This manual, revised for the first time, is designed for use by the pediatric residents and interns created by residents for the UF-Jacksonville Pediatric Residency Program. The recommendations in this manual are specific for the practices in this program. Please understand that this is not a mini-textbook or outline of general newborn care. The purpose of this manual is to assist pediatric house officers by:

a) Providing a guideline for management of patients that require immediate attention
b) Reminders to help them in their daily work

There is little discussion of pathophysiology, pharmacology and infectious disease processes. Certain important and common problems are not covered at all. If at any time you are unsure about the contents provided in this guide, please refer to more comprehensive texts, or contact the on call attending.
Phone Numbers

Dr. Dial 306-4734  Dr. Sharma 306-3979
Dr. Garrison 306-3981  Dr. Banadera 393-5959
Dr. Alviedo 393-4800  Dr. Huddleston 306-3976
NICU 244-5100  Dr. D. Cuevas 306-3974
3N 244-6110  Dr. Tan 498-5350
Step down 244-3330  Dr. Driscoll 393-4414
L&D 244-6127  UF ID 393-6299
Lab 244-6040  Genetics 393-1180
Nemours 697-3600  Dr. Spierre (Rehab) 633-0926
Attending call room 244-3348  Radiology 244-6084
NP/PA call room 244-5109

Schedule

0700 Sign-out from Night call/ on-call PA
1000 Rounds (can be later, attending dependant)
1200 Noon Conference
1400 Radiology Rounds (excluding Thursday)
1900 Sign-out to Night call/ on call PA

Expectations

- If you are on overnight, please double check that your name & pager number are on the white board located by the NICU Secretary and in the Step down unit.
- All orders and notes must have a DATE, TIME, SIGNATURE and DOCTOR NUMBER.
- Be familiar with the standing orders
- Attend afternoon clinic
- If you are called about an infant, evaluate/examine and document a note (why you were called, your examination, your findings, and what actions were taken).
- Document any procedures, updates with family, etc. in Neodata (print & place in chart)

NICU POLICIES & PROCEDURES

Infection Control

- Scrub for 3 minutes to elbow when first coming into the NICU and Step-Down units
- No watches or jewelry, except for a plain wedding band while handling patients
- Please keep sleeves at or above your elbows when handling patients
- Wash hands or use foam before and after handling patients
- Must gown and glove (in addition to hand washing) when handling isolated patients
Patients with NEC should be considered as having a communicable infection with use of isolation technique above until proven Rotazyme negative

NO FOOD/DRINKS are allowed anywhere except the nursing/staff lounge – this includes keeping residents’ computer station clear of food/drinks

Charts, including nursing charting material, must remain at the patient’s bedside

Please follow sterile techniques when introducing central lines/procedures

Chart Entries

- All orders and notes must have a DATE, TIME, SIGNATURE and DOCTOR NUMBER.
- Verbal orders are prohibited except during an emergency such as a resuscitation or when gowned for or performing a sterile procedure.
- All verbal and telephone orders must be signed before leaving for the day.
- All read back orders written by the nurse must be co-signed
- Check all flagged chart pages for need for signature
- Orders “flagged” in chart should be placed face down to comply with HIPPA

Other

- Any baby transferred to normal nursery must have an interim summary on the chart before transfer and either the nursery attending or Patty Williams ANRP notified by telephone prior to transfer
- Patients returning from surgery must have all orders re-written
- Patients transferred to the Step Down unit must have orders re-written
- All medication orders must contain the method of dose calculation ie scheduled doses and, mg/kg; or for infusions: concentration, mcg/kg/min and mL/kg/min.
- TPN, Lipids and large fluid volume infusions should be ordered using the TPN program
  - To access, password is FEEDME
- Remember not to use banned abbreviations, especially the use of “q.d” and trailing zeros on medication or fluid orders (E.g “5” and not 5.0mg )
- All parental consents must be signed on admissions or as close to admission as possible
PULMONOLOGY

Ventilator Terminology

Peak inspiratory pressure (PIP)
- Maximum pressure measured during the delivery of gas during conventional mechanical ventilation.
- PIP reflects the effects of the amount of gas delivered to the lungs in a given breath (tidal volume)

PEEP
- PEEP helps to maintain functional residual capacity (FRC). At the end of expiration, the PEEP exerts pressure to oppose passive emptying of the lung and to keep the airway pressure above the atmospheric pressure. The presence of PEEP opens up collapsed or unstable alveoli and increases the FRC and surface area for gas exchange, thus reducing the size of the shunt

Rate
- Reflects how often a volume of gas in the system is delivered to the infant.

Inspiratory/Expiratory Ratio
- I/E ratio reflects the relationship between time spent in inspiration and time spent in expiration.
- If the I/E ratio is 1:2 with a rate of 60 and the total respiratory cycle is 1 second, inspiration is 0.33 second and expiration is 0.66 second.
- Prolonged inspiration may be associated with more efficient ventilation, optimal arterial oxygenation, a higher risk of air leak, and impending of venous return.
- Prolonged expiration also improves oxygenation, especially in air-trapping conditions

Mean Airway Pressure (MAP)
- Amount of pressure transmitted to the airway throughout an entire respiratory cycle.
- Ways to increase MAP
  - Increase PEEP, or PIP, or Ti, or RR, or Flow

Minute Ventilation (Vt)
- Determines rate of carbon dioxide removal
- Minute ventilation= TVxRR

Amplitude
- Amount of pressure oscillation that occurs around the MAP.
- Increasing the amplitude will increase the TV and therefore decrease PCO2.
- Decreasing the amplitude will decrease the TV and therefore increase PCO2.
Mechanical Ventilation and Respiratory Support

Types of Vents

Synchronized Intermittent Mandatory Ventilation (SIMV)
- Delivers a set number of breaths with a certain amount of pressure each minute, synchronizes with babies’ inspiration attempts
- When ordering SIMV - write for PIP, PEEP, Rate, IT, and FiO2

SIMV/VG
- VG - Volume guarantees a certain tidal volume, usually 6ml/kg per assisted breath. Target tidal volume maintained by the ventilator as the pressure limit varies inversely with lung compliance.
- You still set PEEP and rate

Assist Control
- Not used much in our NICU, you still must set the PIP and PEEP
- The vent assists every breath the baby takes, even if the baby breathes 80 times/min
- Must set a minimum backup rate in case the baby does not breathe at all.
- Reserved for very ill neonates who require very high support

Pressure Support Ventilation (PSV)
- Pressure increases in proportion to inspiratory volume.

