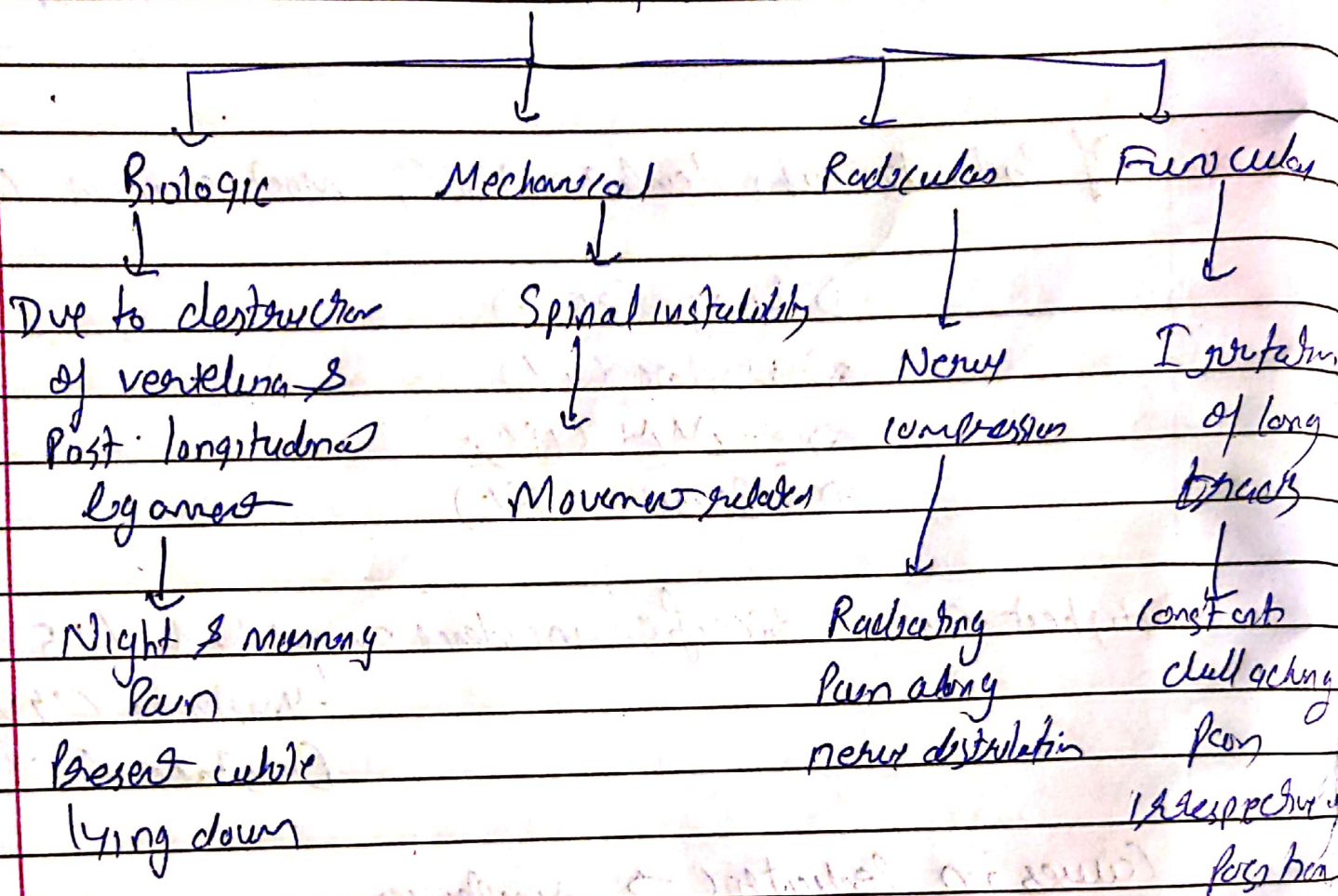


① Pain → Most common feature



- Usually, it exists ~ 8 weeks prior to Dx of MSCC.

- Starts with bony vertebrae, and progresses rapidly once epidural venous plexus is infiltrated.

★ If pain disappears without treatment, it indicates **ADVANCED** destruction of nerve roots causing permanent sensory damage.

- Usually pain is worse at night due to diurnal release of endogenous steroids.

