APPROACH TO HEME/ONC EMERGENCIES

By Kashif Khan (PGY-3)
LEARNING OBJECTIVE

• Identify commonly encountered Hematological and Oncological emergencies

• Order appropriate labs/radiological tests

• Initiate timely treatment

• Early sub-specialty consultations
WHAT IS AN ONCOLOGICAL EMERGENCY

• A clinical condition resulting from a metabolic, neurologic, cardiovascular, hematologic, and/or infectious change caused by cancer or its treatment that requires immediate intervention to prevent loss of life or quality of life.
<table>
<thead>
<tr>
<th>Classifications</th>
<th>Oncologic Emergencies</th>
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</table>
| **Metabolic**    | 1. Hypercalcemia (most common)  
                    2. **Tumor Lysis Syndrome**  
                    3. SIADH (Syndrome of Inappropriate antidiuretic syndrome) |
| **Neurologic**   | 1. **Spinal Cord Compression**  
                    2. Brain metatases/↑ ICP |
| **Cardiovascular** | 1. **Malignant Pericardial Effusion**  
                        2. **Superior Vena Cava Syndrome** |
| **Hematologic**  | 1. Hyperviscosity due to Dysproteinemia  
                        2. **Hyperleukocytosis**  
                        3. DIC (disseminated intravascular coagulation) |
| **Infectious**   | 1. **Neutropenic fever**  
                        2. Septic shock |
CASE - 1
A 60 yo M with PMH of HTN, GERD, depression, who presents with 1 month history of progressively worsening generalized fatigue, weakness, and feeling unwell.

ROS: Positive for dry cough, night sweats, 20 lb unintentional wt loss in last 3 months

Vitals: HR 110/m, otherwise stable

Being an excellent new intern, you perform a full physical and find bilateral cervical and supraclavicular lymphadenopathy and palpable hepatosplenomegaly. Remainder of the exam was unremarkable.

Labs:
CBC/diff: WBC 15k  H/h 7.9/25  Plt 75k  Diff: 85%L 10%N 5%M
RFP: Na138, K 5.6, Cl 105, Bicarb 22, BUN 35, Cr 1.9 (baseline Cr 0.8)
LDH 1205, uric acid 12.6, calcium 7.0, phosphorus 6.5, albumin 3.2
TUMOR LYSIS SYNDROME
TUMOR LYSIS SYNDROME (TLS)

TLS is the result of a massive and abrupt release of cellular contents into the bloodstream after rapid lysis of malignant cells.
TUMOR LYSIS SYNDROME (TLS)

• Seen in high grade liquid tumors like leukemia with leukocytosis, high grade lymphomas, and some solid tumors like small cell lung ca

• Clinical Features: weakness, arrhythmias, paralysis, acute renal failure, tetany, altered mental status, seizures

Diagnosis:

• Laboratory
  - ≥ 2 laboratory abnormalities OR
  - ≥ 25% change in 2 values from baseline value

• Clinical
  - Laboratory diagnosis + end organ damage

HyperKalemia
HyperUricemia
HyperPhosphatemia
HypoCalcemia
TUMOR LYSIS SYNDROME (TLS)

Treatment:

- **Aggressive IV Fluids &/or diuresis (Most important)**
- Manage electrolyte abnormalities
- Rasburicase (Check G-6PD before!)
- Allopurinol (Does not decrease uric acid)
- HD

Prevention:

**Fluids, Allopurinol, Rasburicase**
CASE - 2
A middle aged F with HTN, DM, metastatic breast ca, presents with worsening back pain x 2 weeks. Developed after lifting boxes while moving. More recently has been feeling some RLE numbness and worsening pain.

ROS: Denies fevers/chills, weakness, loss of sensation, bowel/bladder incontinence

Oncologic history: diagnosed with metastatic breast ca 3 yrs ago, last PET-CT 2 months ago showed stable/shrinking osseous mets in L scapula, multiple ribs, thoracic spine (T8), and R femur.

O/E: severe pain on palpation of mid-thoracic spine, strength and sensation intact throughout, normal reflexes, no saddle anesthesia, normal rectal tone

Labs: Unremarkable
MALIGNANT SPINAL CORD COMPRESSION
SPINAL CORD COMPRESSION
SPINAL CORD COMPRESSION

- Majority from:
  - Breast
  - Lung
  - Prostate
  - Lymphoma
  - Myeloma

- About 6-10% of patients with cancer

- Thoracic spine (up to 70%)
SPINAL CORD COMPRESSION

Presentation

- Pain – may not always be present or may be underwhelming
- Weakness
- Sensory deficits: numbness, paresthesias
- Cauda equina syndrome: saddle anesthesia, bowel/ bladder dysfunction, hyporeflexia
SPINAL CORD COMPRESSION

