Lupus Foundation of America, 1991, Rockville, MD, pp. 126-127.
8. Wallace DJ, Does stress or trauma cause aggravate rheumatic disease?, in *Bailliere's Clinical Rheumatology, Exercise and Rheumatic Disease*, 8: 149-159, 1994. Richard Panush and Nancy Lane editors, Bailliere Tindall, London.
9. Wallace DJ, Antimalarial agents and lupus. *Rheumatic Disease Clinics*, in *Systemic Lupus Erythematosus*, WJ McCune editor, 20: 243-263, 1994, WB Saunders, Philadelphia, PA.
10. Wallace DJ, Metzger AL, Ashman RF, Rheumatic diseases, in *Manual of Allergy and Immunology, 3rd edition*, G Lawlor, T Fischer, D Adelman, eds, Little, Brown & Company, 1995, pp 303-352.
11. Silverman S, Gluck O, Silver D, Tesser J, Wallace D, Neumann K, Metzger A, Morris R, The prevalence of autoantibodies in symptomatic and asymptomatic patients with breast implants and patients with fibromyalgia, in *Immunology of Silicones*, M Potter and NR Rose, Eds, Springer-Verlag Berlin, Heidelberg, Germany, pp 317-322, 1996. (Also published as *Curr Top Microbiol Immunol* 1996; 210:317-322.)
12. Wallace D, Metzger AL, Systemic lupus Erythematosus: Clinical aspects and treatment, in Koopman WJ, editor, *Arthritis and Allied Conditions: A Textbook of Rheumatology, 13th Edition*, Williams and Wilkins, Baltimore, MD, 1997, pp 1319-1345.
13. Wallace D, Autoimmune Connective Tissue Disorders, Rakel RE, *Conn's Current Therapy 2000*, WB Saunders, Philadelphia, PA, pp 774-778.
14. Wallace DJ, Severe lupus, in *Rheumatology (3rd Ed)*, MC Hochberg, AJ Silman, JS Smolen, ME Weinblatt, MH Weisman, Mosby, Edinburgh, 2003, pp. 1419-1426.
15. Shinada S, Wallace DJ, Laboratory features of cutaneous lupus, *Cutaneous lupus erythematosus*, Springer, Berlin, 2004, p.p. 311-322.
16. Wallace DJ, Fibromyalgia and the neurobiology of sleep, in *Sleep and Psychosomatic Medicine*, SR Pandi-Perumal, RR Ruoti, M Kramer, Eds, 2007; Informa Healthcare, Abington, United Kingdom, pp. 249-259.
17. Wallace DJ, Systemic lupus erythematosus, *Encyclopedia of Stress*, 2nd edition, G Fink, editor, Elsevier, 2007, Oxford UK, v 3 pp 708-712.
18. Hallegua DS, Wallace DJ, Sleep and fibromyalgia in the elderly, in *Principles and Practice of Geriatric Sleep Medicine*, SR Pandi-Perumal, JM Monti and AA Monjan, Eds, Cambridge University Press 2009, pp 160-166.
19. Wallace DJ, Management of nonrenal systemic lupus erythematosus 2, in *Targeted Treatment of the Rheumatic Diseases*, M Weisman et al Eds, Saunders Elsevier, Philadelphia, PA, 2009, pp 91-107.
20. Goldenberg DM, Wallace DJ, Epratuzumab in SLE, In: *Immunotherapeutic agents for SLE*. Shoenfeld Y, Meroni PL, Cervera R (Eds). Future Medicine Ltd, London , UK , 56–68 (2012).
21. Editor of Up to Date sections since 2011 for: a) Antimalarial drugs in the treatment of rheumatic diseases, b) Diagnosis and differential diagnosis of systemic lupus erythematosus in adults, c) Musculoskeletal manifestations of systemic lupus erythematosus, d) Overview of therapy of the therapy and prognosis of systemic lupus erythematosus in adults, e) Patient education: Systemic lupus erythematosus (Beyond the Basics), Lippincott
22. Wallace DJ, Management of nonrenal and non-central nervous system lupus, in *Rheumatology (5th edition)*, MC Hochberg, AJ Silman, JS Smolen, ME Weinblatt, MH Weisman Eds, Saunders Elsevier, Philadelphia, PA 2014, Chapter 134 pp 1099-1106.
23. Wallace DJ, Fibromyalgia and the neurobiology of sleep, in *Sleep and Psychosomatic Medicine*, SR Pandi-Perumal, Narasimhan, Kramer M, Eds. 2016, pp 127-136, 2nd edition, CRC Press, Boca Raton, FL
24. Wallace DJ, Editor, Special Issue: Biologic Therapies, *Lupus* 2016; volume 25, number 10
25. Wallace DJ, Weisman MH, Clinical features of systemic lupus erythematosus, in *Rheumatology (6th edition)*, MC Hochberg, E Gravalles, A Silman, J Smolen, ME Weinblatt, MH Weisman, Eds, Elsevier, Philadelphia, PA, 2018, Chapter 135, pages 35.1-35.12.
26. Wallace DJ, Systemic and Biologic Agents for Lupus Erythematosus, Springer, Cham Switzerland, 2018, in *Biologic and Systemic Agents in Dermatology*, Paul Yamauchi Ed, 2018, pp 377-390

Organizations and Positions Held:

Lupus Foundation of America

Co-chairman, 1999-2000

Member, Board of Directors 1991-1998, Vice President 1999-2000

National Medical Advisory Board, 1988-

Los Angeles Chapter Medical Advisory Board, Co-Chair 1989-1999

Member, Los Angeles County Medical Association
 Member, California Medical Association
 Member, American Medical Association
 Member, Southern California Rheumatism Society
 Member, Arthritis Foundation Pacific Region
 LA. Metro Committee Chairman, 1989-1994
 Community Services Committee, 1989-1994
 Medical and Scientific Committee, 1989-1994
 Institutional Grants Committee, 1989-1994
 Fibromyalgia Subcommittee, 1988-; Chairman 1990-1995
 Representative, National House of Delegates, 1987, 1990
 Board of Directors, Southern California, 1994-2006

Member, American College of Rheumatology
 Committee on Rheumatologic Practice, 1982-1985
 Lupus Council, 1986-; Chairman 1990-1991
 Nominations Committee 1991-1994
Research & Education Foundation Board of Directors, 1993-1999; Chairman 1995-1999
 Nominating Committee, 2005-2007
 Lupus Abstract Selection Committee 2010-2015
 Research and Education Foundation, National Advisory Council Meeting, 2011-

