

Neonatal Jaundice and Breast Feeding: A Narrative Review

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Abstract. A significant proportion of both term and preterm neonates experience neonatal jaundice. Jaundice is the most common cause of readmission to the hospital in otherwise healthy-term infants. Serum bilirubin levels rise due to red blood cell lysis, which is the main cause of jaundice. The blood carries bilirubin as "unconjugated" bilirubin, mostly attached to albumin. Most term babies have "physiological" jaundice, which doesn't require any special care and can be temporarily alleviated with phototherapy. Neonatal frequently experience unconjugated hyperbilirubinemia, which is typically normal, or the result of breast milk jaundice, which is a benign condition. But neonatal jaundice resulting from underlying cholestasis (and the conjugated hyperbilirubinemia that follows) is invariably pathological and requires immediate medical attention. Severe newborn jaundice is comparatively common in developing nations, and these babies are susceptible to bilirubin-induced brain damage

Highlights:

1. Neonatal jaundice is common, especially in term and preterm infants.
2. Physiological jaundice is benign; phototherapy often provides temporary relief.
3. Severe jaundice risks brain damage, needing urgent care, especially in cholestasis.

Keywords: Neonatal Jaundice, Mothers, Narrative Review

Introduction

One of the most prevalent clinical symptoms among newborns is jaundice. Infants with jaundice will have yellow staining of the skin and sclera, which is a sign of elevated serum bilirubin levels, which lead to bilirubin accumulation in tissues like the skin and mucous membranes. Bilirubin levels of roughly 90 mmol/liter are thought to cause jaundice (1). Although it is more challenging to identify jaundice in infants with darker skin tones, eye examination is an essential component of visual evaluation of jaundice because the sclerae are usually white (2). About 60% of term and 80% of preterm babies get jaundice in the first week of life, and 10% of breastfeeding babies experience it for up to a month (3).

Neonatal Jaundice

Most neonates have jaundice throughout the first week of life. By day three or four, the blood bilirubin level in term newborns reaches a level that may be clinically detected (4, 5). This reaches the normal childhood serum bilirubin level after progressively declining over the following few days. Physiological jaundice is the common term for this. While "breast milk jaundice" was once thought to be the primary cause of persistent unconjugated jaundice (6, 7).

Bilirubin encephalopathy

Babies who are ill or premature are especially vulnerable to the neurotoxic effects of high bilirubin levels. Pathophysiological routes include, but are not limited to, disturbance of energy metabolism, lowering of action potential, disruption of cell membrane function, and disruption of neurotransmitter synthesis (8, 9). "Kernicterus" was the term used to describe the bilirubin buildup in the brain discovered during the autopsy of newborns who had passed away from acute bilirubin toxicity (10, 11). There have been reports linking the brainstem, hippocampus, cerebellum, globe pallidum, and subthalamic nuclei. Regional cerebral blood flow and elevated metabolic activity may be linked to this anatomical tendency. acidosis, hypercarbia, hypoxia, sepsis, and seizures (12, 13).

Acute bilirubin encephalopathy risk is increased by medications that inhibit p-glycoprotein or interfere with bilirubin albumin binding. Less well-differentiated neurons and astrocytes showed higher rates of apoptosis (10, 14). Demonstrating the significance of the brain's developmental stage in connection to bilirubin poisoning. According to a UK surveillance study, one out of every 100,000 live babies develop kernicterus (15, 16).

Etiology of Neonatal Jaundice

Early jaundice

During the first 24 hours of life, clinical jaundice is usually pathogenic and caused by isoimmunization (usually ABO or rhesus incompatibility) or other causes of severe hemolysis (17, 18). An evaluation of the child's and mother's blood groups and rhesus status should be done. Because of passive antibody transmission, anti-D prophylaxis in rhesus-negative women may result in a slightly positive DAT result. The degree of jaundice is not necessarily correlated with a positive DAT test. Furthermore, the

reticulocyte count and blood film are neither very sensitive nor specific in detecting hemolysis in newborns (19, 20).

Prolong jaundice

Prolonged jaundice is defined as persistent clinical jaundice in 2-week-old term neonates and 3-week-old preterm infants. This illness, which usually shows up as unconjugated jaundice in breastfed newborns, is commonly referred to in the community (21, 22). The majority of pathological causes can be ruled out by a clinical examination, feeding history, and stool and urine color. Depending on the findings of the first investigations, the investigations ought to be carried out systematically (23, 24).

Conjugated jaundice

Serum-conjugated bilirubin levels of more than 25 mmol/liter are frequently used to diagnose conjugated jaundice. Even while clinical practice occasionally uses an in situations where the total blood bilirubin level is high, a 10% cut-off figure may provide false comfort. Dark urine and pale, chalky stools may be significant concomitant signs (25, 26). Tests for liver function and a look for clotting problems should be part of the initial inquiries. An ultrasonography of the liver can provide further information when obstructive jaundice is suspected. Further testing can rule out aminoacidemias, glucosemia, sepsis, and congenital infections. Conjugated bilirubin fraction in preterm newborns receiving total parenteral nutrition frequently rises noticeably and then progressively decreases once whole parenteral feeding is discontinued (27, 28).

