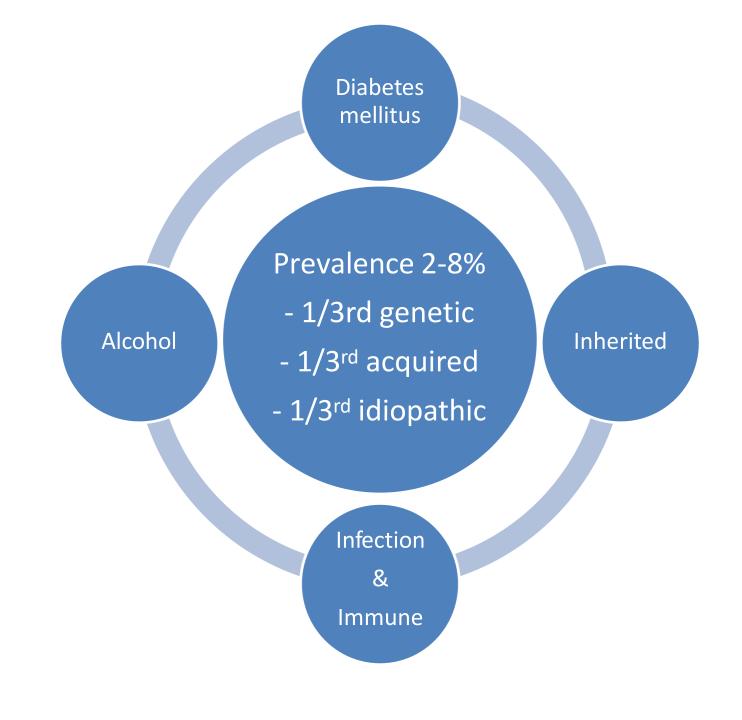
AN APPROACH TO NEUROPATHY

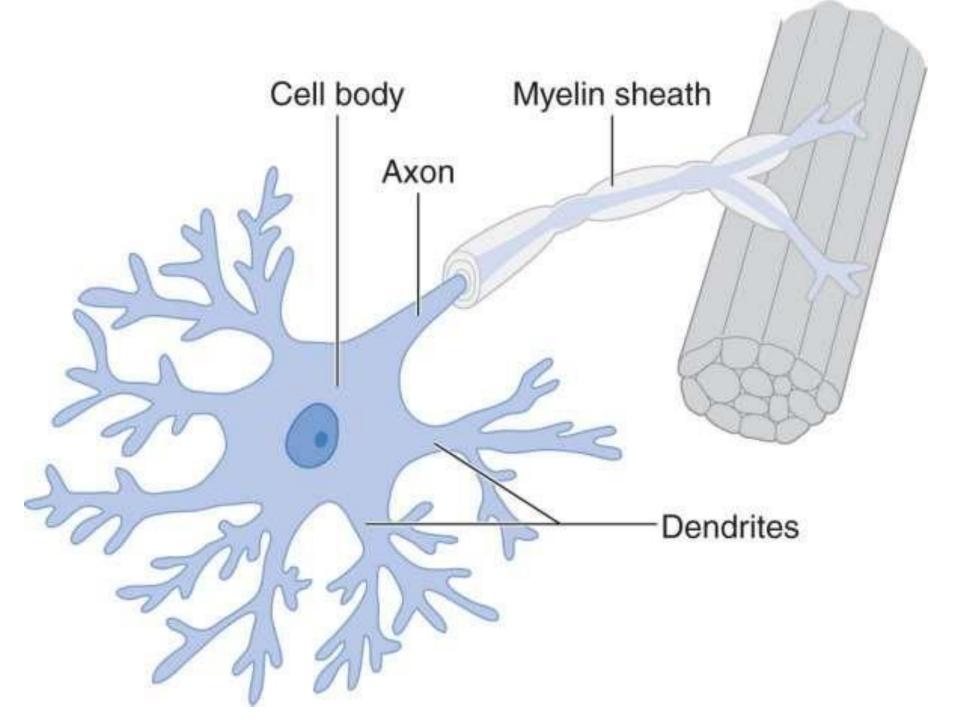
WHAT WE WILL DISCUSS?

