Overview

Iron plays a critical role in early neurodevelopment in the first 1,000 days and is an essential micronutrient during pregnancy, especially in the last trimester. Michael K. Georgieff, MD, discusses how early detection for at-risk infants is crucial in long-term consequences for brain health, and proper screening for maternal iron deficiency is critical. Dr. Georgieff details how to prevent and treat iron deficiency, describes the benefits of iron supplements, and provides recommended daily intake of iron-fortified formula for both formula-fed and breastfed infants.

Target Audience

This activity was developed for pediatric physicians, nurses, nurse practitioners, dietitians, and other healthcare providers who have an interest in newborns, infants and toddlers.

Learning Objectives

At the conclusion of this activity, participants should be better able to:

- Understand how iron deficiency hinders brain development in the first 1,000 days
- Associate early iron deficiency with long-term consequences
- Optimally manage iron deficiency in pregnant women and infants.

Faculty

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This activity was released on December 23, 2020 and is eligible for credit through December 23, 2022.

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**Iron’s Impact on Brain Development**

Michael K. Georgieff, MD: My task today is to talk about nutrition in the first 1,000 days, and to focus specifically on iron. As you just learned in the learning objectives, the first thing we’re going to consider is iron’s impact on brain development.

Before we start, I just want to go back over what I think you’ve already learned about the first 1,000 days. The first 1,000 days turn out to be an important developmental period that is very sensitive to nutrition. Optimal nutrition during this time period is shown to affect fetal growth, and not just fetal growth and development, but also maternal health during the pregnancy and in the postpartum period, and essentially serving as a launching stage for the entire lifespan.

The reason for this is because the central nervous system is developing, of course, from conception onwards. Circuits are being put together during fetal life, particularly during that third trimester, and then continue to a peak in development in that first year postnatally.

We’ve now learned that in fact much of what we see in the postnatal period is driven by proper nutrition or malnutrition in the prenatal period. We’re learning that early intervention is better, and that early might be earlier than we first thought, especially when we think about iron and protein, but really applying to many, many of the nutrients.

You can think of that more as a continuum, interrupted by the process of birth, but still nevertheless a continuum of brain development. More and more studies are showing that it is really important to optimize growth before 12 months, ideally even before 4 months. I want you to think about that in terms of that including pregnancy as well, so fetal life, as well. These all work together to achieve full intellectual functioning later in
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childhood, and even into adult life, as studies in the iron deficiency field have shown.

Iron Is Essential to Mother and Child
- Iron is 1 of 9 nutrients important for healthy pregnancy and infant/toddler development
- Choline
- Folate
- Iodine
- Iron
- Omega-3 fatty acids
- Protein
- Vitamin D
- Zinc
- All these key nutrients should be included in maternal and infant diet
- Failure to provide some of these key nutrients during the first 1,000 days of life can result in a lifelong deficit in brain function
- Strong mother/infant iron relationship that affects status both in utero and in infancy

You can think of this really as an investment in society, and that the cost to society of not having adequate nutrition in those first 1,000 days ends up being loss of intellectual potential, educational loss, job loss, and so on. It's very critical to think in terms of getting nutrition right in those first 1,000 days.

Now, I'm going to be speaking about iron, but I think it's important to recognize that all nutrients are important for both the mother during pregnancy and postpartum, as well as for the child, and especially child brain development.

Essential Nutrients

Some nutrients are more important than others, that is they have a bigger impact if there happens to be a deficiency. These are listed for you on this slide [Slide 3]. You can read them for yourselves. Iron is really one of 9 nutrients that are important for maintaining a healthy pregnancy, which of course means a healthy fetal environment, as well as postnatally for infant and toddler development. All of these key nutrients should be included in the maternal and the infant diet.

Data now show both in clinical studies and in preclinical models, so we understand the mechanisms behind it, that failure to provide some of these key nutrients during the first 1,000 days of life can result in a lifelong deficit in brain function, that is not achieving full potential. The strong maternal-infant relationship as it relates to iron, and so maternal iron status is known to affect fetal iron accretion, and also affects what happens postnatally in terms of what the newborn baby's iron requirements are.

Fetal Iron Accumulation

The discussion in the iron field has been (as I mentioned with the first slide) in order to be successful with maintaining postnatal iron sufficiency, we need to think in terms of fetal iron accumulation. The human, actually all animals, but the human in particular, accretes a lot of its iron during the third trimester, and uses that iron in the first 6 months at a time when dietary iron, that is in breast milk, is fairly low. It's important that fetal accumulation occurs during the pregnancy.

Fetal Accumulation and Iron Needs During Pregnancy
- Iron sufficiency is essential to support a healthy pregnancy
  - Less prematurity
  - Less growth restriction
  - Better maternal outcomes
- Iron is vital for fetal/infant/toddler neurodevelopment
- What happens when you remove iron?
  - Why iron is needed for developing brain
  - What iron acutely does to the developing brain

Moreover, iron sufficiency itself is essential to the mom to support a healthy pregnancy. Iron deficiency during pregnancy, particularly during the first and second trimesters, results in low birth weight, either because of prematurity or because of intrauterine growth restriction. Maintaining iron sufficiency maintains a better maternal outcome. Then postnatally, iron is absolutely vital for infant and toddler neurodevelopment.
We're going to discuss in the next slides what happens when you remove iron, and why iron is needed for the developing brain, and then what does iron deficiency actually do to the developing brain. This is just a slide [Slide 5] to show you that iron, like every other nutrient, has a U-shaped risk curve to it, and that maintaining adequacy or optimum nutrient balance or optimum iron balance is best for the host. In this case, the host being the mom and the fetus.

In this case, the host being the mom and the fetus.

**Slide 5 – U-Shaped Risk for Maternal Iron Status**

You can see to the left of the curve that deficiency impairs host immunity, and, as we're going to talk about, affects brain development negatively. On the other hand, iron overload is also a concern. It's not as frequent as iron deficiency. Iron deficiency is the most frequent micronutrient deficiency on the globe, with about 2 billion people suffering from it. Iron excess can also cause problems both for the pregnancy as well as, potentially, for the fetus.

**Purpose of Nutritional Iron**

What does iron do? Iron is found primarily—there's not a lot of it in the body—but iron is found in iron clusters. That means in enzymes that have activity to produce various proteins, and in hemoproteins. A hemoprotein would be something like hemoglobin or myoglobin that's involved in tissue-oxygen delivery, or cytochromes, which are involved in energy generation. You have to have iron in order to have optimal energetics in the cells.

We'll talk about that shortly. Because of that, it really optimizes organ development and function.