The American Lupus Society (merged with Lupus Foundation of America, 1996)
 National Medical Advisory Board, 1988-1996
 Los Angeles Chapter Medical Advisory Board, 1980-1996
 Los Angeles Chapter Chief Medical Advisor, San Fernando Valley Medical Advisory Board

Member, American Society for Apheresis
 Medical Executive Committee, 1987-1989
 Editor, *ASFA Newsletter*, 1987-1989

United Scleroderma Foundation Board of Directors, 1990-1997
 Scleroderma Foundation, Southern California, Board of Directors, 2001-2004
 American Fibromyalgia Syndrome Association, Medical Advisory Board, 1994-
Board of Directors, Lupus Research Institute, 2000- 2015
 Founder, Lupus LA 2000-
 Co-Chairman, Industrial Relations Council 2012-

Sjogren's Syndrome Foundation Medical Advisory Board, 1997-
 Medical Advisory Board, Maryland Lupus Foundation, 2001-2004
 Examination writer, American Board of Internal Medicine, for Internal Medicine Boards, 1994, 1995
 Member, SLICC (Systemic Lupus International Coordinating Committee), 2003-
 NIH Lupus Biomarkers Committee, 2004-2010
 Member, Lupus Clinical Trials Consortium (LCTC), 2005-2010
 Scientific Advisory Board for lupus, 23andme, 2015-
 NYU Judith and Stewart Colton Center for Autoimmunity, NYU Langone Medical Center, External Advisory Committee, 2014-
 Lupus Research Alliance, Co-chairman, Lupus Industry Council 2016-
 Executive Board, Lupus Clinical Investigative Network, 2016-

Laboratory Experience:

Therapeutic apheresis project, Cedars-Sinai Medical Center, 1977-1997
 Cedars-Sinai Lupus Research Laboratory 1990-
 Summer research fellow, Cedars-Sinai Medical Center, 1968-1970, under Dr. Leon Morgenstern,
 Chairman, Department of Surgery, dealing with wound healing of intestinal anastomoses and synthesizing trypsin inhibitor.
 USC Cancer Virology Laboratory, Dr. Murray Gardner and J. Earle Officer, PhD, 1972-1974, retroviruses and aging.

Publication Review Experience:

Editorial Board, *Journal of Clinical Apheresis*, 1982 -2004
 Editorial Board, *Lupus*, 1997-
 Editorial Board, *Bulletin on the Rheumatic Diseases*, 1998-2004
 Editorial Board, *Arthritis & Rheumatism*, 1998- 2003
 Editorial Board, *Journal of Musculoskeletal Pain*, 1999-
 Editorial Board, *Journal of Rheumatology*, 1999-
 Editorial Board, *Journal of Clinical Rheumatology*, 2003-

Editorial Board, *Future Rheumatology*, 2006-2016
Editor, *Current Opinion in Rheumatology*, Lupus issues, 1994-2000
Editor-in-Chief, *Current Rheumatology Reviews*, 2005-2008
Reviewer for over 50 medical journals

Honors

Globus Award, Best Medical Paper, 1984-5, *Mt. Sinai J Medicine*
Humanitarian Award, Lupus Foundation of America, 1989 and 1991
Best Doctors in the United States, Town and Country Magazine, October, 1989
The American Lupus Society, Lupus Hall of Fame, 1989
"The Best Doctors in America," Woodward/White, Aiken, SC, 1994
Best Doctors in Los Angeles, *Los Angeles Magazine*, 1996
Jane Wyman Humanitarian Award, Arthritis Foundation, 1996
Expert Consultant, Medical Board of California, 1995-
Lupus Foundation of America, Outstanding Service Award, 1997
"Spirit of Hope" Award, Southern California Scleroderma Foundation, 2001
Achievement Award, SLE Foundation, 2002
James R Klinenberg Achievement Award, Arthritis Foundation Southern California Chapter, 2004
Founder's Award, Lupus LA, SLE Foundation, 2008
Top Doctor, US News and World Report, 2011-
Medical Achievement Award, SLE Foundation 2011
Sjogren's Syndrome Foundation, Healthcare Professional Leadership Award, 2012
Lupus Foundation of Northern California, Outstanding Commitment to Treatment and Research, 2016
Los Angeles County Medical Association, Innovation Award for Community Service, 2017
Jane Wyman Humanitarian Award, Arthritis Foundation, 2018

Research Grants:

BSRG Cedars-Sinai Research Grant, \$10,000, 1978-1979,
Apheresis in rheumatoid arthritis
Haemonetics Research Institute Grant, \$25,000, 1980-1981,
Double-blind controlled trial of apheresis in rheumatoid arthritis
Kroc Foundation Grant, \$59,000, 1981-1982,
Apheresis in systemic lupus erythematosus
Haemonetics Research Institute Grant, \$25,000, 1982,
The immunology of apheresis
Parker Foundation Grant, \$76,250, 1983-1985,
Selective immunoadsorption in rheumatoid arthritis
Winthrop Pharmaceuticals Grant, \$5,000, 1989-199,
The role of hydroxychloroquine on lipids
The American Lupus Society Grant, \$28,000, 1990-1992,
An index of lupus literature
Winthrop Pharmaceuticals Grant \$7,500, 1991-1993,
Cytokines and their influence on hydroxychloroquine
Co-Investigator with Dr. Mariana Linker-Israeli, 3 year NIH grant for \$350,000
"Interleukin-6 genetic polymorphisms", 1994-1996
Co-Investigator with Dr. Mariana Linker-Israeli, NIH grant for \$442,670,
"Abnormal IL-6 Production in SLE," Agency Award #2RoLAR4252-04, 1997-1999
American Fibromyalgia Syndrome Association, \$24,600, 1998-1999,
The role of the Th-1/Th-2 axis in fibromyalgia
Food and Drug Administration, RF 412-3324A, \$24,000, 2000-2002
Comparison of IV cyclophosphamide to mycophenolate mofetil for induction therapy of active Class III-IV
nephritis in systemic lupus erythematosus. Co-investigator with M Weisman for Cedars effort
Co-investigator with Dr. Michelle Petri, NIH Grant, Brain Connections: Cognitive function in SLE, National Institute of
Arthritis, Musculoskeletal and Skin disorders, 2003-, Grant number: R01AR4912501, 2004-2008
AROSE: Revising ACR diagnostic/classification criteria for lupus, Lupus Foundation of America, NIH-NIAMS
ARO051871-013, 2004-2009
The role of mycophenolate mofetil in patients with extra-renal lupus, Aspreva, \$62,500, 2006-
A study to investigate pro-inflammatory HDL cholesterol as an indication of risk for atherosclerosis in subjects with

systemic lupus erythematosus, Alliance for Lupus Research, \$72,500, 2006-2008

Co-investigator, Rituxan in the treatment of refractory adult and juvenile dermatomyositis (DM) and adult polymyositis (NIH Grant # N01-AR-4-2273), 2007-2010

