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.

By participating in the Longitudinal Study you will help researchers better understand Takayasu’s Arteritis and work toward developing more effective and safer therapies.

Why participate?

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 Takayasu’s Arteritis 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
- **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
- **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@mtsinaion.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: dlmc@pitt.edu
  - Tel: 412-586-3545
  - Principal Investigator: Larry Moreland, MD
- **University of Utah, Salt Lake City, Utah**
  - Study Coordinator: Jennifer Godina
  - E-mail: Jennifer.Godina@hsc.utah.edu
  - Tel: 801-581-4993
  - Principal Investigator: Curry Koening, MD, MS

The VCRC is a part of the Rare Diseases Clinical Research Network, which is supported by the U.S. National Institutes of Health. Visit us online: www.RareDiseasesNetwork.org/vasculitis