

PMG: Focused On Results.

From timely subject recruitment and patient education, to retention of patient volunteers and the collection of quality data for our clients, we are focused on results. Unlike traditional SMOs, we are an Integrated Site Network (ISN) – a fully integrated network of sites solely dedicated to clinical research. The oversight that PMG provides for the clinical trials process across our network ensures quality, compliance and safety.

The Benefits To You:

- Specialized industry partnership opportunities and access to an experienced team of clinical research talent.
- PMG offers network point of contact for each of the following key areas: feasibility, budget and contract negotiations, recruitment, and project management.
- Our ISN provides effective processes for efficient study start-up, seamless communication, and a cohesive experience.



Our Integrated Site Network

Cary Medical Research
Cary, NC

Crescent Medical Research
Salisbury, NC

East Tennessee Medical Research
Johnson City, TN

Metrolina Medical Research
Charlotte, NC

New Hanover Medical Research
Wilmington, NC

Palmetto Medical Research
Mt. Pleasant, SC

Piedmont Medical Research
Winston-Salem, NC

Triangle Medical Research
Raleigh, NC

TriCities Medical Research
Bristol, TN

Unifour Medical Research
Hickory, NC

Wilmington Medical Research
Wilmington, NC

www.pmg-research.com

Contact Us:

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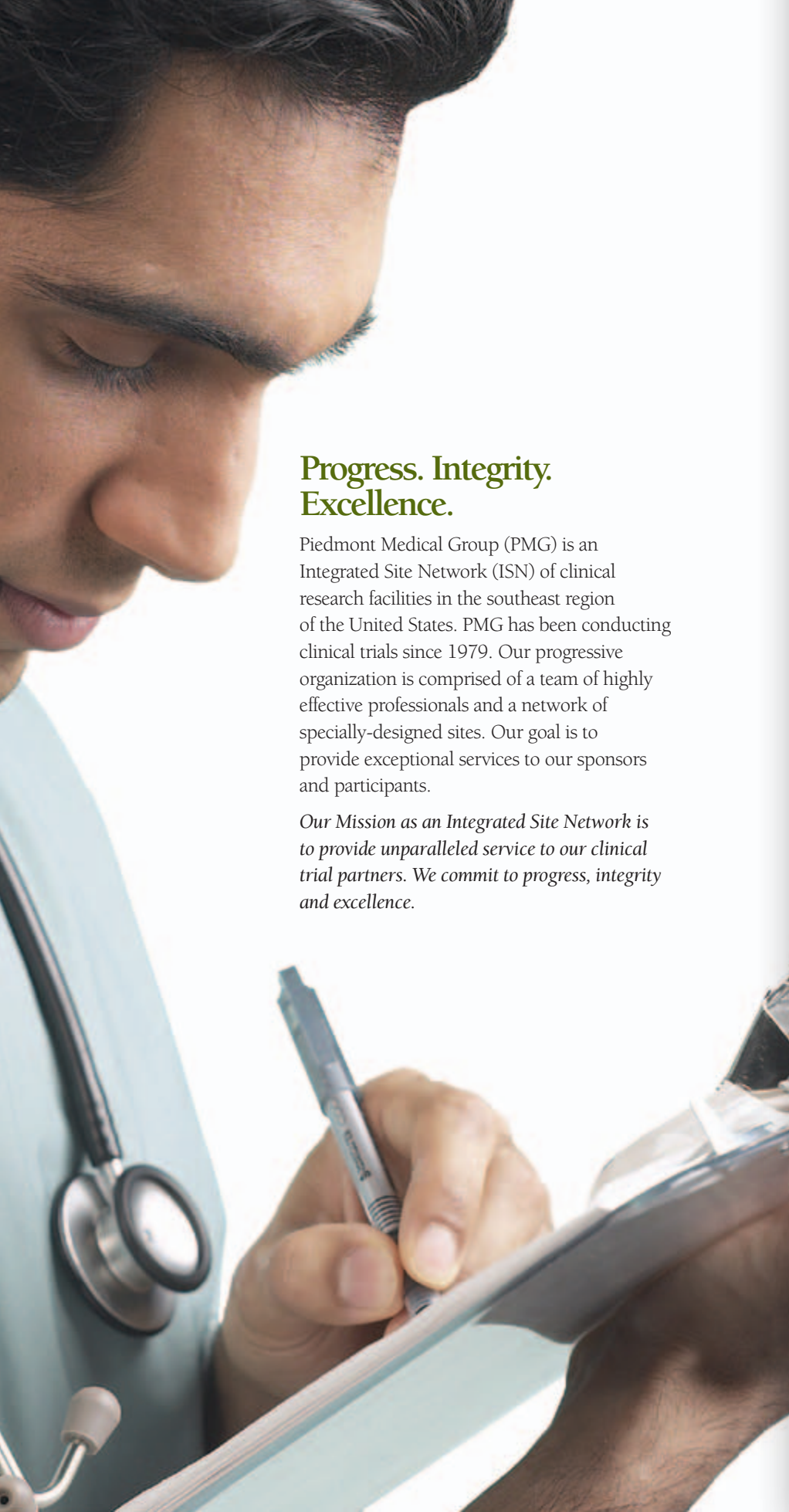
Kathy Cole, Manager of Business Development

T: 919.746.7106



Clinical Trials:
It's what we do.

