



Approach to illness prevention in athletes with spinal cord injuries

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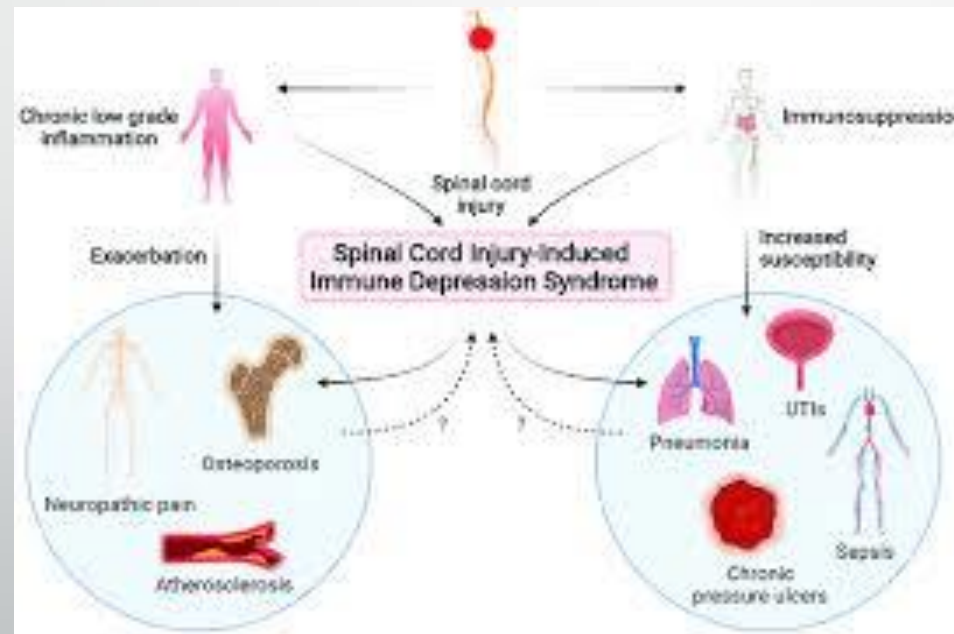
Spinal cord injury (SCI)

- Spinal cord injury (SCI) is a serious medical condition that causes functional, psychological and socioeconomic disorder.
- The goals of rehabilitation and other treatment approaches in SCI are to improve functional level, decrease secondary morbidity and enhance health-related quality of life.



Spinal cord injury (SCI)

- Prevention, early diagnosis and treatment of acute and chronic secondary complications in patients with SCI is critical for limiting these complications.



Long-term complications after SCI

- Respiratory complications
- Cardiovascular complications
- Urinary and bowel complications
- Spasticity
- Pain syndromes
- Pressure ulcers
- Osteoporosis and bone fractures

RESPIRATORY COMPLICATIONS

- Respiratory complications associated with SCI are the most important cause of morbidity and mortality in both acute and chronic stages.
- Forced vital capacity (FVC) and forced expired volume (FEV₁) were normal in patients with low-level paraplegia who had never smoked but they found that both decreased with rising SCI level more prominently in patients with tetraplegia.
- Duration of injury, smoking history, age and body mass index (BMI) effect on this complication.

RESPIRATORY COMPLICATIONS

- Significant decrease in all lung volumes with increasing BMI
- SCI patients have a high prevalence of sleep-related respiratory disorders particularly in para powerlifting athletes.(%25-%45)

RESPIRATORY COMPLICATIONS

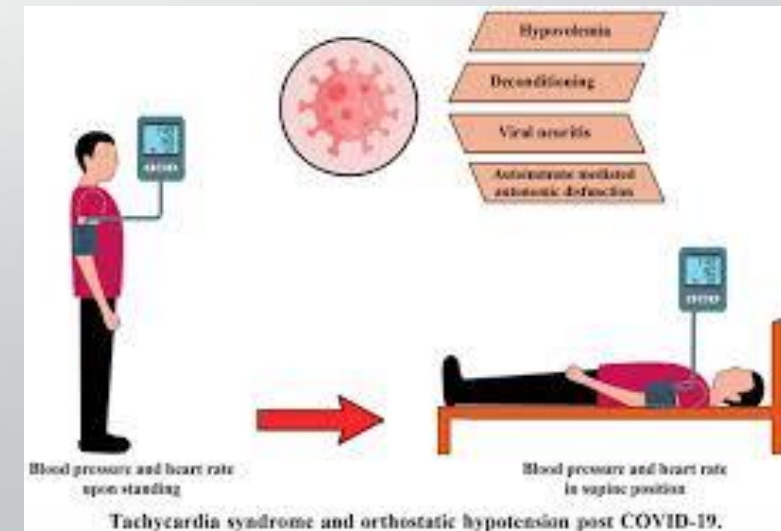
- **Management of respiratory complications include :**
- Positioning and postural changes, breathing techniques, spontaneous cough and cough assistance, suctioning, respiratory muscle training, ventilation techniques and education, vaccination agents for influenza and pneumococcal infections and pharmacological interventions.
- Respiratory muscle training improve respiratory function & improve performance(core box).