Think of iron deficiency in some way as being a metabolic brownout. Not a complete blackout. It won’t kill cells, but a brownout in which the cells are not optimally functioning. It's important in brain development—literally every organ development. It’s also important in immune function.

I've listed in this slide [Slide 6] the ways in which iron affects brain development and function, so you can see neurogenesis, myelination, cellular energetics, neurotransmitter metabolism, because iron is found in the enzymes that synthesize dopamine, serotonin, and norepinephrine. It also regulates growth factors.

**Iron: Critical Nutrient for the Developing Neonate**

Iron is found in iron-cluster (eg, hydroxylases) and hemoproteins

- Tissue oxygen delivery (hemoglobin, myoglobin)
- Optimizes organ development and function
- Immune function
- Brain development and function
  - Neurogenesis
  - Myelination
  - Cellular energetics
  - Neurotransmitter metabolism (monoamines, glutamate)
- Growth Factors

**Iron Affects Brain Function**

We can drill down on that a bit to talk about what happens to the brain in terms of function when you consider iron's role. For example, in myelination there are enzymes that are involved in the synthesis of the fatty acids that are found in the myeline coating. Myeline, of course, is there to make your brain work faster. When we take a look at an iron deficient condition, we see slower speed of processing of the brain.

I mentioned those hemoproteins, so iron is involved in cellular energetics, because hemoproteins like the cytochromes are involved in ATP [adenosine
tri-phosphate] generation by the mitochondria. You need a lot of energy to grow any organ. You particularly need a lot of energy to grow the brain, and specifically its structural development. What I mean by that is during those first 1,000 days, first you have neurogenesis—the birth of the neurons—and you have migration of the neurons.\\(^6\)\\(^7\) Those are all very energy consuming kind of activities.

**Iron's Role in Brain Function**

- Myelination → Speed of processing
- Cellular energetics → Structural development (dendrites, synapses)
- Monoamine metabolism
  - Serotonin
  - Dopamine
  - Norepinephrine systems
  - Dopamine and norepinephrine can affect motor control, sleep cycles and activity, and learning and memory
- Gene regulation → Synaptic plasticity

*Slide 7 – Iron's Role in Brain Function*

Starting at about 28 weeks gestation, we get the complexity of the neurons, that is the dendrites as they branch, and the synapses, which are the connections between the neurons forming. There's a tremendous explosion of that, starting in the last trimester and carrying on through the first couple of years of life. There's your first 1,000 days.

Again, that process is extremely energetic. It takes a tremendous amount of energy, so not surprisingly you need glucose, protein, other substrates for energetics, but you also need iron to convert that through the cytochromes into ATP. Then I had mentioned the monoamines. They are synthesized by iron containing enzymes. Iron deficiency causes deficits in serotonin and dopamine and norepinephrine. Changes in those can affect motor control, sleep cycles and activity, and learning and memory.

**Iron and Gene Expression**

Then a very interesting field that is coming to light is that iron also has a direct role in regulating gene expression through epigenetics. We're not going to get into great detail on it, but you should be aware, at least, that there are iron-containing enzymes in histone, in demethylases that work on histones. That helps or that affects how genes are regulated, and specifically genes that are involved in synaptic plasticity, like growth factors, like brain-derived neurotrophic factor. Just be aware that iron has direct effects on brain function.

**Timing of Iron Intake**

Now, one of the most important principles here is the role of timing. This is one of my favorite slides [Slide 8]. What we see here is a map of brain development across the lifespan.\\(^8\) The first thing you'll notice about this slide is on the x-axis—a lot of the x-axis is devoted to the first 1,000 days. It's marked there with that yellow or orange dotted line that shows where 2 years of age is. It's kind of like everything that's happening early. You can see the waves of activity of the visual cortex, and the receptive language areas, and even the frontal cortex that does higher cognitive function, all ramping up before 2 years of age.

*Slide 8 – Iron's Role in Brain Function: Timing is Important*

If you have a nutrient deficiency, like iron deficiency, that affects any of these processes, you're going to have significant effects when they occur prior to 2 years of age, as opposed to perhaps lesser effects.
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that happen, say, in the teenage years or beyond. We've circled here for you the major areas that are developing that are iron dependent, including that synaptogenesis I talked about, so dendritic arborization and synaptogenesis, myelination, and then rapid hippocampal development.

Not surprisingly, when you look at studies of what is the effect of fetal iron deficiency or iron deficiency in the first 2 years, the affects you see are exactly in those domains.

What might some of these sequelae be, and what has been shown? Well, there's really 2 literatures here. One has to do with the neurodevelopmental sequelae of fetal and neonatal iron deficiency, so that's one peak time when iron deficiency happens, and the other, which is the much larger body of literature, is on infant and toddler iron deficiency.9,10

Iron Deficiency Consequences

If we go over here to look at the neurodevelopmental consequences of having been iron deficient as a fetus and a newborn, this was discovered about 20 years ago that babies who are born with low iron stores—so not having gotten their full complement of iron in the third trimester—that when you follow those kids out to 5, 6 years of age, they had poor school performance, and particularly in areas that involve cognition like math, arithmetic and so on.10

Moving closer to the actual time point of iron deficiency, we now have techniques where we can test the function of babies' abilities in the newborn period and look at it as a function of their iron status. Low cord blood ferritin—so ferritin is the storage molecule for iron—again, low iron stores in the newborn period cause poor recognition memory, poor ability of the baby to differentiate something novel from familiar, which is a very standard way of testing recognition memory.9

If you remember, I said that iron deficiency also affects the monoamines. Babies who are iron deficient and born to iron-deficient anemic mothers have been shown to have an altered temperament.11 They're less responsive to their mother's cue. They are not able to bond as well with their mother. These lower levels of hemoglobin and serum iron are related to higher levels of negative emotion, and lower levels of alertness and soothability. You get this dyad where the mom is iron deficient and the baby is iron deficient, and they're not responding to each other's cues. There's very nice work by Dr. Ted Wachs on that subject.11

I mentioned that iron deficiency affects bioenergetics, and we think that—and specifically neuroenergetics—and we think that's really what confers that long-term risk to neurodevelopment. Plenty of preclinical studies show that iron deficiency compromises the mitochondrial function.3,5,6 The mitochondria cannot generate as much ATP. The oxygen consumption rates of neurons that are iron deficient are cut in half by iron deficiency, and therefore there is less energy available for constructing the brain, making the synapses happen.