- DEFINITION
- TYPES OF NEUROPATHY
- HISTORY AND EXAMINATION
- APPROACH
- INVESTIGATIONS

DEFINITION

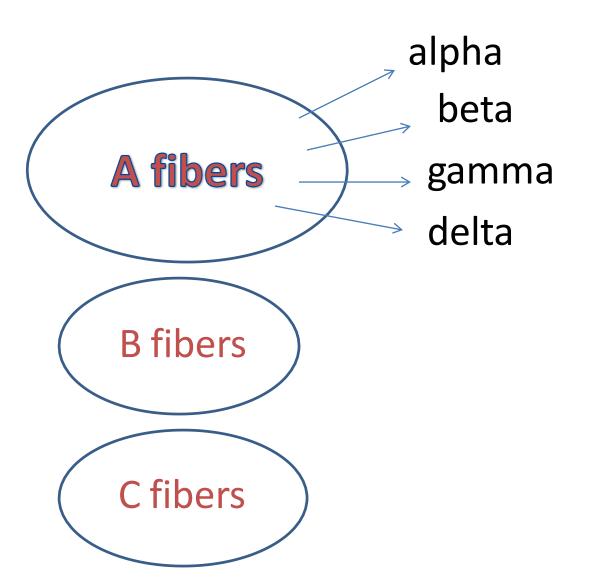
 Functional disturbance or pathological change in peripheral nervous system





TYPES OF NERVE FIBERS

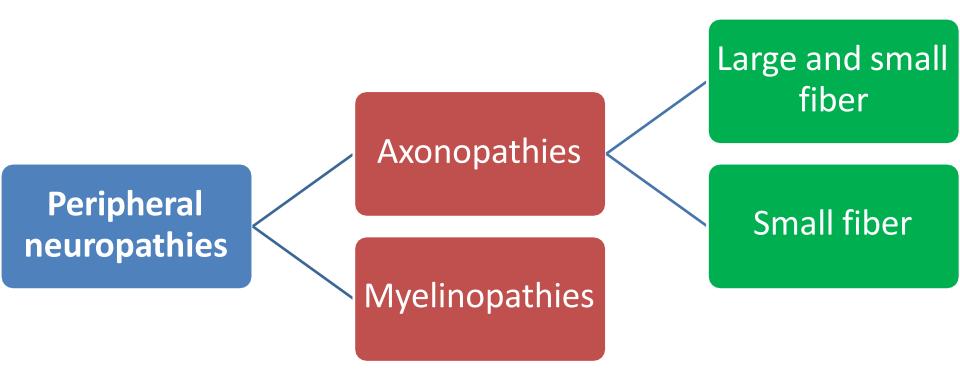
(Erlanger/Gasser classification)



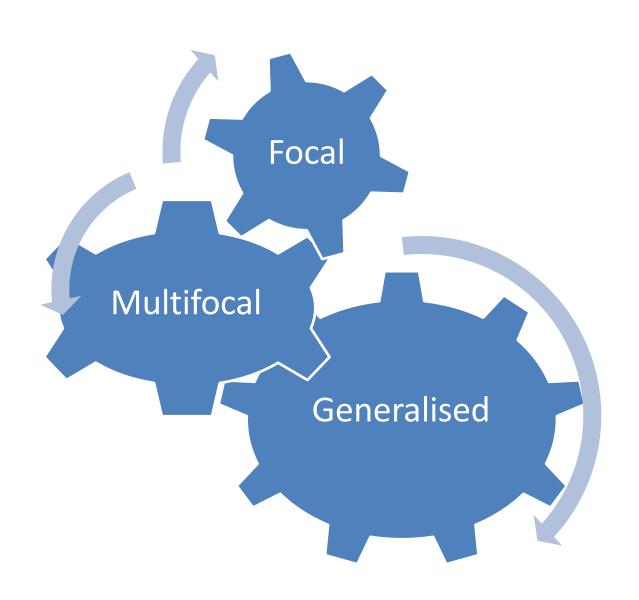
Erlanger / Gasser classification of nerve fibers

Fiber types	Function	Avg. fiber diameters (µm)	Avg. cond. Velocity (m/s)
Αα	Primary muscle spindle afferents, motor to skeletal muscle	15	100 (70-120)
Αβ	Cutaneous touch and pressure afferents	8	50 (30-70)
Αγ	motor to muscle spindle	5	20 (15-30)
Αδ	Cutaneous temperature and pain afferents	<3	15 (12-30)
В	Sympathetic preganglionic	3	7 (3-15)
С	Cutaneous pain afferents sympathetic postganglionic	1	1 (02-2)

CLASSIFICATION OF PERIPHERAL NEUROPATHIES



IS THIS PERIPHERAL NEUROPATHY?