CARDIOVASCULAR COMPLICATIONS

- **Thromboembolism** and **autonomic dysreflexia(AD)**.
- orthostatic hypotension (OH)
- impaired cardiovascular reflexes
- Reduced transmission of cardiac pain
- loss of reflex cardiac acceleration
- Cardiac atrophy with tetraplegia due to loss of left ventricular mass
- pseudo-myocardial infarction

Orthostatic hypotension

- It is defined as a decrease in systolic blood pressure of 20 mmHg or more, or a reduction in diastolic blood pressure of 10 mmHg or more, when the body position changes from supine to upright after 3 min .(30-20-10)
- low level of efferent sympathetic nervous activity and the loss of reflex vasoconstriction after SCI.
- 21% and cervical injuries had the highest prevalence



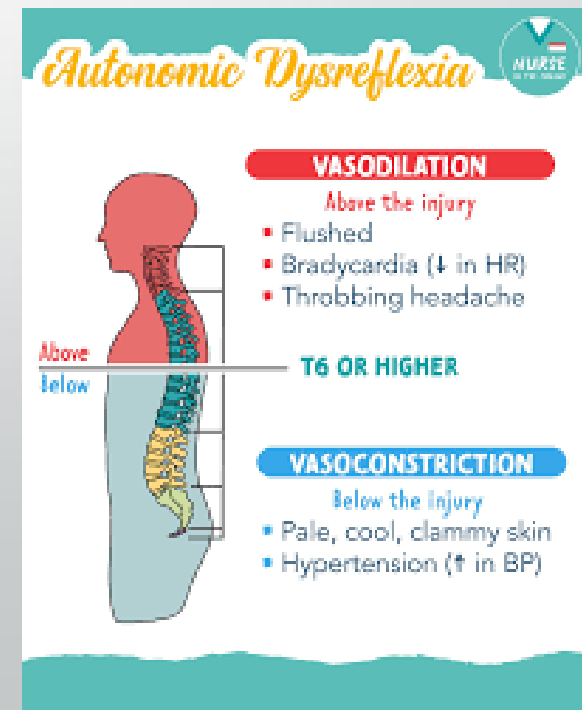
Management of OH

- Management of OH includes application of pressure stockings and abdominal binders, adequate hydration, gradual progressive daily headup tilt and administration of pharmacological agents (salt tablets, midodrine, fludrocortisone, dihydroergotamine, ephedrine or L-DOPS).

####Doping####

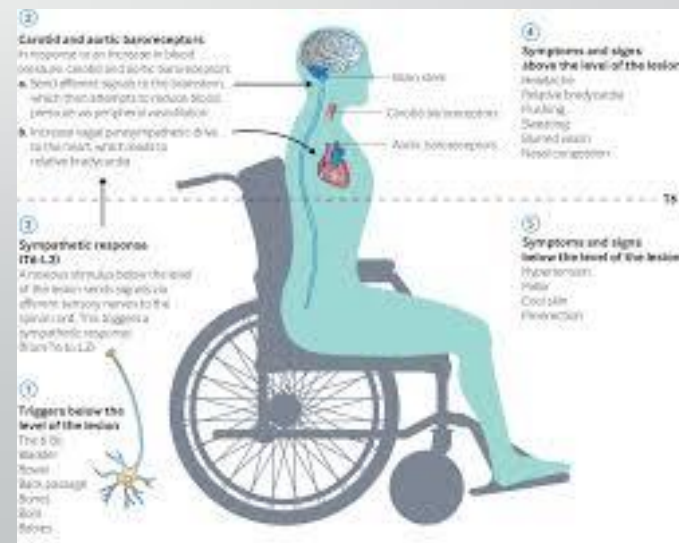
Autonomic dysreflexia

- Autonomic dysreflexia (AD) is a well-known medical emergency.
- It generally occurs in patients with SCI at levels of T6 and above.
- Incidence :19%-70%
- Initiated by a noxious stimulus entering the spinal cord below the level of injury.
- **Bladder distension** is the most common triggering factor for AD.
- The second most common triggering for AD is **bowel distension**.
- (more than in para athletes)