Neurodevelopmental Sequelae of Perinatal Iron Deficiency in Term Infants

- **General:** Low neonatal iron stores (<76 mcg/L)
  - Poorer school-age neurodevelopment [11]
- **Hippocampus:** Cord ferritin <40 mcg/L
  - Impaired recognition memory [12]
- **Dopamine:** Iron-deficient infants born to IDA mothers
  - Altered temperament [13]
  - Linear relation between neonatal iron measures and temperament
  - Lower levels hemoglobin and serum iron related to higher levels of negative emotionality and lower levels of alertness and soothability

1. Iron deficiency anemia
7. Iron Deficiency Anemia
Iron Deficiency Affects Bioenergetics

- Iron deficits in neonate → Long-term risks to neurodevelopment
- Compromises mitochondrial and cellular energetics
- More profound during development
  - Total-body oxygen consumption in infants is 3x greater than in adults
  - 60% of total body oxygen consumption is from the neonatal brain (3x greater than in adults)

**Total-Body Oxygen Consumption**

One of the reasons this happens is because the energy consumption of the baby, and the energy consumption of the baby's brain, is so great compared to adults. For example, adults consume about 40 kcal/kg of body weight. A newborn baby consumes somewhere between 85 and 100 cal/kg. That's a term baby. In a preterm baby, that number might be as high as 120 cal/kg. The total body oxygen consumption of a growing organism—and growth is most rapid from conception through the first 1,000 days—is reflected in that high total-body oxygen consumption.\(^3\)\(^5\)\(^6\)

As you all are sitting there listening to this lecture, you are using about 20% of your 40 cal/kg to run your brain. The brain actually utilizes a lot of oxygen even in us. That number is 3-times higher in babies, so fully 60% of that 100 cal/kg is going to the brain at the time of birth. That, again, just simply reflects the tremendous energy demand that growth and development put on metabolism.

**Impact of Early Iron Deficiency on Long-Term Function**

I think we can move on now into what is the impact of early iron deficiency on long-term neurodevelopment. Here, as I said, there is a fairly extensive literature, and we will go through that.\(^1\)\(^3\)

The important thing to remember is that the baby comes with a history. Until about maybe 10 years ago or so, it was thought that toddler iron deficiency—the classic iron deficiency we all learn about in school—was simply due to the lack of dietary iron intake, and that being iron deficient as a newborn was due to a whole different set of circumstances in terms of not getting enough iron. They were really thought of as 2 different fields.

**Iron Deficiency: Acute vs Long-term Effects**

- Cannot fully repair what has occurred early in the course due to iron deficiency
- Results based on different mechanisms
  - Acute effects:
    - Motor control
    - Electrophysiologic abnormalities
  - Long-term effects:
    - Cognitive delays
    - Neurobehavioral abnormalities

Some of the studies that I'm going to show you from Dr. Betsy Lozoff's group is going to hopefully show you that this is really a continuum. That is, a lot of what was ascribed to postnatal iron deficiency, or postnatal lack of iron intake, was already being set up by a lack of fetal iron accretion. Those kids that were studied in the old studies, we had no idea what their iron status was at birth. It took a study by Dr. Lozoff to show that many of those kids who present with iron deficiency at 6, 9, 12 months of age actually had been set up with low iron stores from the beginning.\(^1\)\(^2\)

One of the points I was trying to make in the previous section was that once iron deficiency occurs, our models tell us, and the clinical studies tell us, we cannot fully repair what has occurred early in the course due to iron deficiency, and that's really based on a couple of different mechanisms. One of them has to do with what are called critical periods. When the brain develops, as you saw in
that map of brain development, there are times when there is very rapid activity, lots of changes going on.

**Long-Term Effects of Iron Deficiency**

If you fail to build those areas of the brain correctly during that critical period, it seems there are, or there's now evidence there are long-term residual structural effects. If you didn't build the hippocampus right or the frontal lobe right, those structural deficits continue throughout the lifespan. The other is, I mentioned, epigenetics, so that iron controls gene regulation through histone modification.

Early on in life, you're setting a lot of your genes for their function across the lifespan. If iron deficiency alters how those genes are expressed, that can have lifelong effects. We see with iron deficiency both acute effects, mostly in the motor domain, and electrophysiologic abnormalities, and we see long-term effects, so cognitive delays and neurobehavioral abnormalities that last beyond the period of iron deficiency. What that tells you is we need to be able to diagnose the risk of iron deficiency to the brain early, and we need to replete that as soon as possible, because of these long-term effects.

**Iron Status Factors**

Now, if we look at iron status of the infant, say, at 9 months of age, and I'm grateful to Dr. Lozoff for this slide [Slide 12]—this is in one of her reviews—you can see there are many reasons why a baby might have low iron status, or an infant might have low iron status at 9 months.

The left side of this diagram shows you the pregnancy effects on that status, meaning the **mother's iron status has an impact on the iron status of the baby at birth.** Whether she has high blood pressure, whether she has diabetes during pregnancy, whether she smokes during pregnancy, those all reduce the transport of iron to the fetus. When the baby is born makes a difference, just because iron is accreted by the fetus in that last trimester. The late preterm who is born between, say, 34 and 37 weeks really has not finished getting all of its iron. Their iron needs postnataally are probably double what the term baby's needs are.

Then, whether the baby got delayed cord clamping, (I prefer not to call it delayed cord clamping. It's actually appropriate cord clamping.) When you clamp the cord early, I think we should be calling that premature cord clamping. Throughout the world now there have been multiple studies that show using delayed cord clamping, which is waiting until the cord stops pulsing, gives the baby better iron status postnataally months down the line.

Then after birth, which is more of the right-hand side of the slide, you can see some of the causes of compromised 9-month iron status. That includes a lack of iron intake and inhibitors of iron intake, like coffee and tea and phytates that are found in grains. Children may lose iron because of bleeding due, for example, to parasites, or excessive amount of cow's
Inflammation or repeated infections decreases the absorption of iron. Whereas taking vitamin C or ascorbic acid increases the availability of iron.

Maternal-Fetal Endowment

There are a lot of factors that play in, but what I wanted to emphasize here was the importance of that maternal endowment of the fetus in terms of mitigating against iron deficiency later in life. When we look at the neurobehavioral consequences of the toddlers with iron deficiency, there’s over 40 studies that demonstrate iron deficiency between 6 and 24 months leads to long-term—both short- and long-term—behavioral abnormalities; so, motor and cognitive delays while the baby is iron deficient. These cognitive delays in spite of treatment of those kids, those cognitive delays can be seen 19 to 23 years after iron repletion.¹³

Deficiency Before Anemia

Also, I want to point out that iron deficiency of all organs happens before you see anemia. Anemia is the last stage of iron deficiency. It is what we screen for iron deficiency around the world, and that’s partly because our hemoglobin is a convenient test to do. In fact, by the time you see anemia, the brain has already been affected. Now, in adults, we use something called total body iron status, but that is much less utilized in neonates and young children.