High-Frequency Ventilation
- Goal of HFOV is to reduce barotrauma
- Delivers a very fast in and out oxygen supply - the baby does not take breaths but there is a continuous in and out motion multiple times per minute
- Gas exchange with a kind of double spiral effect - there is a pulse of O2 going centrally down the airway with pulses of CO2 going out up the sides of the airway.
- Main parameters are amplitude which affects volume in and out with each oscillation. Mean Airway pressure (MAP) affects expansion and oxygenation of the lungs and frequency of oscillation (Hz) affecting how many times per minute the exchange occurs
- Oscillators vibrate columns of air and have active exhalation cycles. Typically set at 10-15Hz (600-900 breaths/min)

JET Ventilator
- Used if the infant has a pneumothorax, PIE, over/hyperexpanded lungs, pneumatoceles, and severe meconium aspiration/PPHN
- Allows longer exhalation time vs oscillator (passive exhalation)
- Change: PIP to change CO2, PEEP (oxygenation & expansion), and rate (expansion)
- Back up breaths: to treat and correct atelectasis (can increase oxygenation in RDS)

Starting Pressures for beginning ventilator support
- FiO2: min. to keep SaO2 88%-92% (for premies) and > 95% (for term infants)
- PEEP: 4-6cm water
- PIP: 14-20cm water (14-16 for premature infants, especially <1000 grams, and 16-18 for term infants)
- Rate: 40-60
- I/E ratio: 1:1-1:2
- I-time: 0.35-0.4
Changing Vent Settings

- What affects Tidal Volume?: PIP, PEEP, and AMP

<table>
<thead>
<tr>
<th>Factors</th>
<th>TV</th>
<th>CO2</th>
</tr>
</thead>
<tbody>
<tr>
<td>Inc PIP</td>
<td>Increases</td>
<td>Decreases</td>
</tr>
<tr>
<td>Dec PEEP</td>
<td>Increases</td>
<td>Decreases</td>
</tr>
<tr>
<td>Inc AMP</td>
<td>No change</td>
<td>Decreases</td>
</tr>
</tbody>
</table>

- What affects respiratory Rate?
  - Vent Rate → increase in rate will decrease CO2
  - I: E ratio → decrease I time will increase E time which will decrease CO2

- What affects airway pressure/expansion?

<table>
<thead>
<tr>
<th>Factors</th>
<th>Volume</th>
<th>O2</th>
</tr>
</thead>
<tbody>
<tr>
<td>Inc PIP</td>
<td>Increases</td>
<td>Increases</td>
</tr>
<tr>
<td>Inc PEEP</td>
<td>Increases</td>
<td>Increases</td>
</tr>
<tr>
<td>Inc MAP</td>
<td>Increases</td>
<td>Increases</td>
</tr>
</tbody>
</table>

- What affects O2 concentration?
  - FiO2
  - Change oxygenation by either changing FiO2 or PIP
  - Monitor oxygenation one of three ways
    - SaO2
    - ABG → PaO2 (only accurate on ABG, on VBG or CBG tells you essentially nothing)
    - CXR - count at least 8 ribs

- How to change your PaCO2?

<table>
<thead>
<tr>
<th>Variable</th>
<th>Rate</th>
<th>PIP</th>
<th>PEEP</th>
<th>IT</th>
<th>FiO2</th>
</tr>
</thead>
<tbody>
<tr>
<td>Increase PaCO2</td>
<td>Decrease</td>
<td>Decrease</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>Decrease PaCO2</td>
<td>Increase</td>
<td>Increase</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>Increase PaO2</td>
<td>NA</td>
<td>Increase</td>
<td>Increase</td>
<td>Increase</td>
<td>Increase</td>
</tr>
<tr>
<td>Decrease PaO2</td>
<td>NA</td>
<td>Decrease</td>
<td>Decrease</td>
<td>NA</td>
<td>Decrease</td>
</tr>
</tbody>
</table>

- Formula
  - Remember AMP changes CO2 (increase AMP dec CO2)
  - MAP changes O2
  - Extubate MAP <7 and FiO2<35%

\[
\text{Formula } OI = \frac{\text{FiO2} \times \text{MAP} \times 100}{\text{PaO2}}
\]

OI = oxygen Index, indicative of severe resp distress, >20 need for Nitric Oxide & >30 ECMO

MAP = \( \frac{(Ti \times PIP) + (Te \times PEEP)}{Ti + Te} \) or \( \left( \frac{\text{RR} \times \text{Ti}}{60} \right) \times \left( \frac{\text{PIP} - \text{PEEP}}{\text{Ti}} \right) + \text{PEEP} \)

Blood Gases

<table>
<thead>
<tr>
<th>Type</th>
<th>pH</th>
<th>PCO2</th>
<th>HCO3</th>
<th>Compensation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Metabolic Acidosis</td>
<td>&lt;7.4</td>
<td>&lt;40</td>
<td>low</td>
<td>-BE (hypervent)</td>
</tr>
<tr>
<td></td>
<td>&lt;7.4</td>
<td>&gt;40</td>
<td>high</td>
<td>+BE (hypovent)</td>
</tr>
<tr>
<td>--------------------------</td>
<td>------</td>
<td>-----</td>
<td>------</td>
<td>---------------</td>
</tr>
<tr>
<td>Respiratory Acidosis</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Metabolic Alkalosis</td>
<td>&gt;7.4</td>
<td>&gt;40</td>
<td>high</td>
<td>+BE (hypovent)</td>
</tr>
<tr>
<td>Respiratory Alkalosis</td>
<td>&gt;7.4</td>
<td>&lt;40</td>
<td>low</td>
<td>-BE (hypervent)</td>
</tr>
</tbody>
</table>

Sodium Bicarbonate: investigate cause of metabolic acidosis, e.g. hypovolemia or sepsis, etc., and treat accordingly. (Refer to Pediatrics - Sodium bicarbonate: basically useless therapy. Aschner JL, Poland RL. Pediatrics. 2008 Oct;122(4):831-5.)

