HODGKIN'S LYMPHOMA

Background

- 1. Definitions:
 - Hodgkin's Lymphoma is a malignancy of lymphatic cell line origin
- 2. General Information
 - Hodgkin's Lymphoma
 - Characterized by Reed-Sternberg cells¹
 - Preapoptotic, germinal center B cell that has lost B-cell phenotype
 - "master regulator" of inflammatory process
 - Types²
 - Classical
 - Nodular sclerosing
 - Lymphocyte-rich
 - o Mixed-cellularity
 - Lymphocyte-depleted
 - Nodular lymphocyte-predominant^{3,4}
 - o Uncommon
 - Low grade monoclonal B-cell lymphoma
 - Can transform to diffuse, large B-cell lymphoma

Pathophysiology

- 1. Pathology of Disease
 - Genetic modifications during B-cell development in the bone marrow lead to DNA alterations that may become pathologic
- 2. Incidence, Prevalence (U.S.)⁵
 - Incidence: 3.1/100,000 men, 2.6/100,000 women
 - Bimodal incidence pattern⁴
 - Peak between ages 15-34
 - Peak in ages above 60
 - ~ 8800 new cases diagnosed annually
 - Prevalence: ~ 167,000 (2008)
- 3. Risk Factors⁶
 - Associated factors include familial, viral disease and immune suppression
- 4. Morbidity / Mortality
 - Morbidity
 - Generally treatment-related (see below)
 - Mortality
 - 5-year survival rate greater than 90%³

Diagnostics^{4,6}

- 1. History
 - "B symptoms"
 - Unexplained fever
 - Night sweats
 - Recent weight loss

- o Adenopathy
- 2. Physical Examination
 - Painless lymphadenopathy
 - Freely moveable with rubbery consistency
 - Hodgkin's
 - Predictable contiguous spread
 - Splenic involvement ~25% of the time
- 3. Diagnostic Testing
 - o Biopsy
 - Large surgical specimen
 - Excisional lymph node biopsy
 - Inguinal, axillary nodes generally avoided due to increase likelihood of revealing "reactive changes"
 - Fine Needle Aspiration unreliable⁷
 - Bone marrow aspiration
 - Unlikely to be positive in absence of B symptoms or subnormal blood cell line counts (under 0.5%)
- 4. Laboratory evaluation
 - Obligatory
 - CBC, ESR, glucose level, alkaline phosphatase, lactate dehydrogenase (LDH,) liver enzymes, albumin, TSH
 - o Compulsory
 - Hepatitis B, hepatitis C, HIV screening
- 5. Diagnostic imaging
 - o Mandatory
 - Chest x-ray, chest and abdominal CT scan
 - Optional
 - PET scan can detect disease not seen with other imaging modalities
- 6. Other studies
 - o Mandatory
 - Cardiac function test
 - Pulmonary function test
 - Optional
 - ENT consult
 - Reproductive consult
- 7. Diagnostic Criteria
 - Lymph node or extranodal tissue biopsy showing "Reed-Sternberg" cells within appropriate cellular environment for diagnosis
 - Staging (based on involved sites, lymph nodes affected on one or both sides of diaphragm, bulky disease, contiguous or disseminated extranodal involvement, presence of "B" symptoms)
 - Stage I
 - Single lymph node region or one extranodal site
 - Stage II
 - Two or more lymph node regions, same side of diaphragm (II)
 - Local extralymphatic extension plus one or more lymph node regions on same side of diaphragm (IIE)

- Stage III
 - Lymph node regions, both side of diaphragm (III)
 - If accompanied by local extralymphatic extension (IIIE)
- Stage IV
 - Diffuse involvement of one or more extralymphatic organs or sites
- Subclassification
 - A absence of "B" symptoms
 - B Presence of at least one of the following:
 - 1. Wt loss (> 10% from baseline 6 months prior to staging)
 - 2. Recurrent fever (> 100° F)
 - 3. Recurrent night sweats

Differential Diagnosis

- 1. Key Differential Diagnoses
 - Malignancies: Leukemias, Metastases of Unknown Primary
 - Infectious Disease: Cat Scratch Disease, Cytomegalovirus, HIV, Mononucleosis
 - Miscellaneous: Sarcoidosis, Kawasaki's Disease
- 2. Extensive Differential Diagnoses
 - Medications: Allopurinol, Atenolol, Captopril, Carbamazepine, Hydralazine, Penicillins, Phenytoin, Primidone, Quinidine, Trimethoprim/Sulfamethoxazole, Sulindac
 - Serum Sickness