- Obtain a good history and neurologic exam

- MRI (CT Myelography)

- Steroids: dexamethasone 10mg IV STAT then 4mg q6

- Time is money! ortho/ neurosurgery, radiation oncology

- Pain control

- Primary determinant of the efficacy of therapy is the patient's neurologic status at time treatment
CASE - 3
75 yo M, heavy smoker, presents with 4 weeks of SOB, worsening non-productive cough, and 20 lb weight loss. Developed neck swelling last week, which worsens on bending forwards. ROS - negative

Vitals: Stable

O/E: R mid-lung crackles, no wheezing/stridor, swelling of neck and right upper extremity, distension of superficial anterior chest and neck veins, no focal neuro deficits

Labs: unremarkable
SUPERIOR VENA CAVA SYNDROME
SUPERIOR VENA CAVA SYNDROME (SVC)

- Most cases are not a true emergency
- Majority of cases are due to Lung Ca or NHL (intrathoracic malignancies)
- Dyspnea (most common). Facial edema, arm edema, distended veins, facial plethora, cough, airway etc.

**Diagnosis:**
- CT/ MRI
- Histological Dx
SUPERIOR VENA CAVA SYNDROME (SVC)

Treat underlying cancer!

Endovascular stents/ Radiotherapy

Supportive care:

- Head elevation, Diuretics
- Avoid high volume fluid infusion through upper extremities
- Anticoagulation
- Steroids
  - Severe Airway Obstruction
  - Lymphoma
CASE - 4
55Y F with AML presents with profound fatigue and 1 week of SOB. Accompanying family members report she initially complained of a headache and dizziness, and since then has started acting confused and been more somnolent.

Vitals: T:100.6, PR: 115/m, BP: 110/76, RR: 26/m, 80%RA

Pertinent exam: A&Ox1 (wrong date and location), confused, strength and sensation intact, ataxic gait, bilateral lung crackles

Labs:
**CBC:** WBC 88K with 78% blasts, H/h 10/30  Plt 3k
RFP: 140 5.0 108 24 20 1.2 100
LEUKOSTASIS

• Increased blood viscosity impedes blood flow, and local hypoxemia is worsened by high metabolic activity of cells and cytokine release

• **Symptoms: (CNS/Eyes/Lungs)**
  Pulmonary: hypoxia, interstitial/alveolar infiltrates
  Neurological: headache, dizziness, ataxia, confusion, somnolence, blurry vision

• **Management:**
  Rapid cytoreduction with chemotherapy
  Consider hydroxyurea or leukapheresis if unable to give chemo
**LEUKOSTASIS**

WBC counts (X 10⁹/L) as indication for leukapheresis in hyperleukocytosis

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<thead>
<tr>
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<th>Symptomatic</th>
<th>Asymptomatic</th>
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<tbody>
<tr>
<td>AML</td>
<td>&gt; 50 000</td>
<td>&gt; 100 000</td>
</tr>
<tr>
<td>ALL</td>
<td>&gt; 150 000</td>
<td>&gt; 300 000</td>
</tr>
<tr>
<td>CML</td>
<td>&gt; 150 000</td>
<td>No</td>
</tr>
<tr>
<td>CLL</td>
<td>&gt; 500 000</td>
<td>No</td>
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<tr>
<td>APL</td>
<td>No</td>
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CASE - 5
A 70 yo M with metastatic colon ca (finished cycle #3 FOLFOX ten days ago), who presents with fevers for 2 days at home. He states he checked his temperature at home and it has ranged from 99 to 102 °F.

ROS: Unremarkable

Vitals: T: 38.7°C, PR: 105/m, BP: 115/75, RR: 16/m, SP02 97% RA

Exam: Mild oral mucositis, Mediport c/d/i, lungs clear, abdominal exam benign, otherwise no focal findings

Labs:
CBC/diff: **WBC 2.0k** H/h 8.5/28 Plt 85k Diff: 68%L **20%N** 9%M 3%E
RFP: WNL
FEBRILE NEUTROPENIA
Infection in a neutropenic patient is an emergency

**Pathogenesis:**

- Breeches in host defenses (breakdown of mucosal barriers)
- Immune system suppression
- Majority of cases of neutropenic fever thought to be caused by bloodstream seeding from GI tract flora

Infectious source: 30%.
FEBRILE NEUTROPENIA

Diagnosis:

ANC < 500 or ANC < 1000 with expected nadir < 500 over next 48 hours

+ T: 38°C for > 1 hour or T > 38.3°C once

Next Steps:

• Is patient HDS? Stable for floor?

• Examine patient: any localizing symptoms? Any role for imaging?