Focal

Entrapment Neuropathies

Meralgia paresthetica Myxedema Rheumatoid Amyloid Acromegaly Hansen's Disease

Multifocal

- Diabetes Mellitus
- Vasculitis
- SLE, PAN
- HIV

MOTOR

- GBS and CIDP
- Diphtheria and Botulism

AUTONOMIC

- Alcoholism
- Amyloidosis and DM

SENSORY

Small fiber

- Leprosy
- > DM

Large fiber

- **→** Paraneoplastic
- Cisplatin and other chemotherapeutics

AXONAL

- Insidious
- Glove and stocking pattern
- Preservation of all DTRs except ankle jerk
- Recovery in months to years
- More Residual deformity
- e.g- Vasculitis, Toxins,
 Metabolic

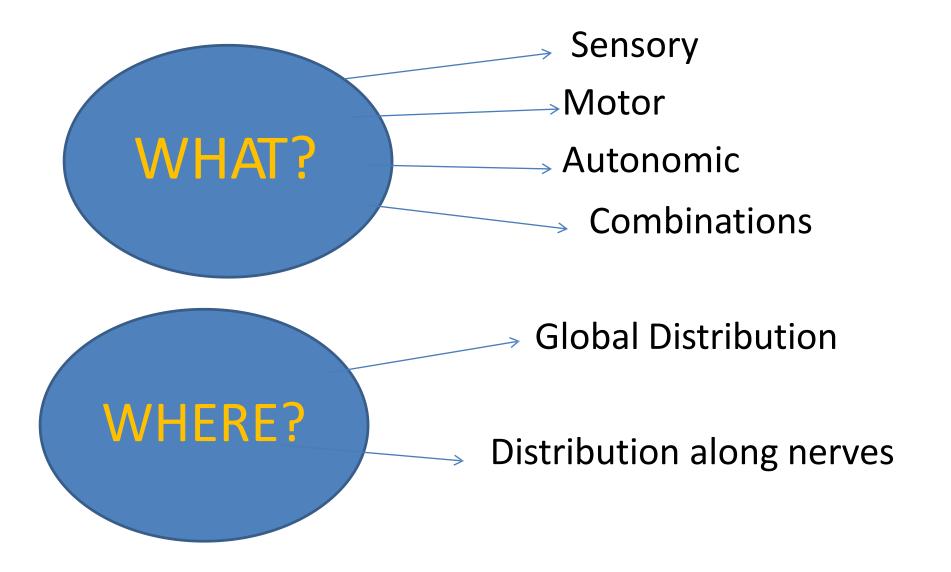
DEMYELINATING

- Generally Acute
- Minimal sensory loss
- Loss of all DTRs
- Rapid recovery
- Residual deformity minimal
- e.g- GBS, CIDP

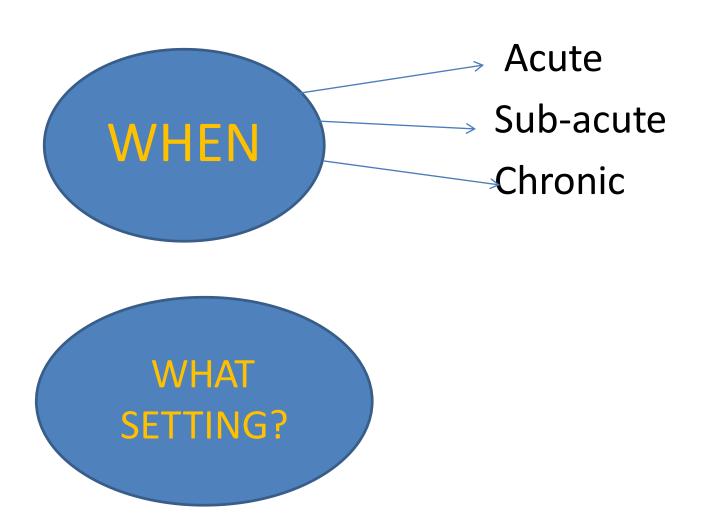
HISTORY

- Ethnicity and globalisation
- Dietary History
- Family History
- Drug History(including abused drugs)
- Concurrent systemic Illness
- Toxins exposure
- Behavioural related(HIV, Hepatitis C, Nutritional)
- Vaccination history

