

Approach to Neurological Examination

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Basics of examination

- Neurological examination starts during the history taking itself.
- Hypotheses about lesion localization, neurological systems involved, and pathology of the disorder can be formed once the history is complete.
- The neurologist then uses the examination findings to confirm the localization of the lesion before trying to determine its cause.
- The full neurological examination is too lengthy to perform in practice.
- This approach should confirm, refute, or modify the initial hypotheses of disease location and causation derived from the history.
- Both the presence and absence of abnormalities may be of diagnostic importance.

Basics of examination

The screening neurological examination is designed for quick evaluation of the mental status, cranial nerves, motor system (strength, muscle tone, presence of involuntary movements, and postures), coordination, gait and balance, tendon reflexes, and sensation.

We couldn't perform a neurologic exam since the patient could not cooperate.

The initial question is whether the disease is in the brain, spinal cord, peripheral nerves, neuromuscular junctions, or muscles.

Screening Neurologic Exam

- More complex functions are tested first; if these are performed well, then it may not be necessary to test the component functions.
- Eg- the patient who can walk heel to toe (tandem gait) does not have a significant disturbance of the cerebellum or of joint position sensation.
- Patient who can do a pushup, rise from the floor without using the hands, and walk on toes and heels will have normal limb strength when each muscle group is individually tested.
- Asking the patient to hold the arms extended in supination in front of the body with the eyes open allows evaluation of strength and posture.
- It also may reveal involuntary movements such as tremor, dystonia, myoclonus, or chorea. A weak arm is expected to show a downward or pronator drift.
- Repeating the maneuver with the eyes closed allows assessment of joint position sensation.

Screening Neurologic Exam

- Of importance, the screening neurological examination may miss important neurological abnormalities.
- Moreover, what is abnormal in young adults may be normal in the elderly. Loss of the ankle reflex and loss of vibration sense at the big toe are common findings in patients older than 70 years.
- Isolated deviation of the tongue or uvula to one side and minor asymmetries of reflexes or sensation – soft signs
- At the end of the neurological examination, the abnormal physical signs should be classified as definitely abnormal (hard signs) or equivocally abnormal (soft signs).


Screening Neurologic Exam

- Loss of pain and temperature sensation on one half of the body, excluding the face, indicates a lesion of the contralateral spinothalamic tract in the high cervical spinal cord.
- A left sixth nerve palsy, with weakness of left face and right limbs, points to a left pontine lesion.
- A left homonymous hemianopia indicates a lesion in the right optic tract, optic radiations, or occipital cortex.

Mini-Mental State Examination (MMSE)

Patient's Name: _____ Date: _____

Instructions: Ask the questions in the order listed. Score one point for each correct response within each question or activity.

Maximum Score	Patient's Score	Questions
5		"What is the year? Season? Date? Day of the week? Month?"
5		"Where are we now: State? County? Town/city? Hospital? Floor?"
3		The examiner names three unrelated objects clearly and slowly, then asks the patient to name all three of them. The patient's response is used for scoring. The examiner repeats them until patient learns all of them, if possible. Number of trials: _____
5		"I would like you to count backward from 100 by sevens." (93, 86, 79, 72, 65, ...) Stop after five answers. Alternative: "Spell WORLD backwards." (D-L-R-O-W)
3		"Earlier I told you the names of three things. Can you tell me what those were?"
2		Show the patient two simple objects, such as a wristwatch and a pencil, and ask the patient to name them.
1		"Repeat the phrase: 'No ifs, ands, or buts.'"
3		"Take the paper in your right hand, fold it in half, and put it on the floor." (The examiner gives the patient a piece of blank paper.)
1		"Please read this and do what it says." (Written instruction is "Close your eyes.")
1		"Make up and write a sentence about anything." (This sentence must contain a noun and a verb.)
1		"Please copy this picture." (The examiner gives the patient a blank piece of paper and asks him/her to draw the symbol below. All 10 angles must be present and two must intersect.) 
30		TOTAL

(Adapted from Rovner & Folstein, 1987)

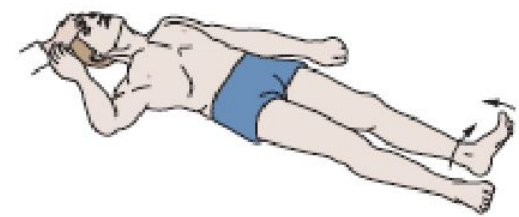
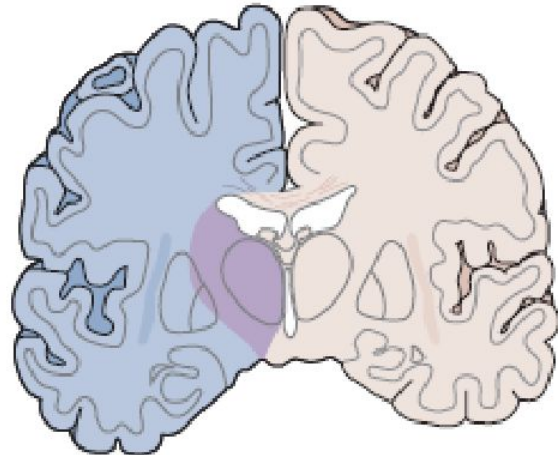
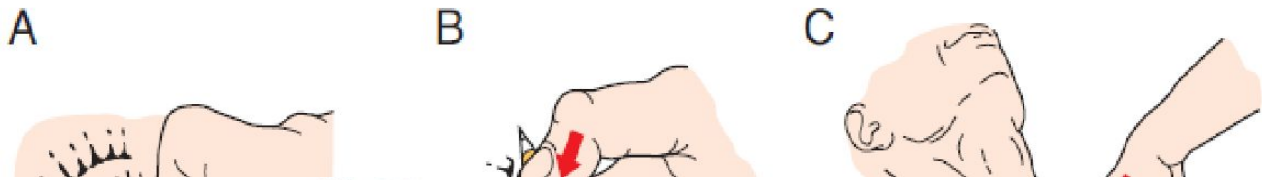
Examination



