The Texas Newborn Screening Program

For Your Baby's Health

The Texas Newborn Screening Program saves and improves hundreds of lives each year through early detection of genetic disorders that leads to timely intervention. The program is expanding to screen for more genetic disorders than ever, which means better health outcomes for Texas newborns.

TEXAS NEWBORN SCREENING

Newborn Screening Disorders

Organic Acid Disorders

PROP, MUT, IVA, 3MCC, HMG, MCD, GA1, Cbl A, B, C, D ßKT, MAL, 2MBG, 3MGA, 2M3HBA, and IBG

Fatty Acid Disorders

MCAD, SCAD, LCHAD, VLCAD, MSCHAD, CACT, CUD, TFP, GA2, MCAT, DE RED, CPT I and II

Amino Acid Disorders

ASA, MSUD, HCY, PKU, ARG, MET, H-PHE, TYR I, II, and III, BIOPT-BS, BIOPT-REG, CIT I and II

Endocrine Disorders

CH and CAH

Hemoglobin Disorders

Hb S/ß TH, Hb SS, and Hb S/C Various Hemoglobinopathies

And Other Disorders

SCID, CF, BIOT, GALT, CCHD, Hearing Loss and T-cell related lymphocyte deficiencies

To view full list and descriptions, visit

https://www.dshs.state.tx.us/newborn/screened_disorders.shtm

TEXAS Department of State Health Services