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



## ChiLDReNLink: BASIC

	Form 29 Change in Diagnosis
	Please identify the subject's primary diagnosis (choose only one):     Please identify the subject's primary diagnosis (choose only one):      Please identify the subject's primary diagnosis (choose only one):
Β1	<ul> <li>Hepatitis C</li> <li>Choledochal cyst</li> <li>Alpha1-Antitrypsin deficiency</li> <li>Hereditary tyrosinemia</li> <li>Hereditary fructose intolerance</li> <li>Storage diseases (i.e. Niemann-Pick type C, Gaucher's, GSD type IV, cholesterol ester), specify:</li> <li>Cystic fibrosis</li> <li>PFIC 1, 2, or 3</li> <li>Alagille syndrome</li> </ul>
	<ul> <li>Bile acid synthetic disorder</li> <li>Operable extrahepatic biliary atresia and choledochal cyst</li> <li>Galactosemia</li> <li>Cholestasis, indeterminate</li> <li>Other, specify:</li> </ul>
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

## ChildrenLink

	Study ID:
	Date of Visit:
	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:
	0
	• Yes

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