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 and featuring a panel of 390+ genes, Revvity's newborn sequencing workflow is powering the next frontier of research.

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

The human genome at a glance:

3 billion DNA base pairs

~20,000 genes total*

6 feet tall human DNA stretched out flat

*Estimated number of genes that produce proteins

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
- Customizable workflow
- Hands-on NGS expertise

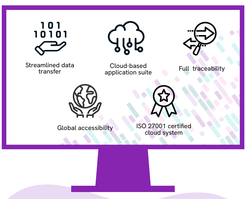
Multiple starting points

- Adopt full solution (DIY)
- End-to-end service
- Research report service
- Data generation service

Complete software suite

- LIMS
- Analysis
- Interpret

Software designed for the lab



Key specifications

- Software engineered by laboratory specialists
- Compatible with diverse laboratory setups
- Runs on dried blood spot samples, whole blood, buccal swaps and saliva

Analysis and interpretation

>390: number of genes we test for

17 research areas

Carefully curated panel

>87K : pre-curated variants with written interpretation

Revvity lab service

Over **40K** samples processed

Validated by **CLIA, CAP, NY State, ISO15189** standards

5 Revvity NGS labs globally

Why Revvity?

> 40 years experience, the leaders in newborn screening

> 75 years providing dynamic research lab solutions

190: number of countries empowering academic & translational research

See how your lab can step into sequencing.

Learn more

For research use only. Not for use in diagnostic procedures.

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