

Plan

Monitor

De-risk

HIT ID

HIT TO LEAD

LEAD OPTIMIZATION

ID CANDIDATE

PRECLINICAL

IND

PHASE I

PHASE II

Operational Best Practices with Proven R.O.I.

Resource

Running an early stage biotech startup is essentially managing capital and optimizing time and resource investment.

Scientific acumen and operational rigor are two equally critical pillars to keep the company moving. Scientific acumen tells you to invest in the right designs and candidates. Operational rigor makes you execute with precision, reduce resource waste and maximally extend your runway.

For founders who are scientists, the most effective approach is to combine your scientific intelligence with proven operational practices.



Learn about the Kaleidoscope platform
<https://kaleidoscope.bio>

Plan

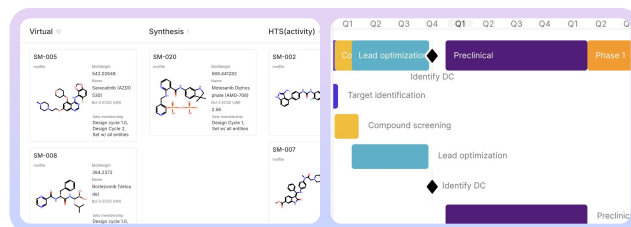
Plan all scientific work with clear expectations, and ensure actionable data and optimized decision making.

Best Practices

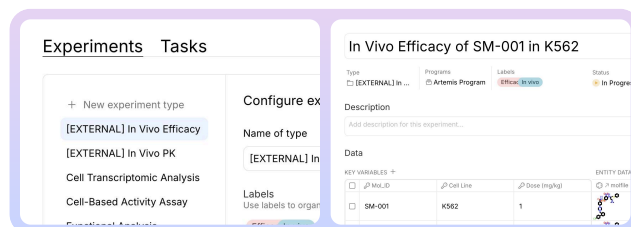
- 1 Document and communicate each stage of your R&D effort clearly with timelines, spending and ownership.
- 2 Plan both CRO and in-house work with clear criteria for completion.
- 3 Forecast experimental work by types and define input analytes and conditions.
- 4 Set clear expectation for data outputs from all R&D efforts (in-house and CRO).
- 5 Ensure all results (data) are driving decisions with clear criteria. Every piece of data need to serve the purpose of advancing or pivoting your program.
- 6 Optimize your decision making and minimize analysis paralysis, by focusing on the most critical data evidence among your candidates.

Software Needs

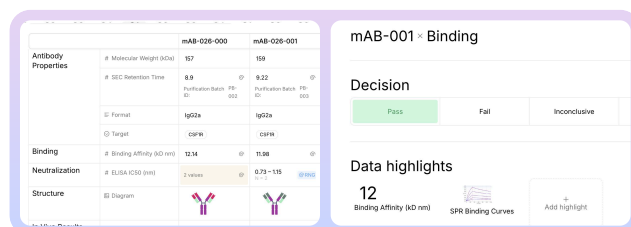
Captures plan details and renders plan visualization by time, type and other properties



Documents attributes of all experiment types, and consistently generates data placeholders



Surfaces data to directly enable decision making and program advancement



Monitor

Track your spending (time and capital) against in-house and CRO results; adjust/pivot sooner than later

Best Practices

1



Ensure all in-house and CRO results (data) are directly viewable from your project and experiment planner.

2

Promptly review data and make critical decisions upon experiment results becoming available.

Software Needs

Enables data review directly in application without need for copy/paste via slides or docs

	Binding	In Vivo PK	Cell L Efficacy	mAB-028-000	mAB-028-001	mAB-028-002
mAB-001	Pass	Pass	Pass	# Molecular Weight (kDa) 157	159	147
mAB-005	Pass	Pass	In Prog	# SEC Retention Time 8.9	9.22	8.5
mAB-006	Pass	Pass	In Prog	Purification Batch ID 1001	Purification Batch ID 1003	Purification Batch ID 1005
mAB-010	Pass	Inconclusive		E Format IgG2a	IgG2a	IgG2a
mAB-015	Pass	Fail		G Target CSF1R	CSF1R	CSF1R
mAB-020	Pass	Fail		# Binding Affinity (nM) 1214	1198	1757
				# SLISA IC50 (nM) 2 values	0.73 - 115	0.63
				E Diagram		

Case studies



How: Plan work, review data and drive decisions in Kaleidoscope without data copying or slide preparation

Therapeutic target: immunology and inflammation

ROI: > 1 day per week saved and reinvested



Ewan Taylor
CSO, Co-founder



How: Monitor and optimize compound progress from design to synthesis to decision, via improved in-house and CRO workflows