- Mental Status Assessed while recording the history
- **Cranial nerves**
- CN I Should be tested in all persons who experience spontaneous loss of smell, in patients suspected to have Parkinson disease, and in patients who have suffered head injury
- CN II Each eye:
 - Visual acuity with glasses/contacts
 - Visual fields by confrontation
 - Swinging flashlight to detect relative afferent pupillary defect
 - Fundoscopy
- CN III, IV, VI:
 - Horizontal and vertical eye movements (saccades, pursuit, vestibulo-ocular reflex)
 - Pupillary symmetry and reactivity
 - Presence of nystagmus or other ocular oscillations
- CN V Pinprick and touch sensation on face, corneal reflex Jaw strength
- CN VII Close eyes, show teeth

Examination

- CN VIII Perception of whispered voice in each ear or rubbing of fingers; if hearing is impaired, look in external auditory canals, and use tuning fork for lateralization and bone-versus-air sound conduction
- CN IX, X Palate lifts in midline, gag reflex present
- CN XI Shrug shoulders
- CN XII Protrude tongue



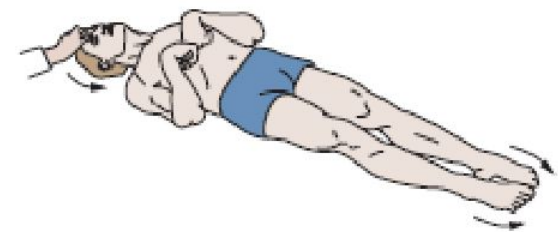
Eye Response

- 4 = eyes open spontaneously
- 3 = eye opening to verbal con
- 2 = eye opening to pain
- 1 = no eye opening

Motor Response

- 6 = obeys commands
- 5 = localizing pain
- 4 = withdrawal from pain
- 3 = flexion response to pain
- 2 = extension response to pain
- 1 = no motor response

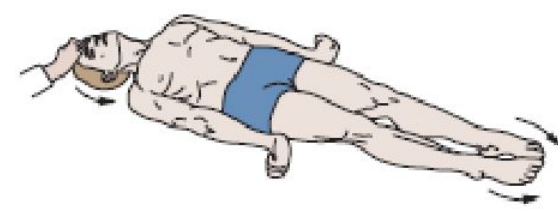
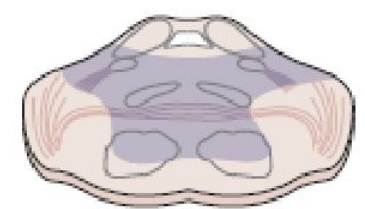
B Upper midbrain damage



