

Study ID: \_\_\_\_\_  
 Date of Visit: \_\_\_\_\_



**ChiLDReNLink: BASIC**

**Form 29 Change in Diagnosis**

Please identify the subject's primary diagnosis (choose only one):

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- Biliary atresia
- Idiopathic neonatal hepatitis
- Hepatitis due to CMV (CMV inclusion disease on liver biopsy or systematic congenital CMV infection)
- Hepatitis due to Rubella (IgM positive, other features)
- Hepatitis due to Herpes (IgM positive)
- Hepatitis due to Toxoplasmosis (IgM positive, other features)
- Hepatitis B
- Hepatitis C
- Choledochal cyst
- Alpha1-Antitrypsin deficiency
- Hereditary tyrosinemia
- Hereditary fructose intolerance
- Storage diseases (i.e. Niemann-Pick type C, Gaucher's, GSD type IV, cholesterol ester), specify:
- Cystic fibrosis
- PFIC 1, 2, or 3
- Alagille syndrome
- Bile acid synthetic disorder
- Operable extrahepatic biliary atresia and choledochal cyst
- Galactosemia
- Cholestasis, indeterminate
- Other, specify:

B1

B2 List other diagnoses at this time (check all that apply):

- None
- Hepatitis due to CMV (CMV inclusion disease on liver biopsy or systematic congenital CMV infection)
- Hepatitis due to Rubella (IgM positive, other features)
- Hepatitis due to Herpes (IgM positive)
- Hepatitis due to Toxoplasmosis (IgM positive, other features)
- Hepatitis B

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- Hepatitis C
- Choledochal cyst
- Alpha1-Antitrypsin deficiency
- Hereditary tyrosinemia
- Hereditary fructose intolerance
- Storage diseases (i.e. Niemann-Pick type C, Gaucher's, GSD type IV, cholesterol ester), specify:
- Cystic fibrosis
- PFIC 1, 2, or 3
- Alagille syndrome
- Bile acid synthetic disorder
- Operable extrahepatic biliary atresia and choledochal cyst
- Galactosemia
- Cholestasis, indeterminate
- Other, specify:

Checking "Yes" to this question indicates that the current questionnaire or task has been completed with all available information. It will be removed from the Task list, but will remain available from the iTask through the CENSUS.

- Z1 This questionnaire or task has been completed with all available data:
- - Yes