Worldwide Clinical Trials

# Where Personalized Attention Meets Global Scale in Oncology Clinical Development

Your mission is fighting cancer. Our mission is helping you achieve that.

Successful oncology clinical development starts with a partner you can trust that has the global resources you need to execute your study. At Worldwide Clinical Trials, we work alongside you from first-in-human through to registration studies, leveraging our global reach to drive your study forward.

Although we are global, our teams are local. Our therapeutically aligned oncology experts:

- Are dedicated to supporting oncology clinical development
- Live locally in your area
- Are available and accessible to you
- Have deep experience with local regulations, standards of care, and nuances of the European region
- Personalize their approach to your needs and the needs of your trial

## Meet Your Team

Worldwide's senior European subject matter experts have a combined 130+ years of experience supporting oncology development and are available to support you throughout your study.

### Tracey Maranthal, BSc

President of Oncology Business Unit, US

- 20+ years of experience in clinical research and operations expertise, including 15 years in oncology research
- Leads Worldwide's Oncology Business Unit, supporting and guiding operational teams in achieving project success for oncology sponsors of all sizes

### Mireille Cantarini

Senior Medical Director, Medical Affairs, UK

- 25 years of experience in global oncology clinical development
- Provides medical monitoring across all phases, from first-in-human to pivotal registration studies

### Matt Cooper, PhD

Executive Director, Therapeutic Strategy Lead, Oncology, UK

- 25 years of experience in the life science industry conducting trials across all phases and 22 years in oncology clinical development
- Previous experience working for sponsors, sites, and the NHS and has extensive experience in site management and expanded use of oncology therapies

### Jelena Vukotic, MD, MBA

Senior Medical Director, Medical Affairs, Serbia

- 14 years of experience, including 6 years in global oncology clinical development
- Provides medical monitoring across all phases and has previous experience working for sponsors

### Tim Demuth, MD, PhD

Executive Medical Advisor, UK

- Over 20 years of experience in biotech and pharmaceuticals, with key leadership roles at Morphosys AG and Merck KGAA, including the global approval of Tepmetko and a $2.9B acquisition by Novartis.
- Recognized expert in oncology with extensive experience in all phases of drug development, from preclinical research to regulatory approval

### Virgilio Garcia Lerma

Executive Director, Global Regulatory Strategist, Spain

- 15 years of experience in global oncology clinical development
- Heads the strategic regulatory pre-award team, providing strategic advice on regulatory clinical development, tool development for business strategic growth, training and coaching, and oversight

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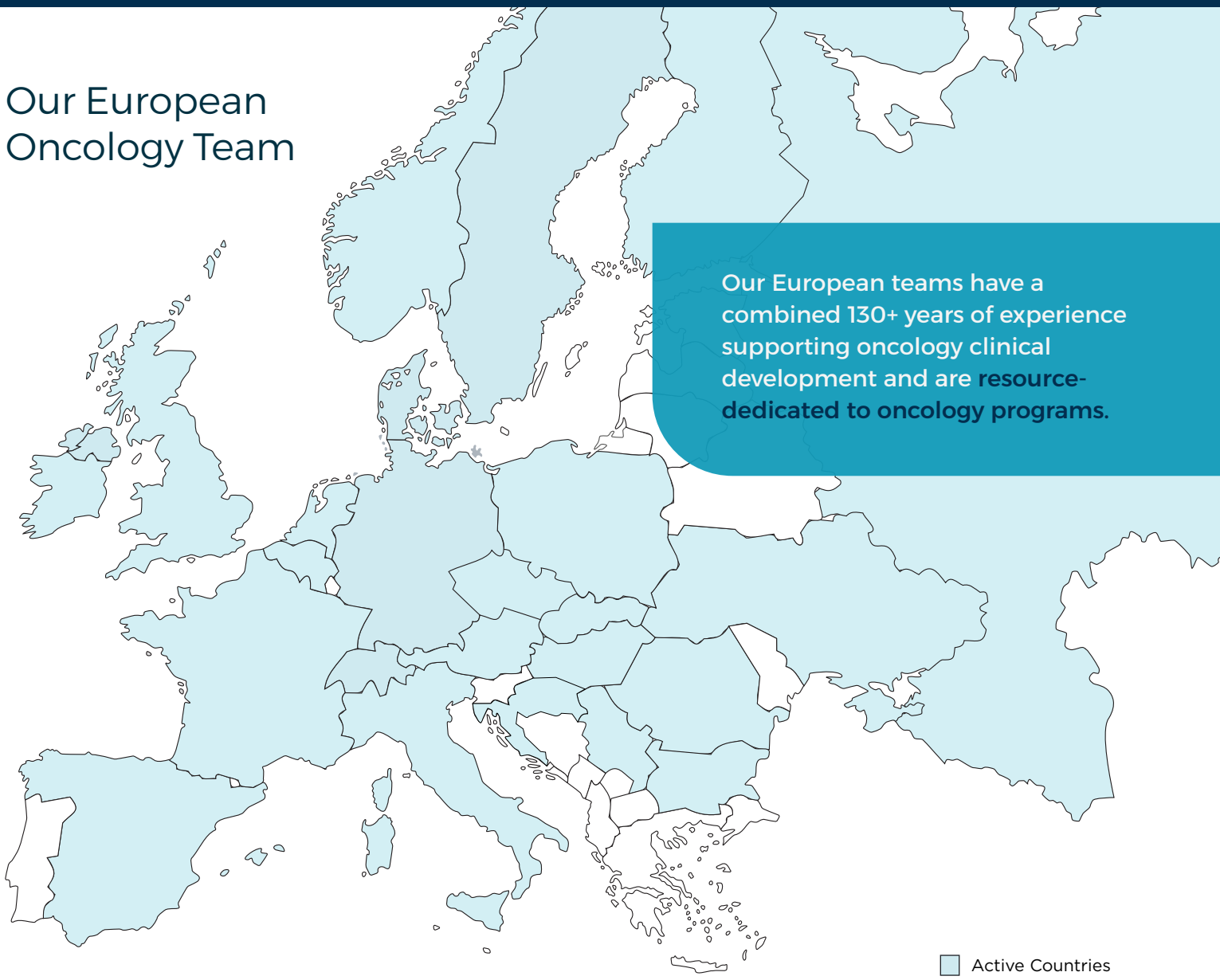
# Global Reach, Local Experts

Our specialized oncology team knows the local standards of care and the regulations in your area, and they have relationships with sites that can support your trial. They are committed and accessible every step of the way, providing a personalized and responsive partnership with the hands-on care that your trial deserves.

## Our European Oncology Team




Our European teams have a combined 130+ years of experience supporting oncology clinical development and are resource-dedicated to oncology programs.

**We're committed to working with sponsors like you to conduct high-quality, global oncology studies. We'd love to talk with you about your oncology program.**

**Contact Us**