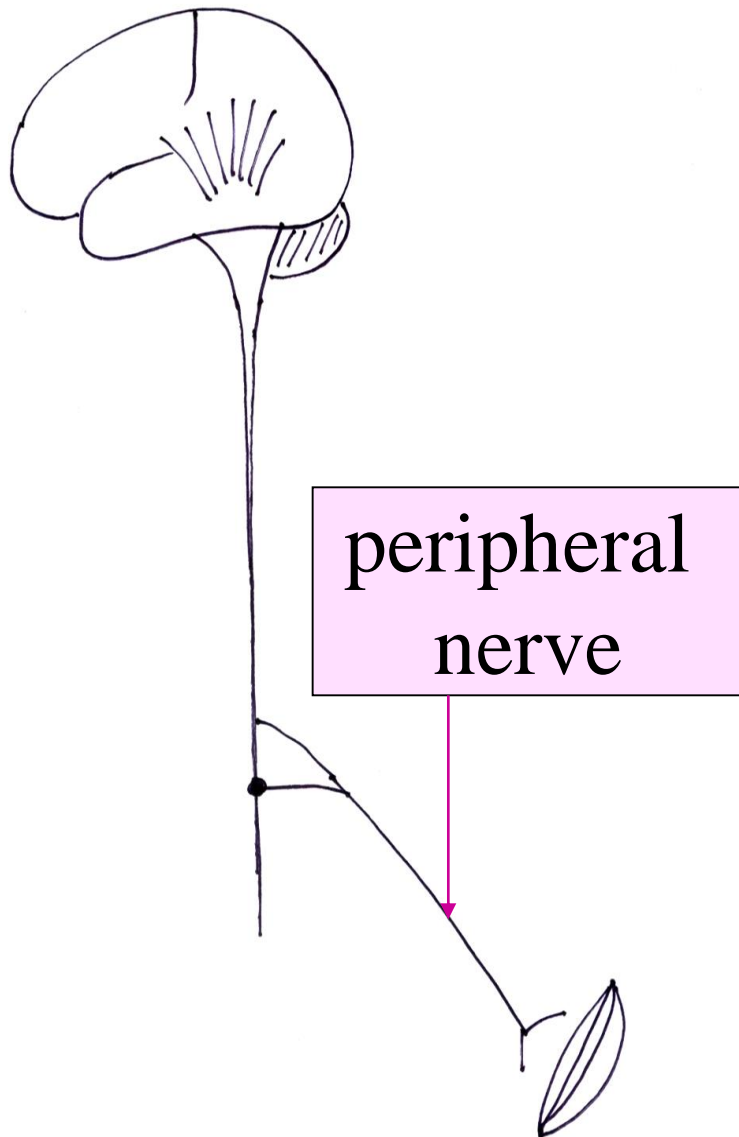


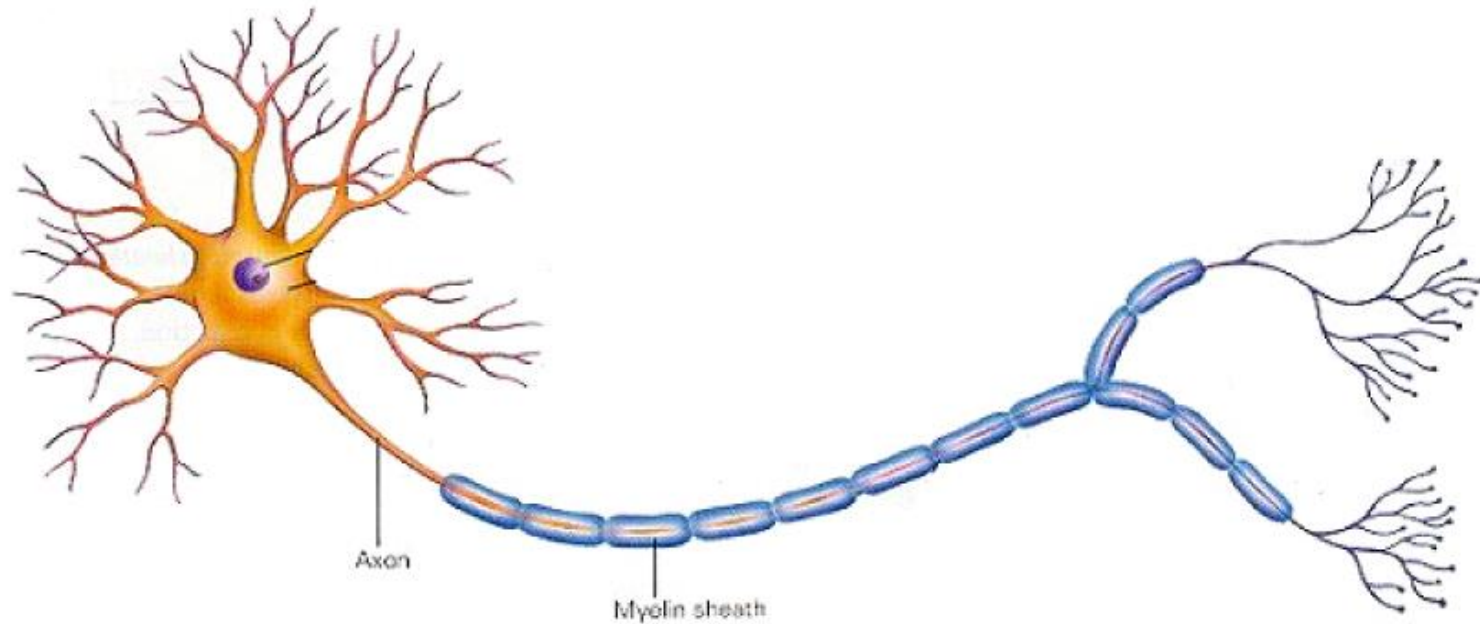
Disorders of peripheral nerves

Peripheral nerve disease



- distal signs
- motor - LMN weakness
- reflex loss
- sensory loss
 - large fibres - JPS, vibration
 - small fibres - pain, temp.
- autonomic

Peripheral nerve disease



- demyelination
 - loss of myelin, axon intact
- axonal degeneration

Peripheral nerve disease - types

- Mononeuropathy
- ‘Mononeuritis multiplex’
- Polyneuropathy

Mononeuropathy

- Trauma
- Entrapment neuropathy
 - median N. - carpal tunnel syndrome
 - ulnar N. - cubital tunnel syndrome
 - lateral cutaneous N. of the thigh -
meralgia paraesthetica
 - tibial N. - tarsal tunnel syndrome

Mononeuritis multiplex

- diabetes
- leprosy
- vasculitis
- connective tissue diseases
- sarcoidosis

Polyneuropathy- causes

- diabetes
- alcohol
- infections - leprosy, diphtheria, HIV, ...
- inflammatory - Guillain-Barre synd., CIDP
- drugs - INAH, metronidazole,
- other metabolic - B12 deficiency, CRF, porphyria, ...
- toxins - lead, organophosphates, ...
- hereditary - HMSN (Charcot-Marie-Tooth dis.)
- paraneoplastic
- other

Polyneuropathy - clinical features

Polyneuropathy - clinical features

- motor
- sensory
- autonomic

Polyneuropathy - clinical features

- bilateral, symmetrical
- distal weakness, wasting
- reflexes - diminished or lost
- numbness, cannot feel texture
 - ‘feeling of mud on my soles’
 - ‘walking on cotton wool’
- pain, ‘pins & needles’, ‘burning’
- ‘glove & stocking’ sensory loss
 - pain, touch, temperature
- JPS loss - sensory ataxia, Romberg’s sign
- thickened nerves - leprosy, HMSN, ...

