Palantir Foundry – Impact Study	Turbocharging re the NIH	esearch at	Q HEALTHCARE
CHALLENGE	The NIH wanted to enable precision medicine by better understanding how genetic and other factors impact drug efficacy. But the scale and intricacy of data from High-Throughput Screening (HTS) robots, genome sequencers, mass spectrometers, and other instruments made it extremely challenging. Scientists had to painstakingly harmonize data from many sources, and informaticians had to do substantial pre-processing work on the data. Existing tools were originally developed with informaticians in mind, and their complexity prevented researchers from accessing, analyzing, and publishing their findings in a timely way.		
SOLUTION	The NIH uses Foundry to integrate and harmonize scientific data from dozens of internal and external sources. Processing, normalizing, and analyzing this data in Foundry allows new modes of informatician / biologist collaboration via:		
	Direct connection to online and offline experiments	Data accessibility for researchers	Traceability and privacy protections
	Online and offline experiments	Researchers independently	Data provenance is fully

IMPACT

- NIH researchers have uncovered insights in hours that used to require months of data gathering and manipulation.
- One lab predicted novel drug combinations for use in an oncology setting, which were then experimentally validated in vitro. In vivo follow up is currently underway for several of these combinations. If successful, this will lead to a clinical trial.
- A different lab validated a gene signature for drug response found in internal experiments. They compared observations from public genomic data with the findings from clinical data, increased their confidence in the initial finding, and defined precise experimental follow-ups.

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