Autonomic dysreflexia

- **Management:**
- Prevention
- non-pharmacological
- Placing the patient in an upright position to take advantage of any orthostatic reduction in blood pressure.
- loosen tight clothing.
- Blood pressure is controlled at least every 5 min



Autonomic dysreflexia

- If arterial blood pressure is 150 mmHg or greater, pharmacological management should be initiated.
- Antihypertensive agent: Nifedipine and nitrates
- captopril, terazosin, prazosin, phenoxybenzamine, Prostaglandin E2 and Sildenafil.

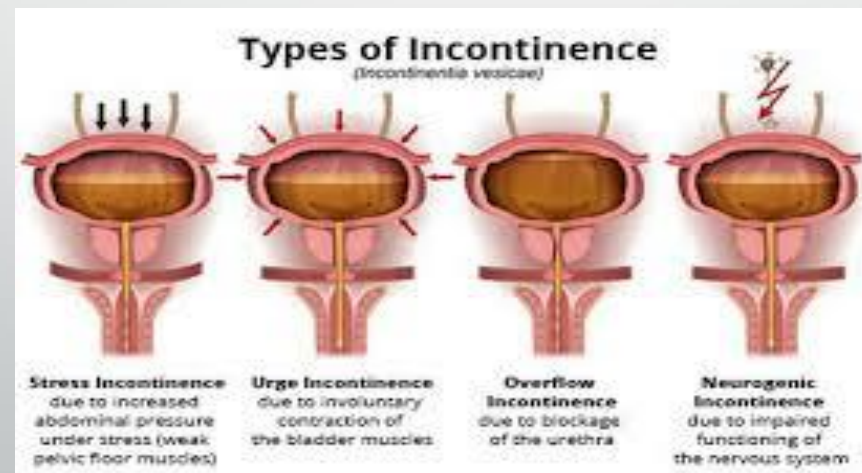
URINARY AND BOWEL COMPLICATIONS

- **Bladder dysfunction:**
- neurogenic bladder
- Decrease psychological and social well-being of the patient.
- Bladder function: the cerebral cortex, the pontine micturition center and the sacral micturition center.
- **Urodynamic evaluation** is essential to provide a precise diagnosis and treatment



Bladder dysfunction

- Treatment methods for neurogenic bladder can be categorized into two groups: therapy to facilitate bladder emptying and therapy to facilitate filling or storage of urine.
- **Clean intermittent catheterization (CIC)**
- **Permanent indwelling urethral or suprapubic catheter** (*for acute phase*)



Crede maneuver

- Don't recommended for bladder emptying in the long-term because vesicoureteral reflux, hernia, rectogenital prolapse and hemorrhoids.
- pharmacological interventions (anticholinergic medications, α -blockers, botulinum toxin)



neurogenic bowel

- **Two main types of neurogenic bowel presented as upper motor neuron (UMN) bowel syndrome and lower motor neuron (LMN) bowel syndrome.**

SPASTICITY

- Spasticity affects 70% of patients with SCI and causes considerable disability for many.
- **light to moderate spasticity** may have a **positive impact** on functional activities, including standing, transfers and ambulation!!!
- Severe spasticity may contribute to increased functional impairment



SPASTICITY

- Additionally, it contributes to better peripheral circulation, thereby **avoiding edema** and reducing the risk of deep **vein thrombosis**.
- Exacerbating factors (such as urinary tract infection, constipation, ingrown nails, pulmonary Infection, pressure ulcers).
- Drug agent: baclofen, tizanidine, botulinum toxin, benzodiazepine, dantrolene sodium, gabapentin and pregabalin.

