



In The Name Of God
Post Discharge Follow-Up
Of High Risk Infants

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A Case Of High Risk Infant

- *A 42 days old preterm infant with GA 29 weeks & birth weight 1100 grams is discharged from the hospital.*
- *What Are Your Follow Up Recommendations?*

The High-Risk Infant

- *The preterm infants (**Extremely preterm, Very preterm**) are the vast majority of high-risk infants.*
- *The World Health Organization (WHO) defines preterm infants as*
 - ***Extremely preterm infants: ≤ 28 wk gestational age ****
 - ***Very preterm infants: 28 - 31 wk, 6 days gestational age ****
 - ***Moderate to Late preterm infants: 32 – 36 wk, 6 days***
- *Classification based on birthweight:*
 - ***Extremely low birthweight (ELBW): birthweight <1000 g ****
 - ***Very low birthweight (VLBW): birthweight <1500 g ****
 - ***Low birthweight (LBW): birthweight <2500 g***

High Risk Infants With Birth weight >1500 grams or gestational age of ≥ 32 weeks and one of the following conditions:

- Infants with **intrauterine growth retardation**
- **Asphyxia** during or before childbirth
- **Severe persistent unstable condition** in infancy (hypoxia, acidemia, hypoglycemia or prolonged hypotension resistant to treatment with pressors)
- **Persistent apnea** requires medical treatment at the time of discharge
- **BPD or CLD**
- **Seizure disorder**
- **Persistent pulmonary hypertension** of the newborn
- **Intraventricular hemorrhage** (grade ≥ 2), periventricular leukomalacia (**PVL**)
- **Polycythemia** and **partial exchange**
- Major congenital anomaly

Follow-up Schedule After Discharge

- *The first follow-up is **24 to 48 hours** after discharge*
- *Next follow-up:*
 - *For babies weighing **1500-2000 gr** are **once a week***
 - *For babies weighing **more than 2000 gr** are **every 2 weeks***
 - *For **four to six weeks** after discharge then **monthly or every two months** (in normal growing infants)*

High Risk Infants Are At Increased Risk For Long-term Complication

- ***Neurodevelopmental Sequelae***
 - *Cognitive delay*
 - *Cerebral palsy (CP)*
 - *Fine and gross motor coordination problems*
 - *Learning disabilities*
- ***Visual and Hearing Problems***
- ***Medical Problems***
 - *Respiratory*
 - *Cardiovascular*
 - *Growth issues*

Respiratory Health

- *Approximately 23% of VLBW infants and 40% of ELBW infants develop **bronchopulmonary dysplasia (BPD)***
 - *BPD: O₂ dependence beyond 28 days with the severity assessed at 36 weeks' postmenstrual age (PMA)*
- ***short and long term respiratory morbidities:***
 - *acute respiratory exacerbations, upper and lower respiratory infections, pulmonary hypertension, cor-pulmonale, growth failure, and developmental delay, asthma-like symptoms, lack of catch-up growth in lung function*
- *commonly, infants with **significant BPD** may be discharged **home** on some combination of **supplemental O₂, bronchodilator, steroid, and/or diuretic therapy.***

Respiratory Health

- *VLBW infants are **4 times** more likely to be **rehospitalized** during the **first year** than are higher birth weight infants; **up to 60%** are rehospitalized at least once **by the time they reach school age**.*
- *Infants with BPD who are discharged home on **O2** are rehospitalized at **twice** the rate during the first 2 years of life compared to those who are not.*

Respiratory Syncytial Virus (RSV)

- *RSV is the **most** important cause of respiratory infection in premature infants, particularly in those with chronic lung disease.*
- *VLBW infants should receive **prophylactic** treatment with **palivizumab** (Synagis) monoclonal antibody.*
- *AAP recommends treatment during RSV season for **at least the first year of life** for*
 - *infants less than 32 weeks' gestation with BPD*
 - *continued in the second year of life for those who have required diuretic or oxygen therapy*

RSV Prophylaxis

- *The method of administering **Palivizumab** to all premature infants under 2 years of age from November to May is once every 30 days in the form of intramuscular injection at a dose of 15 kg/mg for 5 consecutive days.*
- *The **influenza vaccine** is also recommended for **VLBW** infants once they are older than 6 months chronologic age.*

Air Travel

- *Air travel is not recommended for infants with **BPD** because of*
 - *the increased risk of exposure to infection*
 - *the lowered cabin pressure resulting in lower O₂ content in the cabin air*
- *If an infant's partial pressure of oxygen (**PaO₂**) is **≤80 mm Hg**, supplemental O₂ will be needed while flying.*

