Sickle Cell Pain Protocol for BUMC

This protocol is designed for patients who present to the emergency department with pain potentially related to sickle cell crisis.

- 1. Determine if patient has medical emergency other than sickle cell crisis
 - a. Full set of vital signs including temperature and O2 saturation
 - b. Utilize appropriate diagnostics
 - c. Provide any emergent therapeutics necessary
 - d. Address pain management plan upon arrival
 - e. If other emergencies and painful conditions ruled out, proceed with sickle cell pain protocol
- 2. Include standard ED medical assessment for sickle cell crisis.
 - i. Diagnostics
 - 1. CBC, CMP, Retic Count
 - 2. CXR for any cardiovascular or respiratory complaint; EKG as needed
 - ii. If new sickle cell patient, or if something suspicious, consider hemoglobin electrophoresis to confirm diagnosis
- 3. SICKLE CELL PAIN PROTOCOL
 - a. If no vomiting, consider:
 - i. Tylenol 1000mg PO
 - ii. Ibuprofen 600 PO
 - iii. oral morphine 30mg IR
 - b. IV access
 - i. IV access may be problematic in some of these patients.
 - 1. Access mediport as needed
 - 2. Consider IN fentanyl and IM antiemetics while iv pending
 - c. NS fluid bolus for any dehydration
 - i. Then, D5 ½ NS maintenance fluid
 - d. Toradol 10 mg IV
 - e. Dilaudid 2mg in NS 50ml. Infuse over 15 minutes IV q 2 hours prn severe pain x 2 doses. ****NO IV PUSHES
 - f. Ketamine low dose infusion protocol
 - g. Antiemetics
 - i. Zofran 4 mg iv prn nausea
 - ii. Phenergan 12.5-25 mg iv dilution (no iv push)
 - iii. Compazine and Reglan may also be considered
 - h. Benadryl 25 mg PO q 4 hours prn pruritus
 - i. May consider ketamine 0.1-0.3 mg/kg iv

j. NO IV PUSH OPIATES OR PHENERGAN

- 4. If patient with intractable symptoms and unable to be discharged,
 - a. Admit patient