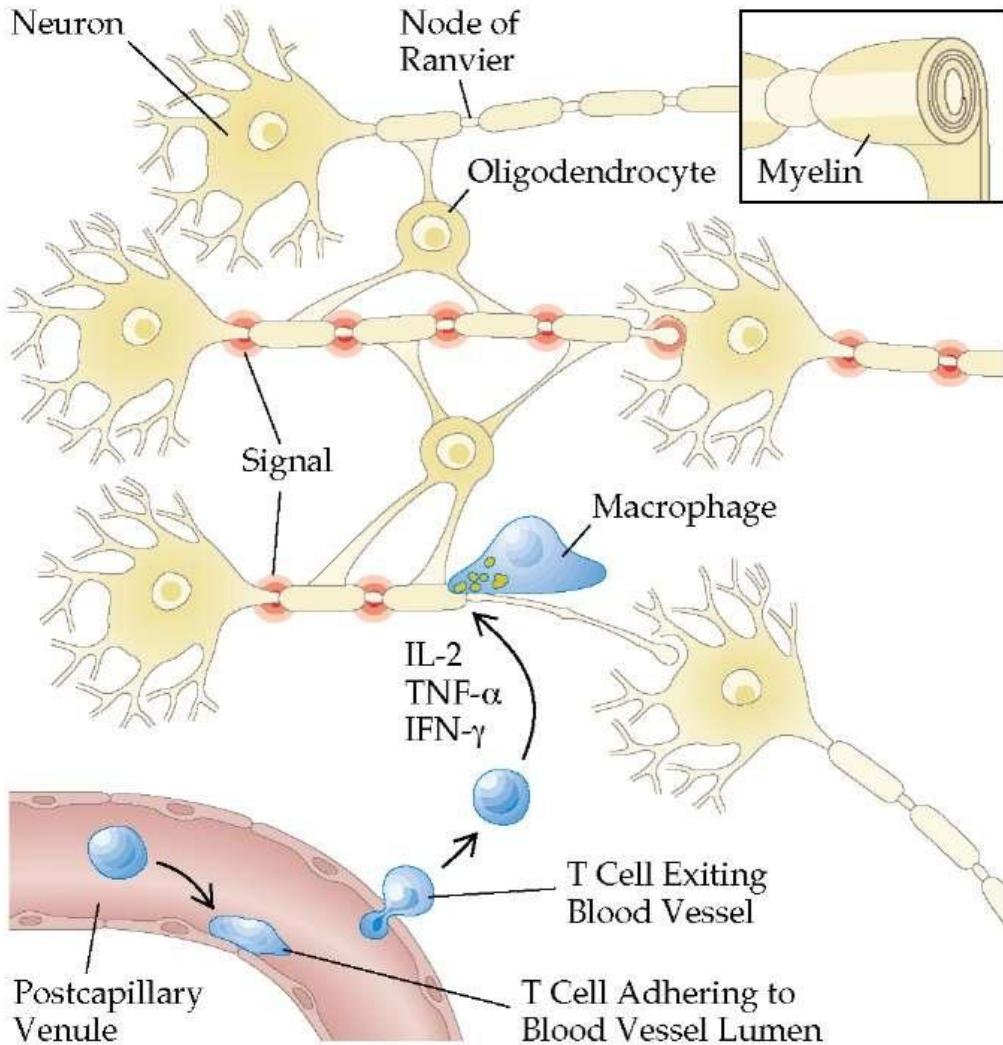
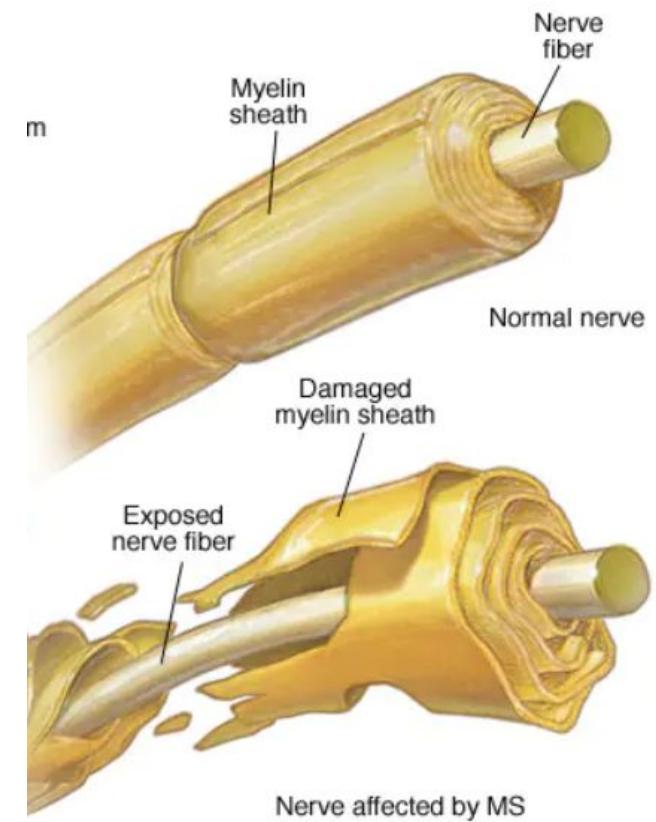

NEUROLOGY

MS, meningitis, encephalitis, intracranial & cerebral abscesses, neurosyphilis, CJD





MULTIPLY SCLEROSIS

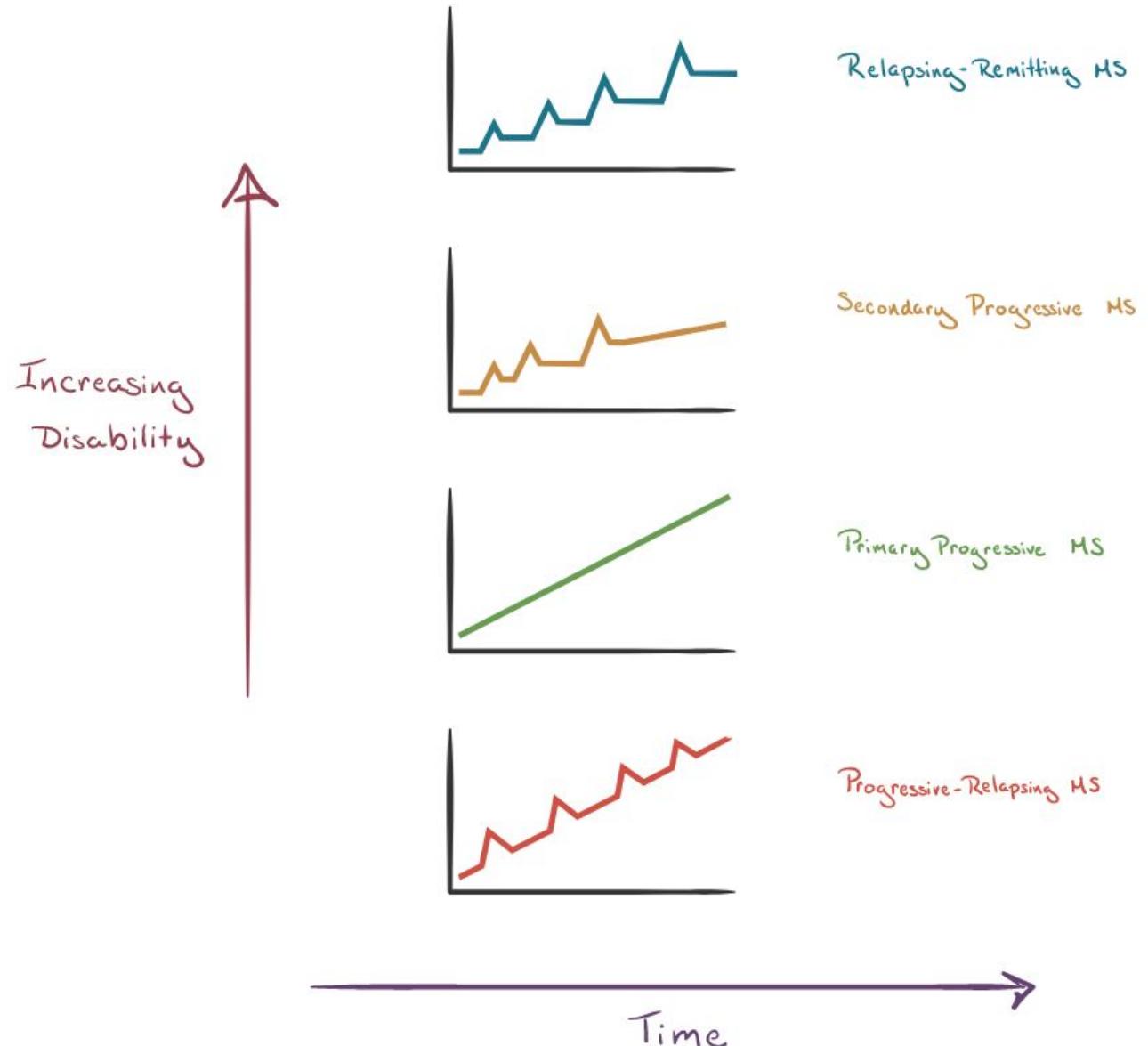


MULTIPLY SCLEROS

Cause is unknown

It's linked to:

- Genetic: female (20-40 years); genes encoding for HLA-DR2
- Infections
- Vitamin D deficiency



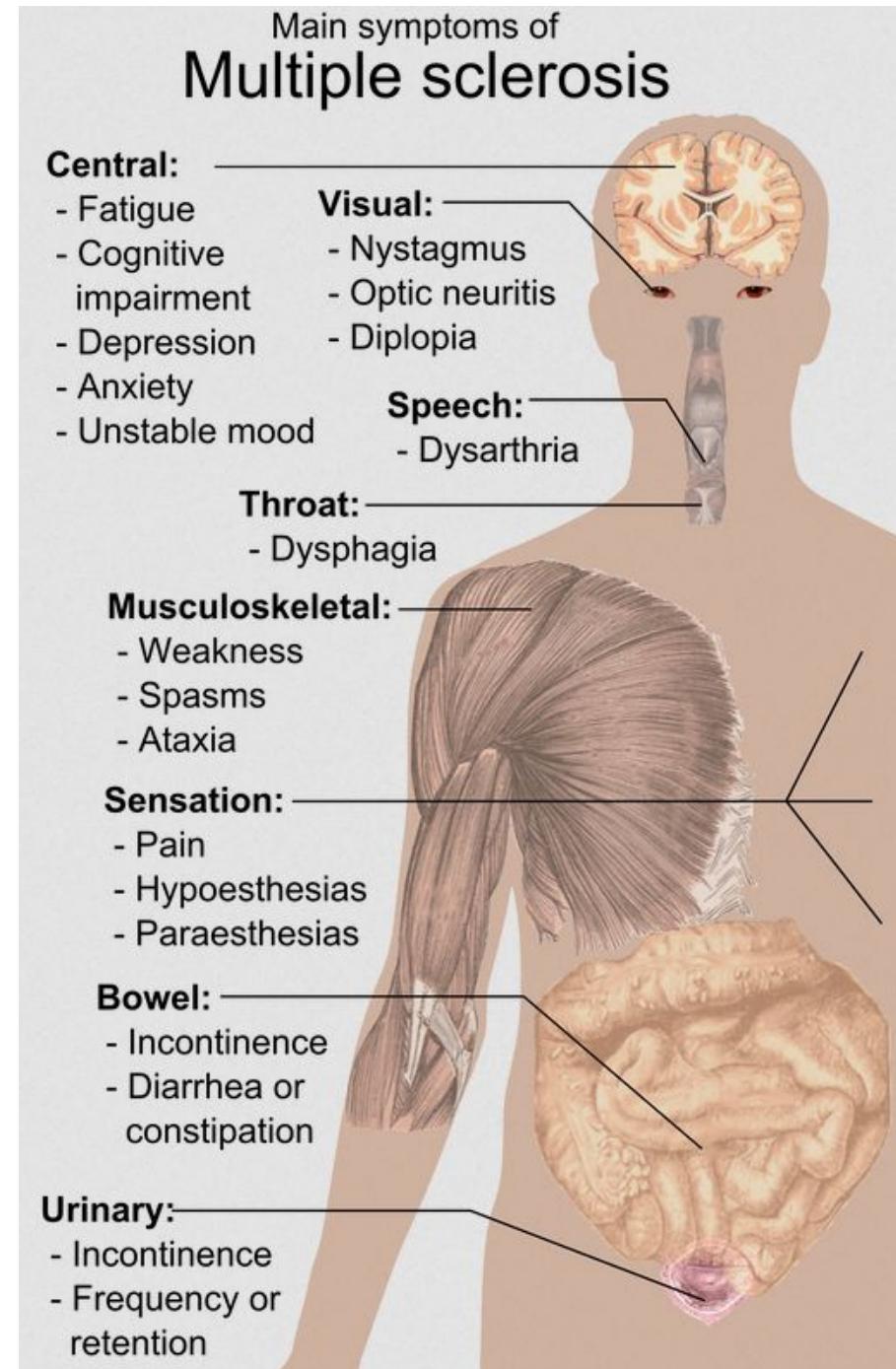
MULTIPLY SCLEROSIS

Charcot's neurologic triad

- Dysarthria
- Nystagmus
- Intension tremor

Specific signs:

