



VASCULITIS CLINICAL RESEARCH CONSORTIUM

The VCRC **Longitudinal Study for Eosinophilic Granulomatosis with Polyangiitis (Churg-Strauss) (CSS)**

What is a longitudinal study?

A longitudinal study follows patients over an extended period of time to look at changes in individuals.

Information is collected on a regular basis and studied by research scientists to find new ways

to track disease and predict who will respond to different treatments, to study the genetics and causes of vasculitis, to understand how best to treat patients, and much more.

The Longitudinal Study is not a drug study but it may lead to development of new treatments.

Why participate?

By participating in the Longitudinal Study you will help researchers better understand Churg-Strauss Syndrome and work toward developing more effective and safer therapies.

What does this study involve?

If you are eligible and decide to participate in this study, you will be asked to come to a clinical center on a regular basis to meet with a physician, to answer medical questions, and to provide blood and urine samples. The study will last for several years.

Who qualifies?

You must have a confirmed diagnosis of Churg-Strauss Syndrome and be willing to be seen for study visits at one of the VCRC study sites.

What is a study visit?

During a study visit, you will meet with a study physician, a study coordinator, and a research nurse. Medical information will be collected and a physical exam conducted. The research nurse will collect blood and urine specimens from you.

Where will the study take place? How do I participate?

If you are interested in participating in this study or would like additional information, please contact a study coordinator at the clinical center most convenient for you:

Boston University School of Medicine, Boston, Massachusetts

Study Coordinator: Naomi Amudala

E-mail: namudala@bu.edu

Tel: 617-414-2512

Principal Investigator: Paul A. Monach, MD, PhD

Cleveland Clinic Foundation, Cleveland, Ohio

Study Coordinator: Katie Gartner

E-mail: gartnek@ccf.org

Tel: 216-445-1397

Principal Investigator: Carol A. Langford, MD, MHS

Mayo Clinic College of Medicine, Rochester, Minnesota

Study Coordinator: Sara Biorn

E-mail: biorn.sara@mayo.edu

Tel: 507-284-4862

Principal Investigator: Ulrich Specks, MD

St. Joseph's Healthcare, Hamilton, ON

Study Coordinator: Sandra Messier,

E-mail: smessier@stjoes.ca

Tel: 905-522-1155 Ext. 35873

Principal Investigator: Nader Khalidi, MD

Mount Sinai Hospital, Toronto, ON

Study Coordinator: Julia Farquharson,

E-mail: jfarquharson@mtsinai.on.ca

Tel: 416-586-8616

Principal Investigator: Simon Carette, MD

University of Pennsylvania Philadelphia, Pennsylvania

Study Coordinator: Brian Rice

E-mail: brian.rice@uphs.upenn.edu

Tel: 215-614-4407

Principal Investigator: Peter Merkel, MD, MPH

University of Pittsburgh, Pittsburgh, Pennsylvania

Study Coordinator: Dawn McBride

E-mail: dllmc@pitt.edu

Tel: 412-586-3545

Principal Investigator: Larry Moreland, MD

University of Utah, Salt Lake City, Utah

Study Coordinator: Julieanne Nielsen

E-mail: Julieanne.Nielsen@hsc.utah.edu

Tel: 801-585-0798

Principal Investigator: Curry Koenig, MD, MS