- ► Environmental Health
- ▶ Food Safety
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- Informatics
- Laboratory Systems and Standards
- Newborn Screening and Genetics

Assuring Laboratory Quality

NBS Molecular Resources
Policy and Positions

- Public Health
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 Response
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Assuring Laboratory Quality

Analyte Interference List

SAVED TOPICS: Newborn Screening, NSQAP, Quality Assurance



The Newborn Screening (NBS) Analyte Interference list is a compilation of published and observed analyte interferences of NBS assays. This list can be used as a reference for quality control issues encountered in a NBS system. It is meant to be an interactive list, meaning users are able to add to it as appropriate, and a one-stop site for all known NBS assay interferences.

This page is currently in beta version. Please contact Ruhiyyih Degeberg, ruhiyyih.degeberg@aphl.org, to provide feedback.

Select Interference Category:

- . Infant Conditions
- Infant Treatment
- Maternal Conditions
- · Special Diets
- Specimen Treatment

Nominate Your Program's NBS Interference

If you have observed an NBS analyte interference in your laboratory, or if you know of a published one that you do not see represented on this list, please fill out a form to nominate it for inclusion to this public resource.

CONTACT

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ruhiyyih.degeberg@aphl.org

 Public Health Preparedness and Response

► Research

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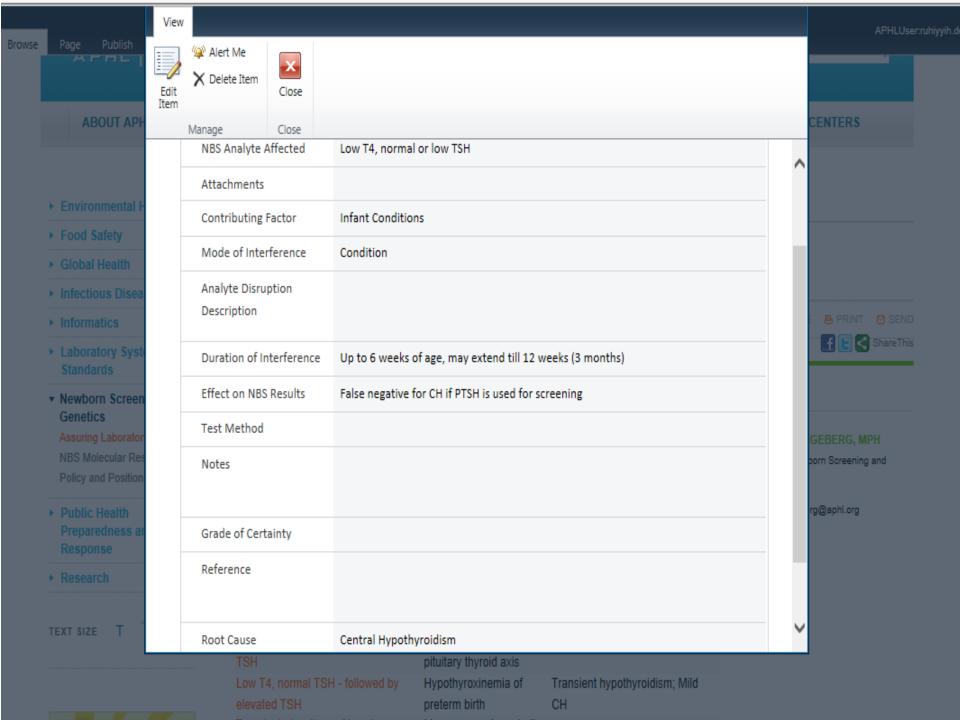