Therapeutic target: oncology

ROI: Meet annual milestone 2 months sooner



Shivam Patel
Head of Data Science



How: Streamline data-to-decision via structured templates and systems auto-sync

Therapeutic target: immunology and inflammation

ROI: Increased confidence and days saved in decision making



Geraldine Paulus
Co-founder

De-risk

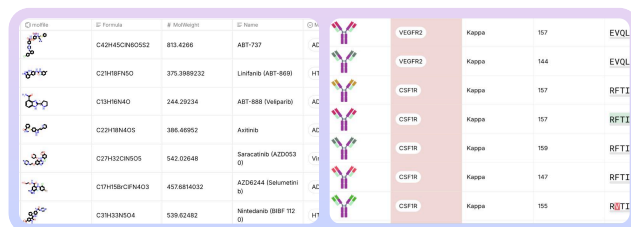
Prevent mislabeling and loss of critical data, unnecessary experiment redo's, and downstream costs of switching systems/tools

Best Practices

- 1 Ensure all designs and modalities are documented with properties in structured tables.
- 2 Enable consistent viewing and actioning via filter views and intuitive searches.
- 3 Document your experimental results as structured data linked to your designs and candidates.
- 4 Minimize use of spreadsheet attachments, or multiple forms of data-to-design association.
- 5 Start early on a future-proof system to build a culture of structured data management.

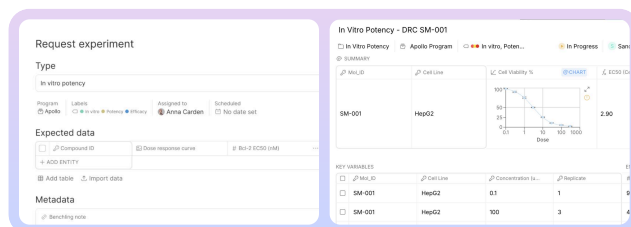
Software Needs

Structurally handles all modalities and their properties with flexibility allowing for innovation



Formula	MultiTarget	Name	Kappa	EVOL
C42H45ClN6O5S2	873.4266	ABT-727	157	EVOL
C21H28FN5O	375.389232	Linfenib (ABT-868)	144	EVOL
C13H18N4O	244.29234	ABT-888 (Velparib)	157	RFTI
C22H18N4O5	386.48952	Avanib	157	RFTI
C27H32ClN6O5	542.52848	Saracatinib (AZD0536)	159	RFTI
C17H18ClFN4O3	457.6814032	AZD6244 (Sunitinib)	147	RFTI
C19H23N5O4	539.62482	Nintedanib (BIBF 112)	155	RFTI

Allows scientists to consistently log results and link them to designs and program stages



Additional resources

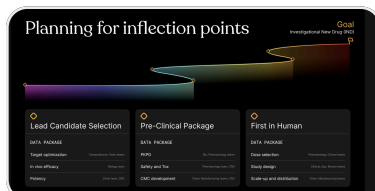
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Planning



Portfolio management in early stage drug discovery – a traveler's guide through uncharted territory ↗

"Management in discovery has to deal with projects covering multiple compounds, sometimes hundreds or even thousands each, that are synthesized, profiled and further optimized"



Plan smartly, execute precisely: the high-stakes game of chasing biotech value inflections ↗

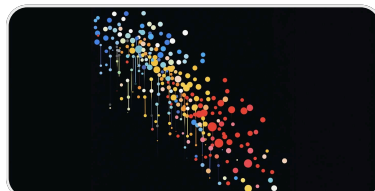
"By knowing the ultimate goal, biotechs are able to understand the data necessary ... and avoid costly pitfalls and ensure they're laser-focused on the right activities rather than chasing irrelevant data or duplicating efforts unnecessarily"

Monitoring



How to improve R&D productivity: the pharmaceutical industry's grand challenge ↗

"resolution of technical uncertainty early in development is necessary to improve R&D productivity. We refer to this as our 'quick win, fast fail' paradigm of drug development".



The 'too early to track' fallacy ↗

"Begin by defining some essentials that you want to track early on, and then add to that list as you scale. This approach allows you to take advantage of the structure you've put in place early on, while also building the muscle and encouraging the behavior of documenting things from the start."

De-risking



Achieving end-to-end success in the clinic: Pfizer's learnings on R&D productivity ↗

"...greater adoption of enhanced, objective, and quantitative methods, including elevating the importance of key scientific quality metrics, has enabled Pfizer to rapidly progress strong programs while stopping weaker programs earlier."



Ending the era of data-blind project managers ↗

"...teams using platforms like Benchling or CDD Vault end up doing double work, manually tracking in their project management systems what's already happening in their ELN. This introduces massive inefficiencies, increases risks of errors, and creates gaps where insights slip through."