① Motor → Motor deficit, as weakness & spasticity is the earliest sign

- Early symptoms - leg heaviness & difficulty in climbing stairs
- Most pt have paraparesis as level of compression is at thoracic spine.
- If higher, it can lead to quadriplegia, resp. failure.
- If extends to lumbar spine, it can cause conus medullaris Syndrome -
 - distal LL weakness
 - Saddle paresthesia
 - Bladder/Bowel overflow.
- Horner's Syndrome (PAMELA) suggests transforaminal progression at cervicothoracic junction & invasion of stellate ganglion.

③ Sensory → Sensory loss is usually ascending in nature

- It is typically at 1-5 levels below actual level of cord damage
- Lhermitte Phenomenon can be seen in cervical spine mets.

Blockage & Bowel retention are namely the sole features & occur late EXCEPT

- Compression at level of conus
- Sacral mets.

Diagnosis - Ix of choice $\rightarrow$ MRI

- Sagittal T1 or STIR sequences of entire spine for screening
- Sagittal T2 weighted sequences done to look for cord compression & lesion within cord itself.

- Sensitivity of MRI - 93-100%.

Specificity $\rightarrow$ 97-100%.

- Whole spine screening should be done to look for assessing other sites as well.

MRI based score $\rightarrow$ ~~Bilsky~~ ^{Bilsky} MSIC score

0 $\rightarrow$ Bone only disease

1 $\rightarrow$ Epidural extension without contact with spinal cord or just spinal cord abutment but no displacement

1a $\rightarrow$ Epidural tumor without thecal sac compression

1b $\rightarrow$ Thecal sac compression but no cord contact

1c $\rightarrow$ Thecal sac compression $\pm$ cord contact

2 → Displaces or compresses Spinal cord without circumferential tumor extension or obliteration of CSF space
(hence, CSF visible around cord)

3 → Cord compression with CSF obliteration.

Grade 2, 3 → High grade.

CT myelography → Done in patients who cannot undergo MRI.

Other use of myelography →

- In patients with indwelling spinal orthopedic hardware

- In pts undergoing treatment planning for radiosurgery or RT.

Management

* If there is no histological evidence of malignancy.
Do NOT start steroid before taking biopsy.

* Steroids → Once biopsy taken, start patient on steroids.

Vecht CT et al study → low dose dexamethasone (10mg) is as effective as high dose (100mg)

Dose → 10mg IV q6h 16mg orally in divided doses

Multiple Prognostic scores to estimate prognosis

1) Tokuhashi Score & Revised Tokuhashi Score

General condition

Number of extraspinal foci

Number of mets in ventral body

mets to internal organs (Resectable or not)

Primary site

Palsy (According to Frankel A, B, C, D)

Highest Score = 15

2) Tominaga score

3) Modified Bauer score - Most favoured score

(A) Score Calculation

Points	Positive Prognostic Factors
1	No visceral mets
1	No lung cones
1	Primary site - Breast, Kidney, Lymphoma (B) < (L) or myeloma
1	Solitary met

(B) Score Interpretation

0-1	→ Supportive care, no surgery
2	→ Short-term palliation, Dorsal Surgery
3-4	→ middle-term control, ventrodorsal Surgery

4) Oswestry Spinal Risk index

5) Spinal instability Neoplastic score (SINS)

- ① Spine location
- ② Pain relief with recumbence or pain with movement
- ③ Bone lesion quality (lytic (mixed) blastic)
- ④ Radiographic Spinal alignment
- ⑤ Vertebral body collapse ($> / < 50\%$)
- ⑥ Posterolateral involvement of ~~vertebrae~~ spinal elements

Score

0-6 - Spine stable

7-12 - Indeterminate - Urgent Surgical consultation

13-18 - Instability - Surgery.

* So, basically, in patient with poor life expectancy, treatment should be along lines of best supportive care or palliative RT