One could misinterpret a normal or even an elevated hemoglobin in a baby as being iron sufficient or perhaps even overload, when in fact it may be just that all of the iron is found in the red cells, for example, due to fetal hypoxia, which would stimulate increased iron into the red cells, but the brain could be iron deficient. Our group has documented that in intrauterine growth restricted babies, infants of diabetic mothers, and babies born to mothers who smoke.

Slide 13 – Neurobehavioral Sequelae of Postnatal Iron Deficiency in Infants

Again, they’re in those domains that I talked about with the cord blood studies, arithmetic, writing, school progress, as well as social and emotional problems; so, anxiety and depression, and social problems and inattention. Those were thought to be dopaminergic effects. These are characteristics of early perturbations of dopamine or monoamines in general and hippocampal dysfunction.
In those cases, you do not see the anemia. It is not iron deficiency anemia; it is actually pre-anemic iron deficiency. You need to know these are associated with neurodevelopmental consequences.\(^3,5,6,7\)

When you look at the toddler literature, and Dr. Lozoff has done this as well, kids who have normal hemoglobin, but who have either low stores or a low MCV (mean corpuscular volume), or some other indication that they are iron deficient—those kids have demonstrable neurodevelopmental abnormalities, particularly in attention and approach.\(^13\)

**Maternal Iron Deficiency Risks**

There are risks to the mom, as well, when she is iron deficient during pregnancy. As I mentioned earlier, there’s a higher risk of low birth weight and preterm birth, smaller placental size, and therefore slower growth of the organs in the first trimester.\(^1\) There’s an impact on fetal growth, with a risk of chronic fetal hypoxia, lower iron stores at birth, which then would give you a risk for running out of those iron stores earlier in the postnatal period. Poor cognitive development, and then cardiometabolic risk later in life.

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**Slide 14 – Nonanemic Iron Deficiency**

**Slide 15 – Risks From Maternal Iron Deficiency**

Dr. Lozoff’s studies have been particularly helpful in terms of understanding the impact of early iron deficiency on motor control. She showed that babies who are iron deficient at 9 months of age have poorer motor control.\(^13\) In the past, as I told you, this was assumed to be due to iron not being accreted postnataally. In a landmark study, or series of studies, she and her group—and [Katy M.] Clark is in her group—have shown that this really stems, to a great deal, from prenatal iron deficiency.\(^14\)

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**Impact on Early Developmental Motor Control**

They did a study that looked at mothers in China, where there is a high rate of iron deficiency. They followed the mom’s iron status throughout pregnancy, measured the baby’s iron status at birth through newborn cord blood ferritin and hemoglobin, and then followed the baby’s iron status and neurodevelopment postnatally. They
were a bit amazed to find out that many of the abnormalities they had previously ascribed to postnatal iron deficiency were actually due to prenatal iron deficiency.

### Assessing Feeding Patterns and Iron Status at 9 Mos

- **Objective** (Clark et al. 2017): Association between breastfeeding and iron status at 9 months of age in 2 Chinese provinces
- Highlights prenatal deficiency
- Odds of ID/IDA at 9 months were increased in BF and MF infants; ID/IDA was common
- Breastfeeding in later infancy identifies infants at risk for ID/IDA in many settings
- Protocols for detecting and preventing ID/IDA in BF infants are needed

<table>
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<tr>
<th>Table. Breastfeeding associated with iron status</th>
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<tbody>
<tr>
<td><strong>Zhejiang (n= 142)</strong></td>
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<tr>
<td>BF infants</td>
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<td>FF infants</td>
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<tr>
<td>Odds of ID/IDA increased in BF and MF infants compared with FF:</td>
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<td>BF vs FF OR</td>
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<td>MF vs FF OR</td>
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**Slide 17 – Assessing Feeding Patterns and Iron Status at 9 Mos**

**Human Milk Low in Iron**

They also assessed the risk to babies' iron status if they were breastfed beyond 6 months of age. This has been well described by Nancy Krebs in Colorado that there is not sufficient iron in human milk to sustain the iron needs of the baby greater than 6 months.15 Exclusive breastfeeding after 6 months of age actually increases the risks of iron deficiency quite dramatically, and you can see that in the table on the slide.14

Some of these neurobehavioral effects, especially the mood and hesitation and wariness, go on beyond the period of early iron deficiency. If I'd showed that slide about 19 to 23 years later, there is still an increased amount of depression and anxiety.8 These are all thought to be monoamine driven.
In this long-term study from China that the Lozoff group did [Slide 19], they looked at optimization or the effect of iron on motor outcomes and also at the optimization of long-term outcomes for intellectual and executive function. This is in the Santos and Lozoff article. Their objective was to assess the relationship between the timing of the iron deficiency, was it prenatal, was it postnatal, was it both, and the duration and severity. What they determined was that more severe iron deficiency in late pregnancy resulted in poorer motor behavior on the infant, and vestibular function at 9 months of age. Whereas more severe iron deficiency in infancy resulted in lower scores for locomotion and overall gross motor. There were differential motor effects based on when the timing of the iron deficiency was. Again, underscored the importance of preventing iron deficiency in the fetus, which really hadn't been thought of all that much before.

<table>
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<th>Prenatal Iron Deficiency and Motor Outcomes (continued)</th>
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<tr>
<td>- Longitudinal follow-up study (Zhang et al 2019)</td>
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<td>9 months (n=107); 18 months (n=109); 5 years (n=114)</td>
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<td>- Children with prenatal ID had significantly lower scores of motor development compared with non-ID children (52.04 vs 54.05 scores, ( \beta = 2.01, P = 0.007 ))</td>
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<tr>
<td>- Children with postnatal ID had similar scores of motor development compared with non-ID children, showing no significant difference (53.07 vs 54.05 scores, ( \beta = 0.98, P = 0.180 ))</td>
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<tr>
<td>- Motor development of children with prenatal ID did not catch up with counterparts without ID by 5 years of age</td>
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The motor development of the children with prenatal iron deficiency didn't catch up with their counterparts even by 5 years of age. Again, you can think in terms of what learning takes place in those first 5 years. Kids are starting to enter kindergarten. They've been through preschool, and so on, and yet there are these lingering effects.

