



ChiLDReNLink: PROBE

Form 14 Diagnosis PROBE

A: VISIT

This form is to be completed at diagnosis or hospital discharge

| | | | |
|----|--|--------------------------|---------------------------|
| A2 | Child was hospitalized: | <input type="radio"/> No | <input type="radio"/> Yes |
| A3 | Child underwent exploratory surgery or portoenterostomy: | <input type="radio"/> No | <input type="radio"/> Yes |
| A4 | Date of diagnosis | ____ / ____ / ____ | |

B: DIAGNOSIS

| | | |
|----|---|--|
| B1 | Please identify the subject's primary diagnosis at discharge: | <input type="radio"/> Biliary atresia <input type="radio"/> Idiopathic neonatal hepatitis <input type="radio"/> Hepatitis due to CMV (CMV inclusion disease on liver biopsy or systematic congenital CMV infection) <input type="radio"/> Hepatitis due to Rubella (IgM positive, other features) <input type="radio"/> Hepatitis due to Herpes (IgM positive) <input type="radio"/> Hepatitis due to Toxoplasmosis (IgM positive, other features) <input type="radio"/> Hepatitis B <input type="radio"/> Hepatitis C <input type="radio"/> Choledochal cyst <input type="radio"/> Alpha 1-Antitrypsin deficiency <input type="radio"/> Hereditary tyrosinemia <input type="radio"/> Hereditary fructose intolerance <input type="radio"/> Storage diseases (Niemann-Pick type C, Gaucher's, GSD type IV, cholesterol ester) (specify): _____ <input type="radio"/> Cystic fibrosis <input type="radio"/> PFIC 1, 2, 3, or 4 <input type="radio"/> Alagille syndrome <input type="radio"/> Bile acid synthetic disorder <input type="radio"/> Operable extrahepatic biliary atresia and choledochal cyst <input type="radio"/> Galactosemia <input type="radio"/> Cholestasis, indeterminate <input type="radio"/> Other (specify): _____ |
|----|---|--|

B: DIAGNOSIS

| | | |
|----|--|---|
| B2 | <p>List other diagnoses at the time of discharge (check all that apply):</p> | <ul style="list-style-type: none"> <input type="checkbox"/> None <input type="checkbox"/> Hepatitis due to CMV (CMV inclusion disease on liver biopsy or systematic congenital CMV infection) <input type="checkbox"/> Hepatitis due to Rubella (IgM positive, other features) <input type="checkbox"/> Hepatitis due to Herpes (IgM positive) <input type="checkbox"/> Hepatitis due to Toxoplasmosis (IgM positive, other features) <input type="checkbox"/> Hepatitis B <input type="checkbox"/> Hepatitis C <input type="checkbox"/> Choledochal cyst <input type="checkbox"/> Alpha1-Antitrypsin deficiency <input type="checkbox"/> Hereditary tyrosinemia <input type="checkbox"/> Hereditary fructose intolerance <input type="checkbox"/> Storage diseases (Niemann-Pick type C, Gaucher's, GSD type IV, cholesterol ester) (specify): _____ <input type="checkbox"/> Cystic fibrosis <input type="checkbox"/> PFIC 1, 2, 3, or 4 <input type="checkbox"/> Alagille syndrome <input type="checkbox"/> Bile acid synthetic disorder <input type="checkbox"/> Operable extrahepatic biliary atresia and choledochal cyst <input type="checkbox"/> Galactosemia <input type="checkbox"/> Cholestasis, indeterminate <input type="checkbox"/> Other (specify): _____ |
|----|--|---|

C: INVESTIGATOR SIGNATURE

| | | |
|----|---------------------------------|---|
| C1 | <p>Investigator Signed?</p> | <p>O No → Done O Yes</p> <p>_____</p> |
| C2 | <p>Date investigator signed</p> | <p>____ / ____ / ____</p> |