

ABM Clinical Protocol #1: Guidelines for Blood Glucose Monitoring and Treatment of Hypoglycemia in Term and Late-Preterm Neonates, Revised 2014

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A central goal of The Academy of Breastfeeding Medicine is the development of clinical protocols for managing common medical problems that may impact breastfeeding success. These protocols serve only as guidelines for the care of breastfeeding mothers and infants and do not delineate an exclusive course of treatment or serve as standards of medical care. Variations in treatment may be appropriate according to the needs of an individual patient.

Purpose

TO PROVIDE GUIDANCE in the first hours/days of life to:

- Prevent clinically significant hypoglycemia in infants
- Appropriately monitor blood glucose levels in at-risk term and late-preterm infants
- Manage documented hypoglycemia in infants
- Establish and preserve maternal milk supply during medically necessary supplementation for hypoglycemia or during separation of mother and baby

Background

Physiology

The term “hypoglycemia” refers to a low blood glucose concentration. Clinically significant neonatal hypoglycemia reflects an imbalance between the supply and utilization of glucose and alternative fuels and may result from several disturbed regulatory mechanisms.¹ Transient hypoglycemia in the first hours after birth is common, occurring in almost all mammalian newborns. In healthy, term human infants, even if early enteral feeding is withheld, this phenomenon is self-limited, without clinical signs, and considered to be part of adaptation to postnatal life, as glucose levels spontaneously rise within the first 24 hours after birth (for some, it is even longer but still physiological).^{2–6} Most neonates compensate for this “physiological” low blood glucose with endogenous fuel production through gluconeogenesis, glycogenolysis, and ketogenesis, collectively called “counter-regulation.” Even in those situations where low

blood glucose concentrations do develop secondary to prolonged intervals (> 8 hours) between breastfeeding, a marked ketogenic response occurs. The enhanced capability of the neonatal brain to utilize ketone bodies provides glucose-sparing fuel to the brain, protecting neurological function.^{3,7–9} The compensatory provision of alternate fuels constitutes a normal adaptive response to transiently low nutrient intake during the establishment of breastfeeding,^{3,10} resulting in most breastfed infants tolerating lower plasma glucose levels without any significant clinical manifestations or sequelae.¹⁰

No studies have shown that treating transiently low blood glucose levels results in better short-term or long-term outcomes compared with no treatment, and in fact there is no evidence at all that hypoglycemic infants with no clinical signs benefit from treatment.^{11,12} Increases in neurodevelopmental abnormalities have been found in infants who have hypoglycemia associated with abnormal clinical signs, especially those with severe, persistent hyperinsulinemic hypoglycemia.^{11–16} Rozance and Hay¹⁷ have delineated the conditions that should be present before considering that long-term neurologic impairment might be related to neonatal hypoglycemia. Transient, single, brief periods of hypoglycemia are unlikely to cause permanent neurologic damage.^{18–21} Therefore, the monitoring of blood glucose concentrations in healthy, term, appropriately grown neonates is unnecessary and potentially harmful to parental well-being and the successful establishment of breastfeeding.^{18–23}

Definition of hypoglycemia

The definition of hypoglycemia in the newborn infant has remained controversial because of a lack of significant

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TABLE 1. POPULATION LOW THRESHOLDS: PLASMA GLUCOSE LEVEL⁴⁰

Hour(s) after birth	≤ 5 th percentile plasma glucose level
1–2 (nadir)	28 mg/dL (1.6 mmol/L)
3–47	40 mg/dL (2.2/mmol/L)
48–72	48 mg/dL (2.7 mmol/L)

correlation among plasma glucose concentration, clinical signs, and long-term sequelae.^{10,24,25} An expert panel convened in 2008 by the U.S. National Institutes of Health concluded that there has been no substantial evidence-based progress in defining what constitutes clinically important neonatal hypoglycemia, particularly regarding how it relates to brain injury.²⁶ Multiple reviews have concluded that there is no specific plasma or blood glucose concentration or duration of low blood glucose level that can be linked to either clinical signs or permanent neurologic injury.^{17,25,27} In addition, blood glucose test results vary enormously with the source of the blood sample, the assay method, and whether whole blood, plasma, or serum glucose concentration is determined. Plasma or serum glucose concentrations are 10–15% higher than in whole blood.^{28,29}

