INDIANA GENOMICS & NEWBORN SCREENING What is Newborn Screening?





Every baby born in Indiana is required by state law to complete newborn screening. Babies with disorders may look and act like healthy newborns, but may have a medical condition that could cause serious illness, developmental delay, disability or even death.



Newborn screening is a set of 3 screens that identifies over 50 disorders. A simple blood screen called a heel stick, a pulse oximetry screen for critical congenital heart defects (CCHDs) and a hearing screen for hearing loss will be completed.



There is treatment available for the disorders screened and most babies who are identified early can grow up to be healthy.



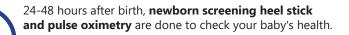
To see a complete listing of all the conditions included on the Indiana newborn screening panel, visit the Indiana Genomics and Newborn Screening Program website.

Genomics and Newborn Screening Program Indiana Department of Health www.NBS.in.gov phone 888.815.0006 Early Hearing Detection & Intervention Program (EHDI) Indiana Department of Health www.Hearing.in.gov phone 317.232.0972





Newborn Screening, Step-by-step



CONGRATULATIONS, BABY IS BORN!

Each spot is smaller than the size of a dime.

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At the NBS lab, special equipment screens the blood spots. Your baby's blood is tested for over 50 disorders.

Your health care provider will give you the results when they are available.

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A **hearing screening** will be completed before your baby goes home.

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A few drops of blood are taken from baby's heel and placed on a newborn screening card with baby's pulse oximtery and hearing screen results and sent to the NBS Lab.

NORMAL RESULTS

If everything looks ok, the results are sent to your baby's health care provider. *Ask for your baby's results at the first newborn visit!*

ABNORMAL RESULTS

If there might be a problem, the lab will call your baby's health care provider. Screening can only tell if your baby **MIGHT** have a health problem; we won't know for sure until your baby has more testing. Your baby's health care provider will talk to you about what needs to happen next.

What happens to blood spot cards?

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After screening, the dried blood spots are stored for 6 months and then destroyed if opted out of storage. If you chose to store the blood spots, they will be stored for 3 years and then destroyed.



Families may benefit from having their baby's blood spot card stored. If a child gets sick, the blood spot can give an idea if something at birth made the child sick. A blood spot may also be used to identify a missing or deceased child, paternity, or if the parents request information for future pregnancies.



To make sure the screenings and equipment work the way they should, quality control may be done at the laboratory with the blood spots in storage.

By law, newborn screening is required; however, you may refuse the screening by completing a religious refusal form, which can be found on the Genomics and Newborn Screening Program website.



Stored blood spots can also be used for research. The blood spots are de-identified, meaning the baby's name, date of birth, and any other identifying information is removed.



No research will be done on your baby's blood spot without your written consent. If you would like to change your baby's blood spot storage or have other requests, please complete the appropriate form that can be found on the Genomics and Newborn Screening Program website or ask your health care provider for more details.

A keepsake for you & your baby
Baby's Name
Baby's Due Date Date of Birth
Timea.m./p.m. Weight Length
Mother's Health Care Provider Image: Constraint of the second s
Heel Stick (blood test) Date
 □ Pulse Oximetry for CCHD □ Pass (Right hand and foot over 95%) □ Echocardiogram
Hearing Screen for hearing loss Ask for your baby's newborn Passed both ears screening results at your first Referred for further testing newborn visit with your health Date care provider!
Time
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