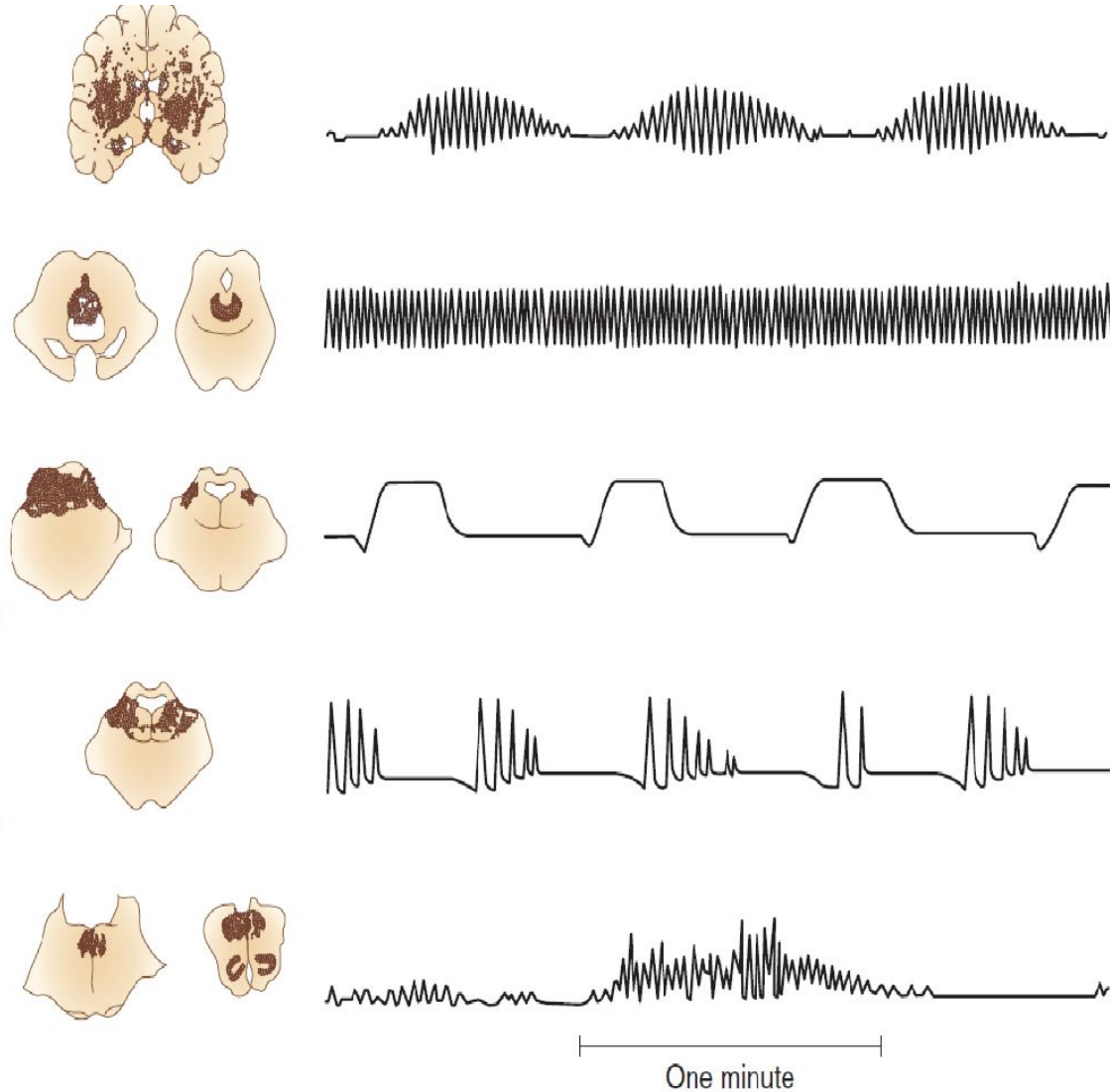
Verbal Response

- 5 = oriented
- 4 = confused
- 3 = inappropriate words
- 2 = incomprehensible sounds
- 1 = no verbal response

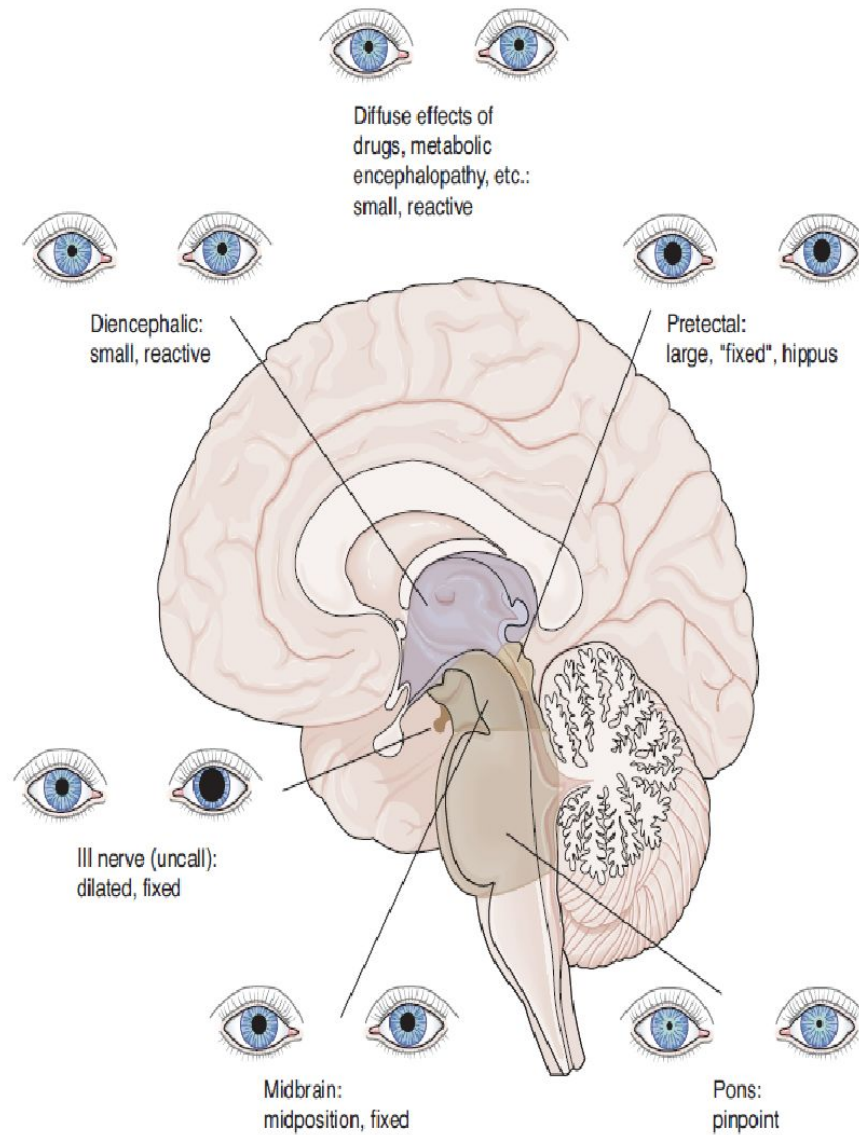
C Upper pontine damage



Patterns of breathing



Pupillary examination



- (i) Coma with meningism (with or without intact brainstem function and lateralizing signs)
- (ii) Coma with signs of focal brainstem dysfunction
- (iii) Coma with intact brainstem function and lateralizing signs
- (iv) Coma with intact brainstem function, no meningism, and no lateralizing signs

Coma with meningism

- Infection: meningitis, encephalitis, malaria, HIV disease
- Vascular: subarachnoid haemorrhage

