

Newborn Screening ACT Sheet

[Absent/Reduced Galactose-1-Phosphate Uridyltransferase (GALT)] Classic Galactosemia

Differential Diagnosis: Duarte variant galactosemia.

Condition Description: Galactosemia refers to a group of disorders which are caused by an inability to metabolize galactose, a sugar found in lactose. Classic galactosemia results from an inherited deficiency of the galactose-1-phosphate uridyltransferase (GALT) enzyme, causing elevated galactose and galactose-1-phosphate. If treatment is not initiated early, life threatening complications can occur. The Duarte variant refers to a diminished ability to break down galactose in childhood.

You Should Take the Following *IMMEDIATE* Actions:

- Inform family of the newborn screening result
 - Ascertain clinical status (poor feeding, vomiting, lethargy, jaundice). Discontinue breast feeding and/or cow's milk formulas and initiate non-lactose based feedings with a soy formula.
 - Consult with pediatric metabolic specialist the same day.
 - Evaluate the newborn (jaundice, poor feeding, vomiting, lethargy, bulging fontanel, and bleeding). If any of these findings are present or if the newborn is ill, transfer to a hospital for further treatment in consultation with the metabolic specialist.
 - Initiate confirmatory/diagnostic testing and management, as recommended by specialist.
 - Provide family with basic information about classic galactosemia, including dietary management.
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- Report final diagnostic outcome to newborn screening program.

Diagnostic Evaluation: **Red blood cell GALT activity:** Complete or near-complete deficiency in classic galactosemia; partial reduction of normal activity with the Duarte variant. **Red blood cell galactose-1-phosphate (gal-1-P):** Elevated in both classic and the Duarte variant in patients consuming lactose. Red blood cell assays are not valid following transfusion. **Molecular genetic testing** may be required to confirm the diagnosis.

Clinical Considerations: Classic galactosemia presents in the first few days of life and may be fatal without treatment. Signs include poor feeding, vomiting, jaundice and may include lethargy and/or bleeding. Neonatal *E. coli* sepsis can occur and is often fatal. The treatment is the avoidance of dairy products and other foods containing galactose, and the administration of soy-based formulas. Symptomatic neonates will require emergency supportive measures. Considered a benign condition, there is no standard accepted management for the Duarte variant; some practitioners restrict high galactose foods in early childhood.

Additional Information:

[How to Communicate Newborn Screening Results](#)

[Gene Reviews](#)

[Medline Plus](#)

[Condition Information for Families- HRSA Newborn Screening Clearinghouse](#)

Referral (local, state, regional, and national):

[Find a Genetics Clinic Directory](#)

[Genetic Testing Registry](#)

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State and Other Resources

State Newborn Screening Program

Metabolic (Bloodspot) Screening, Montana's Newborn Screening Programs
406-444-0984, dphhs.mt.gov/ecfsd/cshs/newbornscreeningprograms/

Short-term Follow-up Coordinator - Crystal Fortune, MT Laboratory Services Bureau
406-444-0930 or 800-821-7284 cfortune@mt.gov

Genetics/Metabolics Consultants

Shodair Children's Hospital Medical Genetics Staff will contact you with recommendations.
Metabolic Clinic: 406-444-1099; Clinics held across Montana: 406-444-1016
shodair.org/about-shodair/contact/

If emergency consultation with a metabolic specialist is necessary, contact One Call (24/7) at Children's Hospital Colorado (800-525-4871) and ask to speak to the metabolic specialist on call.

Information for Clinicians and Families

Montana Medical Home Portal (see also the Parents & Families section)
mt.medicalhomeportal.org/newborn/galactosemia

Parent/Family Support

Galactosemia Foundation
www.galactosemia.org

National Resources (with web addresses)

Additional Information

How to Communicate Newborn Screening Results
www.hrsa.gov/sites/default/files/hrsa/advisory-committees/heritable-disorders/Resources/achdnc-communication-guide-newborn.pdf

Gene Reviews
www.ncbi.nlm.nih.gov/books/NBK1518/

Medline Plus
medlineplus.gov/genetics/condition/galactosemia/

Condition Information for Families-HRSA Newborn Screening Clearinghouse
newbornscreening.hrsa.gov/conditions/classic-galactosemia

Referral (local, state, regional and national)

Find a Genetics Clinic Directory
clinics.acmg.net

Genetic Testing Registry
www.ncbi.nlm.nih.gov/gtr/