APPROACH TO NEUROPATHY



APPROACH TO NEUROPATHY



PHYSICAL EXAMINATION

- 1st step
- (PNS vs CNS
- CNS- speech, diplopia, ataxia, CN involvement/ myelopathy >> bowel, bladder involvement
- Single or multiple nerve root or peripheral nerve plexus
- PNS- peripheral nerve roots vs plexus
- Fundoscopy → optic pallor
 - leukodystrophies and vitamin B12 deficiency

PHYSICAL EXAMINATION

- Motor examination
 - -fasciculations or cramps, or loss of muscle bulk
- **Tone** → normal or reduced
- **Deep tendon reflexes** → reduced or absent.
- Bilateral foot drop > steppage gait
- **Proximal weakness** inability to squat or to rise unassisted from a chair

PHYSICAL EXAMINATION

Respiratory rate and vital capacity → GBS

Mees' lines Arsenic poisoning

HOW TO TEST?



ELECTRODIAGNOSTIC TEST



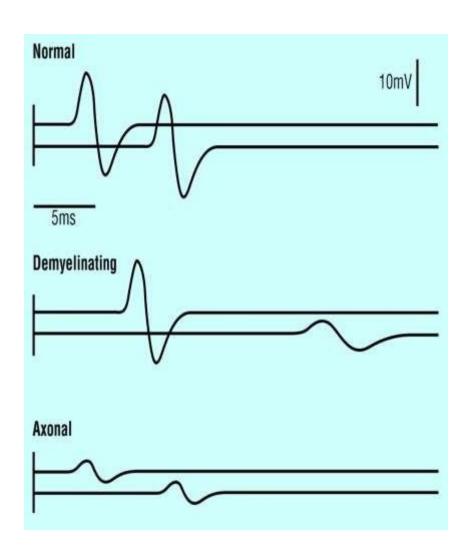
 Cofirmation of presence of neuropathy

Small or large fiber involved

 Motor, Sensory or mixed

Axonal Vs
 Demyelination

ELECTRODIAGNOSTIC TEST



- Velocity of conduction
- Distal latency
- Conduction block
- Temporal dispersion
- F wave latency

NERVE BIOPSY

- If diagnosis is in doubt even after lab and electrodiagnostic findings
- Vasculitis, Amyloid neuropathy, Leprosy, CIDP, Inherited disorders of myelin
- Sural nerve → M/C
- Superficial peroneal nerve alternative; allows simultaneous biopsy of the peroneus brevis muscle
- Combined nerve and muscle biopsy → Vasculitis

LEPROSY

Tuberculoid → Patch of superficial sensory loss

Lepromatous > Widespread invasion of cutaneous nerve

- Tendon reflex Preserved
- No autonomic neuropathy

VITAMIN B12 DEFICIENCY

 Spinal cord, Brain, Optic nerve and Peripheral nerves all affected

Sub-acute Combined degeneration of cord

• Visual impairment — Optic neuropathy

DIABETES MELLITUS

Diabetic Ophthalmoplegia

Autonomic Neuropathy Acute Diabetic Mononeuropathy

Multiple Mononeuropathy and Radiculopathy

Thoracoabdominal Radiculopathy

Distal Polyneuropathy

HYPOTHYROIDISM

Neurological endemic cretinism >> Proximal limb and truncal rigid-spastic motor disorder

Myxedematous → No spastic rigidity
Loss of reflexes,loss of vibration, position and touch

Sporadic → Delayed tendon reflexes

ALCOHOLISM

- More common in our settings
- Slow and insidious progression
- Distal and symmetrical involvement
- Positive sensory symptoms
- Autonomic neuropathy coexistent
- Multi-factorial damage

THANK YOU