- Uhthoff's sign
- Lhermitte's sign





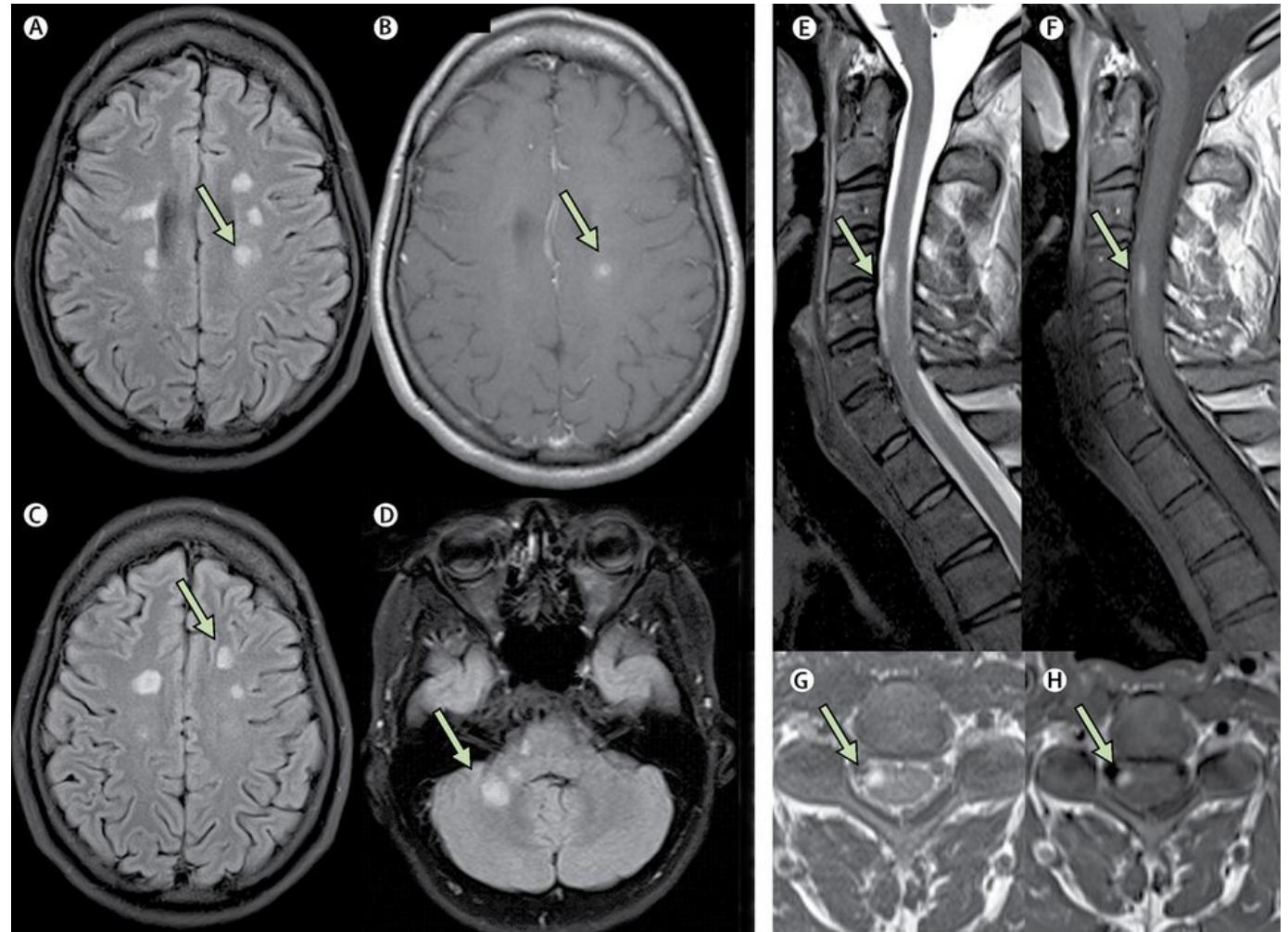
LHERMITTE'S SIGN

Electric shock sensation
which occurs with neck
flexion and often radiates
down the spine

MULTIPLY SCLEROSIS

Diagnosis

- MRI
- Cerebrospinal fluid
- Visual evoked potential



MULTIPLY SCLEROSIS

Treatment

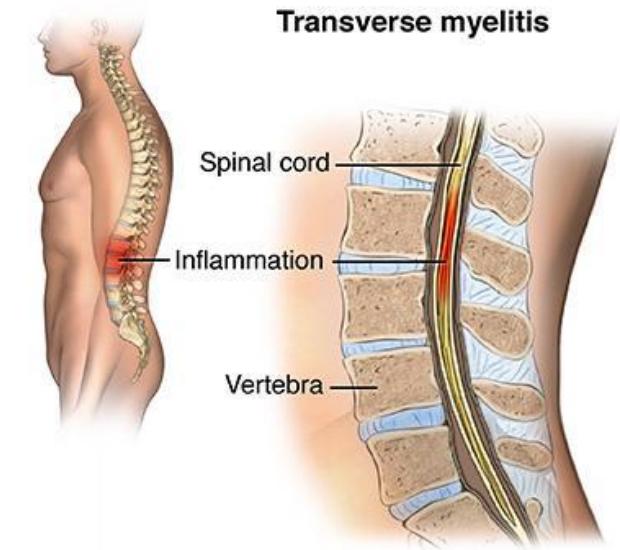
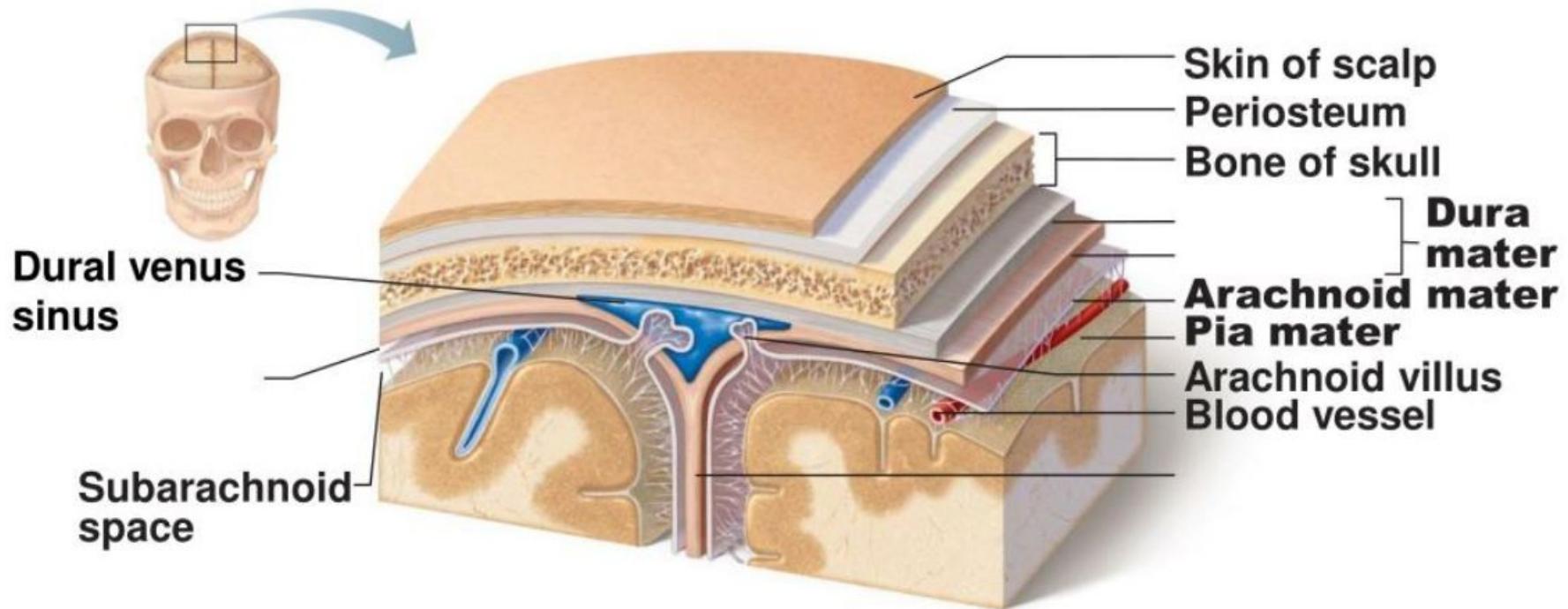
RRMS

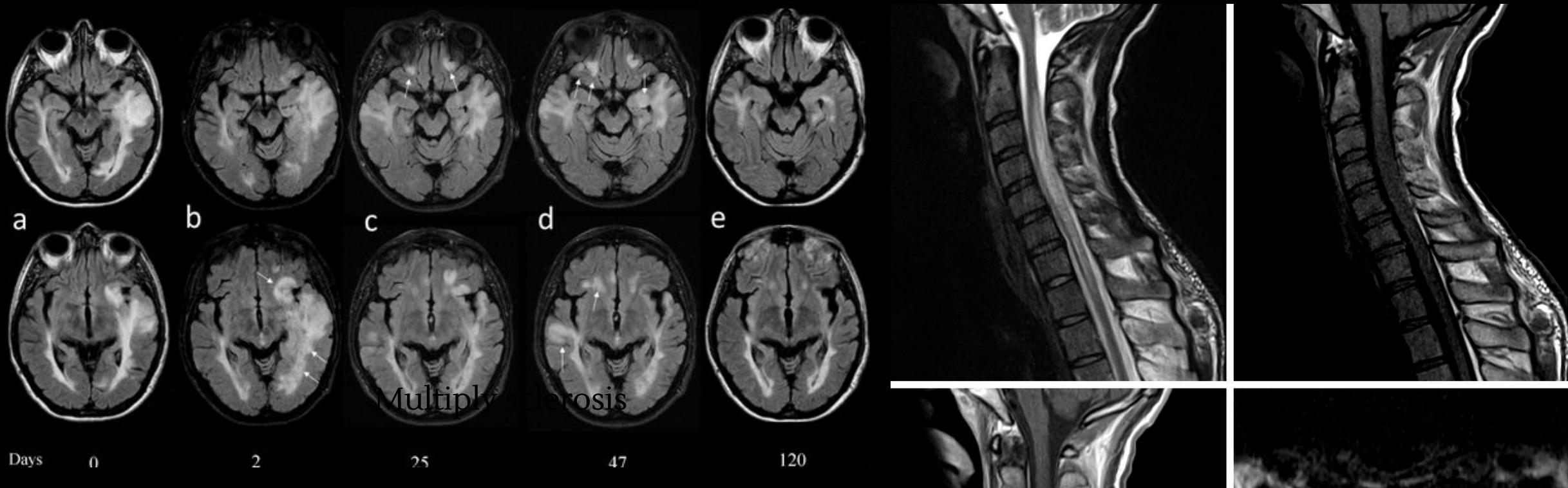
- Corticosteroids, cyclophosphamide, intravenous immunoglobulin
- Plasmapheresis
- Immunosuppressant: recombinant b-IFN

