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:

~20,000 genes total* **3** billion DNA base pairs 6 feet tall human DNA stretched out flat



*Estimated number of genes that produce proteins

Learn more

Newborn sequencing with Revvity

A solution for every step

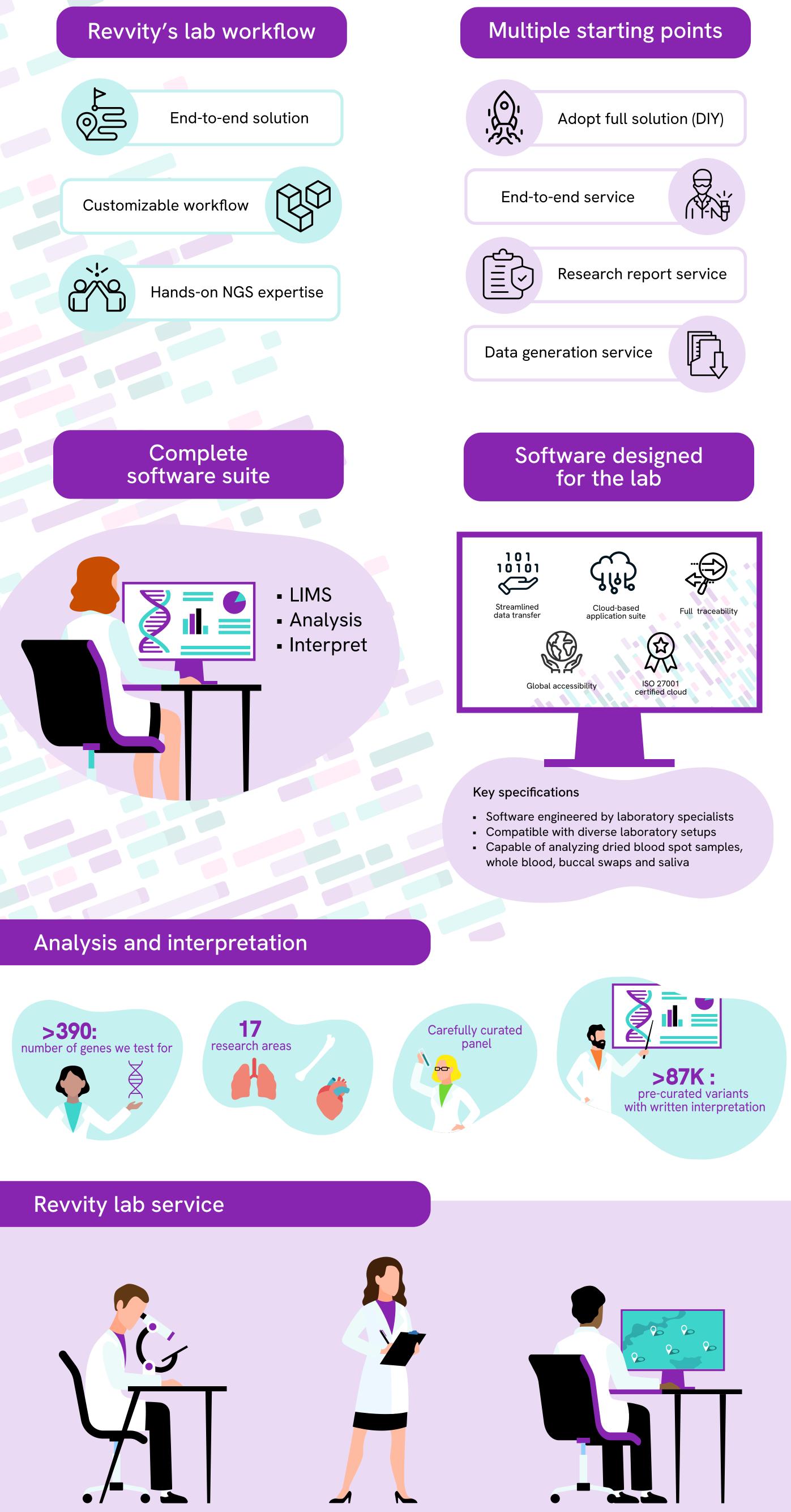
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From dried blood spot card to final data analysis.





Over 40K samples processed



Validated in accordance with 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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