### MANAGING IRON DEFICIENCY

Let's now spend the last part of the talk talking about managing iron deficiency. I think you've been hearing me harp on this idea of starting with a full tank, ensuring maternal iron sufficiency. Prevention starts prenatally. We need to be smarter about assessing maternal iron status and being sure that she doesn't have gestational conditions, like hypertension or glucose intolerance, or smoking, that are going to keep us from being able to load the fetus prenatally to protect against postnatal iron deficiency.

In that same longitudinal cohort, now published by Zhang, they looked at the kids at 9 months and then out to 18 months and 5 years, just to see, again, whether these were long lasting effects. Children with prenatal iron deficiency had significantly lower scores on motor development out at 5 years of age; whereas children with postnatal iron deficiency had similar scores of motor development when they followed them up later.
Now, when we say that, it does not mean that if you load prenatally, you don’t have to worry about postnatal iron. We do. There still is an additional role for iron. It's really a combination of the 2 that's necessary to maintain iron sufficiency throughout these first 1,000 days. Iron deficiency rates vary based on where you live in the world, how you assess the iron status. If you look at the literature, many people use simply hemoglobin to assess iron status. There's a lot of anemia out there that is not due to iron deficiency. The World Health Organization has started to recognize that we need to use a secondary test, for example, ferritin, along with the hemoglobin, to make sure we are truly diagnosing iron deficiency. When you do that or you use saturation of transferrin, or serum transferrin concentrations, you will see the rate of iron deficiency in India and in sub-Saharan Africa approaches 80%, and is highest among multiparous women, presumably because of short interpregnancy time periods.

Iron deficiency prevalence also varies among toddlers by sex and by age and by race and ethnicity. In the US, non-Hispanic blacks and Mexican Americans have the highest prevalence of iron deficiency according to the NHANES database.

**Pregnancy and Iron Deficiency**

I think it's safest to say that pregnancy is an iron deficiency event waiting to happen, if it hasn't already happened. Many moms are entering into pregnancy already with marginal iron status. Using some of the tests, that number may be as high as 40% or 45%. They're not necessarily anemic, but they have compromised either iron stores or are marginal iron status.

What happens during pregnancy is an expansion of mom's blood volume, so that of course requires more iron to fill that hemoglobin that goes into that blood volume, iron needs in the placenta, and then the iron needs of the fetus. That average net pregnancy related loss of iron—or not really loss of iron, but deficit that needs to be made up—is about 740 mg. About a gram of iron would ensure, if it was acquired during pregnancy, [that it would] maintain maternal iron balance and support the fetal and placental development.
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Importance of Iron in Last Trimester

- Iron requirements increase in each trimester
- Maternal hepcidin concentrations are suppressed in second and third trimesters, facilitating an increased supply of iron in healthy pregnancies
- Iron supports fetoplacental development
- **Definitely, Mom has a negative iron balance**
  - Due to Mom's expanding blood volume and iron needs
  - Avg net pregnancy-related loss of iron ~740 mg
  - ~1 g of iron must be acquired during pregnancy to preserve maternal iron balance and support fetoplacental development

Numbers on maternal iron deficiency are hard to get because of the inconsistency of the tests that we use. As I mentioned, anemia is the last stage of iron deficiency. If we’re waiting for that, you’re going to get a pretty low rate of iron deficiency in pregnant women, maybe 16%, 18%. If you look at more sensitive measures of iron, like transferrin saturation percentage, that number goes up quite a bit. We also know that intrauterine growth restriction due to hypertension, which is the most common cause of intrauterine growth restriction in the US, those babies (50% of them) have low iron stores at birth. The iron is just simply not transferred because of the placental insufficiency.

Screening for Maternal Iron Deficiency

- 16–18% of pregnant women are iron deficient
  - This rate (16–18%) is high for pregnant women
  - 50% of infants with IUGR have low iron stores at birth
  - Increased rate of IUGR, results in babies born with lower iron storage
  - Screening alone is not sufficient
  - Guidelines for maternal and neonatal screening and treatment are inconsistent
  - **NEONATE:** Screening should center on biomarkers that index brain health, not hematology

One of the areas that's of most interest is that we eventually want to get to a point where we can screen neonates, and toddlers for that matter, on what the iron is doing for the brain health, as opposed to just screening the hematology. Again, if you just use hemoglobin, you will have already missed the brain iron deficiency. What metrics, what markers can we start using that tell us when the brain is at risk?

**Oral Intolerance Issues**

During pregnancy, the standard is to use oral iron supplementation, but a lot of moms really hate taking iron, and up to 70% report significant gastrointestinal side effects. We know that when they take the iron, it increases the serum hepcidin, which means they are responding, and that leads to decreased absorption.

Iron is very tightly regulated. When you are iron deficient, your hepcidin is low, and you take up a lot of iron. When you ingest the iron as a medication or in your food, then your serum hepcidin will increase, and that down-regulates absorption.

**Oral vs Intravenous Iron Fortification**

- Oral iron intolerance
  - Up to 70% report significant gastrointestinal side effects
  - Oral ingestion reported to increase serum hepcidin leading to decreased absorption (which is the appropriate response)
  - IV may be appropriate
    - When oral iron is ineffective or if/when harmful
    - If anemia is severe (<8 g/dL) in second trimester
  - Evidence (Auerbach et al 2017) reports IV iron safe and effective in second and third trimesters

There are proposals now to look at whether IV [intravenous] iron, particularly in high-risk populations, might be a way to keep the mom adequately loaded, and to ensure fetal loading. These are just trials that are going on now. A recent article just came out showing the safety of it.
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Maternal Dietary Source or Iron Supplements

- Dietary recommended nutrient intakes for women
  - Pregnant: 27 mg/day
  - Lactating: 9 mg/day
- Common dietary sources:
  - Heme sources: Fish, meat, poultry, seafood
  - Non-heme sources: Fortified cereals, nuts, seeds, spinach
- Note: maternal iron status greatly impacts the fetus, however, maternal iron intake does not affect breastmilk.

Iron Food Sources

When we look at the dietary recommended nutrient intakes for Mom during pregnancy, it's about 27 mg per day. During lactation, it's about 9 mg per day. She can get that generally through foods. I think we always think in terms of food sources being better than supplements, but in the common dietary sources, you get more bang for the buck in terms of iron when you use heme sources like fish, meat, and poultry, as opposed to non-heme sources, like grains, which can contain a fair amount of iron, but where there may be phytates that block the absorption of iron.

The maternal iron status does affect the fetal iron status, particularly when she's profoundly iron-deficient. Once a mom's ferritin is below about 13.5, or once a pregnant woman's hemoglobin is less than 10, the fetus starts to get less iron.