If life expectancy > 2 months, then some definitive management can be planned

MNOP Algorithm

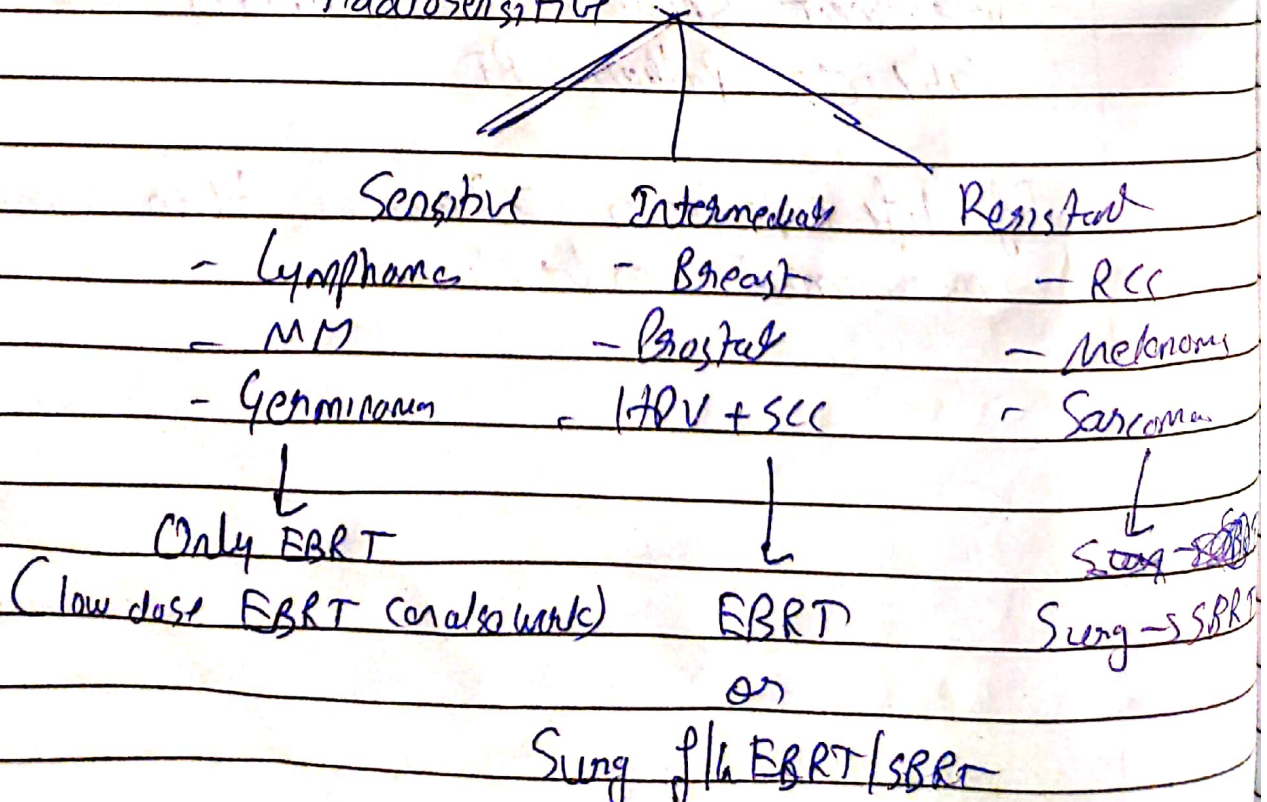
(1) Mechanical $\rightarrow$ If spine is unstable $\rightarrow$ fix it.

Primary RT should NOT be given upfront

(2) Neurologic $\rightarrow$ Grade 0 - no intervention
Grade 1-3 - Intervention needed based on histology & radiosensitivity.

pt with acute decompensation < 72hrs will benefit more from decompression.

(3) Oncologic $\rightarrow$ Whether tumor is radioresistant or radiosensitive



④ Preferred treatment

- ① Surgery - Role of surgery is in
- Decompression
 - Obtaining biopsy
 - Removal of peripheral disease to allow SBRT
 - Local control when RT can't be given.

- The best outcomes are shown with circumferential decompression.

Most versatile approach is Transpedicular which allows for 360° decompression in all flexion/extension.

- ② EBRT - Most common dose - 8 Gy 1#
- 20 Gy 5#
 - 30 Gy 10#

SCORE 1 study → Multifraction EBRT (30-40 Gy) improves 14% local control

BUT in pt with very poor P.S., there is no benefit of local control.

SCORE 2 study → 30 Gy/10# vs 20 Gy/5# in pt of poor P.S. Noninferiority trial.

Further trial built on SCORE 2 compared 20 Gy/5# with 8 Gy 1# - No significant difference.

3) SBRT $\rightarrow$ Stereotactic Body RT.
Delivering high dose to tumor while sparing adjacent tissues.

Much more local control, even for radioresistant tumors.

Very little role left for it, as it can only be given in grade 0.
It needs 1-3mm space between cord & tumor.

However, it can be given in an adjuvant setting

4) Other therapies - Chemo for M12
- Targeted therapies
- Radionuclide - Ra 223 dichloride for painful bone mets

Performance status