Give if adequate ventilation; very seldom indicated
Goal BE < -10
Give not more than 1 meq/kg/hr, followed by ABG after 1hr

Formula
\[
\text{mEq Na HCO}_3 = \frac{\text{BE} \times 0.3 \times \text{wt (kg)}}{2} \quad (\frac{1}{4} \text{ correction})
\]
\[
\text{mEq NaHCO}_3 = \frac{\text{BE} \times 0.6 \times \text{wt (kg)}}{2} \quad (\frac{1}{2} \text{ correction})
\]

Metabolic Alkalosis

<table>
<thead>
<tr>
<th>CO2</th>
<th>Oxygen</th>
<th>DDx</th>
</tr>
</thead>
<tbody>
<tr>
<td>Low</td>
<td>High</td>
<td>Over ventilation, air bubbles, hyperventilation</td>
</tr>
<tr>
<td>High</td>
<td>Normal/High</td>
<td>Obstructed ET, ET down right main stem bronchus, pneumothorax, PDA, permissive hypercapnea</td>
</tr>
<tr>
<td>High</td>
<td>Low</td>
<td>Pneumothorax, improper ET position, PDA, atelectasis, lung disease</td>
</tr>
<tr>
<td>Normal</td>
<td>Low</td>
<td>Agitation, pneumothorax, improper ETT position, atelectasis, pulmonary HTN, pulmonary edema</td>
</tr>
</tbody>
</table>

Criteria for use of Palivizumab (Synagis)

**Indication**
- Indication for the prevention of serious lower respiratory tract disease caused by RSV
- Potential Candidates for Palivizumab

<table>
<thead>
<tr>
<th>Patient Group</th>
<th>Age at Start of RSV Season</th>
</tr>
</thead>
<tbody>
<tr>
<td>Premature, no chronic lung disease or congenital heart disease</td>
<td></td>
</tr>
<tr>
<td>- ≤ 28 weeks gestation age</td>
<td>≤ 12 months</td>
</tr>
<tr>
<td>- 29-32 weeks gestational age</td>
<td>≤ 6 months with 1 additional risk factor</td>
</tr>
<tr>
<td>- 32-35 weeks gestational age</td>
<td>≤ 6 months with 2 additional risk factor</td>
</tr>
<tr>
<td>Chronic Lung Disease</td>
<td>≤ 2 years</td>
</tr>
</tbody>
</table>
Hemodynamically significant congenital heart disease ≤ 2years
Other high-risk children with pulmonary or immune function ≤ 2years

**risk factors:** child care, school-age siblings, exposure to environmental air pollution, congenital abnormalities of the airways, or severe neuromuscular disease

**Dosage**
- 15mg/kg IM
- Given every month during RSV season

**Apnea**
- >20seconds breathing cessation
  - Central- no diaphragm activity
  - Obstructive- upper airway obstruction with diaphragm activity
- Treatment: Caffeine (see dosing in pharmacology guide)

**Bronchopulmonary Dysplasia (BPD)**
- Due to arrested lung development resulting from interference with alveolarization and vascularization
- >36wk, still requiring oxygen at 28days
- If O2 sats <92% and if hypoxia develops chronically, it can cause pulmonary HTN (cor pulmonale)
- Keep CO2>30 on AGB, if less then PVL (periventricular leukomalacia) may develop

**Chronic Lung Disease (CLD)/ Bronchopulmonary Dysplasia (BPD)**
- Supplemental oxygen requirement at 28-30 days of life or 36 weeks postmenstrual age.
- Occurs due to chronic and constant lung injury with ongoing repair and healing of the injury.
- Also caused by lung immaturity, oxygen toxicity, and barotrauma/volutrauma (high PIP and PEEP, pneumothorax, and PIE)

**Meconium Aspiration Syndrome (MAS)**
- Typically occurs in a post term infant who becomes hypoxic in utero. Fetal asphyxia causes the anal sphincter to relax and colonic peristalsis ensues expelling meconium into the amniotic fluid. A second episode of asphyxia occurs, during which the infant makes gasping respiratory movements, causing meconium to enter into the oropharynx and lung.
  - Etiology
    - Atelectasis: due to blockage of smaller airway causing air trapping and alveolar collapse
    - Chemical pneumonitis: inflammatory response to meconium
- Inhibits surfactant function
- Increases pulmonary vascular resistance

  o Diagnosis
    - History
    - CXR: air trapping, hyperexpansion, and hyperinflation; bilateral diffuse, coarse, patchy infiltrates

  o Treatment
    - During labor: Amnioinfusion
    - Delivery: intubate and tracheal suction in depressed infants (*do not stimulate baby prior to this!*)

- Anytime there is a history of thick meconium at delivery with respiratory distress, obtain histopathology on the placenta. Meconium staining of placental can determine length of time (how long prior to delivery) the baby released meconium and was in distress.

**Air leaks**
- Types: pneumomediastinum, pneumothorax, pneumopericardium, or subcutaneous emphysema
- Pulmonary Interstitial Emphysema (PIE)
  - Occurs when free air is released from ruptured alveoli
  - Diagnosis
    - CXR
    - Fiberoptic probe may reveal hyperlucency of affected side
  - Treatment
    - Elevate head 30-40 degrees (this decrease the work of breathing)
    - Give 100% oxygen
    - Needle aspiration and chest tube placement
      - Lateral approach- 4th or 6th intercostals space
      - Superior approach- 2nd-3rd IC space or just lateral to midclavicular line

**Acute Respiratory Distress Syndrome (RDS)**
- Defined- disease of immature lung anatomy and physiology.
- Pathology-
  - Anatomically the preterm lung is unable to support oxygenation and ventilation due to insufficiently developed alveolar saccules, which causes a deficient surface area for gas exchange. The volume of surfactant is insufficient to prevent collapse of unstable alveoli. Because the alveoli collapse with each breath, normal FRC is not established.
  - Increase WOB due to poorly compliant lung and insufficient oxygenation and ventilation requiring increased energy output. Increased WOB results in hypoxemia and academia which cause constriction of pulmonary vascular (arterial) musculature, severely limiting pulmonary capillary blood flow. Without adequate pulmonary capillary blood flow, the type II pneumocytes become deficient in the precursor material necessary for production of surfactant.
  - Diagnosis
    - CXR- show reduced lung volumes, air bronchograms (visible outlines of air-filled bronchi due atelectasis), reticulogranularity, and lung opacification.
  - Treatment
    - Surfactant
    - Correct academia
- Reduce hypoxemia- by maintaining BP and Hct, maintain neutral thermal environment.