In spite of that, and in spite of how common iron deficiency is, universal iron supplementation is currently not recommended. The reason is there are concerns that if you supplement people who are indeed iron-sufficient, that there is a risk to doing that.

Perhaps an increased risk of gestational diabetes. And so there has been some conferences where people have been trying to sort out who should actually receive the iron. Just be aware that universal iron supplementation, the US Preventive Task Force (USPSTF) does not endorse routine iron supplementation during pregnancy. Now, having said that, I think most obstetricians still do put moms on vitamins with iron.

Iron Supplements Reserved for Those at Risk: Pregnancy

- Universal supplementation versus targeted populations
- US Preventive Services Task Force stated there was insufficient evidence to advocate routine iron supplementation during pregnancy
- European Food Safety Authority concluded iron supplementation during pregnancy should be reserved for those at risk or with documented iron deficiency
- Need:
  - Data indicate ~2-3% of pregnant women in US experience IDA
  - ID estimated prevalence 16%
  - The estimate of nonanemic ID is likely an underestimate; may be as high as 45%

After birth, the question is where does the baby get the iron? I told you that is really from the stored iron, so adequate fetal loading, as well as the cord clamping, as well as the small but highly bioavailable amount of iron that is in mom's milk, which is no longer sufficient after 6 months of age. After 6 months, there's this potential gap between human milk and the dietary requirements. By the way, this is true for all of the divalent metals, so that's zinc and copper, as well.

Human Milk vs Dietary Requirements

- After 6 mos, Mother's breastmilk is no longer sufficient as a source for iron or any divalent metal (zinc, copper)
- >6 mos, potential gap between human milk and dietary requirements—highest for iron and zinc
- Infant's iron requirements exceed intake starting at 6 months of life
  - 4-6 months of age, internal stores depleted
  - Iron requirements increase
  - Additional iron support needed from infant formula, complementary foods, or iron supplements

Iron Supplementation

Slide 26 – Maternal Dietary Source or Iron Supplements

Slide 27 – Iron Supplements Reserved for Those at Risk: Pregnancy

Slide 28 – Human Milk vs Dietary Requirements
The infant’s iron status really starts to need supplementation somewhere certainly after 6 months of age, and maybe as early as 4 months of age. When we look at the recommendations for daily iron intake for an infant, again, it can be completely food based.\(^{25}\) This does not have to be medication supplementation.

Zero to 6 months, that 0.27 is reflecting just how much is in breast milk. Then in the second half of the first year, 11 mg per day, and the 1–3 years old more like 7 mg per day.\(^{26,27}\) The reason that number goes down is because the growth rate goes down, and the organ development tax is less.

### Table 2 – Recommended Daily Intake of Iron for Infant

<table>
<thead>
<tr>
<th>Infant Age</th>
<th>RDA Recommended Intake</th>
</tr>
</thead>
<tbody>
<tr>
<td>0–6 months</td>
<td>0.27 mg/day</td>
</tr>
<tr>
<td>7–12 months</td>
<td>11 mg/day</td>
</tr>
<tr>
<td>1–3 years old</td>
<td>7 mg/day</td>
</tr>
</tbody>
</table>

The AAP recommends that babies be breastfed for more than 4 months.\(^{28,29}\) They actually recommend exclusive breastfeeding for the first 6 months. We know that the iron content in human milk is low, and so supplementation needs to happen around 6 months of age. The AAP also recommends iron-fortified formula. How much iron should be in that formula is actually highly debated. I'm going to show you a slide showing the difference between what's done in Europe and what's done in the US.

### Iron-Fortified Formulas

In the US, the iron-fortified formulas have about 10–12 mg of iron per liter. Here's the comparison [Table 3]. On the left, we see AAP recommendations; on the right, we see the European Society for Pediatric Gastroenterology and Nutrition group.\(^{29}\) To be honest, babies who are well-endowed from fetal life will stay completely sufficient in terms of iron on a formula as low as 4 mg/L. The US continues to put out formulas that are 10–12 mg/L.

### Table 3 – Recommendation for Iron-Content Formula or Iron Supplementation of Exclusively Breastfed Infants

<table>
<thead>
<tr>
<th>Age</th>
<th>AAP Committee on Nutrition</th>
<th>ESPGHAN Committee on Nutrition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Formula-fed infants</td>
<td>10–12 mg/L formula</td>
<td>4–8 mg/L</td>
</tr>
<tr>
<td>0 to 6 months</td>
<td>10–12 mg/L formula</td>
<td>No specific recommendation</td>
</tr>
<tr>
<td>6 to 12 months</td>
<td>10–12 mg/L formula</td>
<td>4–8 mg/L</td>
</tr>
<tr>
<td>Exclusively breastfed infants</td>
<td>1 mg/kg per d as a supplement</td>
<td>No recommendation</td>
</tr>
<tr>
<td>&gt;4 months</td>
<td>1 mg/kg per d as a supplement</td>
<td>No recommendation</td>
</tr>
</tbody>
</table>

Is there a risk to that? Well, there might be, because, again, this issue of giving iron to people who are already iron sufficient... might there be consequences?

### When to Supplement Iron

There's one study from Lozoff's group that was done in Chile that showed iron-sufficient kids who were randomized to a high iron formula had poorer neurodevelopment at 10 years of age, whereas in that same study the kids who were iron deficient,
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who got the iron-fortified formula, did better on their outcomes.\textsuperscript{30} It's important to recognize that there may be side effects to giving iron to people who are already iron sufficient.

Timing of Fortification

- Prevention of IDA in infancy is important for brain development
- Timing of nutrient fortification emphasizes fetal loading\textsuperscript{(1)}
- Risks to supplementing iron-sufficient children
- Potential consequence of mistimed or excessive iron
  - Long-term outcomes studies in Chile\textsuperscript{(2)}
    - At 10 years, n=473 assessed (56.6%) [ClinicalTrials.gov NCT01166451]
    - Low-iron group (mean, 2.3 mg/L) compared with iron-fortified group (mean, 12.7 mg/L) scored lower on every 10-year outcome\textsuperscript{(2)}

Potential Risks of Iron Supplements: Infants

- Altered microbiome
  - In iron-replete children (12–35 months), most iron supplements are not absorbed and could promote a more pathogenic microbiome with resulting diarrhea
- Note: US has not sanctioned routine iron supplementation with concern of supplementing kids who do not need it.
- Continued research needed for well-informed public policy to determine who will benefit from iron supplementation
- What amount will provide benefit or may cause adverse outcomes?