Coma with signs of focal brainstem dysfunction

- Herniation syndromes
- Intrinsic brainstem dysfunction
- Vascular: vertebrobasilar occlusion, dissection, haemorrhage, arteriovenous malformation
- Mass lesions: posterior fossa tumours, abscesses, tuberculosis.
- Traumatic brain injury
- Advanced metabolic/ toxic encephalopathy
- Others: central pontine myelinolysis, brainstem encephalitis, leukoencephalopathy

Coma with intact brainstem function and lateralizing signs

- Vascular:
 - Infarction - ischaemic, thromboembolic, or hypoperfusion
 - Haemorrhage - epidural, subdural, subarachnoid, and intracerebral
 - Vasculitis
 - Venous thrombosis
- Traumatic brain injury
- Infection: brain abscess, subdural empyema, Creutzfeldt–Jakob disease, malaria, HIV disease, endocarditis
- Cerebral neoplasm

Coma with intact brainstem function, no meningism, and no lateralizing signs

- Alcohol
- Drugs and toxins: sedatives, anaesthetic agents, opioids, amphetamines, barbiturates, salicylates, organophosphorus poisoning, carbon monoxide, methanol, lead, cyanide
- Seizure: convulsive and non-convulsive status epilepticus, post-ictal states
- Hypoxic: ischaemic encephalopathy
- Endocrine: hypo and hyperthyroidism, Addison's disease, hypopituitarism, diabetic ketoacidosis, hyperosmolar non-ketotic coma
- Metabolic: hypo and hyperglycaemia, sepsis, electrolyte disturbance—hypo and hypernatraemia, hypercalcaemia, hepatic failure, encephalopathy, renal failure
- Respiratory: hypoxaemia, hypercarbia
- Other: porphyria, Reye's syndrome, mitochondrial disease, inborn errors of metabolism NMDA-receptor antibody encephalitis

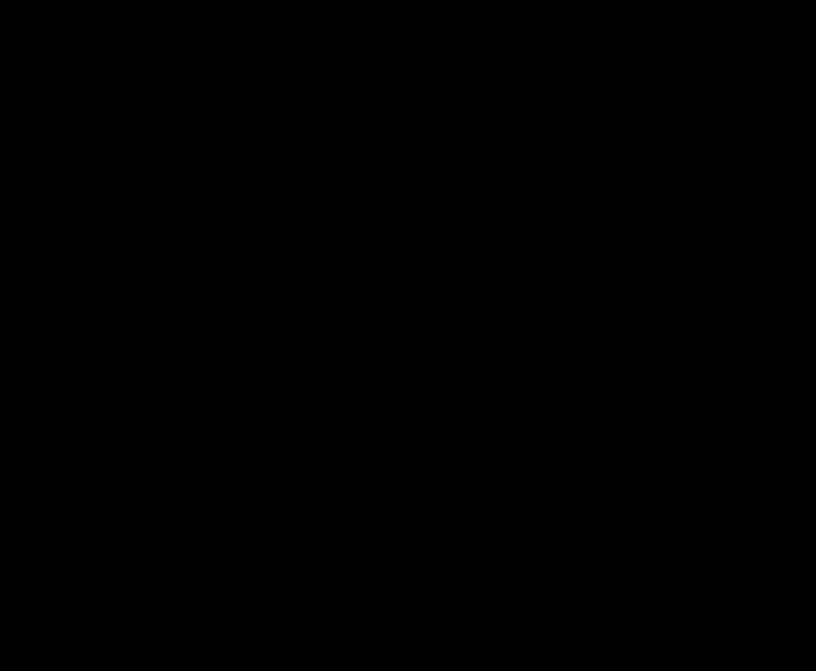
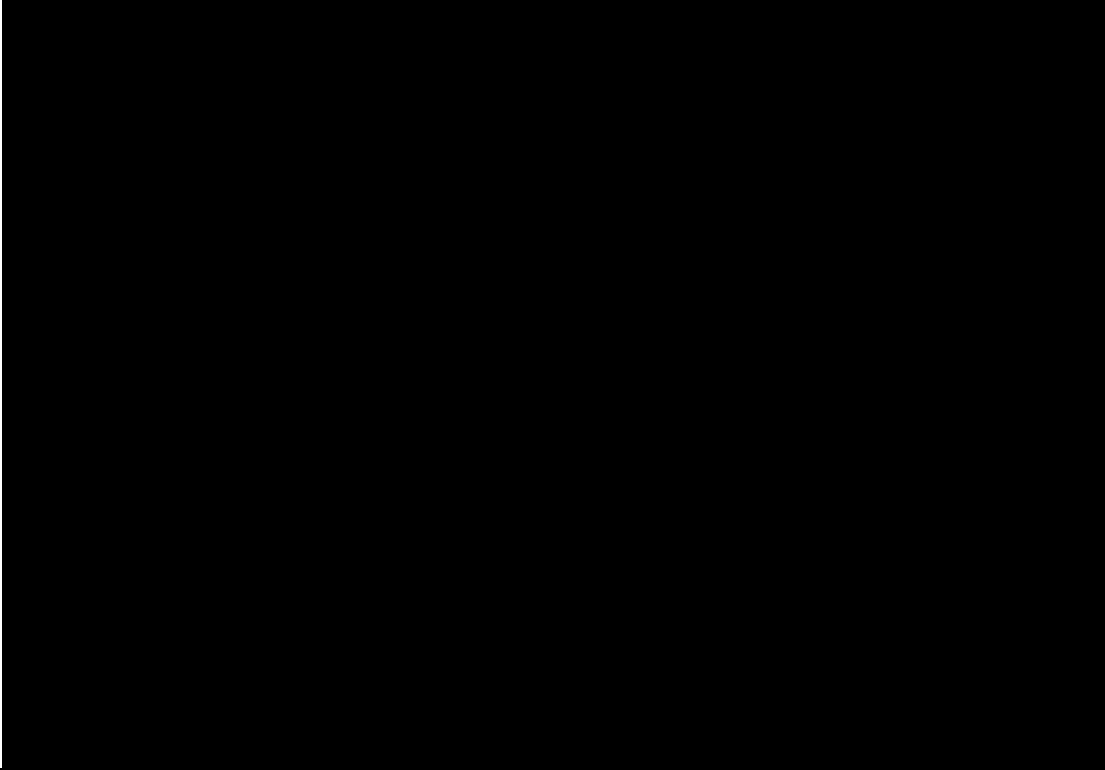