**Transient Tachypnea of Newborn (TTN)**
- Etiology- seen in term or near term infants born via C-section
- TTN is due to lack of gradual compression of the chest that eliminates some fluid during a normal vaginal delivery. This causes a delay in reabsorption of normal lung fluid and lung fluid accumulates in peribronchilar lymphatics and the bronchovascular spaces, causing an “obstructive” lung disease. Accumulation of interstitial fluid interferes with the forces that hold the bronchioli open, causing collapse and air trapping.
- Dx: CXR- show hyperexpansion with streaky infiltrates radiating from the hilum
- Tx: supplemental oxygen

---

**Nitric Oxide Treatment Protocol for Improvement of Survival Without BPD**
- **Eligibility**
  - birth weight/GA groups
  - 500-799 grams: on NCPAP or ventilator
  - 800-1000 grams: if on ventilator
  - 1001-1250 grams: if on ventilator and GA< 28 0/7 weeks
  - postnatal age 7-14 days
  - Respiratory severity score (RSS=MAPxFiO2) ≥2
- **Exclusions**
  - lethal congenital anomalies
  - intent to withdraw care
  - bilateral grad 4 IVH
  - Target SpO2
  - recommend 88-94%
- **Treatment**
  - initiate iNO at 20ppm for 96hrs; obtain methemoglobin level at 24hrs
  - wean iNO to 10ppm for 168 hours (7 days)
  - wean iNO to 5ppm for 168 hours (7 days)
  - wean iNO to 2ppm for 168 hours (7 days)
  - discontinue iNO
  - Note: treatment continues even if on nasal CPAP or nasal cannula

---
NUTRITION

Total Parenteral Nutrition (TPN)
- Source of nutrition for those who cannot eat adequately
  - Contains a mixture of dextrose, protein, electrolytes, vitamins, and minerals.
  - Usually the yellow IV bag hanging at the bedside
- Route
  - Peripheral IV limits
    - Dextrose: 12.5% (12.5 g/100 mL)
    - Calcium: 12 meq/L or 300 mg/100 ml
    - Potassium: 60 meq/L
    - Protein: 2-3%
  - Central IV (PICC, Broviac/Hickman Port)
    - Dextrose: maximum of 25%
- Labs (general guidelines – always subject to change based on individual cases)
  - Prior to starting TPN, check a triglyceride level
    - Goal is to have triglycerides ≤ 150 mg/dL
  - Daily chemistry (BMP)
    - Check magnesium, phosphate, triglycerides q Monday & Thursday
- Formulas
  - Determine Kcal/kg/day
    - \[ \text{Kcal in 24hr period} = \frac{\text{Kcal/kg/24hr}}{\text{Weight (kg)}} \]

<table>
<thead>
<tr>
<th>Weight/Age</th>
<th>Fluids</th>
<th>Labs</th>
</tr>
</thead>
<tbody>
<tr>
<td>&gt;1500grs/ Term</td>
<td>DOL 0-80-90mL/kg/d DOL 1-100-120 mL/kg/d DOL 2-140-150 mL/kg/d</td>
<td>q AM BMP DOL 2 or 3 Tbili/Dibili</td>
</tr>
<tr>
<td>1000-1500gm</td>
<td>DOL 0-80-100mL/kg/day DOL 1-100-200 mL/kg/d DOL 2-120-160 mL/kg/d</td>
<td>At 12hrs of life: BMP DOL 1 Tbili/Dibili q AM BMP, Hct</td>
</tr>
<tr>
<td>800-1000gm</td>
<td>DOL0-100mL/kg/d DOL 1-130 mL/kg/d DOL 2-150 mL/kg/d</td>
<td>BMP q8-12hrs x 48hrs, then DOL1 Tbili/Dibili q AM BMP, Tbili</td>
</tr>
<tr>
<td>600-800gm</td>
<td>DOL0-120 mL/kg/d DOL 1-130-50 mL/kg/d DOL 2-150-170 mL/kg/d</td>
<td>BMP q8hrs x 48hrs, then q AM BMP, Hct, Tbili</td>
</tr>
<tr>
<td>&lt;600gm</td>
<td>DOL0-120mL/kg/d DOL 1-150 mL/kg/d DOL 2-180-200 mL/kg/d</td>
<td>BMP q8hrs x 48hrs, then q AM BMP, Hct, Tbili</td>
</tr>
</tbody>
</table>

Note, it takes 50-75 kcal/kg/day to maintain weight in a thermo-neutral environment

GOAL: 120 kcal/kg/day
<table>
<thead>
<tr>
<th>Standard Electrolytes</th>
<th>Starting</th>
<th>Advance</th>
<th>Weight/Age Based</th>
<th>Max</th>
</tr>
</thead>
<tbody>
<tr>
<td>Carbohydrate</td>
<td>5-8 mg/kg/day</td>
<td>1-2 mg/kg/day</td>
<td>ELBW→6-8 mg/kg/min</td>
<td>Glucose concentration, no more than +/- 2%</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Preterm 4-6 mg/kg/min</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Term 4mg/kg/min</td>
<td></td>
</tr>
<tr>
<td>Protein (watch BUN, Cr)</td>
<td>2-3 gr/kg/day</td>
<td>0.5-1 g/kg/day</td>
<td>Peripheral line-30g/L</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Central line-4g/L</td>
<td></td>
</tr>
<tr>
<td>Fats (20% IL) (watch triglycerides)</td>
<td>Term: 2 g/kg/day</td>
<td>0.5-1g/kg/day</td>
<td>4 g/kg/day</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Preterm: 1 g/kg/day</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sodium</td>
<td>0.8 mEq/kg/day</td>
<td>1-2mg/kg/day</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Start if Na &lt;135 Goal 2-4mEq/kg/day</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Potassium (watch UOP)</td>
<td>DOL 2 when K&lt;3.5 1-4mEq/kg/day</td>
<td>1g/kg/day</td>
<td>4mEq/kg/day</td>
<td></td>
</tr>
<tr>
<td>Calcium</td>
<td>DOL 1 with 2.5mEq/kg/day Goal 1.5-2.5mEq/kg/day</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mg</td>
<td>0.25-0.5 mEq/kg/day</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Phosphate</td>
<td>0.5-1.5 mEq/kg/day</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Fluid Requirements**
- First 24hrs of life
  - Start 10W or D7.5 with AA & calcium at 80 kg/day in term infants and 100 mL/kg/day in preterm infants
  - D10W= 10g/100mL, therefore 1mL D10W= 100mg Glucose (1g Dextrose=3.4kcal)
- Max Volume
  - 150-160 mL/kg/day (>150mL/kg/day is associated with CLD and PDA)
  - 130mL/kg/day for infant’s with CLD
  - VLBW infant may need more fluids due to larger surface area, therefore more insensible loss

**To wean off TPN (general guidelines)**
- Begin enteral feeds on DOL 3 if stable at 20ml/kg/day (20% total volume) q 3hrs
  - Please note: this is a GENERAL rule of thumb, more mature infants can be started on enteral feeds on DOL 1
- Advance enteral by 20-30 mL/kg/day
- Decrease TPN rate accordingly to maintain overall fluid balance

