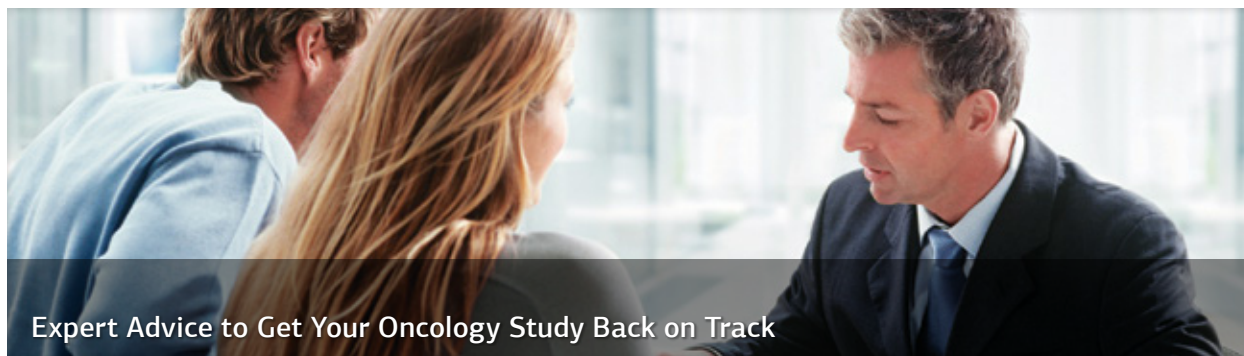




# TROUBLED STUDY ANALYSIS & RECOMMENDATIONS



Expert Advice to Get Your Oncology Study Back on Track

## ONCOLOGY STUDIES ARE COMPLEX

Conducting clinical trials with cancer patients typically takes more time than conducting studies with healthy volunteers. The complexities of oncology studies increase the likelihood of study timelines getting off track, which leads to increased costs and lost time to market for sponsors.

## OUR OFFERING

If your study is off track, Medelis' oncology clinical operations team and members of our Medical Advisory Board can review and assess all of your study elements. We'll identify the key issues preventing your study from meeting your timelines, and develop a study rescue plan focused on resolving current bottlenecks and study challenges.

If you're unsure whether your CRO or clinical team is performing efficiently, we're able to analyze your study in confidence, so you can gain a fresh perspective about the real causes of your study delays.

Getting a study back on track requires an in-depth understanding of:

- The best investigators and sites for a specific protocol and indication
- How to recruit the right patients
- The necessary skills and oncology experience of the CRAs and project managers
- The current regulatory environment
- The most effective method for the collection, organization and submission of the study data

Our analysis provides you with a complete solution to regain lost time and momentum. In virtually every case, the Medelis team has been able to reverse almost all problems, and regain most of the lost time, with minimal additional expense to the client.



## VALUE

By having Medelis review your trial details and provide our recommendations to get your study back on track, you gain a fresh perspective on how to alleviate trial challenges.

Getting your trial back on track allows you to:

- Recover lost time
- Avoid the extra costs of an extended timeline
- Gain the data that you need, faster
- Reduce your time to market

## ABOUT MEDELIS:

As a single-source provider for clinical trial planning, trial management, and drug development planning, Medelis delivers the best total solution—and all the support required—for biotechnology and pharmaceutical companies to optimize their return on clinical research investments. And we accomplish that in the quickest, most responsive way possible.

## OUR EXPERIENCE:

Medelis' medical founders, team physicians and advisors are internationally-recognized oncology thought leaders. They organize and chair sessions at national and international oncology meetings and serve as officers and board members of professional oncology organizations, including The American Association for Cancer Research and The American Society of Clinical Oncology.

Our clinical operations team and medical personnel have critical first-hand experience with the complex issues often associated with oncology clinical trials, and they have handled most targets in oncology, which means that highly experienced resources will be working on your study.

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