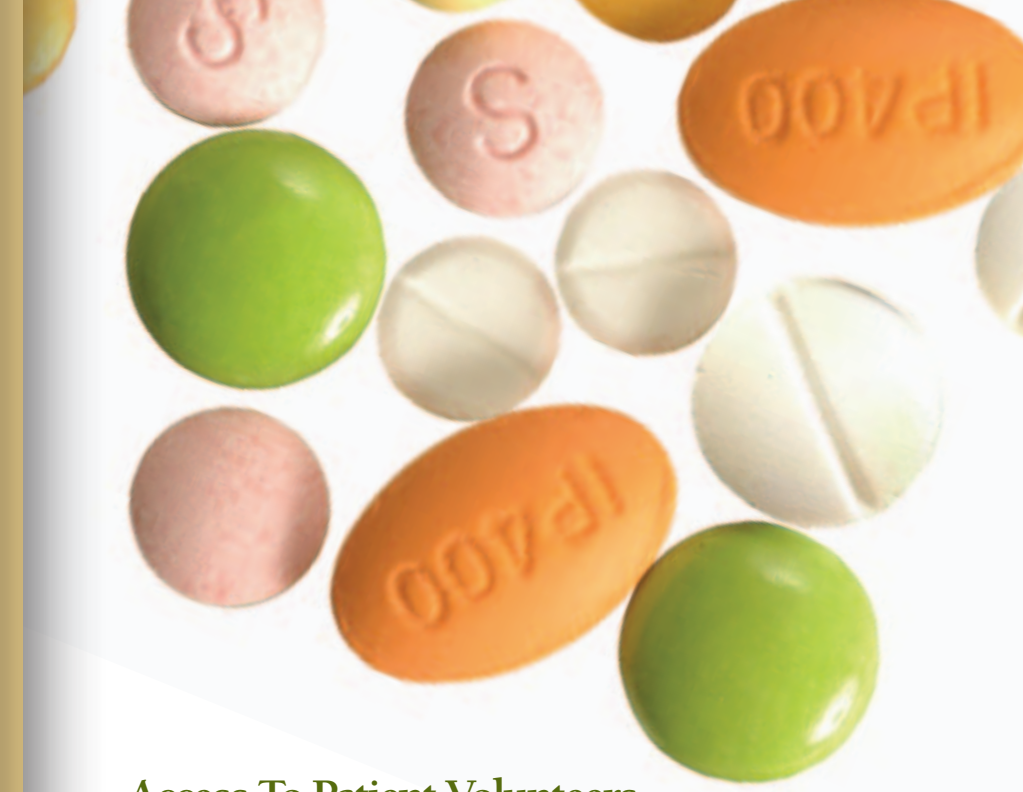




Progress. Integrity. Excellence.

Piedmont Medical Group (PMG) is an Integrated Site Network (ISN) of clinical research facilities in the southeast region of the United States. PMG has been conducting clinical trials since 1979. Our progressive organization is comprised of a team of highly effective professionals and a network of specially-designed sites. Our goal is to provide exceptional services to our sponsors and participants.

Our Mission as an Integrated Site Network is to provide unparalleled service to our clinical trial partners. We commit to progress, integrity and excellence.



Access To Patient Volunteers

- Database with 170,000 potential volunteers.
- Several large, multi-specialty physician practices utilizing EMR systems with access to 1 million active practice patients.
- Network average of 120% for fulfillment of enrollment goals.
- Extensive base of volunteers (50% previous PMG trial participants).
- Staff of professional recruitment specialists.

Quality & Results

- Premier training program and internal QA staff.
- Integrated SOPs developed to ensure compliance with GCPs, subject safety and data integrity.
- Proprietary CTMS includes patient participant, sponsor and study database (5,500 past studies).
- Ability to use Central and Local IRBs.
- Average regulatory document turnaround of 3 business days.
- Contract finalization within 10 business days.
- Master Clinical Trial Agreements.
- FPFV internal requirement of 7 business days.
- TempGuard devices for continuous temperature monitoring of investigational drug.

Staff & Investigator Expertise

- Collectively, PMG sites have 140 years of experience conducting Phase I-IV clinical trials.
- 140 full time staff and 120 Investigators.
- Staff of professional recruitment specialists.
- Integrated team of feasibility specialists.