**Enteral Feeds**
- Criteria for oral (PO) feeds
  - > 1500 grams
  - Suck/swallow reflex (> 32-34wk)
  - Regular stools/ing pattern, good bowel sounds
- No UAC, no NG with bile, no presssors
- **Start feeds**
  - Use breast milk if available (may need to be fortified to meet caloric needs)
  - Formula should be 20kcal with iron
  - 10-20 mL/kg/day q 3hrs
- **Increase Feeds**
  - Advance PO feeds by 20% per day.
    - Be cautious, aggressive feeding can result in NEC
    - Goal is usually between 150-160 mL/kg/day
  - Breast fed infants need to suck well for 10 mins prior to advancing
  - Advance PO attempts as follows: 1PO attempt/day to 1 PO attempt/shift to 2-3 PO attempt/day, PO every other feed, to PO every feed *(again, this is a general guideline)*
- **Weight Gain**
  - If gaining weight well on 24kcal formula, consider changing to a 22cal formula 3-4 days prior to discharge to ensure continued weight gain
- **Human Milk Fortifier (HMF)**
  - May add when tolerating > 100mL/kg/day
  - Add 1pkt/50mL to 22kcal/oz or 1pkt/25mL to 24kcal/oz
  - HMF increases Ca so if patient has hypercalcemic, add MCT oil to increase calories
  - MCT oil can also be used for patients with CLD
- **Supplements**
  - Once on full formula feeds, add:
    - 2-4mg/kg/day of Iron BID
    - Multivitamin 1mL daily
- **Formula**
  - | Formula | Rate | Caloric Value |
  - |-----------------|------------|----------------|
  - | 20kcal/oz | 180mL/kg/day | 120kcal/kg/day | 30mL= 0.67kcal |
  - | 24kcal/oz | 150mL/kg/day | 120kcal/kg/day | 30mL= 0.8kcal |
  - After tolerating 100-140mL/kg/day, can change to 24kcal/oz formula
  - Remember 24kcal/oz formula is ISOtonic and ISOsmolar like 20kcal/oz
  - 27kcal/oz formula is HYPERosmolar, so use with caution
- **Residuals**
  - Often re-feeding regimen is attending dependent
  - Good rule of thumb
    - If < 10% total feeds → refeed and continue feeding
    - If 10-30% → subtract and refeed
    - If >30% → hold feeds and evaluate infant *(always consider NEC)*
  - If >20mL worry about intestinal obstruction, if >60mL almost definite intestinal obstruction
  - If the infant has continued large residuals, look at how the baby is positioned *(R side down will decrease emptying, while L side down will incease emptying)*
GASTEROENTEROLOGY

Gastrochisis vs Omphalocele

<table>
<thead>
<tr>
<th></th>
<th>Gastrochisis</th>
<th>Omphalocele</th>
</tr>
</thead>
<tbody>
<tr>
<td>Location</td>
<td>Right of umbilicus</td>
<td>Central</td>
</tr>
<tr>
<td>Umbilical Cord</td>
<td>Intact</td>
<td>Umbilical cord elements</td>
</tr>
<tr>
<td></td>
<td></td>
<td>surround the sac and join</td>
</tr>
<tr>
<td></td>
<td></td>
<td>above the extrusion</td>
</tr>
<tr>
<td>Protective Sac</td>
<td>Full-thickness defect, no sac</td>
<td>Protective sac is preserved</td>
</tr>
<tr>
<td>Liver and Spleen</td>
<td>Remain within the abdominal</td>
<td>Partially extruded in large</td>
</tr>
<tr>
<td></td>
<td>cavity</td>
<td>defects</td>
</tr>
<tr>
<td>Associated Congenital</td>
<td>Rare</td>
<td>Common</td>
</tr>
<tr>
<td>Anomalies</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Treatment

- Fluid/Temperature regulation- monitor for hypothermia
- Wrap abdomen in wet dressing to protect and cellophane
- Place NG tube to decompress the stomach
- Intubate to reduce abdominal pressure
- Start broad spectrum antibiotics
- Obtain stat KUB
- Immediate Surgical Consult

Duodenal Obstruction

- Present- vomiting and abdominal distension, may have history of polyhydraminos
- Dx: KUB- double bubble sign (x ray to the left)
- Tx: NPO, IVF, NG decompression, and surgical repair

Proximal GI Obstruction

- Vomiting is prominent and distention less prominent
- Dx: KUB and cross table, UGI with SB follow through
- Tx: NPO, IVF, NG decompression, surgical repair

Distal GI Obstruction

- Present- abdominal distention, absence or limited passage of stool, feeding intolerance
- Dx: KUB with cross table reveal multiple loops of dilated bowel, contrast enema may reveal colonic atresia, microcolon, or meconium plug
- Tx: depends on cause
Meconium ileus
- Present - abdominal distension, feed intolerance - vomiting, passage of few (inspissated meconium) or no stool
- Dx: KUB with cross table. Barium enema reveal microcolon
- Tx: Enemas, may need surgery

Necrotizing Enterocolitis (NEC)
- Present - feeding intolerance with increased gastric residuals, abdominal distention, heme + stools, abdominal wall discoloration. Hypotension, hypothermia
- Labs - thrombocytopenia
- Classification
  o IA - suspected NEC
  o IB/IIA - suspected Definite, mildly ill NEC
  o IIB - definite moderately ill
  o II A - advanced severely ill, intact bowel
  o IIIB - advanced, severely ill, perforated bowel
- Tx: NPO, NG decompression, KUB, draw a CBC, test stool for occult blood, serial abdominal exams, Surgical consult

<table>
<thead>
<tr>
<th>Diagnostic Category</th>
<th>Suspected NEC</th>
<th>Definite NEC</th>
<th>Advanced NEC</th>
</tr>
</thead>
<tbody>
<tr>
<td>Systemic illness</td>
<td>Temperature &amp; glucose instability, increased episodes of apena/brady</td>
<td>Mild to moderate symptoms</td>
<td>Shock like picture with possible hypotension</td>
</tr>
<tr>
<td>Abdominal symptoms</td>
<td>Increased gastric residuals, abdominal distention, occult or grossly bloody stools</td>
<td>Abdominal distension &amp; tenderness, absent bowel sounds</td>
<td>Marked distension, peritoneal signs</td>
</tr>
<tr>
<td>Labs</td>
<td>Metabolic acidosis, thrombocytopenia</td>
<td>Metabolic &amp; respiratory acidosis, DIC</td>
<td></td>
</tr>
<tr>
<td>Imaging</td>
<td>Normal to mild distention; bowel wall thickening; possibly a fixed dilated loop</td>
<td>Pneumatosis intestinalis; portal venous gas</td>
<td>Pneumoperitoneum if perforated</td>
</tr>
</tbody>
</table>
HEMATOLOGY