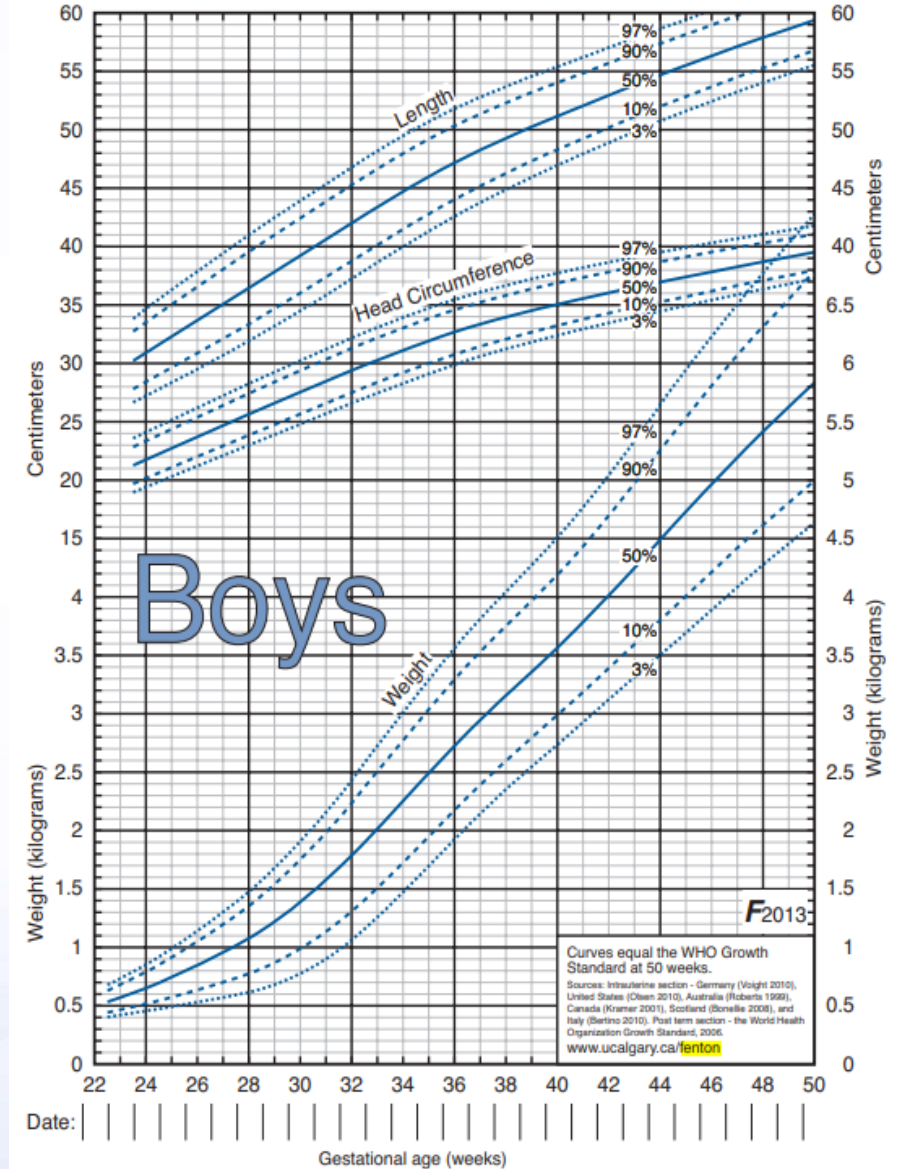
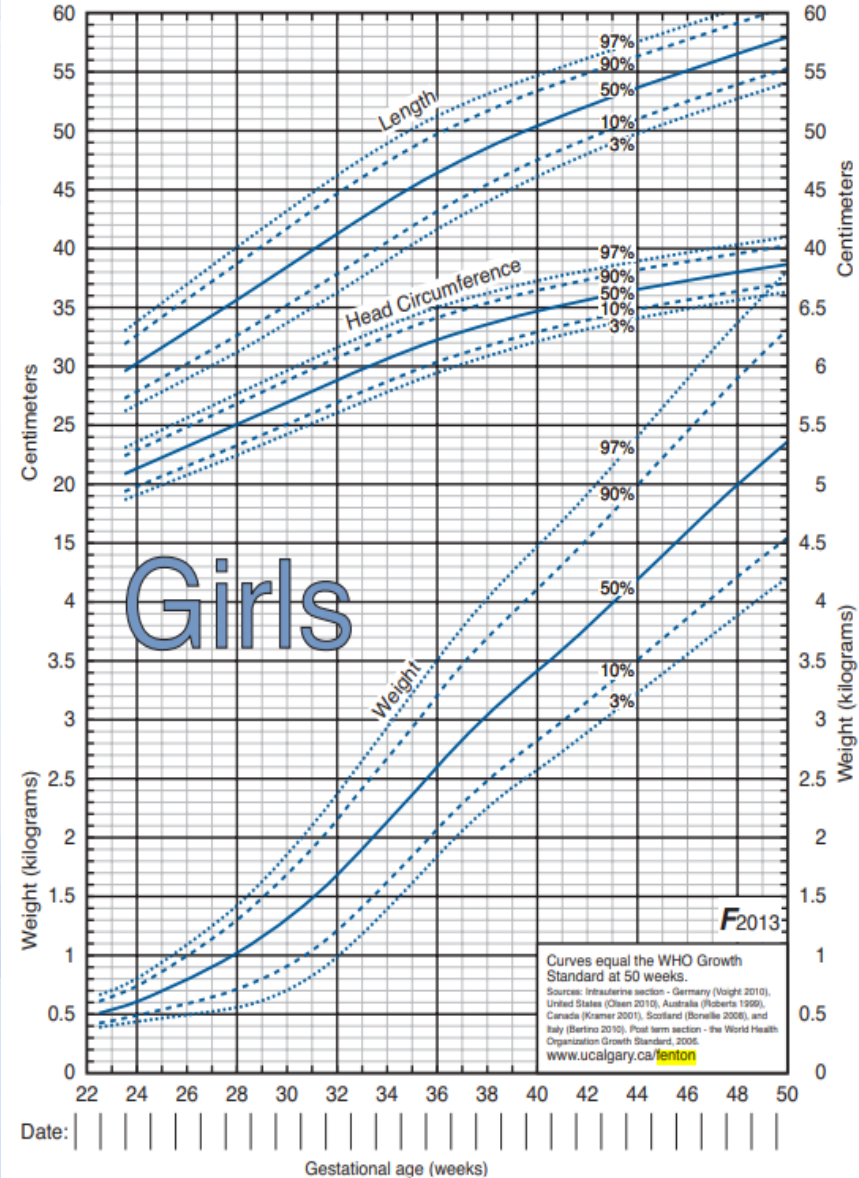
Immunizations