Therapeutics⁶

- 1. Classic Hodgkin's Lymphoma
 - Limited Stage
 - Brief Chemotherapy followed by Radiotherapy
 - 2 or 3 cycles of ABVD (Adriamycin, Bleomycin, Vincristine, Dacarbazine) followed by 30 Gy involved-field radiotherapy (IF-RT)
 - o Intermediate Stage
 - 4 cycles of ABVD followed by 30 Gy IF-RT
 - Other more intensive regimens being evaluated for healthy patients under age 60
 - Advanced Stage
 - Chemotherapy alone
 - 6 to 8 cycles of ABVD (SOR:C)⁴ or 8 cycles of BAECOPP escalated(Bleomycin, Etoposide, Adriamycin, Cyclophosphamide, Vincristine, Procarbazine, Prednisone, G-CSF)
 - BAECOPP escalated should not be used in patients over 60 years old due to increased toxicity
 - Radiotherapy only for large residual masses
 - Relapsed Classic Hodgkin's Lymphoma
 - High-dose chemotherapy followed by Autologous Stem Cell Transplantation (ASCT)

- Refractory or Relapsed Classical Hodgkin's
 - Primary cause of death if diagnosed within 10-15 years after initial treatment⁴
 - High dose chemotherapy followed by autologous stem cell transplant treatment of choice
 - Salvage regimens given to reduce tumor size and mobilize stem cells prior to high-dose chemotherapy and autologous stem cell transplant
 - No treatment standard for patients relapsing after high-dose chemotherapy and stem cell transplant
- 2. Nodular Lymphocyte Predominant Hodgkin's Lymphoma (NLPHL)
 - Stage IA without risk factors
 - 30 Gy IF-RT alone
 - o Other Stages
 - Treated same as Classic Hodgkin's Lymphoma with exception of stage IA
 - Relapsed Nodular Lymphocyte-predominant Hodgkin's Lymphoma
 - Obtain biopsy in those suspected of relapse (transformation to aggressive Non-Hodgkin's Lymphoma needs to be excluded)
 - Localized relapses of NLPHL can be treated with rituximab alone
 - Advanced relapses require aggressive salvage therapy + rituximab
- 3. Hodgkin's Lymphoma Response Evaluation
 - Early/Intermediate Stage: initiated at completion of chemotherapy prior to radiotherapy
 - Advanced Stage: initiated after four cycles of chemotherapy
 - Final staging after completion of therapy
 - Physical exam, lab analysis (CBC with diff, ESR and blood chemistries) and CT scans mandatory (controversial)
 - Early Stage recurrence rate 10% + 80% of recurrences diagnosed by patient or examining provider⁸
 - Advanced stage: FDG-Positron Emission Tomography (PET) helps identify poor-risk individuals
 - Therapy should not be based on results until further clinical trials conducted
 - False-positive PET scan must be excluded
 - Response after relapse is measured in same fashion as initial response

Follow-Up

- 1. Return to Office
 - Follow up protocols vary:
 - European Society for Medical Oncology (ESMO) recommends⁶:
 - Every 3 months for 1st 6 months, then every 6 months until 4th year, yearly thereafter:
 - History, Physical Exam, Laboratory analysis (CBC with differential, ESR, Chemistries), Chest imaging (CT or CXR) (SOR:C)⁴
 - \circ TSH after neck irradiation (1,2, and 5 years out) (SOR:C)⁴