Benefits of Iron Interventions: Infants

- Cai et al 2017 meta-analysis of four RCTs; n=511 infants
- Iron supplementation in exclusively breastfed infants
- Beneficial effects
  - Hematologic parameters
  - Cognitive development
- Significant increase in Bayley psychomotor developmental indices in later life (MD = 7.00; 95% CI; 0.99–13.01)

Who Is At Risk

On the other hand, if you look at the deficiency data, nutrients are really not consumed sufficiently by many children in the US. When you look at the NHANES data from 2001 and 2016, and look at the table [Slide 33], you can see that certain risk groups have high rates of iron deficiency at 1-2 years of age and 1-3 years of age.\textsuperscript{34} I think what that tells you is at least you need to be screening and knowing who is at risk for iron deficiency.
Slide 33 – NHANES Data on Iron Deficiency

We already went through most of this on that diagram I showed you of what determines the infant’s iron needs at 9 months, but just to review: the gestational age, whether there was adequate fetal loading or complications of pregnancy that prevented that; whether the baby got umbilical cord clamping or not; whether the baby grew rapidly. Here rapid growth actually increases the rate of iron deficiency, because the blood volume has to increase. And then duration of exclusive breastfeeding, particularly if it goes on beyond 6 months of age.25

Slide 34 – Older (6 mos) Infants’ Iron Needs

When we look at what the needs are of the baby, and, again, the iron can be provided by complementary foods or by medicinal iron supplementation, the infant between 6 and 12 months needs about 11 mg per day, and between 12 and 24 months needs about 7.28,29 I want to point out that not all food sources are equal in terms of bioavailability. Electrolyte-type iron that is found in infant cereals is far less bioavailable than iron that is found in meats, or any kind of heme iron that has a higher bioavailability.

Slide 35 – Iron Needs of Toddlers

There are some challenges in addressing iron imbalance. The question is who are we going to supplement? I think the challenges are really made more difficult because of no established cutoffs for iron repletion or iron excess, particularly in children, based on that total body iron.25,35 Nevertheless, I think it's important that there's nutritional counseling.

Slide 36 – Challenges Addressing Iron Imbalance

If 14% of children between 1 and 2 years of age are iron deficient, it's up to people taking care of those children (parents but also obviously health care
practitioners), to be informed and to support policies and programs that get them the iron that they need.

That risk of iron deficiency is not equal throughout the pediatric lifespan, specifically the times when you're most at risk is in the newborn period from lack of fetal loading, during infancy and toddlerhood, so let's say from 6 months to about 24 months, and then again during the teenage years, particularly in girls once their menses have started.36

Nutritional Counseling

- Importance of nutritional counseling
  - 14% of children aged 1 to 2 years are iron deficient
- All women and toddlers would benefit from programs and policies that support adequate nutrition
- Risk of ID is not equal throughout the pediatric lifespan
- Pediatricians need to be aware that the newborn, toddler, and adolescent are at highest risk and should be aware of factors that increase those risks

Slide 37 – Nutritional Counseling

I hope I've convinced you that iron is critical for early neurodevelopment, and that poor motor development has been reported both with newborns with fetal iron deficiency, as well as infants and toddlers with postnatal iron deficiency, and that they all probably represent a connected spectrum. That the brain is particularly susceptible to iron deficiency because of its high oxygen consumption rate, and all of those processes that are taking place that are iron-dependent. And that early detection is important, meaning before anemia sets in, in order to provide the best prevention.

Currently the AAP recommends iron-fortified formulas for formula-fed babies, but, of course, the AAP and we recommend breastfeeding as the way that babies should be fed.3,36 Even more so, we need to be aware of what the kid's endowment was, whether they got cord clamping, and then the supplementations that would start, based on the AAP, at about 4 months of age.

QUESTION & ANSWER

Editor's Note: This is a transcript of audience questions together with presenter responses from the December 15, 2020 audio webcast.

Is there any correlation between blood lead levels and iron levels in children with cognitive delay?

Georgieff: That's a great question. Thank you for asking that. It turns out that iron deficiency increases the lead burden, if there is lead to be grabbed from the environment. The reason for that is they use the same transporter from the gut into the body, and from the body into the brain. There is a transporter that takes up iron in the gut called divalent, so 2+: divalent metal transporter. It doesn't care whether it's iron, zinc, or lead. When you're iron deficient, you up-regulate, because you want to grab every bit of iron that is in the diet. If there's no iron there though, it'll take any divalent metal, and that includes lead. Lead toxicity is worse in iron deficiency.

What tests should we use in infants older than 6 months?

I think you're still going to be using hemoglobin. That is what the AAP recommends.3,36 What we're trying to do in the field is to add a second test. If I had one that I would suggest, it would be serum ferritin. Nothing gives you a low serum ferritin other than iron deficiency. It reflects low iron stores and tells you that you are teetering on the edge of your organs becoming deficient, including the brain. But there's no physiologic consequence of having a low ferritin. Nothing gives you a low ferritin, though, other than iron deficiency.

The reason it's not used more is, 1, it requires a bit of blood, and 2, it's an acute phase reactant, which
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means that when you are infected—and what infant and toddler isn't—it goes up. So, you might be fooled by a normal ferritin even though the baby might be iron-deficient or have low stores, because it'll be falsely elevated. That's the risk.

Recently the WHO has put out a bulletin saying we should be using ferritin in addition to hemoglobin to diagnose nonanemic iron deficiency.36

Is iron from a supplement considered heme or nonheme?

Iron from supplements, the medicinal supplements like iron sulfate, iron citrate, all of those, those are nonheme iron.

For infants in neonatal intensive care, do you recommend routine assessment of ferritin as that secondary marker for iron status?

Yes. Absolutely. We do that in my NICU. We start at about 2 weeks of age, because it's about that time that their hemoglobins are dropping. In the NICU, when babies are sick, we are phlebotomizing them for blood tests. That phlebotomy induces a total body decrease in iron, because you're taking out hemoglobin. Every gram of hemoglobin you take out, that's 3.5 mg of iron right there.

We typically phlebotomize and then transfuse back, meaning we're always to the negative, and then transfusing back if we choose to transfuse. We tolerate pretty low hemoglobin. We have been checking ferritin starting at 2 weeks of age. When we see the ferritins going down, we increase the iron dosing by 1–2 mg/kg and repeat the test about 2 weeks later. Then we keep checking them every 2 weeks.

For a breastfed late-preterm infant who had delayed cord clamping, would you still recommend routine iron supplementation?