Click on analyte of interest ____

Infant Conditions

NBS ANALYTE AFFECTED	ROOT CAUSE	EFFECT ON NBS RESULTS
Low T4, normal TSH, delayed rise of	Immature hypothalamic-	False negative for CH
TSH	pituitary thyroid axis	
Low T4, normal TSH - followed by	Hypothyroxinemia of	Transient hypothyroidism; Mild
elevated TSH	preterm birth	CH
Transient elevations of tyrosine,	Liver enzyme immaturity	
methionine, and galactose,		
occasionally other amino acids		
Low T4, elevated TSH	lodine deficiency	Transient hypothyroidism
Low T4, elevated TSH, elevated	Acute illness	Transient hypothyroidism
immunoreactive trypsinogen (IRT)		
Elevated IRT	Hypoxia	False positive for CF primary screen
Elevated tyrosine, methionine,	Liver disease	
galactose, depression of		
biotinidase enzyme activity		
Elevated 17-OHP (?), amino acids,	Renal immaturity; renal	
organic acids	disease	
Lower biotinidase activity levels	Preterm	
inversely related to gestational age		
Acid α-glucosidase (GAA) (Pompe Disease) and other LSD's	High hematocrit	False negative
Varies (most if not all analytes,	Baby screened too early	False negative and false
notably 17-OHP, TSH, IRT)	after birth	positive results
Lower biotinidase activity levels	Preterm	
inversely related to gestational age		
Low T4, normal TSH and FT4	TBG (thyroxine-binding globulin) deficiency	False positive for CH
Low T4, normal or low TSH	Central Hypothyroidism	False negative for CH if PTSH i used for screening
Elevated C3	Hyperbilirubinemia	False positive MMA/PA
Transient elevations of tyrosine	Vitamin C deficiency	Transient neonatal tyrosinemia
Butyrylcarnitine (C4)	Mild form of glutamate formiminotransferase deficiency	False positive for SCAD or IBD deficiency
GALT activity	G6PD Deficiency	False positive for Galactosemi
DNA, PCR	Elevated hematocrit?	Sample failure for CF, SCID an
		other molecular methods
DNA, PCR	Immunoglobulin G in	Sample failure for CF, SCID an
	plasma	other molecular methods

- · Maternal Conditions
- · Special Diets
- · Specimen Treatment



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► Research	
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Partner	0
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Maternal Conditions

NBS ANALYTE AFFECTED	ROOT CAUSE	EFFECT ON NBS RESULTS
Low T4, high TSH	Hyperthyroidism treated with PTU and Methimazole	Transient hypothyroidism
None	Radioactive iodine treatment during pregnancy before 8 weeks gestation	Hypothyroidism
Low T4, high TSH	Radioactive iodine treatement during pregnancy after 8 weeks gestation	
Low or normal 17-OHP	Steroids: prednisone, betamethasone/dexamethasone	False negative for CAH
Elevated 17-OHP	Maternal Congenital Adrenal Hyperplasia	False positive result
Elevated phenylalanine	Maternal PKU or moderate hyperphenylalaninemia (uncontrolled)	Transient hyperphenylalaninemia
Elevated C50H, low free carnitine	Maternal 3-MCC deficiency	False positive 3-MCC
Elevated even-chain acylcarnitines	Fatty liver of pregnancy or HELLP syndrome	False positive, associated with LCHAD
Low carnitine levels	Maternal carnitine deficiency	False positive for CUD
Elevated propionylcarnitine (C3)	Maternal vitamin B12 deficiency	False positive PA/MUT/Cbl / B/Cbl D, E
Low free carnitine, elevated C5DC	Maternal GA-1	False positive for CUD
	Hypothyroidism treated with carbamazepine	
Elevated TSH	Autoimmune (Hashimoto's) thyroiditis	Mild CH or Transient hypothyroidism

- Special Diets
- Specimen Treatment

NBS ANALYTE AFFECTED	ROOT CAUSE	EFFECT ON NBS RESULTS
17-OHP	Blood collection with EDTA	False positive for CAH
TSH, T4	Blood collection with EDTA	False negative for CH
Immunoreactive trypsinogen (IRT)	Blood collection with EDTA	False negative for CF by IRT
Malonylcarnitine (C3DC)	Use of Sani-Wipes	False positive for C3DC
Phenylalanine (Phe)	Benzocaine skin treatment prior to collection	False elevation of phenylalanine, false positive for PKU
DNA, low TREC	Blood collection with heparin	Sample failure for CF, SCID and other molecular methods
Prevent extraction of analytes/degrade protein markers	Heat and/or Humidity	False-positives and false-negatives for many disorders (see Analyte Disruption Description field)
Isovalerylcarnitine (C5)	Nipple-fissure cream containing neopentanoate esters	Elevated C5