Co-investigator, The use of abatacept in lupus nephritis (NIH Grant # N01A115416), 2008-2014
HHSN268200700036C Langford (PI) 09/13/10 – 08/14/12

Concurrent Pilot Studies in Giant Cell Arteritis and Takayasu Arteritis to Examine the Safety, Efficacy, and Immunologic Effects of Abatacept (CTLA4-IG) in large Vessel Vasculitis
The purpose of this study is to determine the effectiveness and safety of the medication abatacept in giant cell arteritis or Takayasu's arteritis.
Role: Co-Investigator

Canadian Arthritis Society Clarke/Bernatsky (PI) 05/05/09 – 12/31/11
Lymphoma Risk in SLE: A Consequence of Immune Suppression or Stimulation?
The purpose of this research is to help identify SLE patients at highest risk for lymphoma and provide guidance regarding the appropriate use of ISDs in both inducing and maintaining remission in SLE.
Role: Co-Investigator

Wallace (PI) Improving the Recognition, Diagnosis, and Referral Patterns of Patients with Systemic Lupus Erythematosus Through Enhanced Care Coordination and Practice Efficiencies – A QI/PI CME Demonstration Project for the Cedars-Sinai Medical Group, France Foundation, 2012- , \$150,000
BASJ02-A randomized, double-blind, placebo controlled trial of Baminercept, a omphotoxin-B receptor fusion, protein, for the treatment of primary Sjogren's syndrome, 2012-2016, Autoimmune Centers of Excellence

Subject: Introduction and Contact Information for the Protocol, "A Phase Ib Study of Milatuzumab Administered Subcutaneously in Patients with Active Systemic Lupus Erythematosus (SLE)," Submitted by Daniel Wallace MD, Cedars-Sinai Medical Center, Los Angeles, California in Support of Proposal, "CD74 Immunotherapy of Systemic Lupus Erythematosus," William Wegener MD, Immunomedics, Incorporated, Morris Plains, New Jersey, Proposal Log Number PR121764, Award Number W81XWH-13-1-0392, HRPO Log Number A-17786, 2014-

NIH R34A114453; Planning grant for 'Mesenchymal Stem Cell Therapy for Active Systemic Lupus Erythematosus' (7/15/14-6/30-15), Co investigator

5P50AR060804-03; Oklahoma Sjogren's Syndrome Center of Research Translation (7/1/13-6/30/15), Co-Investigator

U34 AR067392 Olsen & Karp (MPIs) 02/01/15-01/31/17
NIH/NIAMS
Hydroxychloroquine Treatment for Prevention of Systemic Lupus Erythematosus
Planning grant for a multicenter, placebo-controlled trial of hydroxychloroquine in incomplete lupus patients to determine whether this can ameliorate, delay or prevent progression to SLE.
Role: Co-investigator

LR170141 (Department of Defense); Subject: "Inflammation and Metabolic Reprogramming of Lupus Monocytes – Mechanism of the Pathobiology of Lupus Cardiovascular Disease"
PI: Caroline Jefferies (2018 -
Role: Co-Investigator

Past Drug Study Collaborative Protocols:

Eli Lilly, Benoxaprofen, 1977-1979 in RA

Syntex, Naproxen, 1980-1982 in RA

CIBA-Geigy, Diclofenac, 1985-1987 in RA

Winthrop, Hydroxychloroquine-methotrexate, 1988-1992 in RA

Rorer, Calcitonin in Fibromyalgia, 1989

Sandoz, Sandostatin in Fibromyalgia, 1990-1991

Smith-Kline, Namebutone for Osteoarthritis, 1991-1992

Genelabs, Dehydroepiandrosterone (DHEA) for Lupus, 1994-1997, an open label and double blind trial

La Jolla Pharmaceuticals, LJP 394 for Lupus Nephritis, 1996-1999, Double-blind trial

Immunex, Double-blind trial of TNF-alpha (Enbrel) for Rheumatoid Arthritis, 1997-1999

Boehringer-Ingelheim, Meloxicam for Rheumatoid Arthritis, 1998-1999
Boehringer-Ingelheim, Meloxicam for Osteoarthritis, 1998-1999

Merck, MK-0966 for Rheumatoid Arthritis, 1998

Immunex, Open-label trial of TNF-alpha (Enbrel) for Rheumatoid Arthritis, 1997-

Biogen, Anti-CD40L therapy for lupus nephritis, 1998-1999

Smith Kline, SB 217969/Clenolixib for Rheumatoid Arthritis, 1998-1999

La Jolla Pharmaceuticals/Abbott, LJP 394 for Lupus nephritis, open label #90-11, 1998-2000

Merck, MK-0966 for osteoarthritis, 1998-1999

Smith Kline, SB 217969/Clenolixib (Open Label), for rheumatoid arthritis 1998-1999

Roche, Randomized, double-blind trial of Ro32-3555 (Trocade) for rheumatoid arthritis, 1998-2000

Amgen, A randomized, placebo controlled double-blind, multicenter, dose-finding study to evaluate the safety and efficacy of weekly administration of PEGylated recombinant methionyl human soluble tumor necrosis factor receptor Type I (PEGsTNF-RI) in patients with rheumatoid arthritis, 1999-2000

Merck, An active comparator and placebo controlled, parallel group, double-blind 52 week study to assess safety and efficacy of MK-0966 in rheumatoid arthritis patients, 1999-2000

Merck, Randomized, double-blind multicenter study to evaluate tolerability and effectiveness of rofecoxib (MK-0966) 25 mg/d vs naproxen 500 mg bid in patients with osteoarthritis, 1999-2000

