my Newborn

myNewborn accurately detects more than 390 diseases that can affect newborns in the first years of life.

Diseases are classified into different groups including:

- **Metabolic**
  - E.g. Familial Hypercholesterolemia
- **Endocrine**
  - E.g. Neonatal diabetes mellitus
- **Autoimmune**
  - E.g. X-linked immunodeficiency
- **Deafness**
  - E.g. Usher syndrome
- **Neurological**
  - E.g. Malignant hyperthermia
- **Other**
  - E.g. Cystic fibrosis

How to start?

- The physician prescribes the test.
- Veritas provides a kit to collect the sample.
- Sequencing and interpretation is performed in Veritas accredited laboratory.
- The report is discussed with the parents to stablish next steps according to the results.

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About Veritas

Veritas Genetics, a *LetsGetChecked* company, is one of the world leaders in advanced genetic sequencing and clinical interpretation of the exome and whole genome, driving the transition to personalized and preventive medicine

Using state-of-the-art technologies and the highest safety standards, Veritas Genetics helps individuals, healthcare professionals and institutions worldwide to understand and anticipate genetic risks, enabling more informed and proactive health decisions.

With a focus on innovation and accessibility, Veritas Genetics transforms the way we understand and care for health at every stage of life.

Veritas
a LetsGetChecked company

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my Newborn

Neonatal genomic screening that studies childhood onset actionable diseases

myNewborn provides an early screening for genetic diseases that appear in the first stages of life by giving information for early treatment, preventive or dietary management.

What is myNewborn?

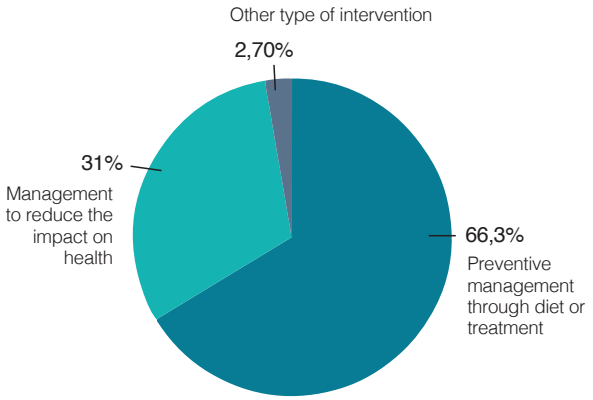
myNewborn is a neonatal screening test that analyses the hild's DNA to determine if any variants related to actionable childhood-onset diseases are present.

Knowing this information makes it possible to adapt medical management to prevent the disease or reduce its impact on the child's health.

What will you get?

- **Analysis of more than 400 genes associated with 390 metabolic and genetic diseases** that appear in the first years of life.
- **The test complements the heel prick test,** improving the clinical utility.
- **In most cases the result is negative which provides tranquility,** while in case of a positive result the information is key for an early management.
- **Most diseases are conveniently treatable** through diet or early intervention

Classification of the pathologies by clinical management



Beyond the heel prick test

Once the child is born, the heel prick test is routinely performed on the newborn. This test is necessary but is limited to a small number of diseases, **myNewborn** allows to expand the diseases studied to offer the most comprehensive neonatal screening.

SIMPLE

Requires a routine blood sample, cord blood or saliva collected with a kit provided by Veritas.

ACCURATE

Analyses the newborn's DNA using advanced sequencing technology which provides reliable results.

PREVENTIVE

myNewborn is for any baby without disease symptoms as part of the medical care.

REALIABLE

myNewborn is based on BabySeq project, which is aimed at to improve medical care for newborns through genomics.

Up to 9.4% of newborns can present genetic variants related to childhood-onset diseases.

Knowing this risk is key to intervene and personalize the newborn's management.

Get the most complete genetic service in the market

veritasint.com