Polycythemia
- Defined as Hct >65%, that is seen in neonates exposed to chronic hypoxia in utero, or a delay in cord clamping
- Risks- hyperviscosity, sludging
- Present- hypoglycemia, ruddy complexion, irritable
- Confirm peripheral stick value with central blood sample
- Treatment is by partial exchange transfusion

Transfusions
- When to Transfuse
  - Hct < 25, asymptomatic with a retic <1%
  - Hct < 30, with:
    - < 35% on Oxy hood,
    - CPAP/Mech vent with MAP<6
    - Significant episodes of apnea/bradycardia (>9 in 12hrs or >2 that require more than moderate stimulation to recover)
    - Persistent HR>180 or RR >80 over 1day
    - <10g/d weight gain over 4days when receiving >100kcal/kg/day
    - Pre-op
  - Hct < 35
    - >35% on oxy hood
    - CPAP/Mech vent with MAP > 6-8
  - Hct< 40
    - On ventilator
- PRBC
  - 15-20mL/kg over 3-4hrs
  - Draw post-transfusion H/H 4hrs after transfusion (especially if no labs scheduled for the morning, or if infant is extremely unstable/critical)
  - If the infant is < 1kg, order “DESIGNATED UNIT” on pink slip held for 35days from 1st draw date
  - 2mL/kg PRBC will increase Hgb by 1g/dL

Anemia of prematurity
- Nadir is usually seen at 4-6wks of life
  - Supplement with iron when on full feeds
    - Some attendings may choose to start Epoepoetin
    - Taper epo dose if Retic >7, and discontinue when infant is > 1800gm
  - Check Hct and Retic q Monday

Platelets
- Plateletphoresis of 15-20mL/kg IV (platelets “drop” in, therefore no time requirement needed on orders)
- Check 4hr post transfusion level (especially if no labs scheduled for the morning, or if infant is extremely unstable/critical)
- Transfuse if <20k or <50k with active bleeding
- Pre-op
OPHTHALMOLOGY

Retinopathy of Prematurity (ROP)

- Occurs when there is disruption of the normal progression of retinal vascular development in the preterm infant which results in abnormal proliferation of blood vessels in the developing retina (neovascularization). ROP develops in 84% of preterm infants < 28wks.
- Etiology
  - Pregnancy complications (HTN, DM, smoking)
  - Supplemental oxygen (oxygen radicals produced)
  - Anemia: decrease in oxygen-carrying capacity, results in increased FiO2 to maintain adequate oxygenation, thus exposing the lung/retina to more oxygen/oxygen toxicity.
  - Apnea/Bradycardia: causes repeated cycles of hypoxia/hyperoxia and hypoxemia/hypercapnia exposing retinal vessels to alternating ischemia and hyperoxic toxicity.
- Degree of retinal vascularization at birth determines an infant’s susceptibility to ROP
- Screen for ROP if the infant has one of the following:
  - Birth weight < 1500 g or <30wk
  - Birth weight between 1500-2000 g with unstable clinical course
  - Exam shall be done at 32 weeks post conception on average
    - If gestational age at birth is between 22-27 weeks, schedule the first ROP exam at 31 weeks
    - If gestational age at birth is 28 weeks, schedule the first ROP exam at 32 weeks
    - If gestational age at birth is 29 weeks, schedule the first ROP exam at 33 weeks
    - If gestational age at birth is 30 weeks, schedule the first ROP exam at 34 weeks
    - If gestational age at birth is 31 weeks, schedule the first ROP exam at 35 weeks
    - If gestational age at birth is 32 weeks, schedule the first ROP exam at 36 weeks
- Pre-exam meds
  - Cyclomydril (cyclopentolate/phenylephrine) 1drop OU Q5min x3, give 1hr prior to exam
  - 0.5% Tetracine at bedside
- Follow up schedule

<table>
<thead>
<tr>
<th>Follow up</th>
<th>Stage I or II with Zone I</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Stage 3 ROP Zone II</td>
</tr>
<tr>
<td>1-2wk Follow up</td>
<td>Immature Vascularization, Zone I, no ROP</td>
</tr>
<tr>
<td></td>
<td>Stage 2 ROP- Zone II</td>
</tr>
<tr>
<td></td>
<td>Regressing ROP, Zone I</td>
</tr>
<tr>
<td>2wk Follow up</td>
<td>Stage I ROP, Zone II</td>
</tr>
<tr>
<td></td>
<td>Regressing ROP, Zone II</td>
</tr>
<tr>
<td>2-3wk Follow up</td>
<td>Immature vascularization, Zone II, no ROP</td>
</tr>
<tr>
<td></td>
<td>Stage 1 or 2 ROP, Zone III</td>
</tr>
<tr>
<td></td>
<td>Regressing ROP,Zone III</td>
</tr>
</tbody>
</table>

- Zones
  - Plus Disease - degree of dilation and tortuosity of the posterior retinal blood vessels
Stages of ROP

Figure from J Bernbaum. Preterm Infants in Primary Care: A Guide to Office Management, 2000.

Stage 1.
Abnormal branching of developing vessels creates a demarcation line between the avascular and vascular retina.

Stage 2.
Vascular retinal engorgement occurs, with dilated and tortuous terminal vessels. The demarcation line becomes a ridge.

Stage 3.
Extraretinal fibrovascular proliferative tissue appears, such as neovascular tufts with hemorrhages or exudates.

Stages 4 and 5.
Peripheral and total retinal detachment, respectively, take place.
CARDIOLOGY

Fetal Circulation and Birth
- In the fetus, pulmonary vascular resistance is high, and pulmonary artery blood pressure > systemic blood pressure, causing blood flow from the main pulmonary artery to travel through the open ductus arteriosus to the descending aorta. A second right-to-left shunt occurs across the foramen ovale in the fetus.
- Pulmonary vascular resistance is high in the normal fetal “hypoxemic” state, which means that pulmonary vascular resistance and pulmonary artery blood pressure decrease as the PaO2 of the neonate increases. At birth, the ductus arteriosus actively constricts in response to the increase in PaO2, eliminating blood flow across the ductus and completing the transition to neonatal circulation.
- During hypoxemia, the pulmonary vasculature vasoconstricts raising pulmonary vascular resistance, thus relaxing the vasculature of the ductus arteriosus and allowing blood flow from pulmonary artery to the descending aorta.