Zeneca, Randomized, double blind placebo controlled, parallel group multicenter trial to assess the analgesic efficacy and tolerability of treatment with multiple doses of 1600 mg ZD6416 bid compared with treatment with placebo in patients with osteoarthritis of the hip or knee, 1999-1999

Biogen C99-1021, An open label, multiple dose study to evaluate the efficacy, safety and pharmacokinetics of BG 9588 (anti CD40 antibody) in subjects with proliferative lupus glomerulonephritis, 1999-2001

Anergen, A phase I double blind, randomized placebo controlled, dose escalation study to evaluate the safety, tolerability and biological activity of a 2 week induction course and 1 maintenance cycle of AG 4263 in subjects with rheumatoid arthritis, 1999-1999

IDEC, Phase II randomized, double blind, placebo-controlled, multiple center, multiple dose, dose finding safety, tolerance and efficacy study of IDEC-131 in patients with active SLE, 1999-2000

Centocor, An open-label trial of anti-TNF chimeric monoclonal antibody Infliximab in patients with active rheumatoid arthritis, 1999-2000

Merck, A randomized, placebo-controlled, parallel group, double-blind study to evaluate the safety and efficacy of rofecoxib 25 mg and celecoxib 200 mg in patients with osteoarthritis of the knee or hip, 2000

Cypress, A post approval, market preference, unblinded general experience study to gather additional information about the ProSORBA column in general rheumatology practice settings, 1999-2000

Searle, Clinical protocol to evaluate the long term safety of valdecoxib in treating the signs and symptoms of rheumatoid arthritis, 1999-2001

Searle, Clinical protocol of a multicenter, double-blind, placebo-controlled randomized, comparison study of the efficacy and safety of 3 valdecoxib doses and naproxen in treating the symptoms and signs of rheumatoid arthritis, 1999-2001

Knoll, A multicenter randomized placebo-controlled Phase II study of the human anti-TNF D2E7 administered as subcutaneous injections in rheumatoid arthritis patients treated with methotrexate, 1999-2001

Knoll, A multicenter Phase II study of the human anti-TNF antibody D2E7 administered as subcutaneous injections in rheumatoid arthritis patients treated with methotrexate, 2000-2001

Amgen, A double-blind extension study to provide treatment with PEGylated recombinant human methionyl soluble tumor necrosis factor Type I (PEG sTNF-R1) to subjects completing trials TNF 980246 and TNF 990136, 2000

Fujisawa, An open-label, long-term study to evaluate the safety of Prograf (tacrolimus) for the treatment of rheumatoid arthritis, 2000-2002

Fujisawa, a randomized double-blind placebo controlled study to assess the efficacy and safety of prograf (tacrolimus) in the treatment of rheumatoid arthritis in patients who have failed one or more disease modifying antirheumatic drugs, 2000-2002

Knoll DE031: A multicenter randomized double-blind placebo-controlled study of the safety of human anti-TNF monoclonal antibody D2E7 in patients with active rheumatoid arthritis, 2000-2008

Knoll, Study of New Onset Rheumatoid Arthritis (SONORA), 2000-2006

Knoll DE013, A prospective multicenter randomized double-blind active comparator-controlled parallel-groups study comparing the fully human monoclonal antibody TNF alpha antibody D2E7 every second week with methotrexate given weekly and the combination of D2E7 and methotrexate administered over 2 years in patients with early rheumatoid arthritis, 2000-2008

La Jolla Pharmaceuticals, LJP 394-90-09, A randomized, double-blind, placebo-controlled, multicenter safety and efficacy trial of LJP 394 in systemic lupus erythematosus (SLE) patients with a history of renal disease, 2000-2002

Centocor, MEDIII Pso-A-1, A multicenter placebo, controlled, double blind, randomized study of anti-TNF chimeric monoclonal antibody (cA2, infliximab) in patients with active psoriatic arthritis, 2000-2002

Bristol-Myers Squibb, IM 101-101, A multicenter, randomized, double-blind, placebo controlled study to evaluate the safety and efficacy of intravenous infusions of BMS-188667 given monthly in combination with sub-cutaneous injections of etanercept given twice weekly to subjects with active rheumatoid arthritis, 2001-2008

Amgen, 2000223, A multicenter double-blind study to evaluate the safety and efficacy of anakinra (r0metHul IL-ra) and etanercept in subjects with rheumatoid arthritis using methotrexate, 2001-

Allergan, 192371-011-01, A multicenter, double masked, randomized, vehicle controlled parallel group study of the safety and efficacy of cyclosporin ophthalmic emulsion used twice daily for 6 months in patients with moderate to severe keratoconjunctivitis, 2001-2002

Vertex, VX00-745-102, A 12 week, randomized, double-blind, placebo-controlled, dose-ranging study of VX-745 in patients with rheumatoid arthritis, 2001-2002

Hoffman-LaRoche, WA15541c, A double-blind, randomized, placebo controlled study to evaluate the safety and efficacy of Ro32-3555 (Trocade) as adjunct to background antirheumatoid therapy, in preventing structural damage in rheumatoid arthritis, 2000-2002

Knoll, DE020, A multicenter 2 year continuation study of the human anti-TNF antibody D2E7 administered as a subcutaneous injection in patients with rheumatoid arthritis, 2001-

Proctor and Gamble, The effect of testosterone patch on activity of systemic lupus erythematosus, 2001-

Pharmacia, 872-IFL-0513-004, Clinical protocol for a double-blind, placebo-controlled, randomized six week comparison study for the efficacy of valdecoxib 20 mg q d and rofecoxib 25 mg q d in relieving the signs and symptoms of osteoarthritis of the knee, 2001-2002

Optime Therapeutics, LEDA.C.001, A phase II study of the safety and efficacy of topical liposome-encapsulated diclofenac analgesic (LEDA) in patients with osteoarthritis of the knee, 2001-2002

Centocor, START Protocol CO1168T41, A randomized, double-blind trial of the safety of TNF-alpha chimeric monoclonal antibody (Infliximab) in combination with methotrexate compared to methotrexate alone in patients with rheumatoid arthritis on standard disease modifying anti-rheumatic background therapy, 2001-2003.

XOMA, HURA 501.02, A Phase II, Double-blind, Placebo-controlled Study to Determine the Safety, Efficacy, Pharmacokinetics and Pharmacodynamics of Efalizumab in Subjects with Moderate to Severe Rheumatoid Arthritis on a Stable Dose of Methotrexate, 2002-2004

Cypress Bioscience, Inc., FMS-021, A Phase II, Multicenter, Double-Blind, Randomized, Placebo-Controlled Study of Milnacipran for Treatment of Fibromyalgia, 2002-2003.