• Cultures STAT (2 sets bld cx peripheral, culture from lines or ports, sputum or stool cx/C Diff or wound cx as indicated), UA, UCx, CXR

• Antibiotics (30-60min)
# FEBRILE NEUTROPENIA

<table>
<thead>
<tr>
<th>Clinical Scenario</th>
<th>Medication</th>
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<tr>
<td>Clinically Stable</td>
<td>Piperacillin/tazobactam 4.5gm Q6h</td>
</tr>
<tr>
<td></td>
<td>Penicillin Allergy</td>
</tr>
<tr>
<td></td>
<td>Aztreonam 2gm IV Q8 + Vanc</td>
</tr>
<tr>
<td>• Suspect cath-related infection OR</td>
<td>Add Vancomycin for empiric regimen</td>
</tr>
<tr>
<td>• Suspected S&amp;S infection OR</td>
<td></td>
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<tr>
<td>• Colonization with MRSA or Penicillin R</td>
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<tr>
<td>pneumococci OR</td>
<td></td>
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<tr>
<td>• Hemodynamic instability OR</td>
<td></td>
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<tr>
<td>• G+ve organism in Blood culture</td>
<td></td>
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<tr>
<td>• If G-ve resistance suspected</td>
<td>Add amikacin 15mg/kg once to empiric regimen</td>
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FEBRILE NEUTROPENIA – LOW RISK PATIENTS

- Certain low-risk patients can be treated at home with PO antibiotics (typically Ciprofloxacin + Augmentin) after initial IV dose and brief observation

- IDSA: anticipated neutropenia ≤ 7 d, clinically stable, ANC > 100, and no medical comorbidities

  - ASCO: MASCC score ≥ 21

<table>
<thead>
<tr>
<th>CHARACTERISTIC</th>
<th>WEIGHT</th>
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<tbody>
<tr>
<td>Burden of febrile neutropenia with no or mild Symptoms^1</td>
<td>5</td>
</tr>
<tr>
<td>No hypotension (systolic BP &gt; 90 mm Hg)</td>
<td>5</td>
</tr>
<tr>
<td>No chronic obstructive pulmonary disease^2</td>
<td>4</td>
</tr>
<tr>
<td>Solid tumor or hematological malignancy with no previous fungal infection^3</td>
<td>4</td>
</tr>
<tr>
<td>No dehydration requiring parenteral fluids</td>
<td>3</td>
</tr>
<tr>
<td>Burden of febrile neutropenia with moderate Symptoms^4</td>
<td>3</td>
</tr>
<tr>
<td>Outpatient status</td>
<td>3</td>
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<tr>
<td>Age &lt;60 years</td>
<td>2</td>
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MALIGNANT PERICARDIAL EFFUSION
Enlarged cardiac silhouette
“Water Bottle” sign

Beat to beat variation in QRS amplitude
MALIGNANT PERICARDIAL EFFUSION

• Can be related to cancer OR chemo/RT/infection/autoimmune

• Clinical manifestations
  Dyspnea, cough, Chest pain, orthopnea, palpitations.
  Exam findings: Beck’s triad (JVD, hypotension, decreased heart sounds), narrow pulse pressure, pulsus paradoxus

Treatment:

• Small/moderate effusions are usually asymptomatic and do not require urgent treatment
• Acute management: drainage with pericardiocentesis
• Prevention of re-accumulation: drainage catheter, pericardial window
• Treat underlying cancer
ACUTE CHEST SYNDROME
ACUTE CHEST SYNDROME

• Vaso-occlusive crises of pulmonary vasculature in patients with sickle cell anemia.

• Leading cause of death in SCD.

• New radio-density on CXR AND any one (T >38.5 °C, >2% drop in SpO₂, CP, cough, wheezing, rales, tachypnea)

• Maintain high suspicion, as some may develop ACS within 48-72 hours after initial pain episode!
ACUTE CHEST SYNDROME

Don’t Forget other D/D

Treatment:
• T&S, adequate pain control, IV access, fluids, oxygen, incentive spirometry, antibiotics, VTE prophylaxis, hematology consult.

• Consider simple Vs exchange transfusion & MICU transfer

• Can use simple transfusion to bridge to exchange transfusion while waiting for MICU bed (does not remove HgbS)
- Primary Oncologist?
- Date of last chemo? (Check on EMR IV chemo)
- What was their last chemo? (know your acronyms)
- Did they get any medications with chemo? (G-CSF)
- What is their previous oncologic course?
- Access for Chemo? (mediport, PICC?)
- Sickle cell crises: check care path in portal and OARRS
- Inform primary oncologist of patient’s admission
REFERENCES


• Symposium on neoplastic hematology and medical oncology - Emergencies in Hematology and Oncology. Mayo Clinic 2017

THANK YOU