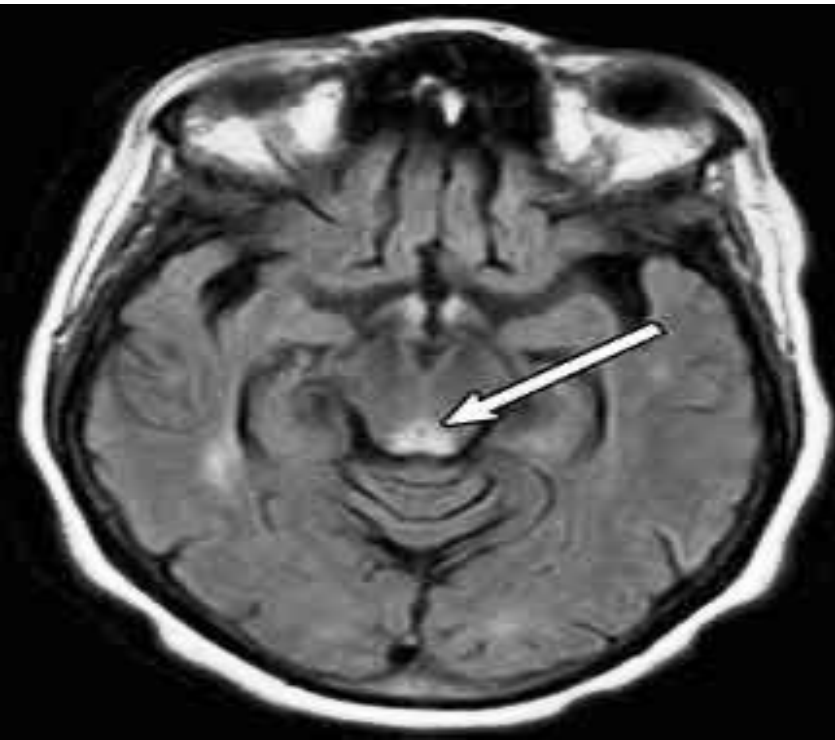
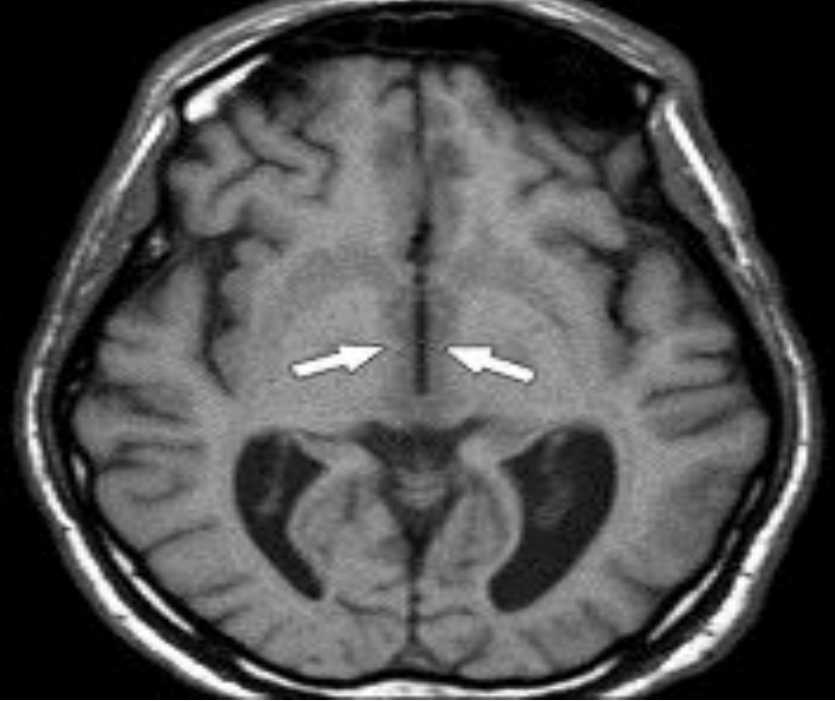
Table 1-1 Cause of Stupor or Coma in 500 Patients Initially Diagnosed as "Coma of Unknown Etiology"*

	Subtotals		Subtotals
I. Supratentorial lesions	101	B. Destructive or ischemic lesions	53
A. Rhinencephalic and subcortical destructive lesions	2	1. Pontine hemorrhage	11
1. Thalamic infarcts	2	2. Brainstem infarct	40
B. Supratentorial mass lesions	99	3. Basilar migraine	1
1. Hemorrhage	76	4. Brainstem demyelination	1
a. Intracerebral	44	III. Diffuse and/or metabolic brain dysfunction	326
(1) Hypertensive	36	A. Diffuse intrinsic disorders of brain	38
(2) Vascular anomaly	5	1. "Encephalitis" or encephalomyelitis	14
(3) Other	3	2. Subarachnoid hemorrhage	13
b. Epidural	4	3. Concussion, nonconvulsive seizures, and postictal states	9
c. Subdural	26	4. Primary neuronal disorders	2
d. Pituitary apoplexy	2		
2. Infarction	9		

