



WHAT?

Guillain-Barré syndrome (GBS)

affects 1:40,000 people each year. While it can affect anyone at any age it is more common in older males.

It is an **autoimmune response to infections** (often *Campylobacter* infection – around 50% of viral).

It is characterised by ascending paralysis, starting with tingling and numbness of the fingers and toes, with decreased or absent reflexes. Symptoms usually progress over a period of days to four weeks, **usually peaking by the 2nd week.**

- Up to 30% of patients will have respiratory muscle weakness and require ventilatory support.
- 1:5 patients will have autonomic dysfunction requiring cardiac monitoring.

WHY?

Diagnosis is largely clinical, however lumbar puncture can be useful. There can be elevated CSF protein levels with a normal WCC. **EEG studies can also aid diagnosis.**

The treatment of GBS is with supportive measures. It is important to remember that these patients require close monitoring and will often require a stay in ITU.

Serial spirometry is used to assess need for intubation and ventilatory support.

All patients with a FVC <20mL/kg require ITU support and will likely need intubation.

Treatment options for GBS are plasma exchange or IV immunoglobulins, these have been shown to be of benefit in RCTs.

HOW?

Guillain-Barré Syndrome (*Life in the Fast Lane*)

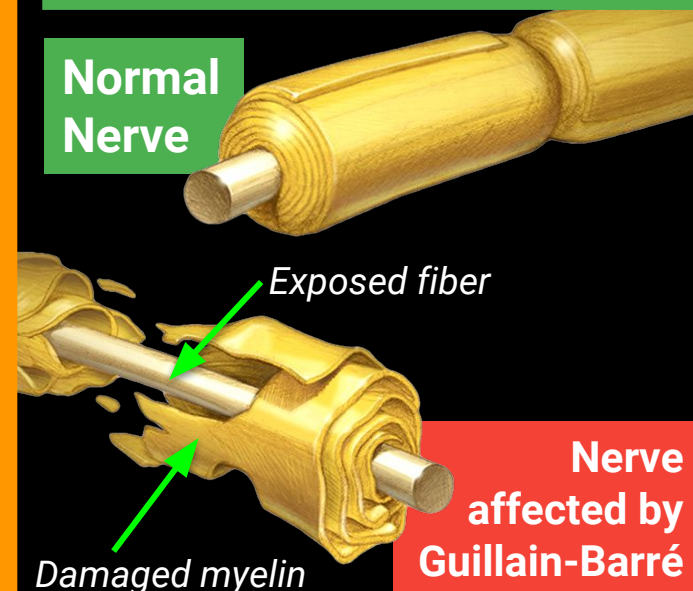
<http://bit.ly/2nDXYbn>

Diagnosis Criteria of GBS (*BMJ*)

<http://bit.ly/2E3jYH5>

GBS Information Guides (*GAIN Charity*) <http://bit.ly/2FOb5xB>

Normal Nerve



Nerve affected by Guillain-Barré