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.
Organic Acid Disorders
PROP, MUT, IVA, 3MCC, HMG, MCD, GAL, 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/B 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