Yes. Absolutely, and there's a great study in Sweden to look at that by Magnus Domellöf.37 I think the first author is Berglund. This was a great study that they did. In Sweden everything can be very controlled, and they have a national registry, and so on. Delayed cord clamping is routine. Breastfeeding is absolutely routine. They took babies who were 34–37 weeks, actually they took babies who were 2,000–2,500 g at birth. So, they were either late preterms (34 or 37 weeks), or else they were term babies who had intrauterine growth restriction.

They randomized those kids to receive no supplemental iron, so 0 mg/kg/day of iron, 1 mg/kg/day, or 2 mg/kg/day. Then they looked at their iron status at 6 months of age, and they looked at their neurodevelopment at 3 years, and at 7 years of age. What they found was a 3- or 4-fold higher rate of iron deficiency in the group that got 0 mg compared to 1 mg or 2 mg. The kids who got 2 mg/kg had less iron deficiency anemia and less nonanemic iron deficiency than the kids that got 1 mg/kg. That's at 6 months of age.37

What was most alarming was that at 3 years of age that group that had gotten 0, had more abnormalities on the Achenbach childhood checklist, particularly in the areas of attention and focus and things that are exactly what we would expect from iron deficiency. Then they looked at those same kids at 7 years of age, and those behavioral problems persisted.

As I mentioned in the talk, it tells you just how important those last 6 weeks might be in terms of accreting your full complement of iron at birth.

From an evolutionary perspective, why do you suppose breast milk is so low in iron?

That's a great question. I didn't design the system, so this is obviously just straight teleology, but I suspect it is because babies are immune-compromised little creatures. They're not very good at fighting infection. Bacteria, and for that matter protozoa, like malaria, love iron. For example, with malaria, an iron-sufficient red cell will replicate malaria a lot faster than an iron-deficient red cell. In
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some sense, iron deficiency is protective against malaria.

So, that young baby who is immune-compromised, or relatively immune-compromised, and is likely to get diarrheal diseases, infectious diseases, why would you feed a large amount of a nutrient, iron in this case, that would feed those pathogenic bacteria? Maybe it would be better, evolutionarily speaking, to load the fetus prenatally with enough iron in the iron stores—so the ferritin, do the delayed cord clamping, have them grow at the moderate rate that a breastfed baby grows, and then you really don't need a lot of iron in the diet itself.

Then you also have developed, or humans have developed a way of protecting the iron from the gut, and that is with lactoferrin as the carrier. The whole system is basically designed to make sure bacteria, that babies become colonized with, aren't pathogenic bacteria.

**Adequate ferritin vs low hematocrit, which value do you treat?**

That's a good question. You're always going to treat a low hematocrit, if it's due to iron deficiency. That's your end stage of the iron-deficiency process. Currently, other than in the NICU, there are not recommendations to treat low ferritins. That said, I think the field will move that way, particularly if the ferritin is going down. If the ferritin is going down, think of ferritin as basically money in the bank. If you're having to spend out your bank account, you're living at too high a rate, and that rate is driven by growth, obviously, in the kids.

I think we will be moving to a point where if we see low ferritins, we're going to... First of all, you can already advise parents to have their kids on more iron bioavailable foods; either foods with more iron in them or foods that have more bioavailability of the iron. If that doesn't solve the problem, I think you're going to see that we're going to be treating those kids with 1 to 2 mg/kg/day of supplemental iron.

**Please comment on iron recommendations for the preterm infant, ie, timing of initiation, dosage, and what is considered excessive.**

Great. Excellent. The starting would be at 2 weeks of age for a couple of reasons. One, your iron needs really go up when you're in your growth phase. Many preemies aren't growing in the first 2 weeks anyway. Secondly, you're waiting for a couple of antioxidant systems to mature. Iron is a potent oxidant stressor, so you want to make sure your antioxidant systems, like vitamin C and E, and so on, are mature. That happens at about 2 weeks of age according to one of the studies from Europe.

When we start iron, we use what the AAP recommends, which is 2–4 mg/kg/day. Based on what the ferritin is doing, we move that number up or down. If it looks like the baby is accumulating ferritin too quickly, meaning the ferritin is going up—so, you're in very, very positive iron balance—we'll cut back on the iron by a mg/kg/day. If the ferritin is dropping, and particularly if it's dropped below 100, then we will increase the iron supplementation by 1 mg/kg/day.

Babies who are on erythropoietin, instead of getting red-cell transfusions, need to be on 6 mg/kg/day.

**Would it be better to give iron supplementation every other day than daily, because of serum hepcidin?**

That's an interesting question. That's not done routinely, but has been proposed, and looks like it's perfectly effective to do that.

**What markers should be used to assess iron stores of brain?**

Ah, there isn't one yet. That's the conundrum. If you divide the body actually into 3 compartments of where iron is, you have the red-cell compartment. That's by far the biggest. Almost 60% of iron in the
newborn is found in the red blood cells, about 55%. You can measure that. That's the hemoglobin. If you know that there's 3.5 mg of iron per g of hemoglobin, and you know how many g of hemoglobin the baby's got, based on the hemoglobin concentration, you can figure that number out.

The second big compartment is the storage compartment, and that's indexed by the ferritin. We can convert. We have nomograms to convert that into the amount of iron. That accounts for 88% of the total body iron of a newborn. But it's that last 12%, which includes the brain iron, that is the functional part. That's the part from which you get the symptomatology and the neurodevelopmental issues, and we do not have a biomarker for that.

The best we can do is to know when the brain becomes at risk based on the other markers. We know that the brain is at risk before the hemoglobin goes down, and the brain is not at risk as long as you have a normal ferritin. That's about as precise as we can be, and there are actually some large studies out there right now trying to come up with new biomarkers for us.

Abbreviations

<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Definition</th>
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<tbody>
<tr>
<td>AAP</td>
<td>American Academy of Pediatrics</td>
</tr>
<tr>
<td>ATP</td>
<td>adenosine triphosphate</td>
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<tr>
<td>Hb</td>
<td>hemoglobin</td>
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<td>ID</td>
<td>iron deficiency</td>
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<td>iron deficient anemia</td>
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<td>IV</td>
<td>intravenous</td>
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<td>IUGR</td>
<td>intrauterine growth restriction</td>
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<td>LBW</td>
<td>low birth weight</td>
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<td>MCV</td>
<td>mean corpuscular volume</td>
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<tr>
<td>NAHES</td>
<td>National Health and Nutrition Examination Survey</td>
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<tr>
<td>USPSTF</td>
<td>U.S. Preventive Services Task Force</td>
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Reference

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