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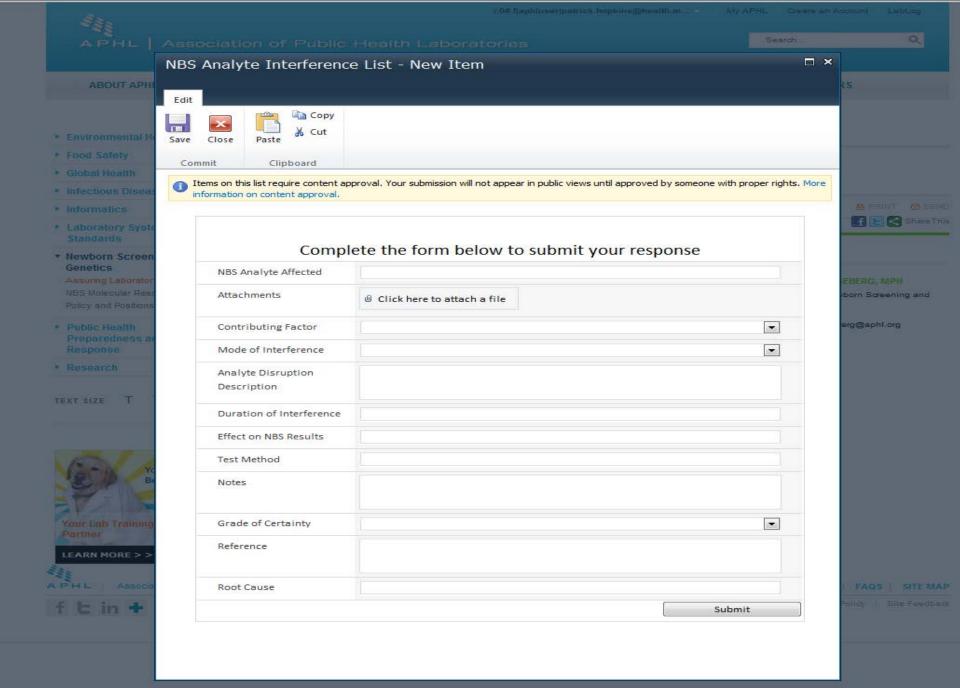
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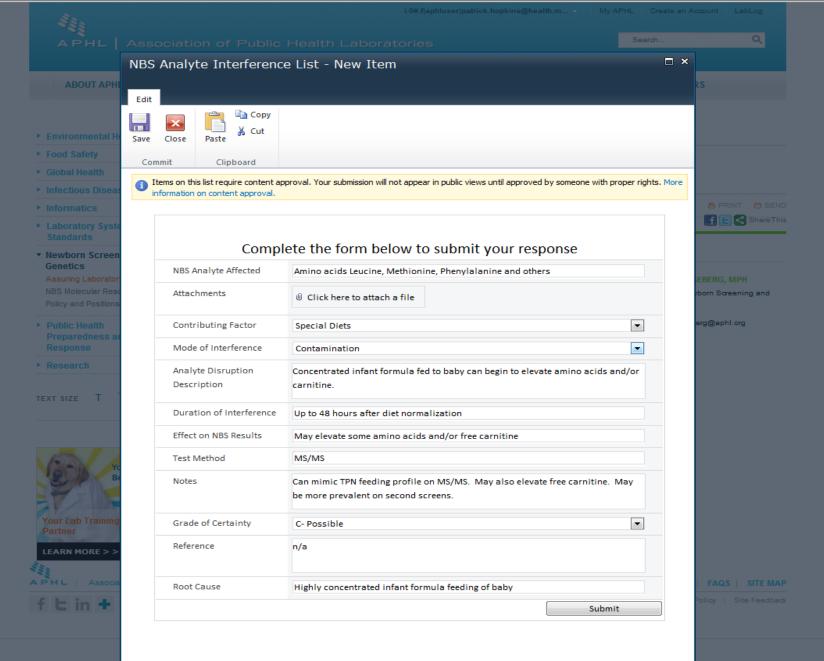
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Degrees of Certainty Assigned by QA/QC Subcommittee

Α	Definite	Well documented – multiple references
В	Probable	Documented – single reference
С	Possible	Not documented, but reported by personal communications or expert opinion
D	Not Likely	Interference is not reproducible or consistent
E	Unknown	No evidence to support or refute interference

NBS Analyte Inteference Site

 http://www.aphl.org/aphlprograms/newbornscreening-and-genetics/qualityassurance/Pages/Analyte-Interference-List

The APHL QA/QC Subcommittee

