

Redefining value in drug development and clinical research

Working with one partner across services unlocks measurable value

A data-driven look at integrated drug development

The Tufts Center for the Study of Drug Development, a nonprofit research group affiliated with Tufts University School of Medicine, analyzed the financial and operational impact of aligning manufacturing, clinical research, laboratory, and supply chain services under a single provider.

To assess value, Tufts developed a model using real-world clinical trial data and industry benchmarks focused on projected outcomes in oncology programs across both monoclonal antibodies (mAbs) and small molecules.

Key inputs to the model



Clinical development costs and durations



Phase-specific success probabilities



Operational efficiencies in integrated trials



Projected peak sales and market exclusivity



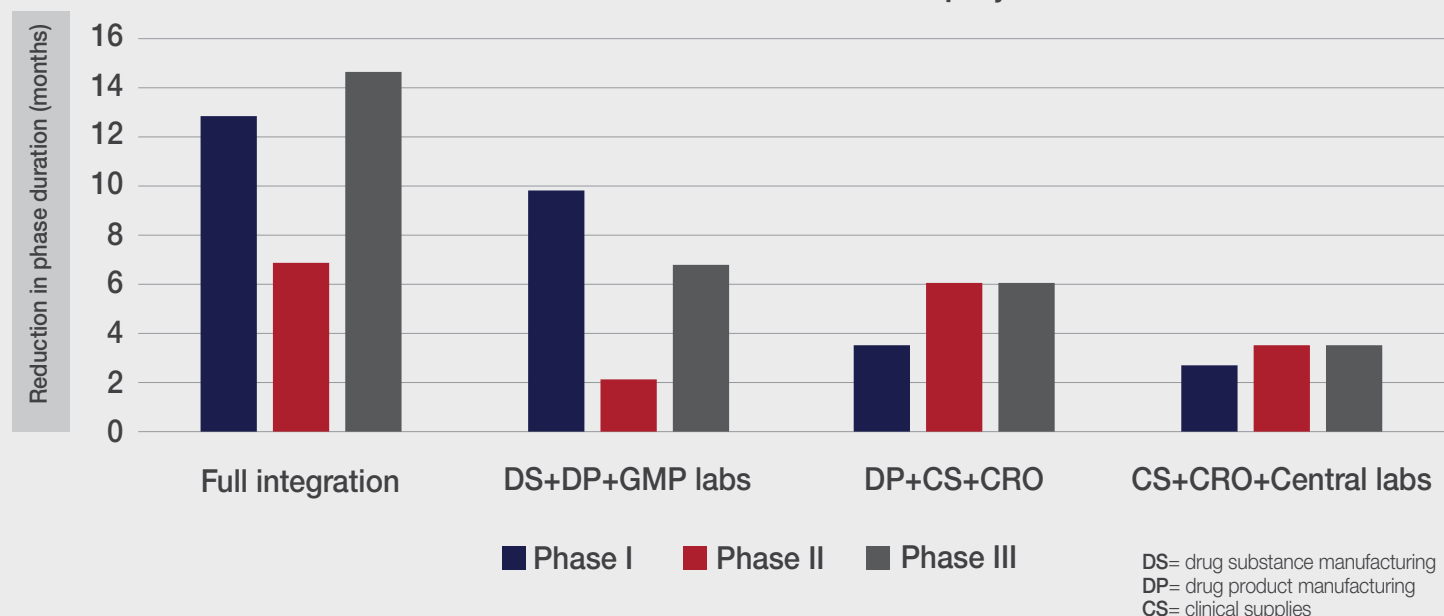
Performance metrics

- **Estimated time savings** based on streamlined execution
- **Expected net present value (eNPV)**, a forward-looking metric reflecting the projected financial return of a portfolio of programs, adjusted for risk and timing
- **Return on investment (ROI)**, the ratio of future returns versus the expected cost of development