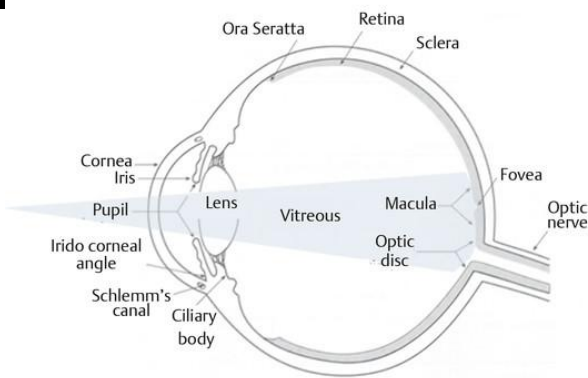
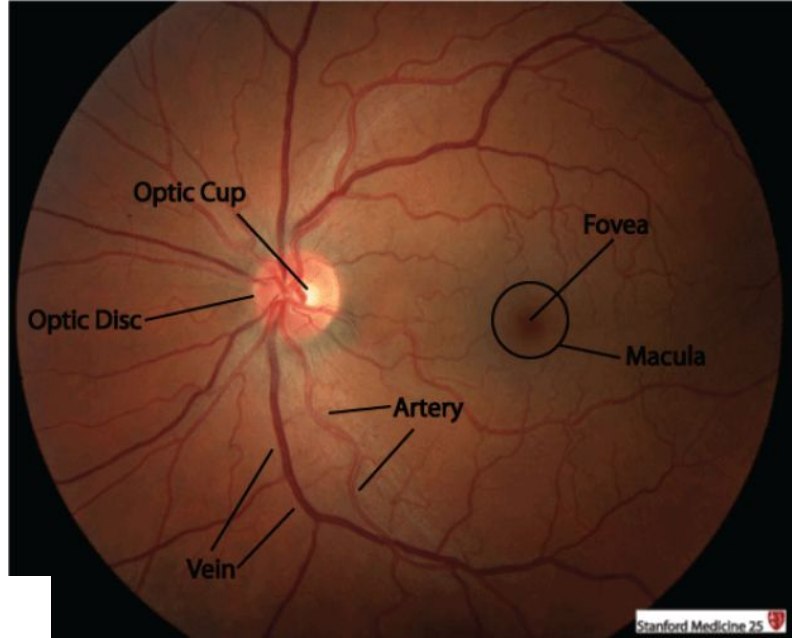
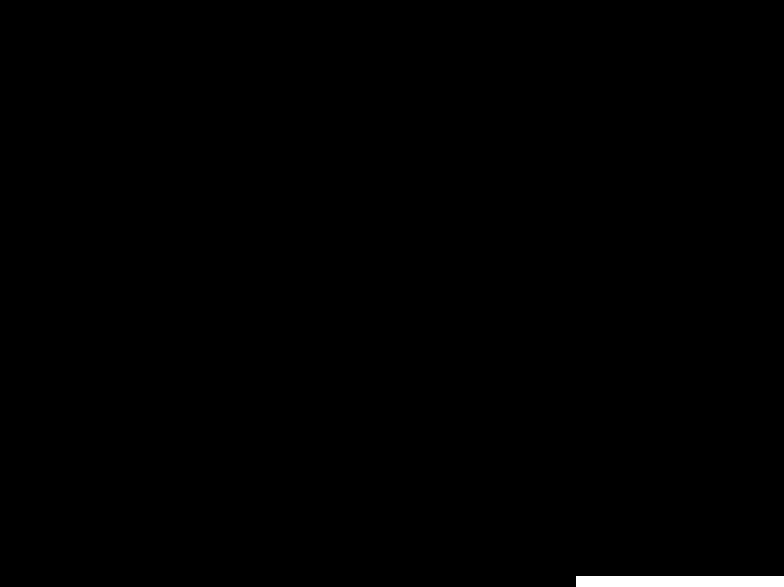
More than half of all cases of coma are due to diffuse and metabolic brain dysfunction

Plum and Posner's diagnosis of stupor and coma, 2007

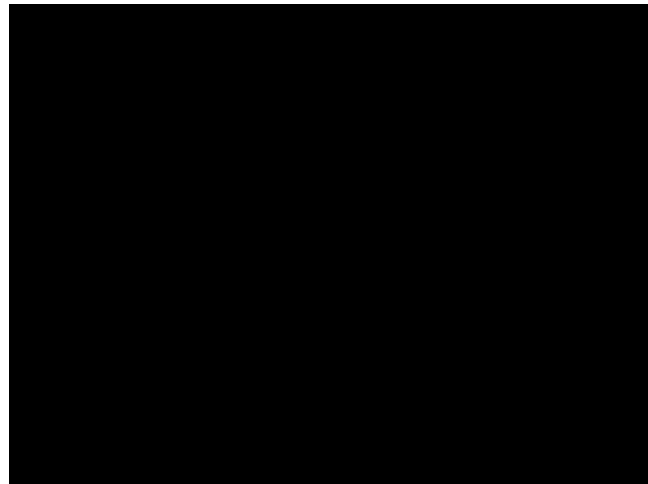
3. Closed head injury	1	8. Remote effects of trauma	0
II. Subtentorial lesions	65	9. Drug poisons	149
A. Compressive lesions	12	10. Ionic and acid-base disorders	12
1. Cerebellar hemorrhage	5	11. Temperature regulation	9
2. Posterior fossa subdural or extradural hemorrhage	1	12. Mixed or nonspecific metabolic coma	1
3. Cerebellar infarct	2	IV. Psychiatric "coma"	8
4. Cerebellar tumor	3	A. Conversion reactions	4
5. Cerebellar abscess	1	B. Depression	2
6. Basilar aneurysm	0	C. Catatonic stupor	2



