Step into newborn sequencing.

Powering the next wave of genomic research.

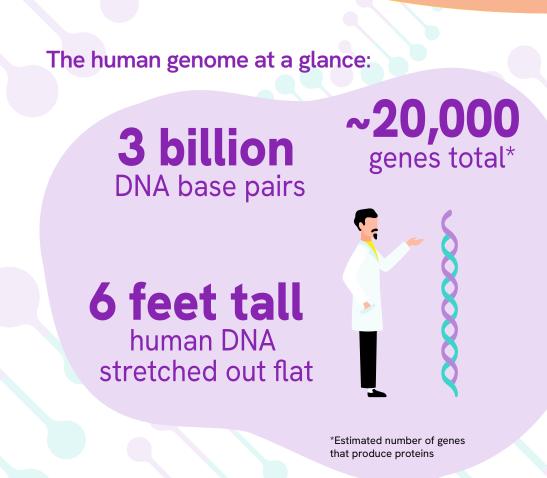


Newborn research stands at the dawn of a whole new era. To date, standard research efforts have documented hundreds of disorders in the context of newborn health. However, less than 80 of them are actively investigated. Now, with the advent of sequencing technology, the limitations have been lifted.

Bringing simplicity to sequencing, Revvity's newborn sequencing workflow features a robust gene panel curated by genomic experts, powering the next frontier of research.

Unleash your laboratory's potential. Explore Revvity's newborn sequencing research worflow today.

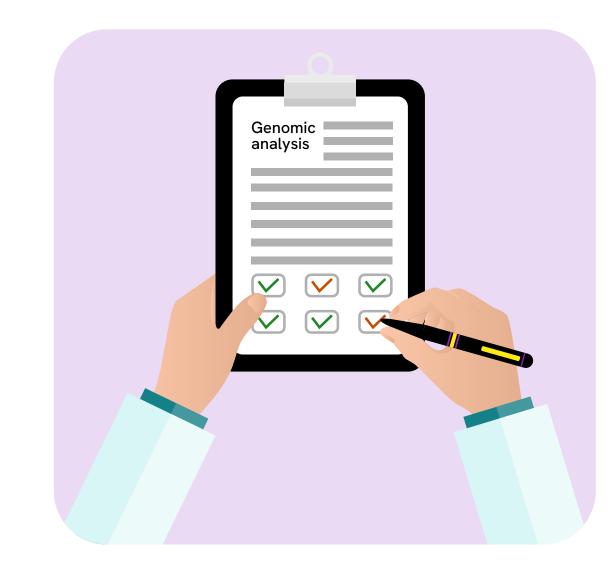
Learn more



# Newborn sequencing with Revvity

## A solution for every step





From dried blood spot card to final data analysis.

## Revvity's lab workflow



End-to-end solution

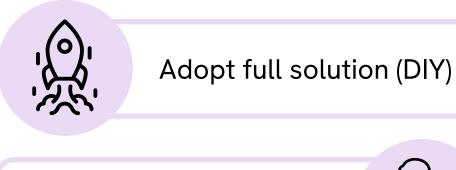
Configurable workflow

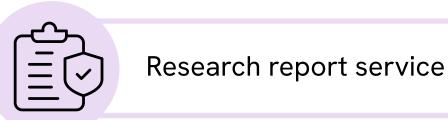




Hands-on NGS expertise

# Multiple starting points





Data generation service

End-to-end service



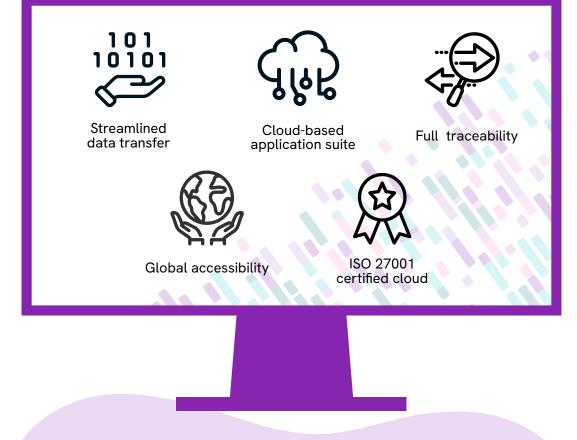
## Complete software suite



- Analysis Interpret

# for the lab

Software designed

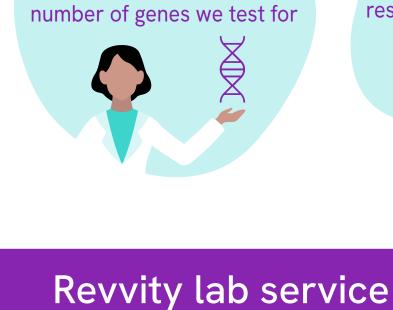


## Software engineered by laboratory specialists Compatible with diverse laboratory setups

**Key specifications** 

- Capable of analyzing dried blood spot samples, whole blood, buccal swaps and saliva

Analysis and interpretation



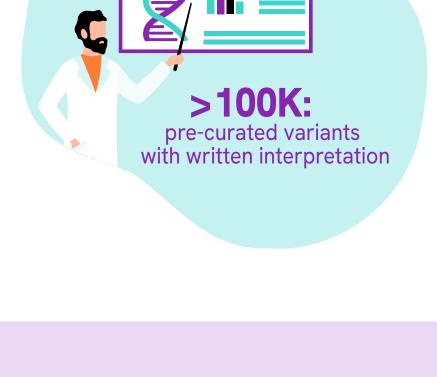
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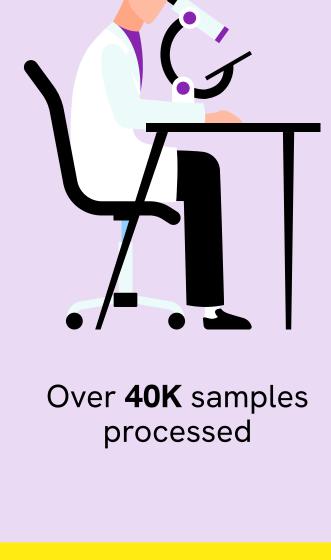


research areas



Carefully curated







Why Revvity?









Learn more

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See how your lab can step into sequencing.