Breastfed, formula-fed, and mixed-fed infants follow the same pattern of glucose values, with an initial fall in glucose level over the first 2 hours of life, followed by a gradual rise in glucose level over the next 96 hours, whether fed or not.^{2,5,6} Artificially fed infants tend to have slightly higher levels of glucose and lower levels of ketone bodies than breastfed infants.^{3,5,18,30–32}

The incidence of “hypoglycemia” varies with the definition.^{33,34} Many authors have suggested numeric definitions of hypoglycemia, usually between 30 and 50 mg/dL (1.7–2.8 mmol/L) and varying by postnatal age.^{2,5,18,24,26,33,35–38} There is no scientific justification for the value of <47 mg/dL (2.6 mmol/L) that has been adopted by some clinicians.^{10,25–27,39} Cornblath et al.¹⁰ summarized the problem as follows:

Significant hypoglycemia is not and cannot be defined as a single number that can be applied universally to every indi-

vidual patient. Rather, it is characterized by a value(s) that is unique to each individual and varies with both their state of physiologic maturity and the influence of pathology

A meta-analysis of studies published from 1986 to 1994 looked at low plasma glucose thresholds in term healthy newborns who were mostly mixed fed (breastfed and formula-fed) or formula-fed. It presented statistical ranges of low thresholds for plasma glucose level based on hours after birth in healthy term infants (Table 1).⁴⁰ The authors specifically noted that given the known lower plasma glucose levels in healthy term breastfed infants as compared with formula-fed infants, the low thresholds for exclusively breastfed infants might even be lower. Table 1 gives recommendations for this timed threshold approach.

This information is translated into guidelines for clinical intervention by the operational treatment guidance of Cornblath et al.¹⁰ As they stated, an operational threshold is that concentration of plasma or whole blood glucose at which clinicians should consider intervention, based on the evidence currently available in the literature (Table 2). It needs to be underscored that the therapeutic objective (45 mg/dL [2.5 mmol/L]) is different from the operational threshold for intervention (36 mg/dL [2.0 mmol/L]), which is different from the population low thresholds in normal babies with no clinical signs or risk factors who do not need to be treated (Table 1). The higher therapeutic goal was chosen to include a significant margin of safety in the absence of data evaluating the correlation between glucose levels in this range and long-term outcome in full-term infants.¹⁰

Given this information, it is clear that routine monitoring of blood glucose in healthy term infants is not only unnecessary, but is instead potentially harmful to the establishment of a healthy mother–infant relationship and successful breastfeeding patterns.^{1,20,22,23,41,42} This recommendation has been supported by the World Health Organization,¹⁸ the American Academy of Pediatrics,^{1,41} the U.S. National Institutes of Health,²⁶ and the National Childbirth Trust of the United Kingdom.⁴³ These organizations all conclude that (1) early and exclusive breastfeeding is safe to meet the nutritional needs of healthy term infants and that (2) healthy term

TABLE 2. OPERATIONAL THRESHOLDS FOR TREATMENT OF PLASMA GLUCOSE LEVELS¹⁰

Infant	Plan/PGL	Treatment
Infant with clinical signs	If <45 mg/dL (2.5 mmol/L)	Clinical interventions to increase blood glucose concentration
Infants with risk factors ^a	Initiate glucose monitoring as soon as possible after birth, within 2–3 hours after birth and before feeding, or at any time there are abnormal signs. If plasma glucose concentration is <36 mg/dL (2.0 mmol/L), close surveillance should be maintained. Intervention is recommended if plasma glucose remains below this level, does not increase after a feed, or if abnormal clinical signs develop.	Clinical interventions to increase blood glucose concentration: at very low glucose concentration (20–25 mg/dL, 1.1–1.4 mmol/L), intravenous glucose infusion to raise plasma glucose levels to >45 mg/dL (2.5 mmol/L) is indicated.

