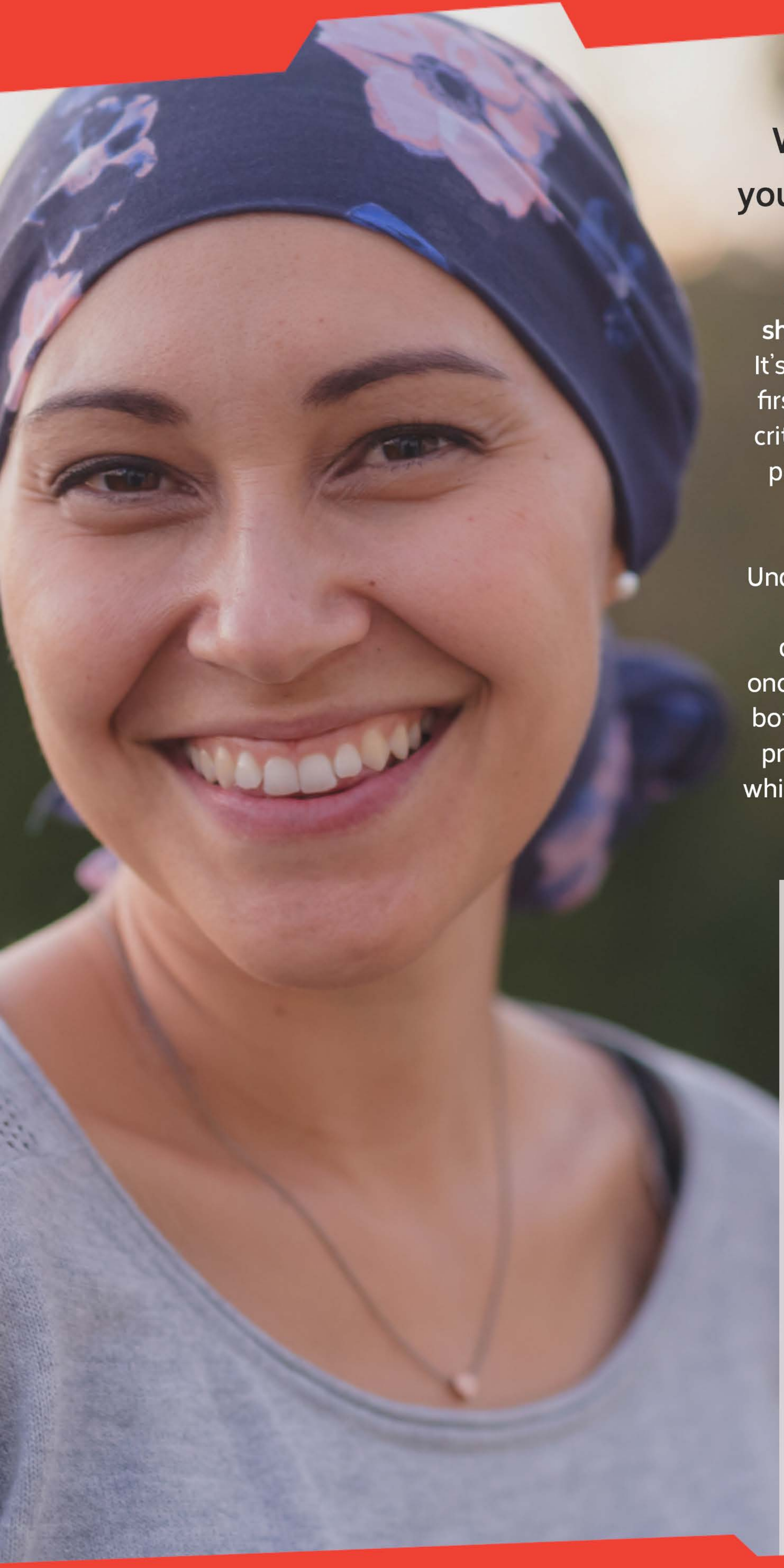
First-In-Human & Early Phase Oncology Trials



When there's a lot at stake, you need a trusted CRO partner

First-in-human (FIH) oncology trials shouldn't be entrusted to just any CRO. It's imperative to partner with one who has first-hand knowledge and expertise in this critical first step where a focus on safety is paramount and each dose and patient is given careful attention.

Understanding that flexibility and speed are also vital to this first stage of clinical development, our resident early phase oncology and rare disease experts can build both detail and flexibility into program and project planning to maximize success, all while mitigating potential issues with dosing protocols and adverse events.

20 YEARS | Cmed

100+ FIH studies

Specialist early phase medical and operations staff

260+ oncology studies
100% oncology experienced project teams

Focused in oncology and rare disease

Full-service CRO for 20+ years in US & EU

Cmed puts your FIH and early phase trials on a path to success

100%
Of Cmed Clinical Staff are Oncology Experts

An expert early phase team

- Clinical Operations
- Clinical Data Management
- Biostatisticians
- Drug Safety
- On-Staff Physicians
- Project Management
- Quality
- Technology

100+
First In Human Trials

260+
Oncology Studies Delivered

Significant experience and solutions to ensure your oncology trial goals are met

- FIH and early phase protocol design, compliance, and delivery
  - Adaptive, Basket, Umbrella, All Comers, Traditional (e,g.,3+3), Innovative (e.g., accelerated titration), Bayesian, etc.
- Investigational product changes/issues
- Data and statistical considerations, challenges, and review
- Safety monitoring
- Enrolment and retention guidance, along with strategies for site engagement
  - Rare patient populations
  - Established site relationships
- Regulatory submission and approvals

Project timelines can be reduced by weeks or months by our experts through protocol adjustments, intelligent use of decentralized services, and real time data visibility via our powerful, holistic technology platform, encapsia®.

- Instant access to all trial data, no matter the source
- Insights that are actionable instantly via live data
- Full compatibility with any mix of EDC, eSource, and third party data sources

FIH and early phase oncology trials benefit from a Cmed partnership

Cmed prides themselves on being a seamless extension to sponsor teams and offer stable, loyal expertise with the right technology, making every clinical trial safer and more efficient, no matter the complexity.

Let's chat and see what our experts can do for your next FIH and early phase oncology trial.

We anticipate. We adapt. We embrace every challenge. We are where you want to be.

Reach out to us today at info@cmedresearch.com