- Testosterone/estrogen in younger patients receiving intense chemotherapy
- CT and other abnormal radiographic tests must be repeated to confirm remission
- Most relapses occur within 5 years of initial treatment
 - 55% suspected on symptom history, 23% found on CXR/CT, 14-18% detected by physical exam⁹
 - Role of Position Emission Tomography in detecting relapse is unclear
 - Usually found by complaint of a new lump⁴
 - Fever, weight loss, night sweats, cough and pain may also be presentation of relapse
- Long-term complications^{4,8}
 - Breast cancer associated with radiation therapy
 - most at risk if radiation received under 35 years of age or if high dose radiation used
 - Long latency period of 10-15 years prior to development
 - Mammogram effective screening tool
 - American Cancer Society recommends yearly Breast MRI in addition to mammography for patients with chest irradiation prior to age 30
 - Premature menopause has protective effect
 - Lung cancer associated with radiation and chemotherapy
 - Older alkylating agents cited as causative in dose dependent manner (procarbazine, mechlorethamine, dacarbazine)
 - Smoking cessation should be emphasized at time of diagnosis
 - Median survival of lung cancer after Hodgkin's Lymphoma patients is under 1 year
 - Other malignancies due to chemotherapy regimens:
 - Acute Myeloid Leukemia 1-3% incidence, usually within 10 years of treatment
 - Older Alkylating agents main risk
 - Wide field radiation (uncommon today)
 - ABVD regimen lowers risk
 - Myelodysplatic Syndromes
 - Cardiovascular complications
 - Coronary Artery Disease (CAD) primary contributor to excess cardiac mortality after therapy
 - Main risk factor mediastinal radiation
 - Doxorubicin also cardiotoxic
 - Cardiomyopathy Doxorubicin
 - Female, Cumulative high dose, Younger age at exposure and increased time from exposure
 - Pericardial disease
 - Valvular abnormalities
 - Conduction disturbances

- Reproductive problems (increase with age at treatment)
 - Ovarian failure higher risk with abdominopelvic radiation therapy and older alkylating agents
 - Infertility
 - Men spermatogenesis affected by alkylating agents and cyclophosphamide
 - ABVD regimen has lowered male infertility
 - Radiation contributes to damage
 - Consider cryopreservation of sperm prior to treatment
- Other Late Complication of Therapy
 - Thyroid abnormalities primarily hypothyroidism
 - Irradiation to neck and upper mediastinum increases risk
 - Risk greatest first 5 years after therapy
 - Can occur 20 years after therapy
 - Evaluate thyroid function 1,2, and 5 years after treatment in those status post neck irradiation
 - Dental problems primarily related to oropharyngeal radiation induced decrease in salivation
 - Preventive dentistry recommended
 - Pulmonary dysfunction
 - Radiation dose and lung volume exposed dependent
 - Bleomycin associated pulmonary fibrosis (common after doses greater than 200 Units per m²)
 - Fatigue related to cardiac, pulmonary, thyroid disease

Prognosis

- 1. Hodgkin's Lymphoma⁶
 - 80-90% of treated patients secure permanent remission and should be considered cured
 - \circ 15% of patients with early disease will relapse¹
 - \circ 1/3 of patients with advance disease will relapse¹

Prevention

1. Post Therapy - annual influenza and appropriate interval pneumococcal immunizations recommended

Patient Education

- 1. The Leukemia and Lymphoma Society
 - o http://www.lls.org/diseaseinformation/lymphoma/
- 2. National Cancer Institute
 - o http://www.cancer.gov/cancertopics/types/hodgkin

References

- 1. DeVita VT, Costa J. Toward a Personalized Treatment of Hodgkin's Disease (Editorial). NEJM 2010; 362(10): 942-3.
- 2. Lee AI, LaCasce AS. Nodular Lymphocyte Predominant Hodgkin Lymphoma. The Oncologist 2009;14:739-751.
- 3. Armitage, JO. Early-Stage Hodgkin's Lymphoma. NEJM 2010; 363(7): 653-62.
- 4. Glass C. Role of the Primary Care Physician in Hodgkin Lymphoma. Amer Fam Phys 2008;78(5):615-22, 625-6.
- 5. National Cancer Institute Surveillance Epidemiology and End Results. <u>http://seer.cancer.gov/statfacts/html/hodg.html</u>. accessed January 26, 2012.
- 6. Eichenauer DA, Engert A, Dreyling M. Hodgkin's Lymphoma: ESMO Clinical Practice Guidelines for Diagnosis, Treatment and Follow-up. Ann Onc 2011; 22(Supp 6): vi55-vi58.
- 7. Ansell SM, Armitage JO. Management of Hodgkin Lymphoma. Mayo Clnc Proc 2006;81(3):419-26.
- 8. Juweid ME, Vose JM. Imaging in Early-Stage Hodgkin's Lymphoma. [editorial] NEJM 2010; 362(10): 962.
- 9. Ng A, et al. Expert Panel on Radiation Oncology-Hodgkin's Lymphoma: Appropriateness Criteria follow-up of Hodgkin's Lymphoma [online publication] American College of Radiology 2010; 10 pages.

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