

Management of increased intracranial tension

Increased intracranial tension is a common neurological problem encountered in cancer patients.

Causes:

- (1) Metastatic brain tumors (commonly from Ca Lung & Melanoma)
- (2) Primary brain tumors [especially those located in the posterior fossa (Medulloblastoma, Ependymoma)]
- (3) Intracranial hemorrhage
 - tumors, eg metastasis from Melanoma, Germ Cell Tumor, RCC)
 - bleeding diathesis
 - trauma
 - vascular causes, eg vasculitis, aneurysms
- (4) Dural sinus thrombosis
- (5) Vasogenic oedema (surrounding tumors/ hemorrhage)
- (6) Medications (eg Vitamin A, ATRA, Tetracyclines, Sulfonamides)
- (7) Cerebral Infections (Toxoplasmosis, HSV, Candidiasis)
- (8) Brain abscess
- (9) Leptomeningeal carcinomatosis

Workup:

- (1) History & Physical examination (neurodeficits, fever, meningism, etc)
- (2) Complete blood count & coagulation profile
- (3) Serum Urea, Creatinine, Electrolytes
- (4) Liver function test
- (5) CT scan/ MRI scan of brain
- (6) Fundoscopy
- (7) Lumbar puncture

Treatment:

- (1) Rest with head and upper trunk slightly elevated
- (2) Corticosteroids-Dexamethasone 8 mg PO./IV QID-to be avoided in case of suspected primary CNS lymphoma
- (3) Osmotic diuretics-Mannitol
- (4) Hyperventilation (for rapid reduction of raised ICP)
- (5) Surgery-for obstructing lesions-resection +/- shunting
 - for hemorrhage-evacuation of clot
 - for brain abscess-drainage
- (6) Stoppage of drug-for drug-induced increase of ICP
- (7) Antimicrobials-for infections
- (8) Thrombolytics-for venous thrombosis
- (9) Transfusion of platelets/ plasma-for bleeding diathesis
- (10) Chemotherapy-for meningeal carcinomatosis & for chemosensitive metastatic tumors (germ cell tumors, SCLC)
- (11) Radiotherapy-for primary/metastatic malignant brain tumors.

Management of Malignant Spinal Cord Compression

Causes -ADULTS: (1) Vertebral metastasis (Ca Lung, Breast, Prostate, Colorectum, Multiple Myeloma)

- (2) Spinal tumors (metastasis-rare, primary-meningioma, Schwannoma)
- (3) Paravertebral tumors (Lymphoma, Neuroblastoma)

Causes-CHILDREN: (1) Neuroblastoma

- (2) Wilms' tumor
- (3) Lymphoma
- (4) Primary vertebral osteosarcoma

Initial presentation of cancer with MSCC: (20%)

- (1) Multiple Myeloma
- (2) Metastasis from SCLC
- (3) NHL

Rare in case of Ca Breast

Common sites of MSCC:

- (1) Thoracic (commonest-65-70%)
- (2) Lumbar (20%)
- (3) Cervical
- (4) Sacral

Non-malignant causes of spinal cord compression in cancer patient:

- (1) Epidural hematoma (coagulopathy)
- (2) Abscess

Differential diagnosis of spinal cord compression:

- (1) Pain-degenerative disease
- (2) Neurodeficit-transverse myelitis

Symptoms of MSCC:

- (1) Pain (commonest)
- (2) Neurodeficits-
 - Motor deficits (occurs earlier than sensory)
 - Cervical → quadriplegia/quadriparesis
 - Thoracic → paraparesis/paraplegia
 - Lumbo-sacral → cauda equine /conus medullaris syndrome (saddle anaesthesia, distal lower extremity weakness, overflow bladder and bowel incontinence)
 - Sensory deficits → hypoesthesia (usually 2-3 segments below the level of the spinal lesion)
 - Autonomic deficits are common (urinary retention, fecal incontinence).