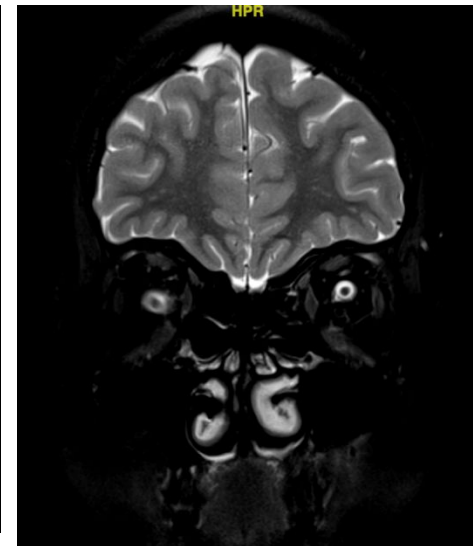
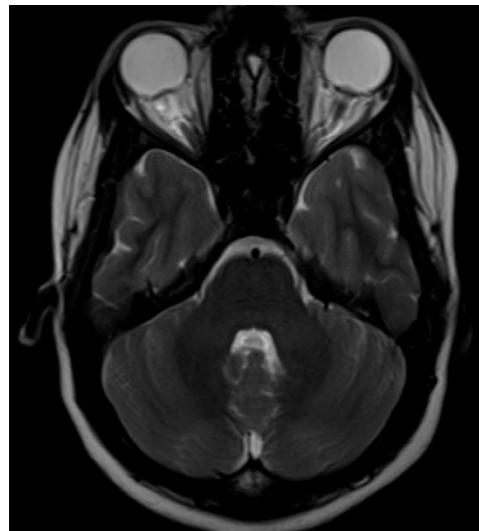
35 year lady, 2 months gestation presented with h/o acute onset of memory disturbances. She was previously diagnosed with hyperemesis gravidarum and managed accordingly.



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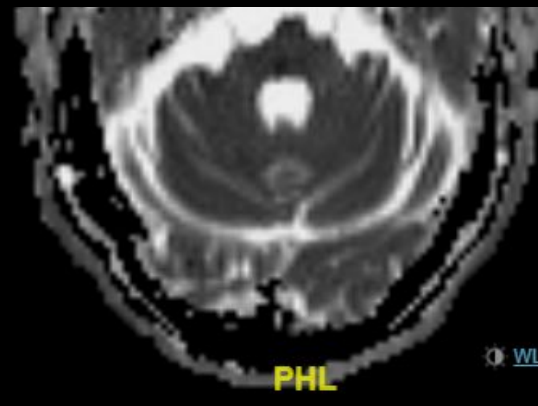
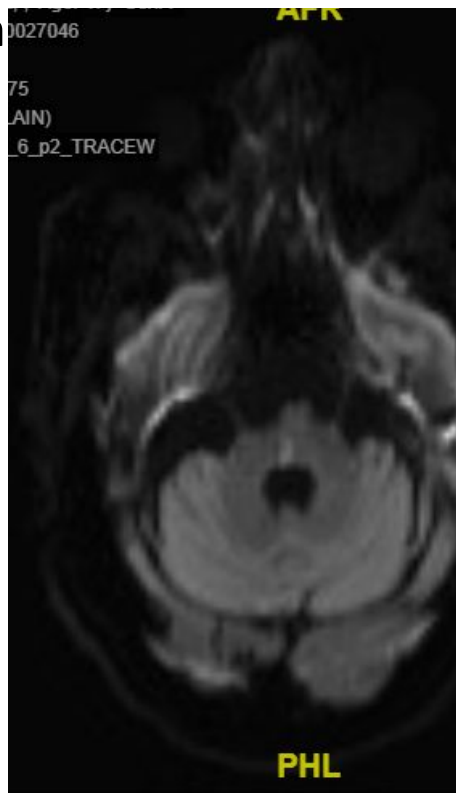


- A 18 year-old woman with a history of obesity and PCOS presented to our emergency department complaining of bifrontal headache for one month associated with nausea, vomiting, transient visual disturbances, and Pulsatile tinnitus.



A 65 year old lady presented with sudden onset of blurring of vision of two days duration.

