

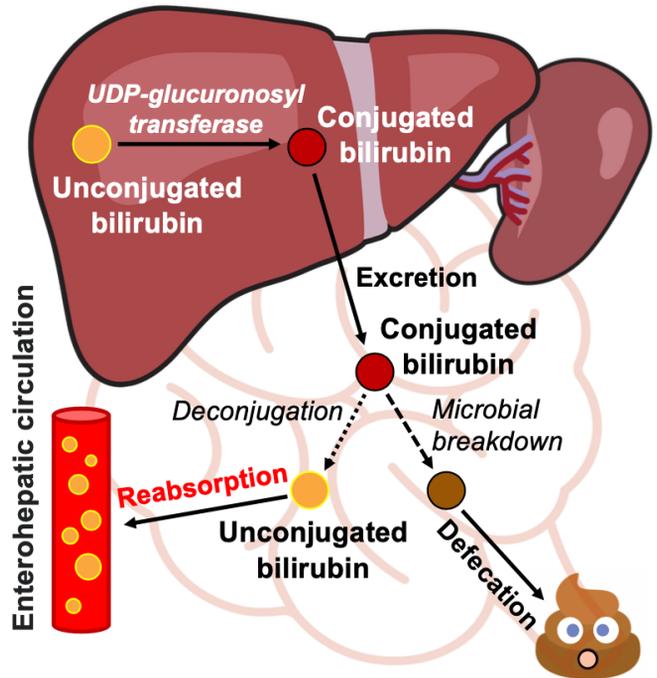


Jaundice: yellowing of the skin, scleral, and mucous membranes from a build up of bilirubin.
60% of term infants and 80% of preterm infants develop jaundice in the 1st week of life, typically an **unconjugated hyperbilirubinemia**.

SCREENING
 All infants in the **first 24 hours** of life:
 • Total serum bilirubin (TSB)
 • Transcutaneous bilirubin

RISK FACTORS

<ul style="list-style-type: none"> • Prematurity • Cephalohematoma • Bruising • Dehydration • Male sex • Maternal age ≥ 25 • Asian or European descent 	<ul style="list-style-type: none"> • Visible jaundice ≤ 24 hours • Visible jaundice before discharge at any age • Exclusive and partial breastfeeding • Sibling with severe hyperbilirubinemia
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UNCONJUGATED HYPERBILIRUBINEMIA – PHYSIOLOGIC JAUNDICE

BREASTFEEDING JAUNDICE	BREAST MILK JAUNDICE	PREMATURITY
<ul style="list-style-type: none"> • Early onset – 1st week after birth • Insufficient milk intake leads to dehydration resulting in hemoconcentration of bilirubin • Fewer bowel movements increases the enterohepatic circulation of bilirubin 	<ul style="list-style-type: none"> • Later onset – after 1st week of life • Bilirubin levels peak during weeks 2-3 of life • Can persist for 3-12 weeks • Cause unknown • It is thought that substances in breast milk interfere with the breakdown of bilirubin 	<ul style="list-style-type: none"> • Occurs in preterm infants (< 37 weeks) • More likely to require phototherapy

MANAGEMENT

<ul style="list-style-type: none"> • Phototherapy (use AAP normograms to determine the need for phototherapy – based on TSB and age in hours) • Continue breastfeeding • Supplemental PO or IV fluids (PO preferred over IV) 	<ul style="list-style-type: none"> • Phototherapy makes bilirubin water soluble by inducing a conformational change • Hyperbilirubinemia is treated to prevent kernicterus/acute bilirubin encephalopathy
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PATHOLOGIC UNCONJUGATED HYPERBILIRUBINEMIA

HEMOLYTIC	NON-HEMOLYTIC
<p>INTRINSIC</p> <ul style="list-style-type: none"> • G6PD deficiency • Hereditary spherocytosis • Thalassemia <p>EXTRINSIC</p> <ul style="list-style-type: none"> • Drugs • Iso-immune (ABO, Rh) • Sepsis 	<ul style="list-style-type: none"> • Sepsis • Hypothyroidism • Cephalohematoma • Gilbert • Crigler-Najjar <p>Work-up: Coombs test, CBC with differential, blood smear, blood culture</p>

CONJUGATED HYPERBILIRUBINEMIA

EXTRAHEPATIC	INTRAHEPATIC
<ul style="list-style-type: none"> • Biliary atresia • Choledochal cysts • Perforated bile ducts • Tumour/mass • Cystic fibrosis • Galactosemia 	<ul style="list-style-type: none"> • Infections: hepatitis, TORCH, UTI, etc. • Drugs: eg. ceftriaxone, sulfonamides, etc. • Genetic/metabolic: eg. Alagille syndrome, etc.

Must rule out biliary atresia!
 Initial investigation: **abdominal ultrasound**