PGL, plasma glucose level.

^aSee Table 3.

infants do not develop clinically significant hypoglycemia simply as a result of a time-limited duration of underfeeding.

Testing methods

Bedside glucose reagent test strips are inexpensive and practical but are not reliable, with significant variance from true blood glucose levels, especially at low glucose concentrations.^{22,38,44–46} Bedside glucose tests may be used for screening, but laboratory levels sent STAT (immediate determination, without delay) (e.g., glucose oxidase, hexokinase, or dehydrogenase method) must confirm results before a diagnosis of hypoglycemia can be made, especially in infants with no clinical signs.^{1,18,22} Other bedside rapid measurement methods such as reflectance colorimetry and electrode methods may be more accurate.^{47–50} Continuous subcutaneous glucose monitoring, as is used in diabetic patients, has been used experimentally in neonates with good correlation with laboratory glucose values but is not currently recommended for screening.^{51,52}

Risk factors for hypoglycemia

Neonates at increased risk for developing neonatal hypoglycemia should be routinely monitored for blood glucose levels irrespective of the mode of feeding. At-risk neonates fall into two main categories:

1. Excess utilization of glucose, which includes the hyperinsulinemic states
2. Inadequate production or substrate delivery^{32,53,54}

Infant risk factors for hypoglycemia are listed in Table 3.^{3,10,18,19,21,30,32,34,53–56}

Clinical manifestations of hypoglycemia

The clinical manifestations of hypoglycemia are nonspecific, occurring with various other neonatal problems. Even in the presence of an arbitrary low glucose level, the physician must assess the general status of the infant by observation and physical examination to rule out other disease entities and processes that may need additional laboratory evaluation and treatment. Some common clinical signs are listed in Table 4.

A recent study found that of the 23 maternal/infant risk factors and infant signs/symptoms studied, only jitteriness and tachypnea were statistically significant at predicting low blood glucose—not even maternal diabetes!⁵⁷ A diagnosis of hypoglycemia also requires that signs abate after normoglycemia is restored (the exception being if brain injury has already been sustained).

General Management Recommendations (Table 5)

Any approach to management needs to account for the overall metabolic and physiologic status of the infant and should not unnecessarily disrupt the mother–infant relationship and breastfeeding.^{1,21} Because severe, prolonged hypoglycemia with clinical signs may result in neurologic injury,^{11,14,15,58} immediate intervention is needed for infants with clinical signs. Several authors have suggested algorithms for screening and treatment.^{1,17,26,27,59} (Quality of evidence [levels of evidence I, II-1, II-2, II-3, and III] is based on the U.S. Preventive Services Task Force Appendix A Task Force Ratings⁶⁰ and is noted in parentheses.)

TABLE 3. AT-RISK INFANTS FOR WHOM ROUTINE MONITORING OF BLOOD GLUCOSE IS INDICATED

Small for gestational age: <10 th percentile for weight commonly cited in the United States; <2 nd percentile cited in the United Kingdom as above this considered small normal ^a
Babies with clinically evident wasting of fat and muscle bulk
LGA: >90 th percentile for weight and macrosomic appearance ^b
Discordant twin: weight 10% < larger twin
All infants of diabetic mothers, especially if poorly controlled
Low birth weight (<2,500 g)
Prematurity (<35 weeks, or late preterm infants with clinical signs or extremely poor feeding)
Perinatal stress: severe acidosis or hypoxia-ischemia
Cold stress
Polycythemia (venous Hct > 70%)/hyperviscosity
Erythroblastosis fetalis
Beckwith–Wiedemann’s syndrome
Microphallus or midline defect
Suspected infection
Respiratory distress
Known or suspected inborn errors of metabolism or endocrine disorders
Maternal drug treatment (e.g., terbutaline, beta-blockers, oral hypoglycemics)
Infants displaying signs associated with hypoglycemia (see Table 4)

^aAs per Dr. Jane Hawdon (personal communication).

