



Improving newborn screening

By participating in **Baby Detect**:

- You allow your child to benefit from a more extensive screening, free of charge
- You participate in the progress in the fight against serious diseases

A few drops
of blood to
diagnose
in time!





Baby Detect is...

What?

Research that aims to demonstrate the possibility of screening at birth for more than 120 serious and treatable childhood illnesses.

Why?

This test makes it possible to identify more than 120 genetic diseases before the symptoms appear in order to treat them before the disease acts.

For whom?

For all newborns after parental consent.

Where?

At the maternity ward, 2 days after birth.

How?

As part of the official newborn screening program, a few drops of blood are taken to screen for 19 serious and treatable childhood illnesses. This is called the Guthrie test. With Baby Detect, we take a few extra drops to screen for over 120 diseases. If you do not hear back from us, it means that no disease included in the program has been detected. In this case, “no news is good news”.

If an anomaly is detected, you will then be contacted as soon as possible by a specialist doctor.

This causes no problems and no additional discomfort for your baby.

For more information:

- Scan the QR-code



- Visit www.babydetect.com

- Speak to your gynaecologist, paediatrician or midwife.