Progressive MS

- Manage symptoms
- Physical therapy
- Cognitive rehabilitation therapy

MENINGITIS ENCEPHALITIS MYELITIS





ENCEPHALITIS MYELITIS

MENINGITIS

Inflammation triggers

- Autoimmune disease
- Adverse reaction to medication
- Infection

Two ways of spreading

Direct spread

- Through overlying skin
- Up through nose
- Anatomical defect

Hematogenous spread

- Through binding to surface receptor
- Areas of damage
- Vulnerable spot

CAUSES

Bacteria

- Newborns: group B streptococci, E coli, Listeria monocytogenes
- Children and teens: Neisseria meningitidis, Streptococcus pneumonia
- Adults and elderly: Streptococcus pneumonia, Listeria monocytogenes

Tick-borne: *Borrelia burgdorferi*

Viruses:

- Enteroviruses, Herpes simplex, HIV
- Mumps, Varicella zoster, Lymphocytic Choriomeningitis

Fungi: *Cryptococcus* genuses, *Coccidioides* genuses

Tubercular meningitis

Parasitic cause: *Plasmodium falciparum*

SYMPTOMS

Meningitis

Headache, fever, nuchal rigidity

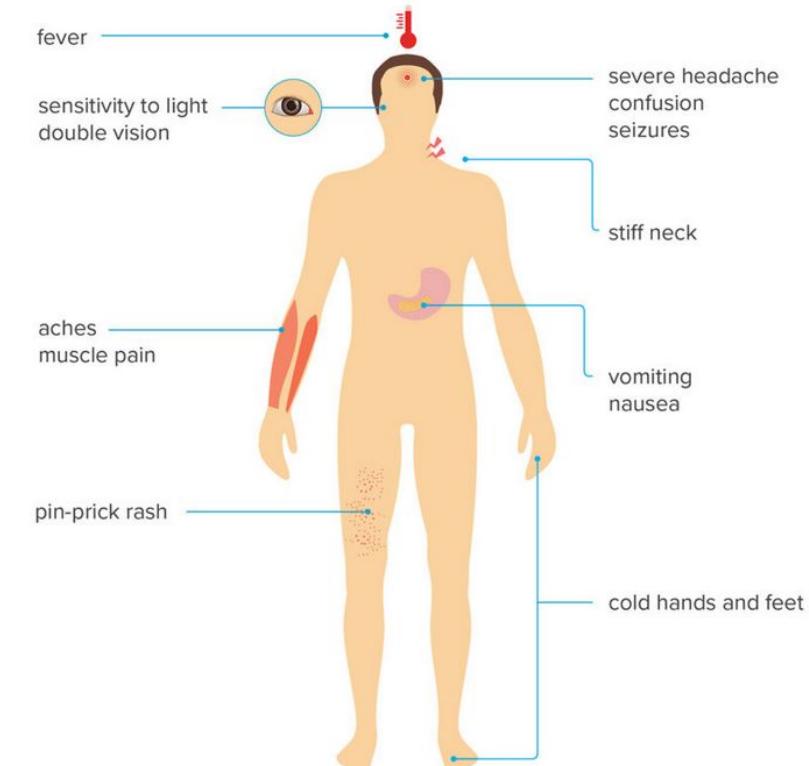
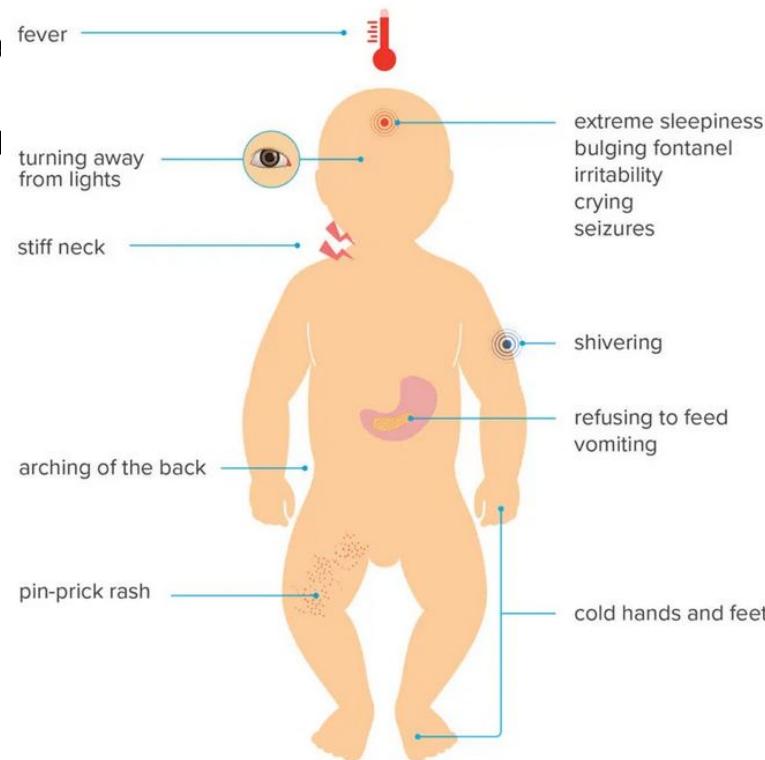
Photophobia and phonophobia

Encephalitis

Fever, altered mental status, seizure or focal neurologic symptoms

Myelitis

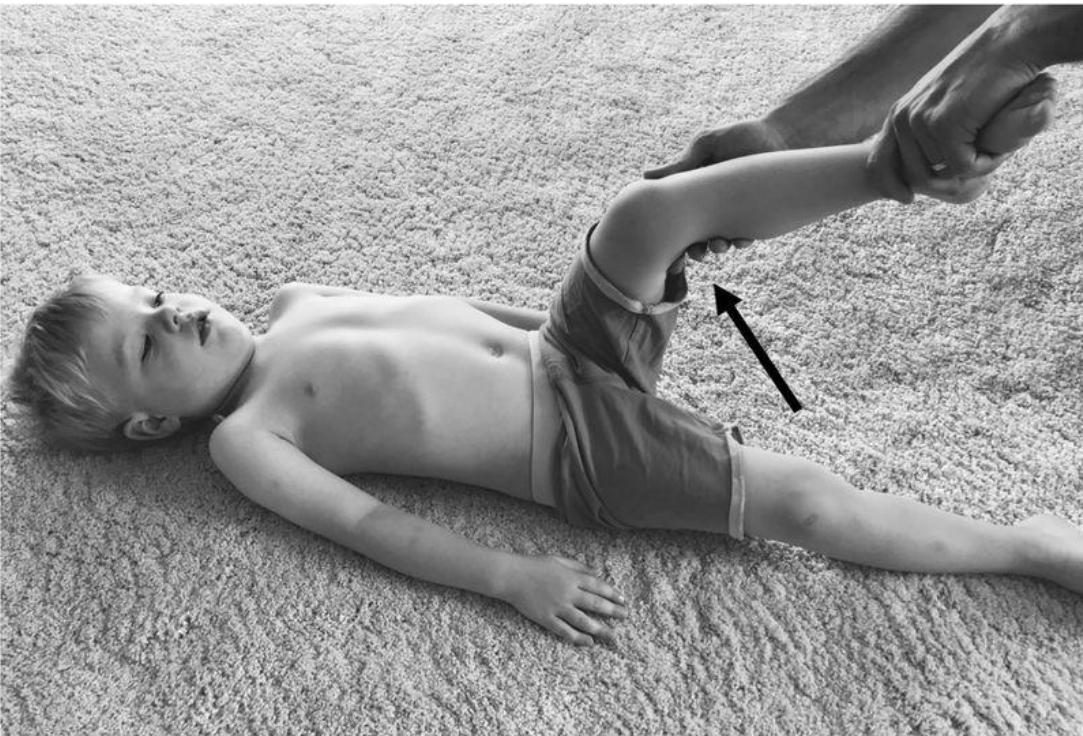
Flaccid paralysis and sensory loss



DIAGNOSIS

OF MENINGITIS

A



Kernig's Sign

B



Brudzinski's Sign

DIAGNOSIS

Lumbar puncture

PCR

Western blot

Thin blood smear

Etiology	Glucose	Protein	White Cell Count	Gram Stain	Bacterial Culture
Bacterial meningitis	Low	High	>100; ↑ PMN	+	+
Aseptic meningitis	Low/NL	NL/High	> NL; variable	-	-
<i>Pre-treated bacterial meningitis</i>	Low	NL/High	>100; ↑ PMN	-	-
<i>Viral meningitis</i>	NL	NL/High	>10-100's; ↑PMN (early) ↑ Lymphs (late)	-	-
Encephalitis	NL	NL	> NL	-	-