- ***VLBW infants should receive their routine pediatric immunizations, with the exception of rotavirus and hepatitis B vaccine.***
- ***The AAP recommends initial vaccination of preterm infants at or following discharge from the hospital if***
 - ***clinically stable and***
 - ***chronologic age between 6 weeks and 14 weeks 6 days.***
- ***Medically stable, thriving VLBW infants should receive the hepatitis B vaccine as early as 30 days.***
- ***If the baby is ready for discharge to home before 30 days of age, it can be given at the time of discharge to home.***

Growth

- *VLBW infants have a high incidence of **feeding and growth problems** for multiple reasons:*
 - ***Increased caloric needs** for appropriate weight gain (severe BPD)*
 - ***Abnormal or delayed oral motor development***
- *Growth should be followed carefully on standardized growth curves (WHO International Growth Curves 2006) using the **corrected gestational age for the first 2 years of life** and then using the Centers for Disease Control and Prevention (CDC) standardized curves.*
- *The Revised **Fenton** growth charts combine intrauterine growth with the WHO chart to construct a growth chart from **22 to 50 weeks' PMA.***

Fenton Growth Chart



Assessment Of Nutritional Status

- *Nutritional assessment is recommended for all infants with gestational age ≤ 34 weeks or birth weight ≤ 1800 grams.*
- *Values indicating impairment in nutritional assessments:*
 - *Weight gain less than 15 to 20 g/kg/day for infants < 2 kg and 25 to 35 g/day for larger infants*
 - *Height increase less than 1 cm per week*
 - *Increase in head circumference less than 0.7 cm per week*
 - *Phosphorus < 4 mg/dl*
 - *Alkaline phosphatase > 900 IU/L*
 - *BUN < 5 mg/dl*