Patent Ductus Arteriosis (PDA)
- Allows blood from right ventricle (RV) and pulmonary arterial system to flow into descending aorta (thereby bypassing the pulmonary circulation). The PDA closes quickly after birth (hours to several days).
- Physical Exam Findings:
  o Holosystolic murmur heard at 2nd or 3rd intercostal space at left sternal boarder
  o Bounding peripheral pulses
  o Hypotension- decreased mean arterial pressure (MAP)
  o Respiratory deterioration- Tachypnea
- Diagnosis
  o ECHO (Gold Standard)
    ▪ Perform pre and post treatment of PDA
  o CXR may show cardiomegaly with increased pulmonary vascularity in large shunts.
- Treatment
  o Indomethacin or Ibuprofen
    ▪ Monitor urine output (UOP) during treatment
  o Respiratory support (may need to be intubated)
  o Fluid restriction
  o A Hct > 40% will decrease L to R shunting

Coarctation of the Aorta
- Localized constriction of the aorta that usually occurs at the junction of the transverse aortic arch and the descending
aorta in the vicinity of the ductus arteriosus. Coarctation can occur anywhere in the aorta from above the aortic valve to the abdominal aorta.

- **Diagnosis**
  - Blood pressures in all four extremities.
  - SBP > 15mmHg in upper extremities vs lower
  - Typically no murmur on exam

- **Treatment**
  - Keep the PDA open with prostaglandin (PG).
  - Stat Cardiology consult
  - Closely monitor pulses, and any signs of CHF such as increased work of breathing (WOB), UOP, and HTN

**Transposition of the Great Arteries**

- Most common cyanotic congenital heart lesion in the newborn period; cyanosis will present within hours after birth
- Aorta arises from the morphologic right ventricle and the pulmonary artery arises from the morphologic left ventricle which means that the two circulations are in parallel (ie independent)
  - Aorta is positioned anterior and to the right of the pulmonary artery
  - Symptoms on presentation will be tachypnea, tachycardia and a second heart sound that is greater in intensity
    - Murmur not always heard unless another lesion is present
  - EKG finding will be RVH (non-specific)
  - Diagnosis
    - CXR: increased pulmonary vascular markings and a narrow mediastinal shadow secondary to a small thymus
      - “egg on side” or “apple on a string”
      - Echocardiogram
  - Treatment
    - Keep the PDA open with prostaglandin E1 (PG).
    - Metabolic acidosis should be corrected
    - Mechanical ventilation may be necessary if pulmonary edema develops in concert with severe hypoxemia
    - Stat Cardiology consult

**Tetralogy of Fallot**

- Common cyanotic heart lesion, 7-10% of congenital heart disease
- Comprised of
  - RV outflow tract obstruction
  - VSD
  - Overriding aorta (detropoitioning)
  - RVH

![Diagram of Tetralogy of Fallot]
Cyanotic due to systemic venous blood shunting across the VSD into the aorta

Tet spell - acute reduction in pulmonary BF, a drop in systemic afterload, and worsening R-L shunt. Squatting allow increase systemic arterial resistance, increase pulmonary blood flow

Murmur - systolic murmur at middle or LLSB (RV outflow obstruction)

- CXR- "boot shaped heart"
- EKG- RAD, RVH
- Presents
  - Infants with mild degrees of right ventricular outflow obstruction may initially be seen with heart failure caused by a ventricular-level left-to-right shunt.
  - Often, cyanosis is not present at birth, but with increasing hypertrophy of the right ventricular infundibulum and patient growth, cyanosis occurs later in the 1st yr of life. It is most prominent in the mucous membranes of the lips and mouth and in the finger-nails and toenails.
  - In infants with severe degrees of right ventricular outflow obstruction, neonatal cyanosis is noted immediately. In these infants, pulmonary blood flow may be dependent on flow through the ductus arteriosus. When the ductus begins to close in the 1st few hours or days of life, severe cyanosis and circulatory collapse may occur.

Treatment
  - PGE1 helpful in neonate with pulmonary outflow obstruction to alleviate cyanosis, surg by 6-12mo,
  - The modified Blalock-Taussig shunt is currently the most common aortopulmonary shunt procedure and consists of a Gore-Tex conduit anastomosed side to side from the subclavian artery to the homolateral branch of the pulmonary artery

Mx: Hct 50-55%, give Beta Blocker until surgery
NEUROLOGY

Intraventricular Hemorrhage (IVH)
- 90% of bleeding events occur in first 72hrs of life. Most commonly seen in premature infants < 1500gm
- Etiology
  - Asphyxia, severe RDS, pneumothorax, hypoglycemia, shock, acidosis
  - Tend to occur in first few hours or days of life
- Classification
  - 0- No bleeding
  - I- Germinal matrix only. Confined to subependymal area
  - II- Hemorrhage filling 50% of lateral (as seen on image)
  - III- Ventriculomegaly and more than 50% of occlusion of lateral ventricles
  - IV- Intraventricular and perenchymal bleeding (other than germinal matrix)
- Symptoms
  - Apnea and bradycardia
  - Pale or cyanosis
  - Weak suck
  - High-pitched cry
  - Seizures
  - Swelling/bulging fontanelle
  - Anemia
- Treatment
  - Indomethacin 0.1mg/kg/dose PO q 24hr x 3days
    - Mechanism of Action: cause cerebral vasoconstriction
    - Decreases PG production which influences platelet function and improves gut/renal perfusion
  - Starting criteria
    - Good urine output (>1mg/kg/hr) with Cr <1.8 and BUN < 30
    - No evidence of active bleeding with ulcer
    - Hct >40
    - No NEC or guiac +stools
  - Repeat HUS 3 days post treatment
  - Depending on HUS, infant may need an MRI prior to discharge

Seizures
- Management:
  - Review birth and FHx
  - Obtain Chem7, ionized Ca, Mg, pH, Blood Cx, HUS, Lumbar puncture with CSF studies
  - Give phenobarbital bolus to stop seizure
INFECTIOUS DISEASES
Please see Red Book for full discussion and management of ID topics

Congenital Syphilis
- Present: snuffles, band keratitis, polymorphic skin rashes, PNA alba, perositis
- Adequate Treatment
  - Mom is adequately treated if she received 3 doses of PCN greater than 30 days before delivery PLUS demonstrates a 4-fold drop in titer, AND baby's titer cannot be higher than Mom’s.
- Inadequate treatment
  - Check serum RPR, CSF for cell count and VDRL, long bone radiographs, and CBC.
  - Start Penicillin G (50,000 U/kg/dose IV q 12hrs for 7 days then IV Q8hr for 3 days)
  - If maternal status is reactive, infant RPR lab must be drawn in 2cc red top tube