TREATMENT

Bacterial: Steroids and antibiotics

Antivirals, antibacterial, antifungals, antiparasitic

Prevention vaccine: Neisseria Meningitidis, Disseminated tuberculosis

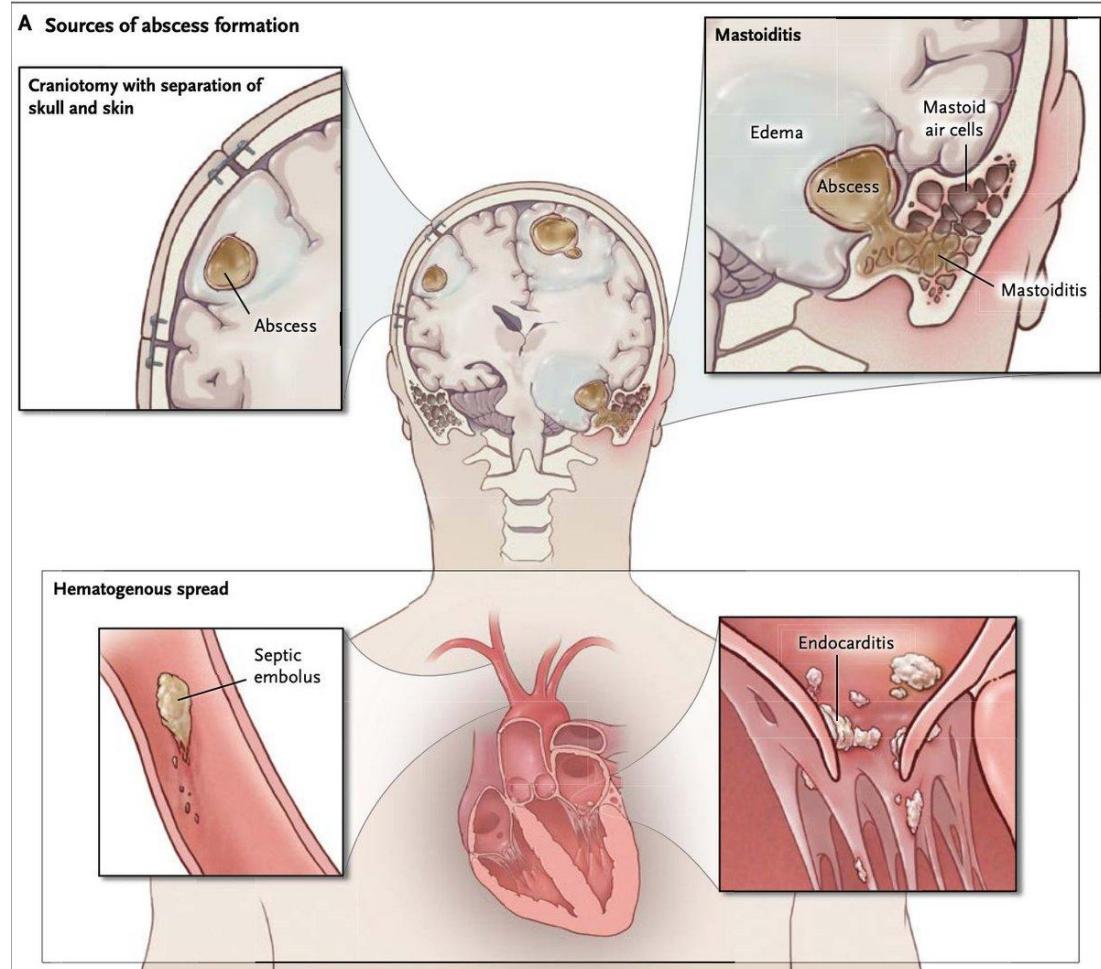
BRAIN ABSCESS

Direct spread

Cause a single brain abscess

Primary infection include:

- Subacute and chronic otitis media and mastoiditis (the inferior temporal lobe and cerebellum)
- Frontal or ethmoid sinuses and dental infection
(the frontal lobe)



BRAIN ABSCESS

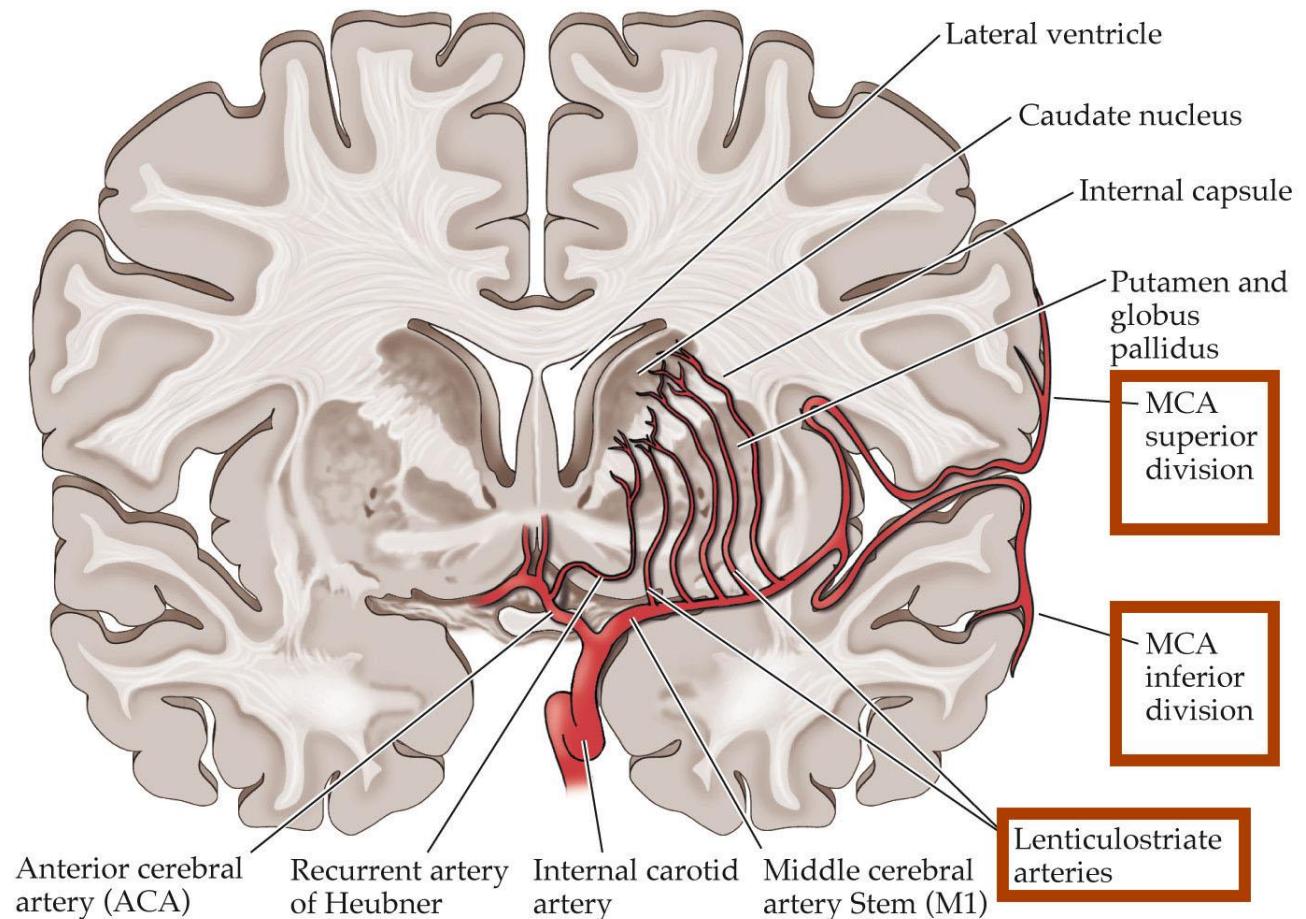
Hematogenous spread

Usually multiply abscess

Most commonly located in the distribution of the middle cerebral artery

Sources:

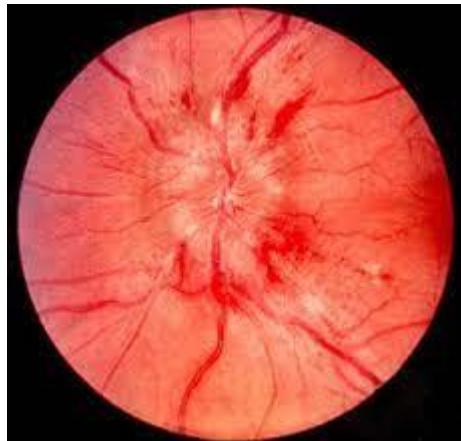
Skin infection, pelvic infection, intraabdominal infection, esophageal dilation, bacterial endocarditis, cyanotic congenital heart disease



BRAIN ABSCESS

Symptoms

- A headache (69% to 70%)
- Mental status changes (65%) lethargy progressing to coma is indicative of severe cerebral edema
- Focal neurologic deficits (50% to 65%)
- Fever (45% to 53%)
- Seizures (25% to 35%).
- Nausea and vomiting (40%)
- Nuchal rigidity (15%)

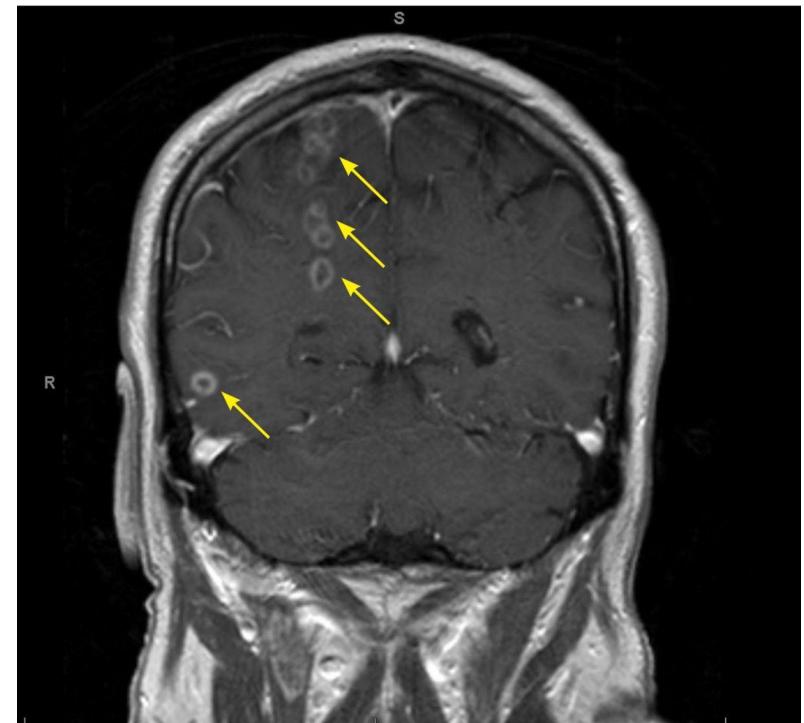


Diagnosis

Clinical: focal symptoms and signs

Papilledema

MRI



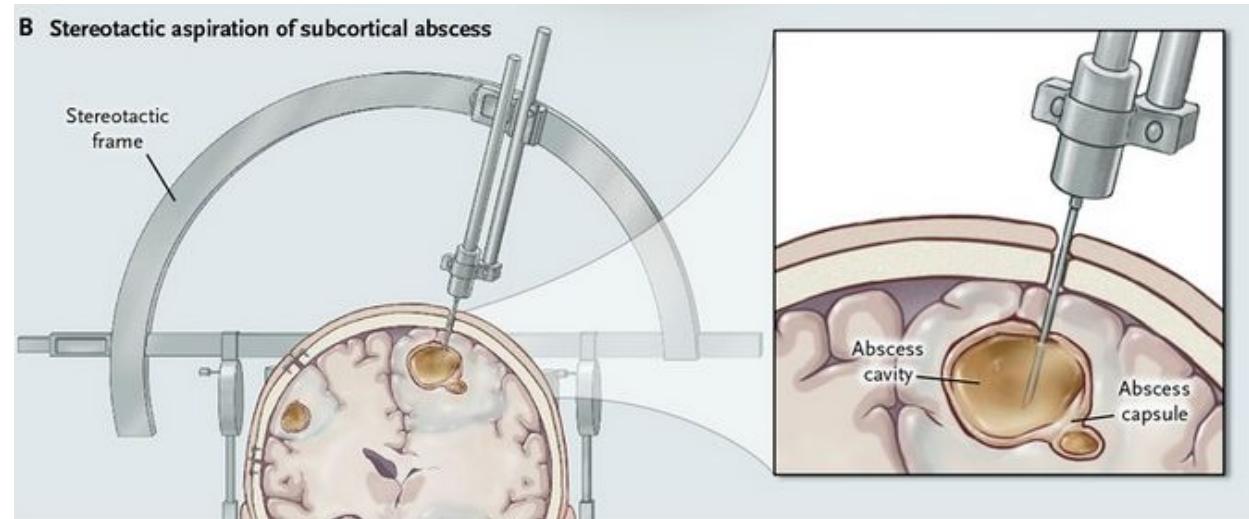
BRAIN ABSCESS

Treatment

- IV antibiotic: PenG + Chloramphenicol or Metronidazole

For MSSA: Nafcillin or Oxacillin

- Surgery
- Aspiration
- Glucocorticoids: dexamethasone



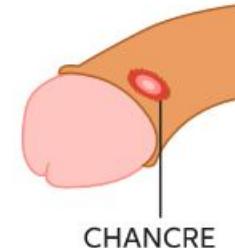
NEUROSYPHILIS

Neurosypilis is caused by *Treponema pallidum*

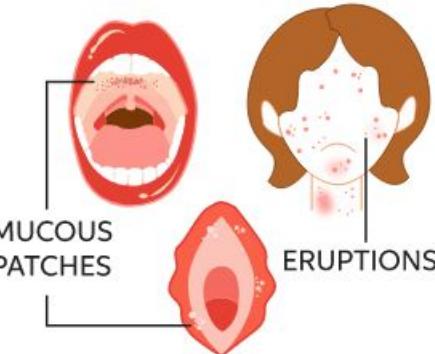
There are different forms of neurosyphilis:

- asymptomatic neurosyphilis
- meningeal neurosyphilis
- meningovascular neurosyphilis
- general paresis
- tabes dorsalis

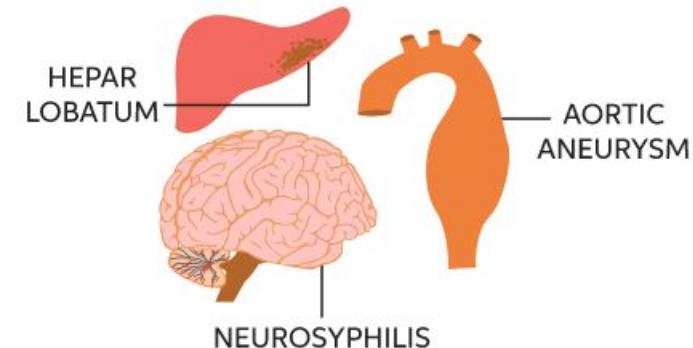
Primary syphilis



Secondary syphilis



Tertiary syphilis



NEUROSYPHYLIS

Early

- Asymptomatic neurosyphilis
- Acute symptomatic syphilitic meningitis: nausea, vomiting, headache, CN 2,4-8 abnormalizes
- Meningovascular syphilis