Immunex Corporation, 016.0034, Rheumatoid Arthritis DMARD Intervention and Utilization Study (RADIUS 1), 2001-2003.
Immunex Corporation, 016.0036, Phase 3 Randomized, Double-Blind, Placebo-controlled Study of 50mg Etanercept (Enbrel®) Administered SC Once Weekly in Patients with Active Rheumatoid Arthritis, 2002-2003

Immunex Corporation, 016.0037, Multicenter, Double-blind, Placebo-Controlled, Randomized Phase 3 Study of Etanercept in the Treatment of patients with Ankylosing Spondylitis, 2002-2003

Amgen, Inc., 20020103 KONTROL, Psychometric Assessment of the Cedars-Sinai Health Related Quality of Life (CSHQ-RA) Instrument in Rheumatoid Arthritis Subjects Receiving Kineret (Anakinra) Therapy, 2002-2003.

Isis Pharmaceuticals, Inc., ISIS 104838-CS7, A Double-blind, placebo-controlled, randomized trial of the safety, efficacy, and pharmacokinetic profile of ISIS 104838 (TNF- antisense oligonucleotide) subcutaneous injections in active rheumatoid arthritis patients, 2002-2004

Scios, Inc., 782.344, A Phase II, Multicenter, Randomized, Double-Blind, Placebo-Controlled, Dose-Escalating Study of SCIO-469 in Patients with Active Rheumatoid Arthritis Receiving Methotrexate, 2002-2004.

Janssen Pharmaceutica Products, L.P., CIS-USA-154, Limited access protocol for the use of oral cisapride in the treatment of refractory gastroesophageal reflux disease and other gastrointestinal motility disorders, 2002-2005

Abbott, A multicenter study of the safety of human anti-TNF monoclonal antibody D2E7 in subjects with active rheumatoid arthritis, 2002-

Amgen, Rheumatoid arthritis DMARD intervention and utilization study (RADIUS 2), 2002-2005

Genelabs, The use of GL701 in the prevention of osteoporosis in patients on corticosteroids with lupus erythematosus, 2003-2005

Cipher Canada, A double-blind, randomized, placebo-controlled, multi-dose, Phase III, parallel group study of Tramadol ER in the relief of signs and symptoms of osteoarthritis of the hip and knee, 2003-2005

Centocor, A multicenter, double-blind trial of anti TNF alpha chimeric monoclonal antibody (Infliximab) for the treatment of subjects with psoriatic arthritis, 2003-2005

Centocor OPPOSITE, Open label, Pilot Protocol of Patients with Rheumatoid Arthritis who switch to Infliximab after incomplete response to etanercept, 2003-2005

Amgen MRI, The use of MRI to Describe and Identify the Early Findings Leading to Foot Erosions in High Risk Subjects with Rheumatoid Arthritis, 2003-

Glaxo Smith Kline, CXA20006, A Multicenter, Randomized, Double-Blind, Placebo and Active-Controlled, Phase 2, Parallel Group, dose-ranging Finding study To assess the Safety and efficacy of GW406381 Administered For 42 days to Subjects with Rheumatoid Arthritis, 2003-2005

IDEC, 102-20/WAI17042, A Randomized, Placebo-Controlled, Double-Blind, Multicenter Study to Evaluate the Safety and Efficacy of Rituximab in Combination with MTX in Subjects with Active Rheumatoid Arthritis Who have had an inadequate Response to MTX and Anti-TNFalpha Therapies 2003-

Novartis, CCOX189A2335, A 13-Week, multicenter, international, randomized, double-blind, placebo-controlled,

parallel-group study of COX189 200mg in patients with rheumatoid arthritis using naproxen 500mg b.i.d. as comparator, 2003-2004

Prometheus, 03-MTX-02, Measurement of Methotrexate and Folate Polyglutamate Levels and MTHFR Polymorphisms in a Cross-Section of Rheumatoid Arthritis Patients to Assess Correlations of Toxicity and Efficacy, 2003-2006

Human Genome Science, LBRA01, A Phase 2, Multicenter, Double-Blind, Placebo-Controlled, Dose-Ranging Study to evaluate the safety, tolerability and Efficacy of LymphoStat-B™ (LSB) in subjects with RA, 2004-2005

Abbott, M02-537, A Multicenter continuation trial for Patients Completing Study M02-518 and M02-570 of the Human Anti-TNF Monoclonal Antibody Adalimumab (D2E7) in Patients with Moderate to Severely Active Psoriatic Arthritis, 2003-2006

Biorad, Collection of Prospective Samples for Investigational Studies of Bio-Rad BioPlex 2200 ANA Screen on the BioPlex 2200, 2004-

SLICC (Systemic lupus erythematosus International Coordinating Committee) Registry for Atherosclerosis, 2003-, Registry for central nervous system lupus, 2003-
Prometheus Imuran SLE: An Open Label Safety and Efficacy Trial of Imuran for Patients with Systemic Lupus Erythematosus, 2004-2006

Wyeth Research: Protocol 3140A1-200-WW: A Double-Blind, Placebo-Controlled, Parallel, Randomized Study to Evaluate the Efficacy and Safety of 3 Oral Dose Levels of TMI-005 in Subjects with Active Rheumatoid Arthritis on a Background of Methotrexate, 2004-2005.

LJP: Protocol 394-90-14: A Randomized, Double-Blind, Placebo-Controlled, Four-Arm Parallel-Group, Multicenter, Multinational Safety and Efficacy Trial of 100mg and 300mg of LJP 394 in Systemic Lupus Erythematosus (SLE) Patients with a History of Renal Disease, 2004-2007.