Causes of pain in MSCC:

- (1) Stretching of vertebral periosteum, damage to posterior longitudinal ligament, damage to synovia of intervertebral joints
- (2) Compression of spinal nerve roots
- (3) Paravertebral muscle spasm

Mechanism of spinal cord compression:

- (1) Direct extension of vertebral bone metastasis into epidural space
- (2) Pathologic fracture of vertebral segment with impingement of bony fragments into epidural space
- (3) Extension from paravertebral mass into epidural space
- (4) Intradural mass lesions
- (5) Direct hematogenous metastasis to epidural space (rare)

Pathology:

- (1) Acute stage → impairment of venous drainage & vasogenic oedema
- (2) Chronic stage → necrosis & demyelination

Workup:

- (1) History & physical examination (including complete neurological examination)
- (2) Routine biochemistry
- (3) Chest X Ray
- (4) USG/ CT scan whole abdomen & pelvis
- (5) MRI scan whole spine
- (6) WBBS
- (7) FNAC from spinal lesion

Treatment:

- (1) Corticosteroids-Dexamethasone 20 mg PO/IV loading followed by 10 mg PO/IV QDS
- (2) Radiotherapy-Most cases are managed by EBRT, Usual dose is 30 Gy/10#/ 2 weeks to an area comprising the vertebra affected with 2 vertebra above and below it. Shorter regimes are also there. Dose is prescribed at 3 cm for cervical cord, 4 cm for thoracic and 5 cm for lumbar cord (alternatively the depth may be assess from the MRI).
- (3) Surgery-antero-lateral decompression. It is only indicated in certain specific instances:
 - (1) Vertebral fracture or instability
 - (2) Known radio-resistant tumor
 - (3) Recurrent lesion following EBRT
- (4) Chemotherapy-only for extremely radiosensitive tumors such as SCLC and NHL.
- (5) Bisphosphonates-used in case of metastatic Ca Breast/Prostate/Myeloma

Management of Superior Vena Cava Syndrome

It comprises the signs and symptoms caused by obstruction to blood flow (intraluminal/ luminal/ extraluminal) through the superior vena cava.

Why does it occur?

Due to the fact that the SVC is a thin-walled compliant blood vessel with scanty muscular tissue in its walls-so it is easily compressed.

Causes: (ADULTS)

Malignant (main) → Ca Lung (80%)(SCLC > NSCLC)

Lymphoma (8%) [NHL (Lymphoblastic lymphoma & DLBCL) > HD]

Mediastinal germ cell tumor

Thymoma

Metastatic mediastinal lymphadenopathy from Ca Breast

Non-malignant (10-20%) → Mediastinal fibrosis (histoplasmosis)

Thrombosis of central venous catheter

Retrosternal goiter

Aortic aneurysm

Causes: (CHILDREN) –rare

Non-malignant (70%) → Iatrogenic (post-surgery for congenital heart disease, ventriculo-atrial shunting, central venous catheterisation)

Benign granulomas

Malignant (30%) → Lymphoma is the commonest cause (70%).

Prognosis: Median survival= 9 years (benign causes)

5 mo (Lung Cancer)

Clinical features:

Symptoms → Dyspnoea, Fullness in face & neck, headache, cough, chest pain, hemoptysis

Signs → Distended veins in neck & chest wall, facial oedema & plethora, cyanosis

Workup:

- (1) History & physical examination
- (2) Complete blood count
- (3) Routine biochemistry
- (4) Chest X Ray
- (5) CECT scan thorax +/- guided FNAC
- (6) Biopsy of any suspicious lymph node
- (7) Thoracocentesis (if pleural effusion)
- (8) Sputum cytology
- (9) Bronchoscopy +/- biopsy
- (10) Mediastinoscopy
- (11) Thoracoscopy
- (12) Thoracotomy

Treatment:

- (1) Medical management → all cases. Bed rest with head end elevated + Salt-restricted diet + Diuretics + Corticosteroids + Moist oxygen inhalation
- (2) Radiotherapy → for malignant SVCO. Whole mediastinum along with primary parenchymal tumor (if present) with margin, should be treated. Dose = 16 Gy/4#/4 days.
- (3) Chemotherapy → for chemosensitive tumors like SCLC and NHL.
- (4) Surgery (bypass grafting) → in case of rapidly progressive SVCO/ refractory SVCO/ SVCO caused by retrosternal goiter or aortic aneurysm.
- (5) PTCA +/- stenting → can be done for both benign & malignant causes
- (6) Thrombolysis (urokinase/streptokinase/TPA) → for thrombosis of central venous catheter (catheter should be removed in all cases)
- (7) Anti-coagulants (Heparin followed by Warfarin) → for thrombosis of central venous catheter