5-6 years after infection

Focal neurologic signs, vasculitis, stroke, transverse myelitis

Late

- Dementia paralytica
10-20 years after infection
Slow cognitive decline, weakness, tremor, pupillary abnormalities, bowel-bladder incontinence
- Tabes dorsalis
15-20 years after infection
Radicular paresthesia, “thunder bolt” pain in limbs, back or face; broad-based, foot-slapping gain, loss of reflexes in lower limbs, Argyll-Robertson pupils

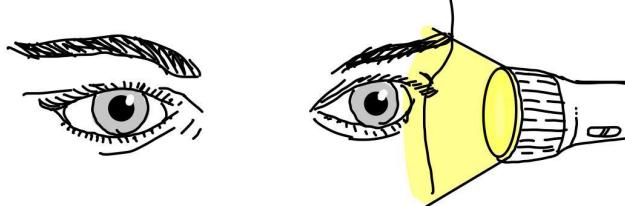
NEUROSYPHILIS

Tabes Dorsalis morphology:

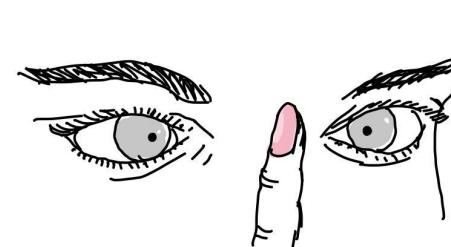
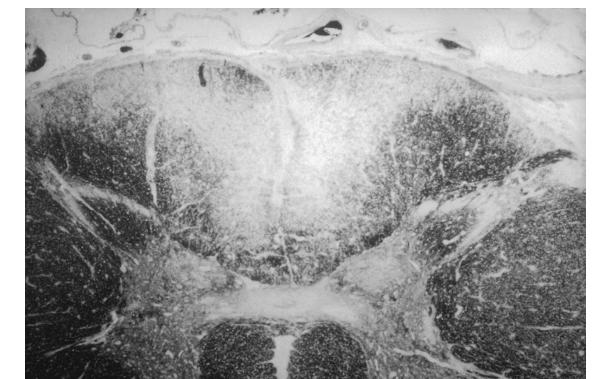
- Dorsal column degeneration
- Orthopedic pain (Charcot joints)
- Reflexes decreased (deep tendon)
- Shooting pain
- Argyll-Robertson pupils
- Locomotor ataxia
- Impaired proprioception
- Syphilis



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Pupils DO **NOT** constrict when exposed to bright light. ("light reflex")



Pupils DO constrict on a near object.
("accommodation reflex")

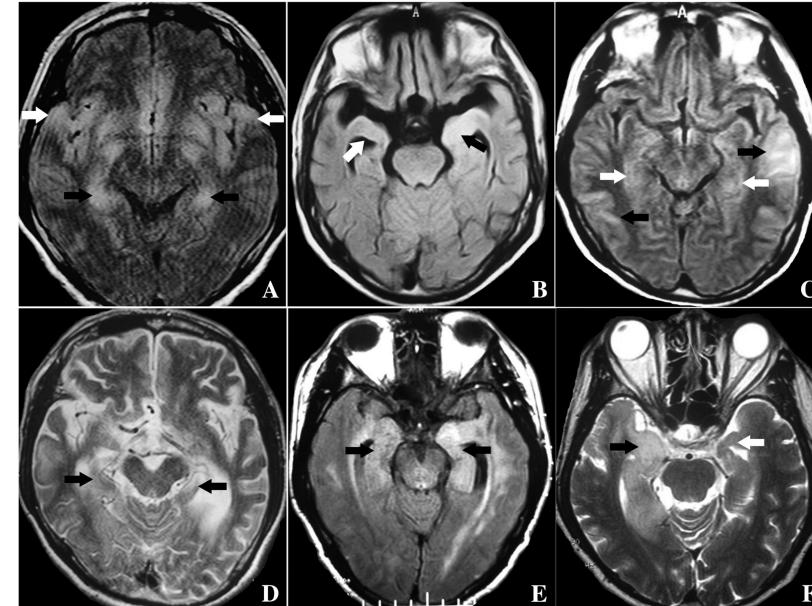
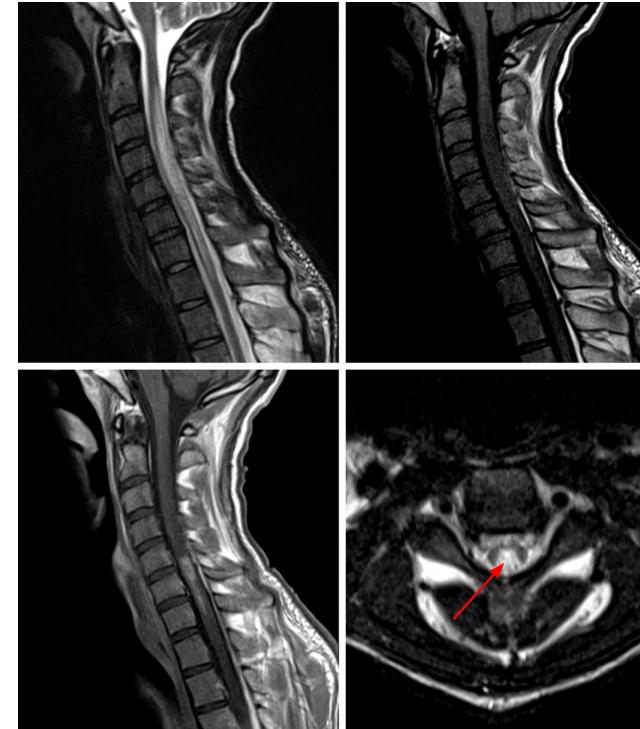
NEUROSYPHILIS

Diagnosis

- Serum nontreponemal tests : RPR, VDRL

Nonreactive in late neurosyphilis

- Serum treponemal test: FTA-ABS, TPA or syphilis EIA
- LP: lymphocytic pleocytosis , high protein, low or NM glucose, reactive csf-VDRL