^bUnnecessary to screen all large for gestational age (LGA) babies. Glucose monitoring is recommended for infants from maternal populations who were unscreened for diabetes during the pregnancy where LGA may represent undiagnosed and untreated maternal diabetes.

Hct, hematorit.

A. Initial management

Early and exclusive breastfeeding meets the nutritional and metabolic needs of healthy, term newborn infants. Healthy term infants do not develop clinically significant hypoglycemia simply as a result of time-limited underfeeding.^{18,19,21} (III)

1. Healthy, appropriate weight for gestational age, term infants should initiate breastfeeding within 30–60 minutes of life and continue breastfeeding on cue, with

TABLE 4. CLINICAL MANIFESTATIONS OF POSSIBLE HYPOGLYCEMIA

Irritability, tremors, jitteriness
Exaggerated Moro reflex
High-pitched cry
Seizures or myoclonic jerks
Lethargy, listlessness, limpness, hypotonia
Coma
Cyanosis
Apnea or irregular breathing
Tachypnea
Hypothermia; temperature instability
Vasomotor instability
Poor suck or refusal to feed

TABLE 5. GENERAL MANAGEMENT RECOMMENDATIONS FOR ALL TERM INFANTS

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- A. Early and exclusive breastfeeding meets the nutritional and metabolic needs of healthy, term newborn infants.
1. Routine supplementation is unnecessary.
 2. Initiate breastfeeding within 30–60 minutes of life and continue on demand.
 3. Facilitate skin-to-skin contact of mother and infant.
 4. Feedings should be frequent, 10–12 times per 24 hours in the first few days after birth.
- B. Glucose screening is performed only on at-risk infants or infants with clinical signs.
1. Routine monitoring of blood glucose in all term newborns is unnecessary and may be harmful.
 2. At-risk infants should be screened for hypoglycemia with a frequency and duration related to the specific risk factors of the individual infant.
 3. Monitoring continues until normal, prefeed levels are consistently obtained.
 4. Bedside glucose screening tests must be confirmed by formal laboratory testing.
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the recognition that that crying is a very late sign of hunger.^{41,61,62} (III)

2. Initiation and establishment of breastfeeding, and reduction of hypoglycemia risk, are facilitated by skin-to-skin contact between the mother and her infant immediately after birth for at least the first hour of life and continuing as much as possible. Such practices will maintain normal infant body temperature and reduce energy expenditure (thus enabling maintenance of normal blood glucose) while stimulating suckling and milk production.^{31,41} (II-2, III)
3. Feedings should be frequent, at least 10–12 times per 24 hours in the first few days after birth.⁴¹ (III) However, it is not unusual for term infants to feed immediately after birth and then sleep quite a long time (up to 8–12 hours) before they become more active and begin to suckle with increasing frequency. They mount protective metabolic responses throughout this time so it is not necessary to try to force-feed them. However, an unusually, excessively drowsy baby must undergo clinical evaluation.
4. Routine supplementation of healthy term infants with water, glucose water, or formula is unnecessary and may interfere with the establishment of normal breastfeeding and normal metabolic compensatory mechanisms.^{3,30,41,43} (II-2, III)

B. Blood glucose screening

Glucose screening should be performed only on at-risk infants and those with clinical signs compatible with hypoglycemia. Early breastfeeding is not precluded just because the infant meets the criteria for glucose monitoring.

1. At-risk infants should be screened for hypoglycemia with a frequency and duration related to the specific risk factors of the individual infant.^{1,19} (III) Monitoring should begin no later than 2 hours of age for infants in risk categories.¹ Hawdon⁶³ recommended blood glucose monitoring should commence before the second

feeding (i.e., not so soon after birth that the physiologic fall in blood glucose level causes confusion and over-treatment). (III)

2. Monitoring should continue until acceptable, prefeed levels are consistently obtained, meaning until the infant has had at least two consecutive satisfactory measurements.⁶³ A reasonable (although arbitrary) goal is to maintain plasma glucose concentrations between 40 and 50 mg/dL (between 2.2 and 2.8 mmol/L)¹ or >45 mg/dL (2.5 mmol/L).¹⁰ (III)
3. Bedside glucose screening tests must be confirmed by formal laboratory testing, although treatment should begin immediately in infants with clinical signs.