Recommendations For Nutritional Impairment

- *In case of the above disorders, the following are recommended:*
 - *Control of feeding and breastfeeding*
 - *Use of breast milk fortifiers*
 - *Use of premature or post-discharge formula*
 - *Use of nutritional supplements*
 - *Treatment of infants with osteopenia of prematurity*
- *Specialized preterm infant formulas with increased protein, calcium, and phosphate (either added to human milk or used alone) should be considered for the first 6 to 12 months of life for infants who have borderline growth.*

Recommendations For Nutritional Impairment

- *If growth failure persists, gastrointestinal pathology such as **severe gastroesophageal reflux disease** or endocrinologic problems such as **growth hormone deficiency** should be considered.*
- *Administering iron supplements in premature babies:*
 - *for **LBW** infants: **2-3 mg/kg** daily from 1 to 24 months old*
 - *For **VLBW** & **ELBW** infants: **3-4 mg/kg** daily from 1 to 24 months old*
- *Clinical evaluation and, if necessary, **screening** by measuring hemoglobin and hematocrit levels **before the age of 6 months***
- *For **iron deficiency anemia**, treatment with **ferrous sulfate** daily at the rate of **4-6 mg/kg** for **at least 3 months**.*

Hearing Follow-up

- *Hearing loss occurs in approximately **2% to 11%** of VLBW infants.*
- *Prematurity increases the risk of both **sensorineural and conductive hearing loss.***
- *All VLBW infants should be **screened** both in the **neonatal period** and **again before 1 year of age.***
- *There is also evidence that VLBW infants are at increased risk for **auditory dyssynchrony** (also called **auditory neuropathy**) and **central auditory processing problems.***

Audiometry In High-risk Infants

- *Clinical conditions leading to hearing impairment in high-risk infants:*
 - *use of **ototoxic drugs** such as **aminoglycosides** and **furosemide***
 - ***Respiratory alkalosis***
 - ***Lack of noise control** in the NICU*
- *Recommended method to evaluate this group of infants:*
 - ***Automated Auditory Brain Stem Response (AABR)***

Screening and follow-up process for hypothyroidism in high-risk infants

- *Check TSH at 3 to 5 days, 2, 4 and 6 weeks*
- *In the first month of birth, $T4 < 5.6$ and $TSH > 5$ mU/L should be treated as primary hypothyroidism until 3-4 years of age.*
- *Levothyroxine dose: 10-15 mcg/kg Max: 50 mcg/ day*
- *Follow up of treatment:*
 - *2 weeks after the start of treatment*
 - *Then monthly in the first 6 to 12 months of life*
 - *Every 2 months for the next 12 months*
- *Ferrous sulfate and soy-based formula interfere with the absorption of levothyroxine*
- *It is recommended to give the medicine to the infant 2 hours before or after breastfeeding*

Follow-up Of Osteopenia Of Prematurity

- ***Osteopenia** often appears a few weeks after birth, but the symptoms of **rickets** (epiphyseal dysplasia and skeletal deformation) usually appear in the **sixth** week of life.*
- *VLBW infants who have had nutritional deficits in calcium, phosphorous, or vitamin D intake are at increased risk for rickets.*
- *Etiology:*
 - *Inadequate intake of minerals and reduced absorption after birth*
 - *The use of Furosemide*
 - *long-term use of corticosteroids*
 - *TPN for a long time*
 - *Renal excretion of phosphorus and vitamin D reduction*
- *Measurement of serum levels of calcium, phosphorus and alkaline phosphatase at **4-5 weeks** and at **3 and 6 months of age**.*

Diagnostic Criteria Of Osteopenia Of Prematurity

- *Serum phosphorus $< 4 \text{ mg/dl}$ and alkaline phosphatase $> 800-1000 \text{ IU/L}$. are helpful in diagnosing osteopenia. Serum calcium level can be normal, high or low.*
- *Treatment : start with 20 mg/kg of elemental calcium and $10-20 \text{ mg/kg}$ of elemental phosphate and $400-1000 \text{ IU}$ of vitamin D daily.*
 - *Max: $70-80 \text{ mg/kg}$ e-calcium and $40-50 \text{ mg/kg}$ e-phosphorus.*
- *Prevention of metabolic bone disease:*
 - *Supplementation of calcium and phosphorus in premature babies weighing less than 1800 to 2000 grams (breastfed)*
 - *In cases of breast milk insufficiency, prescribing premature infant formula at the rate of 120 kcal/kg*

Dental problems

- *VLBW infants are at increased risk for **enamel hypoplasia and discoloration**.*
- *Long-term intubation may result in **palate and alveolar ridge deformation** affecting tooth development.*
- *Initiation of routine **supplemental fluoride at 6 months PMA** is recommended, as is referral to a pediatric dentist in the **first 12 months of life**.*