Scios: Protocol B007: A 24-Week, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Study of the Efficacy of Oral SCIO-469 in Subjects with Active Rheumatoid Arthritis Who are not Receiving DMARDS Other than Hydroxychloroquine, 2004-2006

Orphan Medical, Inc.: Protocol OMC-SXB-26: Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Multi-Center Trial Comparing the Effects of Orally Administered Xyrem (sodium oxybate) with Placebo for the Treatment of Fibromyalgia, 2004-2005

Amgen B cell: An exploratory study to characterize the variability in circulating B cell populations in subjects with systemic lupus erythematosus (SLE), 2004-2006

A Phase IIB MultiCenter, Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Efficacy and Safety of Abatacept in Combination Therapy with Glucocorticosteroids vs. Placebo plus Glucocorticosteroids in the Treatment of Active SLE and the Prevention of Subsequent Lupus Flares, BMS, 2005-2007

A Randomized, Double-Blind, Placebo-Controlled Multicenter Phase II/III Study to Evaluate the Efficacy and Safety of Rituximab in Subjects with Moderate to Severe SLE, Genentech, 2005-

A Multi national, Multi center, randomized, double blind, placebo controlled, multiple dose, four arm study to assess the efficacy, tolerability and safety, of three different doses of Edratide (TV-4710) Subcutaneous injections in SLE patients, TEVA 2005-2007

A phase III, Randomized, Double blind, Placebo controlled, multi center study of Epratuzumab in patients with acute severe SLE Flares Excluding the Renal or Neurologic Systems, Immunomedics, 2005-2006

A Phase 2, Multi-Center, Double-Blind, Placebo-Controlled, Dose-Ranging Study to Evaluate the Safety, Tolerability, and Efficacy of LymphoStat-B™ Antibody (Monoclonal Anti-BLyS Antibody) in Subjects with Systemic Lupus Erythematosus (SLE), Human Genome Sciences, 2005-2007

A Phase Ib, multi-centre, double-blind, placebo-controlled, dose-escalating, single dose study to assess the safety, pharmacokinetics and pharmacodynamics of TACI-Fc5 when administered subcutaneously to patients with SLE, Serono,

2005-2006

A randomized, double-blind, placebo-controlled study to evaluate the safety, tolerability, pharmacokinetics and pharmacodynamics of AMG 623 following multidose administration in subjects with SLE, Amgen , 2005-2006

A Phase 2 study to evaluate the safety, tolerability and activity of fontolizumab (HuZaf) in patients with active rheumatoid arthritis, Protocol ZAF-711, sponsored by Protein Design Labs, Inc (PDL), 2006-2007

Phase 1 Single Ascending Dose Study of the Safety, Pharmacokinetics, and Pharmacodynamics of TRU-015 Administered Intravenously in Subjects with Rheumatoid Arthritis. Protocol 15001, Trubion 2004-

A double-blind, randomised, placebo controlled, dose escalating, multi-center phase I/II trial of HuMax-CD20, a fully human monoclonal anti-CD20 antibody, in patients with active rheumatoid arthritis who have previously failed one or more disease modifying anti-rheumatic drugs. Protocol Hx-CD20-403
Genmab, 2005-2007

A randomized, double-blind, parallel group, international study to evaluate the safety and efficacy of Ocrelizumab compared to placebo in patients with active Rheumatoid Arthritis continuing Methotrexate treatment. Protocol WA20494/ACT3985g, Genentech, 2005- 2007

A Phase III, Multicenter, Open-Label, Continuation Trial of LymphoStat-B Antibody (Monoclonal Anti-BLys Antibody) in Subjects with Rheumatoid Arthritis (RA) who Completed the Phase II LBRA 01. Protocol LBRA99, Human Genome Sciences, 2006-2009

An Exploratory study to characterize biomarker assays in healthy subjects and in subjects with Rheumatoid Arthritis. Protocol 92005637, Amgen 2006-2007

A Phase I, Randomized, Double-Blind, Placebo-Controlled, Dose-Escalation Study to Evaluate Safety and Tolerability of a Single IV Dose of MEDI-545, a Fully Human Monoclonal Antibody Directed Against Interferon Alpha Subtypes, in Patients Who Have Mild System Lupus Erythematosus (SLE) With Cutaneous Involvement. Protocol MI-CP126, MedImmune, 2006-2008

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A Phase III, Randomized, Double-Blind, Placebo Controlled, Multi-Center study of Epratuzumab in Patients with Active Systemic Lupus Erythematosus. Protocol Immu-103-03
Immunomedics, 2005-2006

A Multi Center, open label, continuation trial of lymphostat b antibody (monoclonal anti-blys antibody) in subject with Systemic Lupus Erythematosus (SLE) who completed the phase 2 protocol lbsl02. Protocol LBSL9, Human Genome Sciences, 2006-

Genentech IFN 3958g, A Phase I, Randomized, Double-blind, Placebo-controlled, escalating single and multiple dose study of the safety, tolerability, and Pharmacokinetics of rhuMAB IFNalpha In adults with mildly active SLE, 2007-2009.

UCB SL0006, An Open-Label Re-treatment Trial for Patients Previously Randomized into the SL0003 and SL0004, Randomized, Double-Blind, Placebo-Controlled, Multi-Center Studies of Epratuzumab in patients with SLE, 2007-.

UCB SL0007, A Phase lib, Randomized, Double Blind, Placebo controlled, dose and dose regimen-ranging study of the Safety and Efficacy of Epratuzumab in Serologically-positive Systemic Lupus erythematosus patients with Active Disease. 2008-2009

Medimmune MPI-CP152, A Phase IB, Multicenter, Randomized, Double-blind, Placebo-controlled, dose escalation study with an open label extension to evaluate the safety and tolerability of multiple intravenous doses of MEDI-545, a fully human Anti-Interferon-Alpha Monoclonal Antibody , in patients with Systemic Lupus Erythematosus 2007-2010.

Human Genome Sciences C1056, A Phase 3, Multi-Center, Randomized, Double-Blind, placebo-controlled, 76-Week Study to Evaluate the Efficacy and Safety of Belimumab (HGS1006, LymphoStat-B™), a Fully Human Monoclonal Anti-Blys Antibody, in Subjects with SLE, 2007-2010.

Human Genome Sciences C1066, A Multi-Center, Continuation Trial of Belimumab (HGS1006, LymphoStat-B-), a Fully Human Monoclonal Anti-BLYS Antibody, in Subjects with Systemic Lupus Erythematosus (SLE) who Completed the Phase 3 Protocol HGS1006-C1056 in the United States. 2008-

BMS Lupus Nephritis IM 101-075, A sequential adaptive phase II/III multi-center, randomized, double-blind, placebo controlled study to evaluate the efficacy and safety of Abatecept versus Placebo on a background of Mycophenolate Mofetil and Glucocorticoids in subjects with active Proliferative Glomerulonephritis due to Systemic Lupus Erythematosus (SLE), 2007-

The systemic lupus erythematosus (SLE) activity gene expression (SAGE) study, XDx protocol SL 105, 2007-2009
BMS IM 101-167, A Phase IIIb, Multicenter, Randomized, Withdrawal study to evaluate the Immunogenicity and safety of Subcutaneously Administered Abatacept in Adults with Active Rheumatoid Arthritis, 2008-.