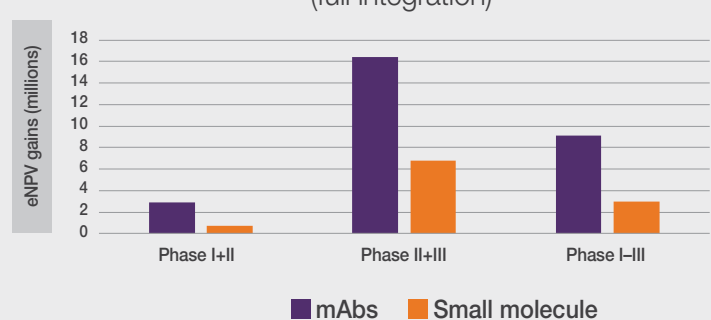
Impact across phases and modalities

The study found that integration consistently reduced drug development time and improved projected value and ROI across multiple phases and drug modalities, with various combinations of services.

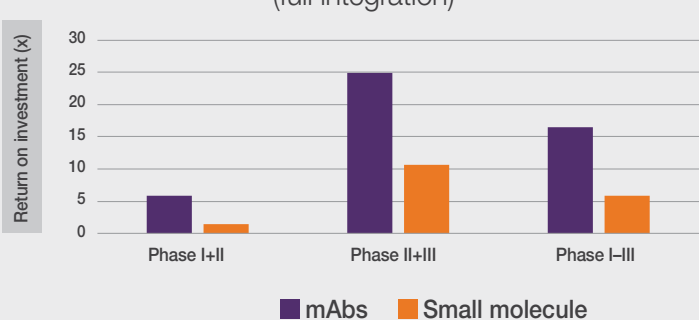
Reduction in drug development timeline using integrated CDMO and CRO services for mAb projects*



Comparison of eNPV gains across scenarios* (full integration)



Comparison of ROI across scenarios* (full integration)



*DiMasi, Joseph, Dirks, Abigail, Getz, Kenneth. "The Net Financial Benefits of Single Vendor Integrated CDMO and CRO Drug Development Services." (2025) Manuscript submitted for peer review and publication. Tufts Center for the Study of Drug Development, Tufts University.

Key findings

Up to
34

months saved through coordinated execution

Up to
\$16.4M

eNPV gains for Phase II + Phase III

Positive return

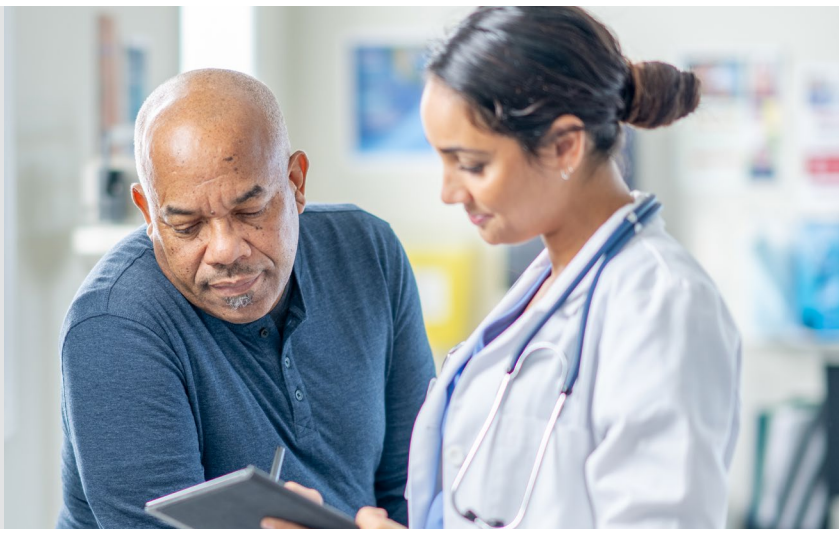
on investment in all phases

Benefits observed across both
mAbs and small molecules

Why CDMO + CRO integration matters

Coordinating key functions through one partner:

- ➔ Reduces handoffs and delays by aligning development functions
- ➔ Improves coordination and decision-making across teams
- ➔ Mitigates risk by proactively identifying roadblocks and enabling greater flexibility to adapt to evolving development needs



White paper

Value reimaged: Unlocking ROI and efficiency in drug development

Is integration the right move for your program?

Learn more about the impact it has across phases and modalities.

[Download the white paper](#)

Learn more about **Accelerator™ Drug Development**

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