Polyneuropathy - clinical features...

autonomic neuropathy

- CVS - postural hypotension
loss of HR & BP variation with Valsalva
- GIT - vomiting (gastroparesis), nocturnal diarrhoea
- GUT - bladder - retention, incontinence
erectile dysfunction
- skin - loss of sweating

Peripheral nerve disease - investigation

- Confirm peripheral nerve disease
 - nerve conduction studies / electromyography
 - demyelination/ axonal degeneration
 - nerve biopsy
- Look for cause

Peripheral nerve disease - treatment

- Symptomatic treatment - peripheral nerve disease
 - Neuropathic pain —
 - anti-depressants - tricyclics, ...
 - anti-epileptics - carbamazepine, gabapentin, pregabalin
 - physiotherapy
- Treat the cause

Guillain-Barre syndrome

GBS - pathophysiology

Antibodies against peripheral nerve myelin or axonal proteins

Disruption of myelin sheath or axonal transport

Impaired nerve conduction

Motor and sensory deficits

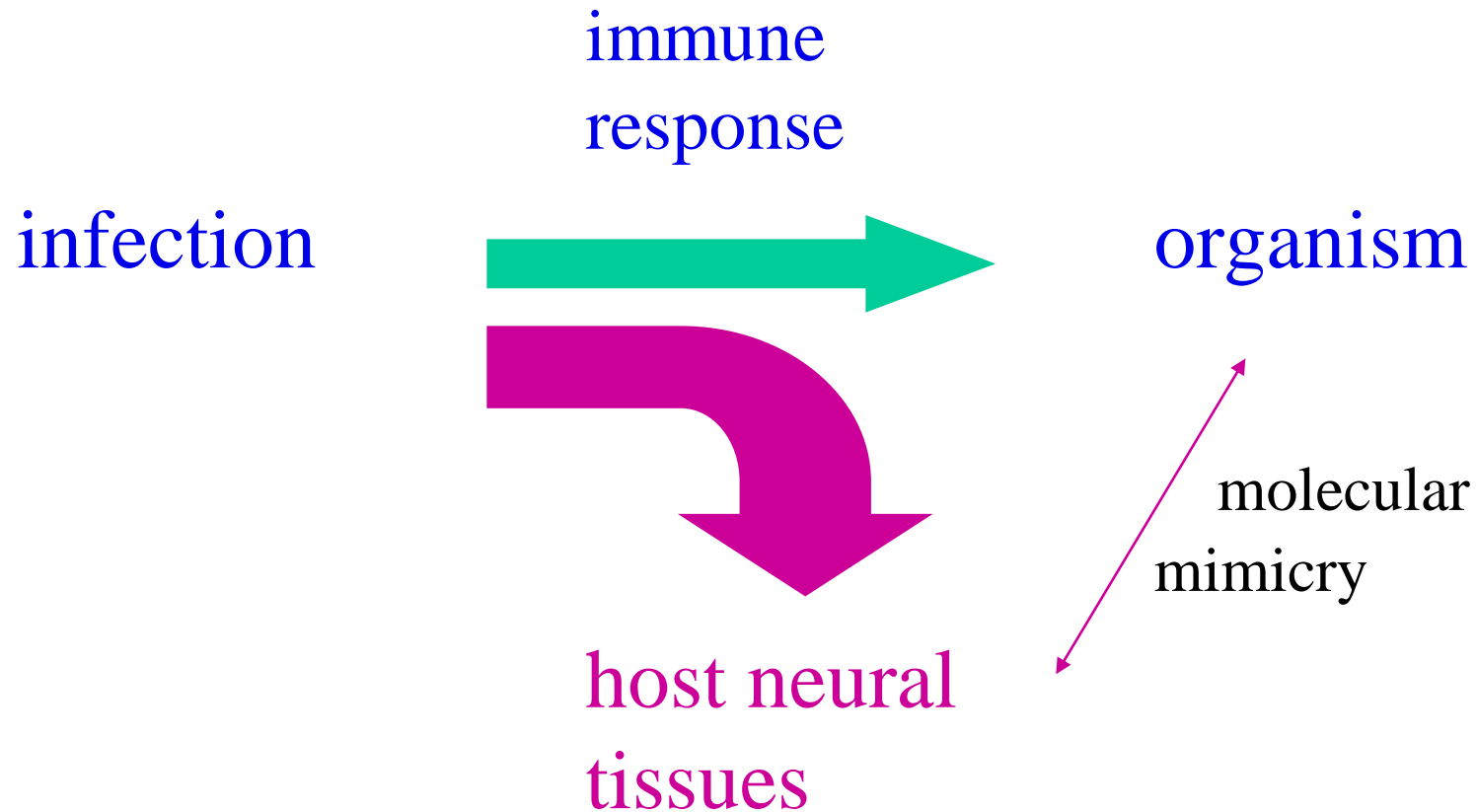
Recovery and remyelination

Supportive care and symptom management

Prognosis and long-term outcomes

Research and clinical trials

GBS - pathophysiology



GBS - pathophysiology

- antecedent events

infection - viral - CMV, EBV, VZV, HIV

Campylobacter, mycoplasma

vaccines

- molecular mimicry

shared antigens between pathogen &
surface glycolipids of peripheral nerves

- immune mediated inflammatory poly-radiculo-neuropathy

GBS - pathophysiology

target epitopes

- Schwann cell membrane or myelin
demyelination - AIDP
- axonal membrane
axonal damage - axonal GBS