Table 5 summarizes these recommendations.

Management of Documented Hypoglycemia (Table 6)

A. *Infant with no clinical signs (absence of clinical signs can only be determined by careful clinical review)*

1. Continue breastfeeding (approximately every 1–2 hours) or feed 1–3 mL/kg (up to 5 mL/kg)¹⁸ of expressed breastmilk or substitute nutrition (pasteurized donor human milk, elemental formulas, partially hydrolyzed formulas, or routine formulas). Glucose water is not suitable because of insufficient energy and lack of protein. Recent reports of mothers with diabetes expressing and freezing colostrum prenatally (beginning at 34–36 weeks of gestation) to have it available after birth to avoid artificial feedings should their infant become hypoglycemic are mixed in terms of association with earlier births, and currently this procedure is not widely recommended.^{64–68} (III)

TABLE 6. MANAGEMENT OF DOCUMENTED HYPOGLYCEMIA

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- A. *Infant with no clinical signs*
1. Continue breastfeeding (approximately every 1–2 hours) or feed 1–5 mL/kg of expressed breastmilk or substitute nutrition.
 2. Recheck blood glucose concentration before subsequent feedings until the value is acceptable and stable.
 3. Avoid forced feedings (see above).
 4. If the glucose level remains low despite feedings, begin intravenous glucose therapy.
 5. Breastfeeding may continue during intravenous glucose therapy.
 6. Carefully document response to treatment.
- B. *Infant with clinical signs or plasma glucose levels <20–25 mg/dL (<1.1–1.4 mmol/L)*
1. Initiate intravenous 10% glucose solution with a minibolus.
 2. Do not rely on oral or intragastric feeding to correct extreme or clinically significant hypoglycemia.
 3. The glucose concentration in infants who have had clinical signs should be maintained at >45 mg/dL (>2.5 mmol/L).
 4. Adjust intravenous rate by blood glucose concentration.
 5. Encourage frequent breastfeeding.
 6. Monitor glucose concentrations before feedings while weaning off the intravenous treatment until values stabilize off intravenous fluids.
 7. Carefully document response to treatment.
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2. Recheck blood glucose concentration before subsequent feedings until the value is acceptable and stable (usually >40 mg/dL [2.2 mmol/L]). If staff is unavailable to check blood glucose and an infant has no clinical signs, breastfeeding should *never* be unnecessarily delayed while waiting for the blood glucose level to be checked.
3. If the infant is simply worn out and not otherwise ill, nasogastric feeds of human milk can be initiated, watching carefully for signs of intolerance or evidence of significant underlying illness. If the neonate is too ill to suck or enteral feedings are not tolerated, avoid forced oral feedings (e.g., nasogastric tube) and instead begin intravenous (IV) therapy (see below). Such an infant is not normal and requires a careful examination and evaluation in addition to more intensive therapy. Term babies should not be given nasogastric feedings. They are much more likely to fight and aspirate.
4. If the glucose level remains low despite feedings, begin IV glucose therapy and adjust the IV rate by blood glucose concentration. Avoid bolus doses of glucose unless blood glucose is unrecordable or there are severe clinical signs (e.g., seizures or coma). If a bolus dose is given, use 5 mg/kg of glucose in 10% dextrose preparation.
5. Breastfeeding should continue during IV glucose therapy when the infant is interested and will suckle. Gradually wean from the IV glucose as the serum glucose level normalizes and feedings increase.
6. Carefully document physical examination, screening values, laboratory confirmation, treatment, and changes in clinical condition (i.e., response to treatment).
7. The infant should not be discharged until reasonable levels of blood glucose are maintained through a fast of 3–4 hours. Monitoring must be recommenced if there are adverse changes in feeding.