Assessment of neonatal jaundice

In a well-lit space, the infant should be examined for visual evaluation of clinical jaundice. Visual evaluation is unreliable, especially in artificial light and after phototherapy has started. Dark-skinned babies may also find it challenging; in these cases, it's crucial to examine the skin, gums, and pinched areas. Serum bilirubin levels must be assessed for any newborn exhibiting clinical jaundice to facilitate care planning. When a baby's conjugated bilirubin fraction is pale, clinical jaundice often manifests visually at a serum bilirubin level between 80 and 90 mmol/l. This condition eventually gets better once all parenteral nourishment is discontinued (29, 30).

Transcutaneous bilirubin meter use has revolutionized community care for jaundiced infants in the United Kingdom in recent years. These devices' bilirubin measurements show a strong correlation with serum bilirubin levels, which can help

prevent needless blood tests and/or hospital admissions. The two measurements differed by an average of 12.7 ± 32.9 mmol/liter, even though there was a high association between serum bilirubin measurement and transcutaneous bilirubin meter measurement (31, 32).

neonatal jaundice Management

Distinguishing the rare youngster with a fast-rising bilirubin level who is susceptible to worsening is the goal of treating neonatal jaundice. Encouraging moms and babies to start breastfeeding as soon as possible (pre-discharge and in the community) can reduce the risk of elevated blood bilirubin levels and avoid readmission to the hospital (33, 34)

Unconjugated Jaundice

Phototherapy

The primary line of treatment for newborns with elevated unconjugated jaundice is phototherapy. It reduces the need for exchange transfusions and more invasive therapies. Phototherapy is only effective when bilirubin penetrates the skin and the blood bilirubin level is higher than 80 mmol/liter (35, 36). The effectiveness of phototherapy depends on the dosage, wavelength, and surface area of the baby's body exposed to the light. The dose will be increased by utilizing more phototherapy devices and positioning them at the very least safe distance from the infant. According to the American Academy of Pediatrics, a phototherapy device must have a spectral irradiance of at least $30 \text{ microW/cm}^2/\text{nm}$ (37, 38).

Bilirubin levels can be effectively lowered with fiberoptic Bili blankets. When used with an overhead unit, they can be a useful method of administering "double" phototherapy and aid in continuing phototherapy during parent cuddles. Since they produce a negligible amount of heat, they can be put near the baby (39, 40). The main way that bilirubin is eliminated is by irreversible photo alteration, which turns it into lumirubin, a structural isomer that dissolves in water and is eliminated along with bile and urine (41, 42).

Pharmacologic therapy

The only drug used in clinical practice for infants with elevated jaundice levels caused by rhesus or ABO isoimmunization is high-dose intravenous immunoglobulin.

Even if it shortens the duration required for phototherapy, hospital stays, and exchange transfusions. are eliminated through urine undergo configurational isomerization to produce less toxic and more water-soluble isomers (43, 44).

Exchange transfusion

The emergence of more advanced, extremely effective phototherapy systems and better prenatal care have greatly decreased the necessity for exchange transfusions in recent years. For babies who don't react, it's a crucial intervention. It is also recommended for newborns who have severe anemia from in-utero hemolysis (45, 46). Arterial damage, and biochemical and hematological abnormalities. Before, during, and following the exchange transfusion, the baby must be continuously monitored, and the hematological and biochemical parameters must be checked (47, 48).

Conjugated jaundice

When treating a baby with conjugated jaundice, determining the cause is essential. The baby must be examined in a surgical center for additional testing and care if a congenital blockage is detected (21, 49). To enhance bile flow, ursodeoxycholic acid, and phenobarbital are utilized, and fat-soluble vitamin supplements are also necessary (50, 51).

Breast Feeding

Hyperbilirubinemia and bilirubin accumulation are the results of delayed or insufficient breastfeeding, which impairs gastrointestinal motility and bilirubin excretion. Meconium passage is delayed as a result of inadequate milk. The production of the mother's milk is triggered when the kid is passed through the vagina during delivery. An infant born via cesarean section has a little delayed milk excretion. Due to the moms' discomfort and anesthesia, feeding is postponed; these kids are more susceptible to jaundice. Regardless of whether they gave birth naturally or via caesarian section, mothers should begin breastfeeding as soon as possible (52, 53).

Several documented factors have been linked to the occurrence of kernicterus, also known as bilirubin encephalopathy, which is a disorder caused by severe jaundice in the newborn with bilirubin deposition in the brain that causes brain damage, potentially leading to cerebral palsy, hearing impairment, visual impairment, or dementia. These factors include a brief hospital stay after delivery, delayed post-discharge appointments, an increased frequency of breastfeeding, and pediatric care

providers' lack of concern about high bilirubin levels in newborns, and delayed recognition and initiation of effective therapy for neonatal jaundice (54, 55). Research indicates that the morbidity rate for kernicterus is 70% and the mortality rate is 10% (56, 57). Most of the difficulties caused by untreated or poorly managed unconjugated hyperbilirubinemia, which are incurable, might be avoided with early detection and effective treatment of newborn jaundice (58).

Conclusion

The most prevalent ailment in neonates that requires medical treatment is newborn jaundice. Once the bilirubin level hits the treatment threshold, most babies respond well to phototherapy, and the majority of these instances start with unconjugated hyperbilirubinemia. Early treatment and attentive monitoring are necessary to prevent bilirubin neurotoxicity in infants with risk factors for developing severe hyperbilirubinemia. Future use of mobile phone cameras as an objective indicator of sclera color may be made easier by current research

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