CARDIOVASCULAR EXPERTISE

imarc
WE'LL EARN YOUR APPROVAL.

IMARC HAS EXPERIENCE IN THESE THERAPEUTIC AREAS:

- 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 CARDIOVASCULAR DEVICES:

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

Conducting Cardiovascular Clinical Trials?

Clinical trials conducted in the cardiovascular 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 Clinical Research Organization (CRO) for your cardiovascular clinical trial is imperative.

What You Need is IMARC Expertise

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

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

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 cardiovascular.

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 experience working on the monitoring team, IMARC provides the level of expertise during cardiovascular clinical trials that 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.

INTERNATIONAL TRIAL

1. Monitored 40+ sites in the US
2. Provided enrollment support
3. Audited each site / followed up on audit findings
4. Provided training and guidance to Japanese monitors
5. Audited 4 Japanese sites and in-country sponsor
6. Monitored 3 sites in Canada
7. Audited European sites and in-country sponsor
8. Conducted targeted AE and device audits for US sponsor
9. Provided audit support for 5 US sites that underwent FDA audit
10. Established presence during FDA inspection of sponsor

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 cardiovascular 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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