- cerebellum, cranial nn. 3 4 6 , dorsal root ganglia
Miller- Fisher syndrome
ataxia, ophthalmoplegia, areflexia

GBS - clinical features

acute onset of weakness and sensory deficit

ascending paralysis

areflexia

autonomic dysfunction

resolves within 4 weeks

diagnosis by exclusion

CSF proteinopathy

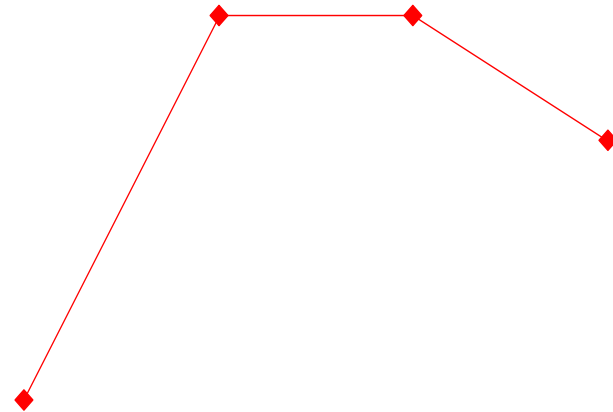
antibodies to gangliosides

GBS - clinical features

- acute flaccid areflexic paralysis
- self - limiting
- 2/3 - trigger event
- rapid progression - upto 4 weeks

plateau

recovery



GBS - clinical features

- acute, symmetric, flaccid motor weakness
 - 2 or more limbs - proximal, ascending
 - cranial nerves – facial, bulbar, ophthalmoplegia
 - respiratory
- areflexic
- sensory - symptoms common, signs few
- autonomic - cardiovascular, bladder, bowel
- pain - 1/3rd severe

GBS - diagnosis

- Clinical
- CSF - high protein (80%)
cells normal
‘protein-cellular dissociation’
- NCS / EMG
demyelination
axonal degeneration
- changes best seen after 10-14 days

Exclude - other causes of
acute flaccid paralysis

Exclude - other causes of acute flaccid paralysis

poliomyelitis

myasthenia

paralytic rabies

porphyria

toxic neuropathy - OP

electrolyte imbalance

diphtheria

botulism

snake bite

GBS - management

- General

monitor — pulse, BP, RR, single breath count,
vital capacity

1/3 - intensive care

- Respiratory support - $VC < 20 \text{ ml/kg}$

* no signs of breathlessness

- Specific treatment - Immunomodulation

plasma exchange, IVIg - equally effective

* steroids - no benefit

- Rehabilitation