imarc

WE'LL EARN YOUR APPROVAL.

CARDIOVASCULAR EXPERTISE

Conducting Cardiovascular Clinical Trials?

Clinical trials conducted in the Cardiovascular (CV) arena are challenging, first and foremost, due to the variety of underlying disease states and conditions that affect a majority of potential trial participants. This is why choosing IMARC as your Contract Research Organization (CRO) for your CV clinical trial is imperative.

What You Need is IMARC's Specialty

IMARC has a strong therapeutic background, as well as a strong regulatory foundation. IMARC also has a powerful knowledge of CV trials along with the know--how to manage the complexities of your trial.

IMARC has experience in a vast majority of these therapeutic trials:

- Abdominal aortic, aortoiliac, or iliac aneurysms
- Thoracic aneurysms
- Coronary disease
- Renal artery disease
- Superficial femoral artery disease
- Heart failure
- Ischemic cardiomyopathy
- Aortic dissection
- Fibrillation (both atrial and ventricular)

And has experience with these CV trials:

- Endovascular grafts
- Balloon- expandable stents
- Drug- eluting stents (coronary, iliac, renal, femoral)
- Vena cava filter
- Defibrillators
- IVDs for diagnosing MI
- IVUS imaging

Get to the Heart of the Matter with IMARC as Your CRO

Monitoring • Auditing • Research & Development • Consulting

To learn more, contact John E. Lehmann, Director of Business Development **440.801.1540**

Choosing IMARC as Your Go-To CRO

IMARC Research, Inc, based in Cleveland, Ohio, was built upon a regulation--focused compliance foundation and has an expansive resume, which touts several therapeutic verticals, including CV.

Across the board, IMARC focuses on quality over quantity and strives to get it right the first time.

Regulatory Focus – Regulation--focused compliance should be at the foundation of your CRO partner. IMARC always works within a regulatory framework, any non--compliance can be identified quickly, and corrective actions are put in place to secure compliance.

Knowledgeable Background – Started by a nurse whose background includes coronary, medical, and surgical intensive care units and with a depth of nursing experience on the monitoring team, the level of expertise provided by IMARC during CV clinical trials will meet your approval.

Puts Patients First – You will want to work with a company that understands the trial requirements, puts human subject protection at the forefront, and will work with the sites to ensure there is no compromising in this regard.

Gets Results – During a recent trial for a drug--eluting stent, IMARC:

- ✓ Monitored 40+ sites in the US
- ✓ Provided enrollment support
- ✓ Audited each site/followed up on audit findings
- ✓ Provided training and guidance to Japanese monitors
- ✓ Audited 4 Japanese sites and sponsor
- ✓ Monitored 3 sites in Canada for registry
- ✓ Audited European sponsor for registry sites/oversight
- ✓ Provided AE and device audits for sponsor
- ✓ Audit Prepped for 5 sites receiving audit
- ✓ Established presence with the sponsor for FDA inspection

The company was happy to report no findings GLOBALLY. If you are interested in experiencing success like this, consider IMARC as your go-to CRO for CV clinical trials.

For more information on how you can help prepare your sites for a better outcome, starting from Day One, please contact John Lehmann at 440.801.1540 or via e-mail at jlehmann@imarcresearch.com.

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