

SHORT TITLE: PROMUS

RESEARCH PROTOCOL TITLE: Element Plus US Post-Approval Study

PRINCIPAL INVESTIGATOR: Matthew R. Wolff, MD

RESEARCH SPONSOR: Boston Scientific Corporation

Clinical Trials.Gov Listing at:

<http://clinicaltrials.gov/ct2/show/NCT01589978?term=promus+element+plus&rank=1>

STUDY PURPOSE

The purpose of this study is to collect more information on the PROMUS Stent system, details of the stent procedure and information about the patients who are treated with the stent. The PROMUS stent is used to treat narrow or blocked arteries in the heart. The study will collect information regarding the medical care and outcome of the patients heart condition over a 5 year period.

ELIGIBILITY CRITERIA FOR SUBJECTS

All patients who are treated with a PROMUS stent will be asked to participate in this study.

SUBJECT PAYMENT

There is no compensation for participating in this study.

RESEARCH LOCATION AND CONTACT PERSON

Lynn Skatrud, RN, CCRC
Meriter Wisconsin Heart
2601 West Beltline Hwy.
Suite 200
Madison, WI 53713

PHONE: 608-417-2168