Follow-up Of Nephrocalcinosis

- *Factors related to kidney calcification include:*
 - ***Hypercalciuria** as the main pathophysiological factor, **hypophosphatemia**, **hypercalcemia**, long-term **oxygen** therapy, **male** gender, treatment with drugs such as **gentamicin**, **vancomycin**, **dexamethasone**, **Lasix** and **methylxanthines***
- *Nephrocalcinosis screening :*
 - ***KUB ultrasound at 36 weeks** (corrected gestational age) or **1 month**; and **4 years old**.*

Intraventricular Hemorrhage Follow-up

- ***Predisposing factors include:***
 - ***prematurity, intrauterine growth restriction, hypothyroxinemia, hypercapnia/hypercarbia, hypothermia, asphyxia, placental or maternal infection, and increased cytokines.***
- ***Most babies with IVH have no clinical symptoms.***
- ***Apnea, pallor, cyanosis, poor sucking, loud crying, seizures, hypotonicity, metabolic acidosis, and shock can be symptoms of IVH.***
- ***50% occur on the first day of birth and 75% of cases occur between the first and third days.***
- ***PVL is usually clinically asymptomatic until neurological complications and white matter damage appear as spasticity in the infant.***

Intraventricular Hemorrhage Follow-up

- *The ecodense phase occurs on days 3 to 10 and the cystic phase on days 14 to 20.*
- *Brain ultrasound should be performed in **ELBW** infants between **days 3 and 5** and in other **VLBW** infants at the **end of the first week** of life.*
- *Repeat brain ultrasound **2 to 3 times a week** (at least once a week).*
- *Follow-up of IVH should be done at **36 to 40 weeks** of corrected gestational age or at the **end of the first month** and before discharge.*
- *Cranial ultrasound is recommended in babies **older than 32 weeks** if there are risk factors such as **asphyxia, pneumothorax** or **neurological symptoms**.*

- ***When is the first screening program for retinopathy of prematurity in preterm babies between 23 and 26 weeks of gestational age?***
 - 1. At 4 weeks of chronological age.***
 - 2. At 30 weeks' PMA.***
 - 3. At 31 weeks' PMA.***
 - 4. At 32 weeks' PMA.***

Follow-up Of Retinopathy Of Prematurity

- *AAP clinical guideline recommendations for ROP screening:*
 - *All babies with a gestational age < 30 weeks or*
 - *Birth weight < 1500 grams*
 - *Babies with a gestational age > 30 weeks or a weight of 1500 to 2000 grams if they have an **unstable clinical condition***
- *Problems causing clinical instability:*
 - *perinatal **asphyxia**, prolonged **hypoxia**, severe **acidosis**, severe **hypoglycemia** or **hypotension** requiring **vasopressor drugs**, **IVH**, **BPD**, repeated **apnea** ,....*

Follow-up Of Retinopathy Of Prematurity

- *The time of the first eye examination:*
 - *In infants with a gestational age ≤ 27 weeks \rightarrow at the age of **31 weeks' PMA***
 - *In infants with a gestational age > 27 weeks \rightarrow **4 to 5 weeks after birth***
- *Eye examination can be done with or without anesthesia.*
- *If anesthesia is required, examinations should be performed in a hospital with a neonatal intensive care unit.*
- *Examination without anesthesia can be done at the bedside or in a clinic equipped with newborn monitoring and resuscitation facilities.*
- *Eye exams are usually continues **every 1 to 2 weeks** until the baby is **at least 38 to 40 weeks' PMA.***

Treatment Of Retinopathy Of Prematurity

- *Laser therapy to ablate the entire peripheral avascular retina (the standard treatment)*
- *Intravitreal injection of anti-endothelial growth factor mainly **bevacizumab** (Avastin)*

Other Ophthalmologic Problems

- ***Refractive errors (Myopia is the most common problem and may be severe)***
- ***Amblyopia (can become permanent if it is not treated before 6 to 10 years of age)***
- ***Strabismus (especially in those with a history of ROP, intracranial hemorrhage, or white matter injury)***
- ***In infants who have had severe ROP including those treated with laser therapy, there is an increased risk of cataracts, glaucoma, late retinal detachment, abnormal color vision development, and visual field deficits.***



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- *Close monitoring until the **retinal vessels** have reached **maturity** (in the **first months of life**)*
- *Subsequent assessments (by **8 to 12 months of age** then **annually** or **at 3 years of age**)*

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