NEUROSYPHILIS

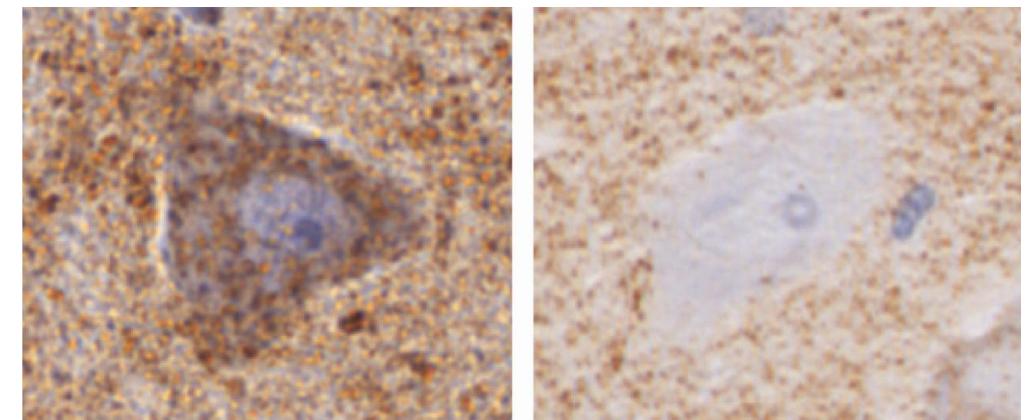
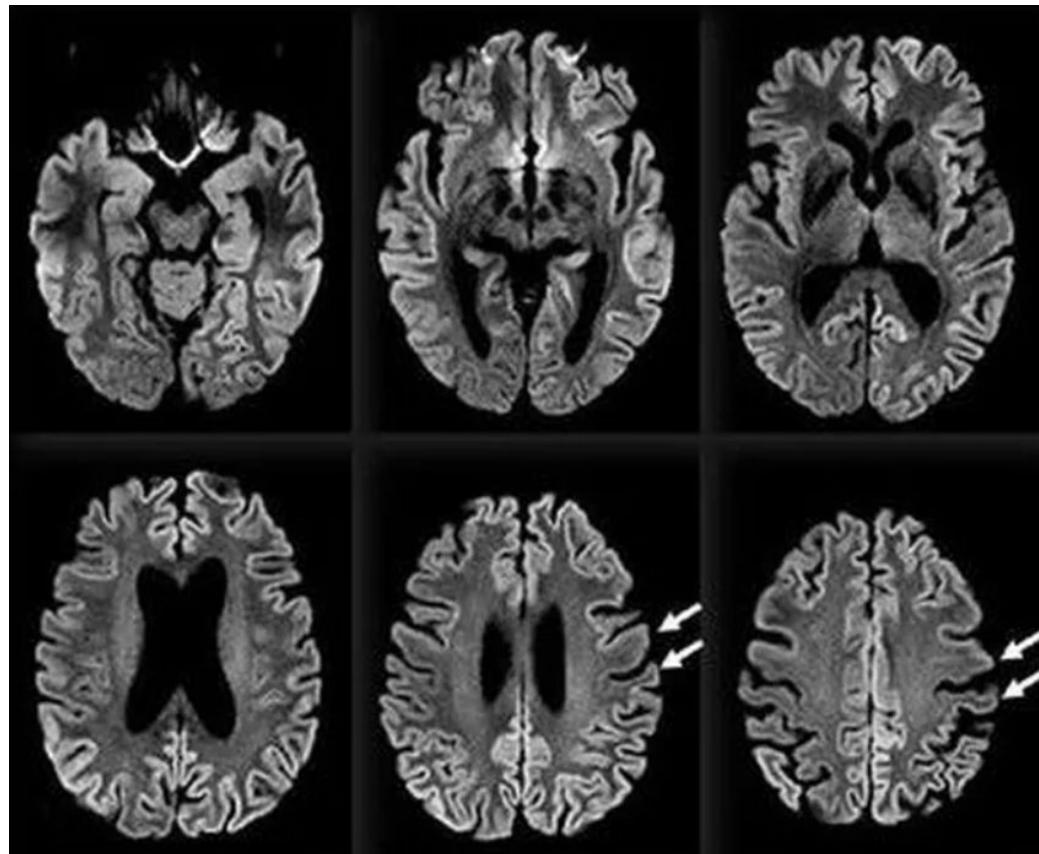
Treatment

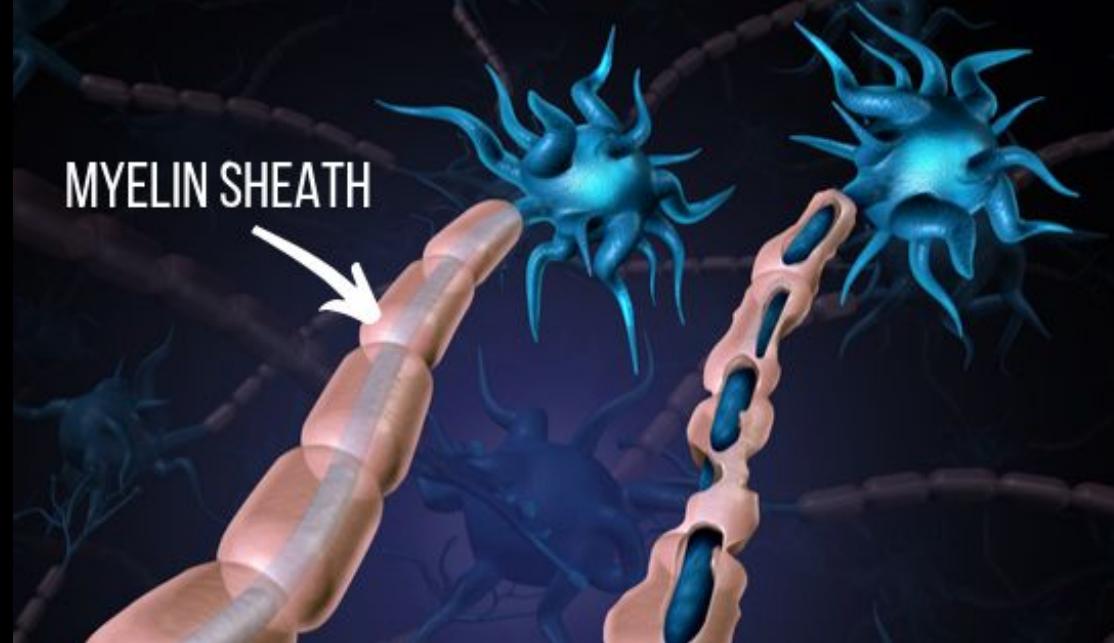
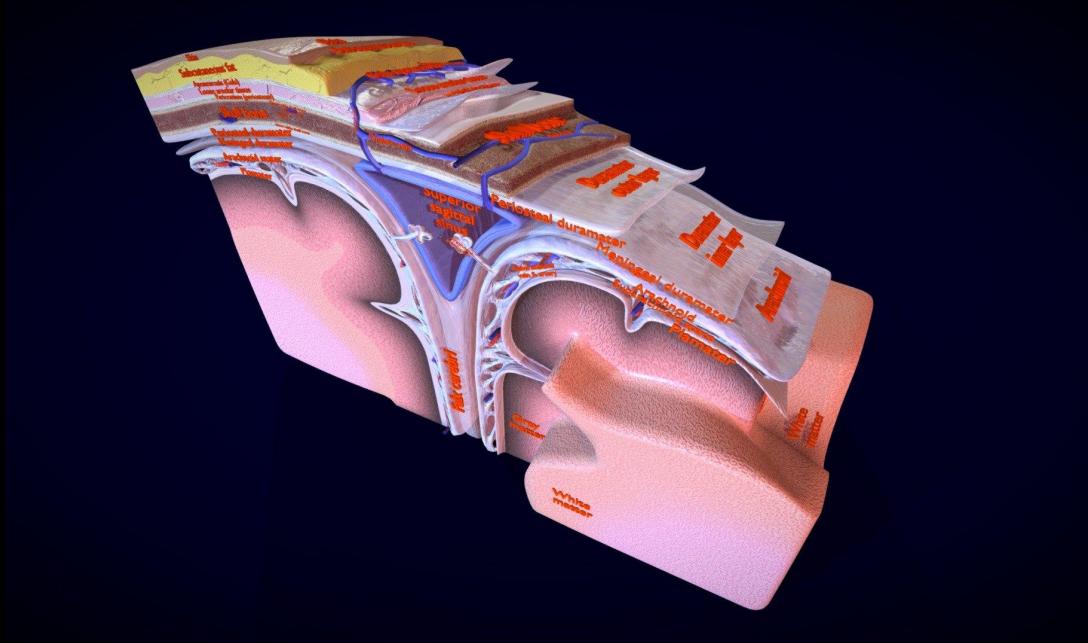
- Aqueous crystalline **penicillin G** (18 to 24 million units per day, administered as 3 to 4 million units intravenous every four hours, or 18 to 24 million units daily as a continuous infusion) for 10 to 14 days, or
- Procaine **penicillin G** (2.4 million units intramuscular [IM] once daily) plus **probenecid** (500 mg orally four times a day), both for 10 to 14 days
- Ceftriaxone 2 g IV daily 10-12 days

CREUTZFELDT-JAKOB DISEASE

- CJD is a neurodegenerative disease with a rapid onset characterized by progressive dementia, myoclonus and also cerebellar, pyramidal and extrapyramidal signs.
- Abnormal prion protein accumulate in the brain and it can cause irreversible damage. It lead to brain atrophy or wasting; cytoplasmic vacuoles in neurons and astrocytes
- Symptoms: fatigue, sleep problems, reduces appetite; dementia, behavior changes and confusion; cerebellar ataxia, aphasia, visual disturbances and motor weakness
- Diagnostic: exclude infection and toxicity. Brain biopsy
- Treatment: no cure

CREUTZFELDT-JAKOB DISEASE





THANK YOU