Novo Nordisk NN8360-3559, A randomized, double-blind, placebo-controlled, single dose-escalation and multiple dose extension trial of NNC 0152-0000-0001 administered i.v. or s.c. in subjects with Systemic Lupus Erythematosus. 2009

Genentech IFN4575g, A phase II, Randomized, Double-blind, placebo-controlled study to evaluate the efficacy and safety of Rontalizumab (rhuMAb IFNalpha) in patients with moderately to severely active Systemic Lupus Erythematosus. 2009-

Amgen (AMG 827) 20070264, A Randomized, Double-blind, Placebo-controlled, Ascending Multiple-dose Study to Evaluate the Safety, Tolerability, Pharmacokinetics, Pharmacodynamics and Efficacy of AMG 827 in Subjects with Rheumatoid Arthritis. 2009- 2010

Crescendo Bioscience, Inc. CR10, Index for Rheumatoid Arthritis Measurement (InFoRM) Study. 2009-

Immune Tolerance Network: Protocol ITN034AI, A randomized, double-blind, controlled, phase II Multicenter trial of CTLA4Ig (Abatacept) Plus Cyclophosphamide vs Cyclophosphamide Alone in the Treatment of Lupus Nephritis. 2009-2010

Array BioPharma Inc. 797-201, A Phase 2, Randomized, Double-Blind, Multicenter, Placebo-Controlled, Parallel-Group Study To Investigate The Safety, Efficacy, Pharmacokinetics and Pharmacodynamics Of 12 Weeks Of Treatment With ARRY-371797 In Patients With Active Ankylosing Spondylitis And Inadequate Response To Conventional Therapy. 2009-2009

UCB C87094, A Phase IIIB, multi-centre study with a 12 week double-blind, placebo-controlled, randomized period, followed by an open-label extension phase to evaluate the safety and efficacy of certolizumab pegol administered to patients with active rheumatoid arthritis. 2008-2010

Roche Laboratories Inc. ML22533/A, An open-label, randomized study to evaluate the safety, tolerability and efficacy of tocilizumab (TCZ) monotherapy or TCZ in combination with non-biologic disease modifying antirheumatic drugs (DMARDs) in patients with active rheumatoid arthritis who have an inadequate response to current non-biologic or biologic DMARDs. 2009 –

SLICC, Lymphoma Risk in SLE: A Consequence of Immune Suppression or Stimulation? 2009-

Lupus Clinical Trials Consortium, Inc., LCTC Lupus Data Registry. 2009-

Cedars Sinai Medical Center, Cross Cultural Spanish Validation of Lupus Pro: A Patient Reported Outcome Measure for Lupus. 2009-

TEVA Pharma - A Phase IIa, Multicenter, Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Safety, Tolerability and Clinical Effect of Laquinimod in Systemic Lupus Erythematosus Patients with Active Lupus Arthritis. PROTOCOL LA-LAQ-202. 2010 - , NCT01085084

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Sanofi Aventis US Inc. - A randomized double blind-placebo controlled dose ranging study to evaluate the efficacy and

safety of SAR153191 in patients with Ankylosing Spondylitis (AS). Protocol Number: DRI11073 – ALIGN. 2010 – 2011

UCB - A phase 3, multicenter, randomized, double-blind, placebo-controlled study to evaluate efficacy and safety of certolizumab pegol in subjects with active axial spondyloarthritis (AS001). 2010 – 2011

UCB. - Phase 3, multicenter, randomized, double-blind, parallel-group, placebo-controlled study to evaluate the efficacy and safety of certolizumab pegol in subjects with adult-onset active and progressive psoriatic arthritis (PsA). 2010 – 2011

Study of Epratuzumab in systemic lupus erythematosus, NCT00383513, 2010

Duke Autoimmunity Pregnancy Registry (DAP Registry), 2010-2012

Studies of B cell abnormalities in Systemic Lupus Erythematosus via MiRNA. 2010 –2011

Concurrent pilot studies in Giant cell arteritis and Takayasu's arteritis to examine the safety, efficacy, and immunologic effects of abatacept (CTLA4-Ig) in large vessel vasculitis. 2010 –

Hp-MMP 9 levels in humans: a pilot study. 2010

GlaxoSmithKline - Lupus Impact Tracker: A Longitudinal Validation Study
Protocol GHO-09-1621 2011-2012

Eli Lilly and Company – A Phase 3, MultiCenter, Randomized, Double-Blind, Placebo Controlled study to evaluate the efficacy and safety of Subcutaneous LY2127399 in patients with Systemic Lupus Erythematosus (SLE), Protocol H9B-MC-BCDS 2011-2012

Teva Pharmaceutical Industries, LLC - A Phase IIa, Multicenter, Randomized, Double-Blind, Placebo-Controlled, Study to Evaluate the Safety, Tolerability and Clinical Effect of Laquinimod in Active Lupus Nephritis Patients, in Combination with Standard of Care (Mycophenolate Mofetil and Steroids)
Protocol LN-LAQ-201 2011 – 2012

UCB, Inc. - A Phase 3, Randomized, Double blind, placebo controlled, multicenter study of the Efficacy and Safety of Four 12-Week Treatment Cycles (48 Weeks Total) of Epratuzumab in Systemic Lupus Erythematosus Subjects with Moderate to Severe Disease (Embody1).
Protocol SL0009 2011- 2012

UCB, Inc. – A Phase 3, Multicenter, open label, extension study to assess the safety and tolerability of Epratuzumab treatment in Systemic Lupus Erythematosus Subjects (Embody 4)
Protocol SL0012 2011 -

Roche - A randomized, double-blind, parallel group study of the safety and effect on clinical outcome of tocilizumab SC versus tocilizumab IV, in combination with traditional disease modifying anti-rheumatoid arthritis drugs (DMARDs), in patients with moderate to severe active rheumatoid arthritis.
Protocol Summacta WA22762 2011 –2014

Study of Lymphoma in Systemic Lupus Erythematosus, SLICC (Systemic Lupus International Collaborative Clinics), 2009-2011