Chlamydia
- Conjunctivitis 5-14 days, PNA by 4-8wks
- Begin treatment with Erythromycin (PO/IV)

Gonorrhea
- Conjunctivitis (2-5 day of life) can cause blindness, sepsis, meningititis
- Full septic work up if mom is positive and untx
- Treatment
  - Rocephin 125mg IV/IM x 1 dose
  - Preterm/LBW Infant: 25-50mg/kg/dose IV/IM x 1 (maximum of 125mg/dose)

Hepatitis B Status
- If maternal status is unknown, infant needs to receive HBV and HBlg prior to discharge
- If maternal status is positive, infant needs to receive HBV and HBlg prior to discharge
- Note
  - Must give HBV within 12hrs of delivery
  - HBlg can be given up to a week of age or at discharge

CMV
- SGA, HSM, jaundice, cerebral calcifications, microcephaly
- Send Urine Cx
- Treatment: Acyclovir +/- CMV Ig

Herpes
- Swab eye, nasopharynx, rectum, CSF, Blood
- Treatment: Acyclovir
OPERATING NEODATA

1. You must first have a username and password setup
2. Double click on the icon that says “Neodata”.
3. With the screen opened, look at the top of the right column. Click on the drop down menu that says “practitioner” and select “newborn”. Next, click on the drop down menu that says “NICU” and select “Nsy”. Finally, click on “service” and this should pull up just the newborn nursery babies.
4. Click on Procedures and select “Clone”. This will give you a drop down menu from which to choose from. Select “Clone all patients”. This will allow you to enter patient information for that day, otherwise you will get a window asking you if you want to “continue or clone”.
5. Once you have the list in front of you. Look at the box that says “All Forms” and click on the drop down menu. Select the type of form you will be currently filling in for the day. Ex “progress note, admission, or discharge”.
6. Now look at the left column on the page and notice that depending on the type of form you have selected your list of entries will also change.
7. Click on each folder and enter the respective data (including all labs). NOTE: when entering vital signs enter ranges whenever available.
8. Make sure to go through each folder, so that you don’t forget to enter any information.
9. Most boxes within each folder contains drop down menus from which you can choose your entries. If your physical exam entry or any other entry is not found on the drop down menu, type it in. At times there will be a window that pops up and asks if you would like to override the entry. Enter “yes”.
10. When you have completed entering all of your information for the day, go to File and select Print. This will open another drop down menu from which to select the type of form you are working with. Please remember if you are doing a progress note, it must be printed out on progress note paper. Admission and discharges can be printed on plain white paper. Please place the admission papers in the “H&P” section of the binder and the discharge papers at the end of the “progress notes”.

Special Instructions for Admissions:

1. When entering an admission click on “Procedures” and look at the drop down menu. Please select “New Patient”. Here you will enter the babies’ last name, their first name (gender), MRN number, and account number.
2. Once you have entered the baby, you may proceed by choosing the “Admission Form.”
3. Again, go down the list of folders and enter all information (including labs).
4. You must enter your diagnoses. You may look at other baby’s charts to help you enter information and to see what kind of information needs to be entered.
5. You must fill in as complete as possible the “Pregnancy”, “Birth”, and “Admit” folders. This is like writing up a delivery note. Again you may look at another baby’s chart to get an idea.

Special Instructions for Discharge:

1. Click on “Procedures” and select “Discharge Form”
2. Enter your daily information. This includes a complete physical and all labs.
3. You must enter your updated assessment for the day under the respective diagnosis box.
4. Diagnoses must be consolidated and finalized before completion of the form.
5. You must enter completion of the anticipatory guidance under the folder that says “Daily”. You may have to go under “progress notes” or “all forms” in order to pull up this folder. At the bottom, where it says “supplemental notes” is where you will enter it.
6. Under diagnosis, in the middle of the page on the left hand side click on the circle that says “Final Comments”. Click on the “Add” button and it will add the entry in the box into the final comments. These comments should be as concise as possible and a synopsis of all of the assessments done during the hospital stay, including the diagnoses that have been resolved. To see the resolved diagnoses, click on the box on the right top corner that says inactive diagnosis (the same for any meds or procedures you may be inquiring about). This is what prints out in the discharge report. NOTE: You must remove all entries under “plan” if the diagnosis has been resolved.

7. Click on the “Discharge” folder and click on the gray “extract” box to the left lower hand corner.

8. Then look at right half of the screen and right click on the box that says “Discharge addendum”. Click on “Macros” and select “Disch Documentation 2”. There will be a line for you to enter the amount of time it took to prepare discharge (> 30 mins).

9. Then click on the folder that says “Summary” and once again click on the gray box in the middle of the screen. This will extract all of your information.

10. If all of your information has been entered including (length and head circumference), you may now print. Depending on the parent, you may print two copies. One to give to the parents for their records or to present to the PCP, so they may make a copy, and the other copy is for the babies’ chart.

Special Instructions for Progress Notes:

1. You must enter your diagnoses. You may look at other babies charts to help you enter information and to see what kind of information needs to be entered.

2. None of the folders should be left without information being entered into it.

3. Please enter you assessment for the day on each diagnoses listed in the box available.

4. You must add any meds that were started for the day under the respective diagnosis.

5. Make sure to keep your entries updated.
SPEECH PATHOLOGY IN THE NICU AT SHANDS JAX

Speech Referrals:

- A’s, B’s, and/or desats associated with po feeding
- Minimal interest with po offers
- Oxygen dependency when initiating nipple feeding
- Absent or weak gag reflex and/or cry
- Excessive gagging
- Inspiratory or expiratory stridor
- Oral aversions/defensiveness
- New onset of feeding difficulty
- Nasal regurgitation
- Cleft lip and/or palate
- Craniofacial anomalies

Infants born before 32 weeks gestation at birth, birth weight less than 1000 grams, and/or present IVH are the most at risk for prolonged transitions to full oral feedings (Shaker & Woida, 2007)

Speech/ Feeding Therapy in the NICU:

- Appropriateness for po feedings
- Non-nutritive suck techniques
- Oral sensory stimulation
- Nipple suggestions/modifications
- Feeding positions
- Techniques to encourage latching, coordination, and rhythmic patterns
- Patent/caregiver education and demonstration

Advancing Nippling Attempts

- Consistent at each stage x 24 hours
- Quality more important than quantity

Writing Orders for Speech & PO Feeding

- Examples:
  - ST Eval and Treat
    - re: po recs
    - re: poor po feeding
  - 1 PO Attempt a day
  - PO attempts TID and with cues

- Cue based feeding
  - 35 weeks and older
  - PO feedings offered when infant showing hunger cues