B. Infants with clinical signs or with plasma glucose levels <20 – 25 mg/dL (<1.1 – 1.4 mmol/L)

1. Initiate IV 10% glucose solution with a bolus of 3 mL/kg and continuous IV treatment at 5–8 mg/kg/minute.
2. Do not rely on oral or intragastric feeding to correct extreme or symptomatic hypoglycemia. Such an infant most likely has an underlying condition and, in addition to IV glucose therapy, requires an immediate and careful examination and evaluation.
3. The glucose concentration in infants with clinical signs should be maintained at >45 mg/dL (>2.5 mmol/L).
4. Adjust the IV rate by blood glucose concentration.
5. Encourage frequent breastfeeding after initiation of IV therapy.
6. Monitor glucose concentrations before feedings while gradually weaning from the IV solution, until values are stabilized off IV fluids.
7. Carefully document physical examination, screening values, laboratory confirmation, treatment, and changes in clinical condition (i.e., response to treatment).

Supporting the Mother

Giving birth to an infant who develops hypoglycemia is of concern to both the mother and family and thus may jeop-

ardize the establishment of breastfeeding. Mothers should be explicitly reassured that there is nothing wrong with their milk and that supplementation is usually temporary. Having the mother hand-express or pump milk that is then fed to her infant can overcome feelings of maternal inadequacy as well as help establish a full milk supply. It is important for the mother to provide stimulation to the breasts by manual or mechanical expression with appropriate frequency (at least eight times in 24 hours) until her baby is latching and suckling well to protect her milk supply. Keeping the infant at breast or returning the infant to the breast as soon as possible is important. Skin-to-skin care is easily accomplished with an IV line in place and may lessen the trauma of intervention, while also providing physiologic thermoregulation, thus contributing to metabolic homeostasis.

Recommendations for Future Research

1. Well-planned, well-controlled studies are needed that look at plasma glucose concentrations, clinical signs, and long-term sequelae to determine what levels of blood glucose are the minimum safe levels.
2. The development and implementation of more reliable bedside testing methods would increase the efficiency of diagnosis and treatment of significant glucose abnormalities.
3. Studies to determine a clearer understanding of the role of other glucose-sparing fuels and the methods to measure them in a clinically meaningful way and time frame are required to aid in understanding which babies are truly at risk of neurologic sequelae and thus must be treated.
4. For those infants who do become hypoglycemic, research into how much enteral glucose, and in what form, is necessary to raise blood glucose to acceptable levels is important for clinical management.
5. Randomized controlled studies of prenatal colostrum expression and storage for mothers with infants at risk of hypoglycemia are important to determine if this is a practical and safe treatment modality.

Summary

Healthy term infants are programmed to make the transition from their intrauterine constant flow of nutrients to their extrauterine intermittent nutrient intake without the need for metabolic monitoring or interference with the natural breastfeeding process. Homeostatic mechanisms ensure adequate energy substrate is provided to the brain and other organs, even when feedings are delayed. The normal pattern of early, frequent, and exclusive breastfeeding meets the needs of healthy term infants.

Routine screening and supplementation are not necessary and may harm the normal establishment of breastfeeding. Current evidence does not support a specific blood concentration of glucose that correlates with signs or that can predict permanent neurologic damage in any given infant. At-risk infants should be screened, followed up as needed, and treated with supplementation or IV glucose if there are clinical signs or suggested thresholds are reached. Bedside screening is helpful, but not always accurate, and should be confirmed with laboratory glucose measurement. A single low glucose value is not associated with long-term neurological abnormalities,

provided the treating clinician can be assured that the baby was entirely well up until the time of the low value. Hypoglycemic encephalopathy and poor long-term outcome are extremely unlikely in infants with no clinical signs and are more likely in infants who manifest clinical signs and/or with persistent or repeated episodes of severe hypoglycemia.

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ABM protocols expire 5 years from the date of publication. Evidence-based revisions are made within 5 years or sooner if there are significant changes in the evidence.

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