SPASTICITY

- **Tizanidine** acts to reduce reflex mechanical responses substantially without inducing comparable changes in intrinsic muscle properties in individuals with SCI.(without change in postural stability).
- **Botulinum toxin** is an injectable medication that acts on the neuromuscular junction. A chemical denervation in intrafusal and extrafusal muscle fibers and its effect is reversible.
- **Benzodiazepines**
- **Dantrolene sodium**(directly on muscle but it tends to cause generalized weakness)

PAIN management

- **Chronic pain is one of the frequent secondary complications for individuals with SCI, with up to 80%.**
- **Chronic pain may lead to functional disability and emotional discomfort.**



PAIN

- Pain types are divided into two main groups: nociceptive (musculoskeletal or visceral) and neuropathic (either above level, at level or below level of injury).

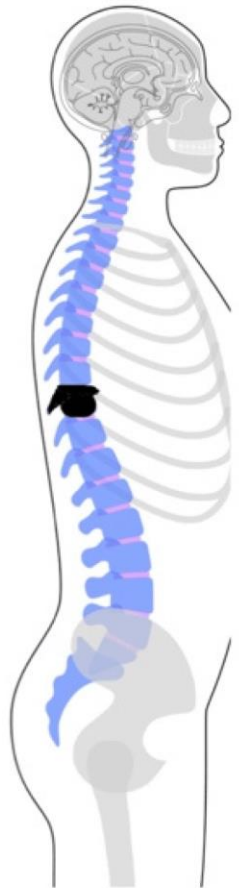
PAIN

- **Nociceptive pain:**
- Shoulder pain due to manually operated wheelchair
- Carpal tunnel syndrome and ulnar nerve entrapment
- Muscle spasm pain in patients with incomplete SCI
- Visceral pain due to irritation or distention of internal organs. This type of pain is reported in 15% of patients with chronic SCI .

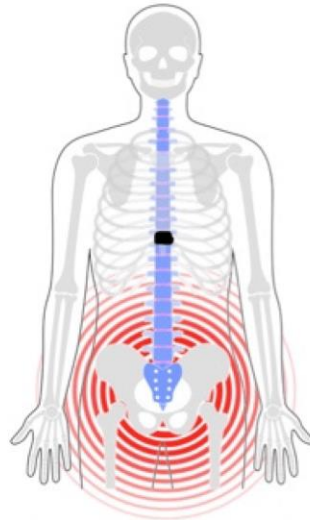


Neuropathic pain

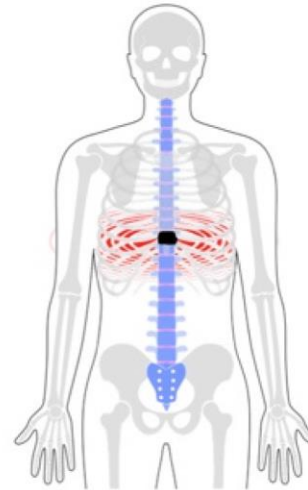
SPINAL CORD INJURY



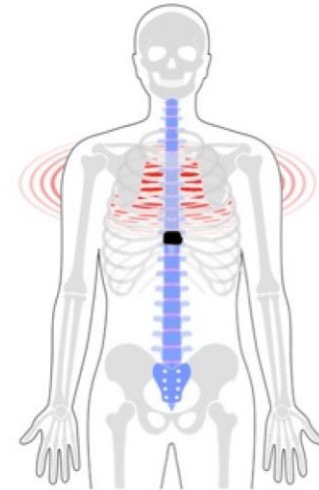
PAIN SENSATION: tingling, burning, compression



NP BELOW THE NEUROLOGIC LEVEL OF SCI LESION



NP AT THE NEUROLOGIC LEVEL OF SCI LESION



NP ABOVE THE NEUROLOGIC LEVEL OF SCI LESION

SENSORY SYMPTOMS ASSOCIATED:

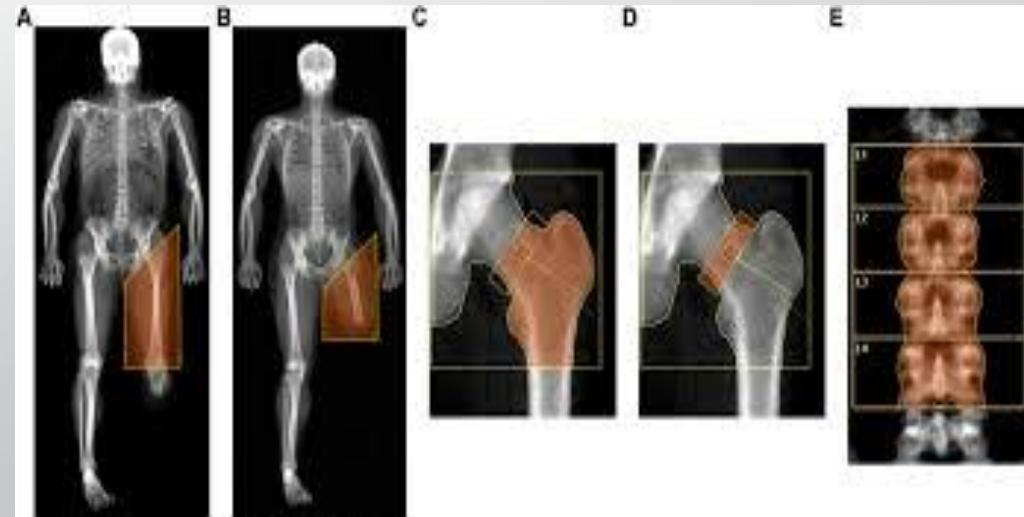
1. allodynia: pain perception in response to innocuous tactile stimuli
2. paraesthesia: abnormal painless sensation
3. dysesthesia: abnormal painful sensation
4. phantom sensations

Pain treatment

- Simple analgesics, non-steroidal anti-inflammatory drugs and opioids.
- **Physiotherapy** (transcutaneous electrical nerve stimulation, acupuncture, spinal cord stimulation) with and without drug.
- Surgical interventions.
- Anticonvulsants ,Antidepressants.

OSTEOPOROSIS AND BONE FRACTURES

- Osteoporosis, a condition characterized by low bone mass and deterioration of the skeletal microarchitecture.
- It occurs rapidly in the first 12-18 mo but continues for several years.

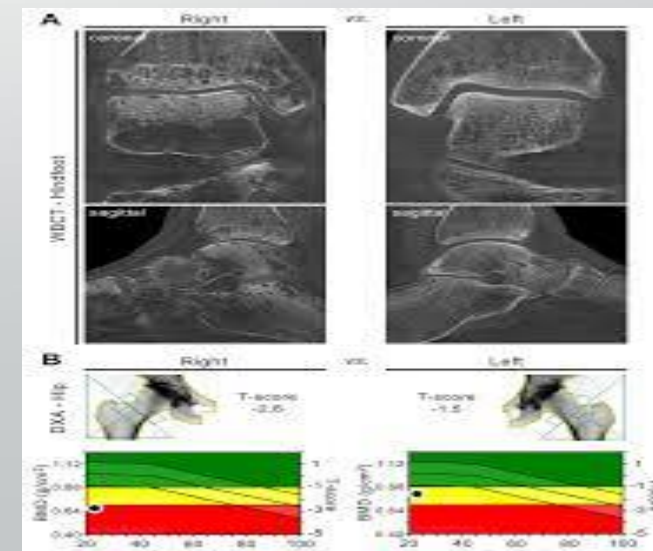


OSTEOPOROSIS AND BONE FRACTURES

- The mechanism of osteoporosis is complex and multifactorial .
- Disuse(regional) may play an important role .
- Non-mechanical factors also appear to be important:
 - *insufficient nutritional support*
 - *disordered Vaso regulation*
 - *hypercortisolism (either therapeutic or stress-related)*
 - *alterations in gonadal function and other endocrine disorders .*

OSTEOPOROSIS AND BONE FRACTURES

- The most common fracture sites appear to be those **around the knee**, such as the distal femur or proximal tibia.
- More severe in patients with complete SCI than with incomplete SCI .
- No standardized treatment guidelines for management of osteoporosis in patients with SCI .
- pharmacological and rehabilitation-oriented approaches .



OSTEOPOROSIS AND BONE FRACTURES

- **Bisphosphonates (strongly inhibit bone resorption).**
 - ✓ Gilchrist *et al* concluded that alendronate 70 mg orally per week for 1 year initiated soon after acute SCI prevents bone loss .
 - ✓ Zehnder *et al* also reported that SCI bone loss was stopped at all measured cortical and trabecular infralesional sites over 2 years with alendronate .
 - ✓ In a recent study, zoledronic acid.

- **Non-pharmacological treatment methods :**



- standing-up.



- orthotically aided walking.



- weight - bearing physical exercises.



- functional electrical stimulation .



- pulsed electromagnetic fields.



Inspiring!

