

Clinical decision makers confirm shift to hybrid and decentralized studies.



73% of oncology executives plan to run a Hybrid or Fully Decentralized Clinical Trial in the next 12 months

up from **49%** from the previous 12 months

Fewer than **2/3** of respondents (65%) are planning a Traditional, Site-Based Clinical Trial in the next 12 months

down from **88%** from the previous 12 months

Trial Type	previous 12 months	next 12 months
Traditional Trial	88%	65%
Hybrid or Fully DCT	49%	73%

The three **biggest perceived challenges** concern speed...

Challenge	Percentage
Patient Recruitment	72%
Study Start-up	55%
Timeline / Delays	54%

Time is of the essence when executing an oncology clinical trial. Delays in getting new drugs to market can cost sponsors upwards of **\$600 million a day**.

TYPES OF CANCERS FOR DCT

Sponsors and CROs are planning to execute Hybrid Trials/DCTs in the next 12 months for a wide range of oncology indications, including:

Cancer Type	Percentage
Lung Cancer	40%
Leukemia / Blood Cancers	37%
Breast Cancer	30%

Plus, Colorectal, Lymphoma, Gastric, Pancreatic, Melanoma, Endometrial, Ovarian, Bladder, Liver, Thyroid, and Kidney and Renal Pelvis cancers.

Top three **perceived benefits** of using Hybrid / DCT tools in oncology trials:

Benefit	Percentage
Increased Patient Retention	67%
Greater Patient Diversity	54%
Faster Patient Recruitment	50%

SURVEY PARTICIPANTS

by Sector

Sector	Percentage
Pharmaceutical / Biotech	60%
CRO	39%
Medical Devices	1%

by Function

Function	Percentage
Clinical Operations	46%
R&D Management	20%
Corporate Management	12%
Other	18%
Outsourcing	5%