



Myelodysplastic Syndrome: Let's build a definition

Myelo – bone marrow







Syndrome

Collection of signs and symptoms associated together

Myelodysplastic Syndrome

- Heterogeneous group of clonal hematopoietic stem cell disorders characterized by ineffective hematopoiesis, progressive pancytopenia, morphologic abnormalities and propensity to transform to AML
- Dysplastic hematopoiesis
 - Impaired differentiation
 - Accumulation of blasts
 - Hypercellular bone marrow in ~90%
- Peripheral cytopenias
- Risk of progression to AML in 25-35%
- Abnormal bone marrow cytogenetics in ~50%
- Cazzola M, Malcovati L. *N Engl J Med.* 2005;352:536-538 Heaney ML, Golde DW. *N Engl J Med.* 1999;340:1649-1660 Hofmann W-K, et al. *Hematol J.* 2004;5:1-8
- MDS Foundation Resource Center. Available at: http://www.mdsresourcecenter.org/



| Factor | Evidence |
|-----------------------------------|--------------|
| Increasing Age | ++++ |
| Male Gender | ╋ |
| Chemotherapy Agents/XRT | +++ + |
| Benzene/Solvents | +++ |
| Smoking | ++ |
| Pesticides/Herbicides/Fertilizers | ++ |
| Ionizing Radiation | + |
| Hair Dye | + |

Bone Marrow Failure: Signs and Symptoms

<u>Anemia</u>

- Fatigue, pallor
- Shortness of breath, decreased exercise tolerance
- Exacerbation of heart failure, angina

Neutropenia

- Active infection (bronchitis, sinusitis, pneumonia, etc.)
- Risk of infections

Thrombocytopenia

- Petechiae, bruising, bleeding
- Risk of bleeding

Performing a bone marrow aspiration





Required Initial Evaluation NCCN (2013) Guidelines

- H&P
- CBC with diff, platelet count, & retic
- Examination of peripheral blood smear
- BM aspirate with iron stain and cytogenetics
- BM biopsy
- Baseline serum EPO level prior to RBC transfusion
- RBC folate and serum B12
- Serum iron/TIBC/ferritin
- Check thyroid function
- Documentation of transfusion history

NCCN Practice Guidelines in Oncology: Myelodysplastic Syndrome v.2.2013

IPSS-R Prognostic Score Values

| Prognostic variable | 0 | 0.5 | 1 | 1.5 | 2 | 3 | 4 |
|------------------------|--------------|-----------|----------|-----|--------------|------|-----------|
| Cytogenetics | Very Good | | Good | | Intermediate | Poor | Very poor |
| BM Blast % | ≤2 | | >2 - <5% | | 5 – 10% | >10% | |
| Hemoglobin | ≥10 | | 8 - <10 | <8 | | | |
| Platelets | ≥100 | 50 - <100 | <50 | | | | |
| ANC | ≥0.8 | <0.8 | | | | | |
| | | | | | | | |



How is MDS treated?

- Supportive Care (transfusions, antibiotics, growth factors, ? iron chelation)
- Hypomethylating agents (azacitidine, decitabine)
- Immunomodulators (e.g. lenalidomide)
- Hematopoietic stem cell transplantation
- Novel Agents/Clinical trials





- If possible, cure me
- If you can't cure me, at least make me live longer and feel better
- If you can't make me live longer, at least make me feel better
- If you can't even make me feel better, then get me another doctor and go back to school...







- Know your risk group
- Know your treatment options, including whether you should be considering stem cell transplant and/or clinical trials
- Know what results are reasonable to expect from your treatment
- Know the potential side effects
- Know about resources (e.g. the LLS)
- Include your caregiver in treatment planning
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