IRBIS (Internal Registry for Biologics in SLE) Phase I, Retrospective data collection, SLICC (Systemic Lupus International Collaborative Clinics), 2010-2012, Phase II and III, 2012-

ACR/EULAR Diagnostic and Classification Criteria for Vasculitis, ACR, EULAR, Vasculitis Foundation, Oxford University, 2011-2012

A study to learn about the safety, effectiveness and effects on the body of abatacept in large vessel vasculitis.
Concurrent pilot studies in Giant cell arteritis and Takayasu's arteritis, Vasculitis Clinical Research Consortium, 2011-2012

Vasculitis Clinical Research Consortium (VCRC) Genetic Repository DNA Protocol, 2011-2016,

A Phase 3/4, Multi-Center, Randomized, Double-Blind, Placebo Controlled, 52 week study to evaluate the efficacy and safety of Belimumab (HGS1066) in Adults subjects of Black Race with Systemic Lupus Erythematosus (SLE), Human Genome Sciences, 2012-

A Randomized, Double-Blind, Placebo-controlled, multiple dose, parallel, Multiple dose-level study to evaluate the safety, tolerability and efficacy of AMG 557 in (SLE) subjects with active Lupus Arthritis, Amgen, 2012- 2014

A Double Blind, Randomized, Placebo-controlled, Multicenter, dose ranging study to evaluate the efficacy and safety of PF-04236921 in subjects with Systemic Lupus Erythematosus, Pfizer, 2012-2013.

OMRF Sjogren's Studies: Gene Expression Profiling in Primary Sjogren's Syndrome

A dose escalation, multi-center study to evaluate the safety, tolerability and proof of mechanism of DV1179 in Subjects with Systemic Lupus Erythematosus, Dynavax, 2012

A Phase 3, multicenter, randomized, double-blind, placebo-controlled study to evaluate the safety and efficacy of subcutaneous LY2127399 in patients with systemic lupus erythematosus (SLE), 2012- 2014 , Lilly , 2012- 2015

A longitudinal observational study of CXCR5, CXCL13 and other biomarkers in patients with lupus and healthy control subjects , Sanofi, 2012-2015

Protocol WA 27893: Prospective, observational safety study of patients with Granulomatosis with polyangiitis (Wegener's) or microscopic polyangiitis treated with rituximab, Genentech 2012- 2014

A Study to Evaluate the Efficacy and Safety of R333 6% Ointment Administered Topically to DLE and SLE Patients with Active Cutaneous Discoid Lesions , Rigel, 2012-2013

A Multicenter, Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Efficacy, Safety, and Tolerability of BIIB023 in Subjects With Lupus Nephritis, Biogen, 2012- 2013

A phase 111, Multicenter, Randomized, Double-blind Placebo-controlled Study to Assess The efficacy and Safety of Tocilizumab in Subjects With Giant Cell Arteritis, Roche, 2013- 2015

Nodality – Characterization of Immune Alterations in Systemic Lupus Erythematosus (SLE) using Single Cell Network Profiling (SCNP) Protocol 2012087 SLE Landscaping, 2013-2014

Ignyta – Molecular Analysis in Biological Specimens from Subjects with Rheumatoid Arthritis (RA) Protocol – IGN-RA104
Ignyta – Molecular Analysis in Biological Specimens from Subjects with Systemic Lupus Erythematosus (SLE) and Non-Lupus Control Protocol – IGN-SLE104, 2013- 2014

A Phase IIb, Multi-Center, Randomized, Double-Blind, Placebo-Controlled, Multidose, 24-Week Study to Evaluate the Efficacy and Safety of Atacicept in Subjects With Systemic Lupus Erythematosus (SLE), 2013-

Protocol IMMU-115-04: A Phase Ib Study of Milatuzumab Administered Subcutaneously in Patients with Active Systemic Lupus Erythematosus (SLE) 2014-

Pharmacokinetic Evaluations of Tabalumab Following Subcutaneous Administration by Prefilled Syringe or Auto Injector in Patients with Systemic Lupus Erythematosus, 2014-2015

An International, Open Label, Randomized Controlled Trial Comparing Rituximab with Azathioprine as Maintenance Therapy in Relapsing ANCA-Associated Vasculitis (RITAZAREM) 2014-

An open-label, Non-randomized, 52-week study to evaluate treatment Holidays and rebound phenomenon after treatment with Belimumab 10mg/kg in Systemic Lupus Erythematosus subjects, 2015-

A 52-week, randomized, double-blind, parallel-group, placebo-controlled study to evaluate the efficacy and safety of a 200 mcg dose of IPP-201101 plus standard of care in patients with SLE, ImmuPharm-Orion-Simbec (Lupuzor), 2015-

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EMR Serono Research & Development Institute, Protocol EMR200527-002 Protocol Title: A Phase Ib study to evaluate the safety, tolerability, PK and Biological Effect of MSC2364447 in systemic lupus erythematosus, 2015-2017

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Protocol I4V-MC-JAHH, Lilly, 2016-, A Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Phase 2 Study of Baricitinib in Patients with Systemic Lupus Erythematosus (SLE)

RSLV-132 Protocol 132-03, Resolve, 2016-, A Phase 2A, Double-Blind, Placebo Controlled Study of RSLV-132 in Subjects with Systemic Lupus Erythematosus

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A Multi-Center, Randomized, Double-Blind, Placebo-Controlled, Parallel Group Phase 2A Study to Assess the Efficacy of RO5459072 in Patients with Sjogren 's Syndrome

2017, Protocol: MS200527-0018, 2017 A Phase II, Randomized, Double-Blind, Placebo-Controlled Dose-Ranging Study To Evaluate the Safety and Efficacy of M2951 in Subjects with Systemic Lupus Erythematosus (SLE)

2017, Aurina Pharma, A Randomized, Controlled Double-blind Study Comparing the Efficacy and Safety of Voclosporin (23.7 mg Twice Daily) with Placebo in Achieving Renal Response in Subjects with Active Lupus Nephritis

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2017, AMPEL, A Randomized, Double-Blind, Active Comparator-Controlled, Crossover Study to Assess the Capacity of Delayed-Release Prednisone (RAYOS®) Compared to Immediate-Release Prednisone to Improve Fatigue and Control Morning Symptoms in Subjects with Generalized Systemic Lupus Erythematosus

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2017, Exagen 17-SLE1 CARE Study, Clinical Laboratory Assessments and Recommendations for Lupus

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