SUBURBAN LUNG ASSOCIATES, S.C.

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| NAME | James M Moss, MD |
|----------------------------|---|
| EDUCATION | Rush Presbyterian/St. Luke's Medical Center PGY- IV & V- Allergy/Immunology Fellow 1988-1990 |
| | University of Illinois- Urbana/Champaign, Illinois PGY- I, II & III- Internal Medicine 1985-1988 |
| | Mt. Sinai Hospital- Chicago, Illinois 1983-1985 PGY- I, II & III- General Surgery |
| | Chicago Medical School, M.D. 1979-1983 |
| PROFESSIONAL POSITION | Suburban Lung Associates. S.C. 7/2006-present 800 Biesterfield Rd, Suite 510 Elk Grove Village IL 60007 |
| | Oakbrook Allergists, S.C. – 2000-2006 120 Oak Brook Center Mall Suite 424 Oak Brook, Illinois 60523 630.574.0460 630.574.0470 fax |
| | Midwest Research Associates- 2000-2003 Medical Arts Associates, Ltd, Moline, Illinois- 1990-2000 Medical Director Student Health Augustana College 1990-1993 Anchor HMO, Oak Park, Illinois-1988-1990 Paxton Community Hospital, Emergency Room, Paxton, Illinois- 1987 Convenient Care Clinic, Carle Clinic, Danville, Illinois- 1986-1988 |
| HONORS | University of Illinois at Urbana/Champain, Illinois Internal Medicine Resident of the Year 1987-1988 |
| CERTIFICATION | Certified; American Board Internal Medicine, 1995; Recertified 2006 Certified; American Board Allergy and Immunology, 99 Recertified 06 |
| SOCIETIES | Illinois State Allergy Society Illinois State Medical Society DuPage Medical Society American College of Physicians American College of Allergy, Asthma and Immunology American Academy of Allergy, Asthma and Immunology |
| LICENSURE | Illinois 036- 070259 |
| PHARMACEUTICAL RESEARCH | C1Q Binding by HIV Infected H-9 Cell Supernatants, |

IVIG Therapy in a patient with hypo-complementemic Vasculitis, A multi-center, double-blind randomized parallel-group study investigating the clinical effect of L858674 in patients with seasonal allergic rhinitis, a pilot study the fall season. Merck. 2000

A phase 2/3 open-label, prospective multicenter study of the pharmacokinetics, efficacy, safety, and tolerability of Immune Globulin Subcutaneous, (human) CE 1200 in subjects with primary immune defiency. Aventis. 2001.

An observational study of the epidemiology and natural history of asthma; Outcomes and treatment regimens (Tenor). Genentech 2001.

An observational study of epidemiology and natural history of asthma; outcomes and treatment regimens. (Tenor) Genentech. 2001.

A multi-center, double-blind, randomized, parallel-group study investigating the clinical effect of Montelukast in patients with seasonal allergic rhinitis over a 4- week treatment period-fall 2001. Merck. 2001

A multicenter, randomized, controlled, open-label study to evaluate the safety of Omalizumab in moderate to severe, persistent asthma subjects already treated with other therapies (ALTO). Protocol 2143g. Genentech, 2001.

Open-label extension study II of Xolair (Omalizumab) in moderate to severe, persistent asthma subjects who completed study Q2143g. Protocol 2461g. Genentech. 2002.

A phase III double-blind, double-dummy, parallel-group, multicenter, placebo-controlled, efficacy and safety study of ciclesonide MDI 400 uG/day, 800 uG/day (ex-valve) and Flovent MDI (fluticasone propionate) 880 uG/day (ex-actuator) administered twice daily for 12-weeks in the treatment of severe persistent asthma in adolescents and adults. Aventis. 2002.

Protocol By217/FK1 021:12 weeks treatment with 125 ug roflumilast versus 250 ug roflumilast versus placebo in patients with asthma. Byk Gulden. 2002.

Boehringer Ingelheim "205.416 A Phase III randomized, double-blind, placebo-controlled, parallel grouptrial to evaluate efficacy and safety of tiotropium inhalation solution delivered via Respimat® inhaler (5 $\mu g/day$) over 48 weeks as add-on controller therapy on top of usual care in patients with severe persistent asthma"-2008

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