O/E- her visual acuity was normal. Eye movements were examined

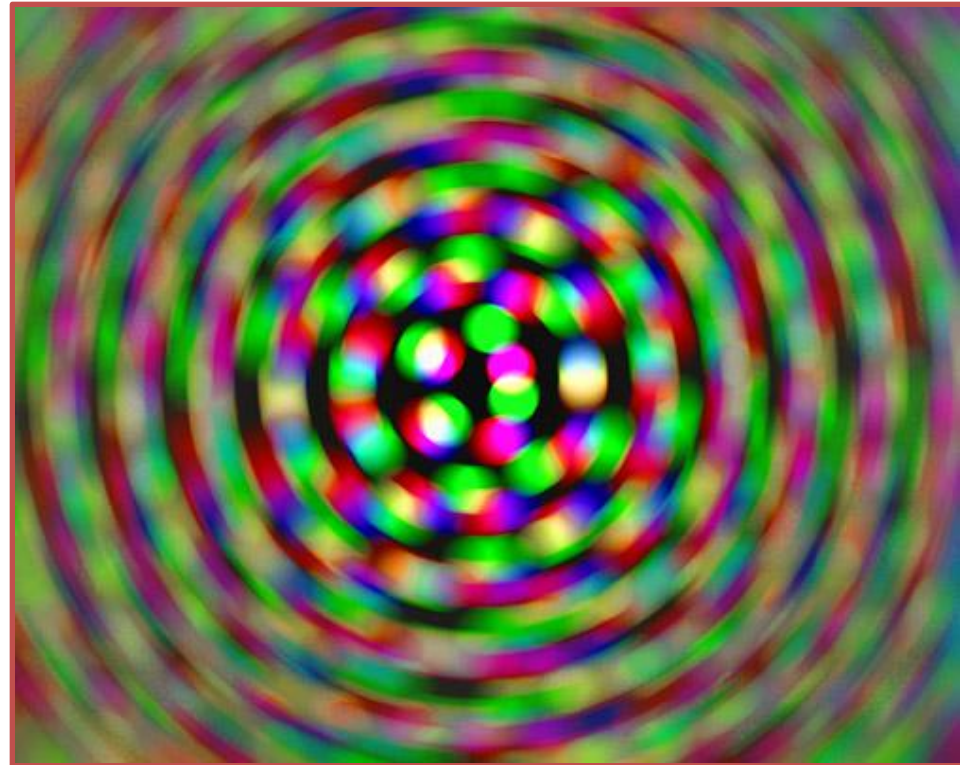


Optic Neuritis and MS Do

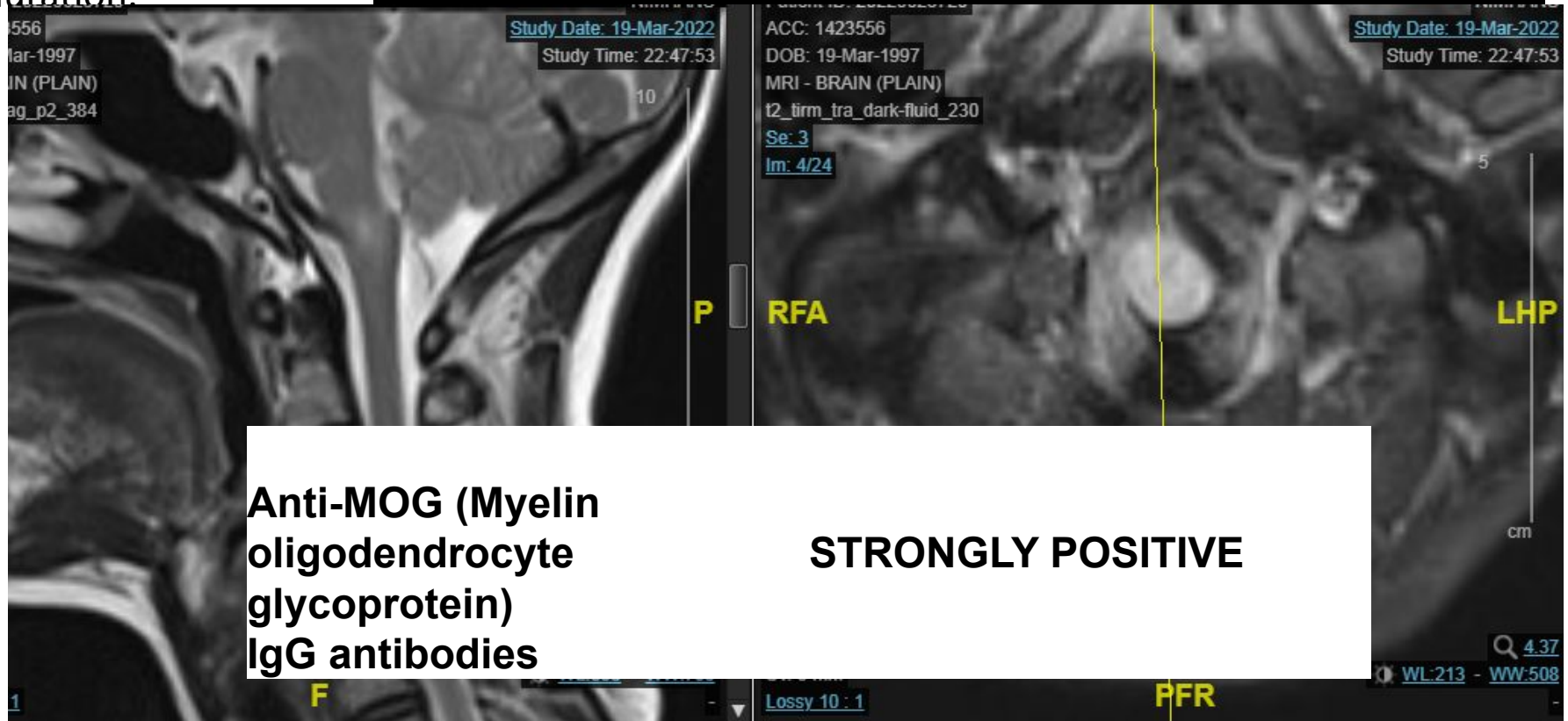


Oscillopsia

- Both nystagmus and saccadic dyskinesia move the fovea off target, which results in a decrease in vision.
- They also cause oscillopsia, an **illusion of motion in the stationary environment**.



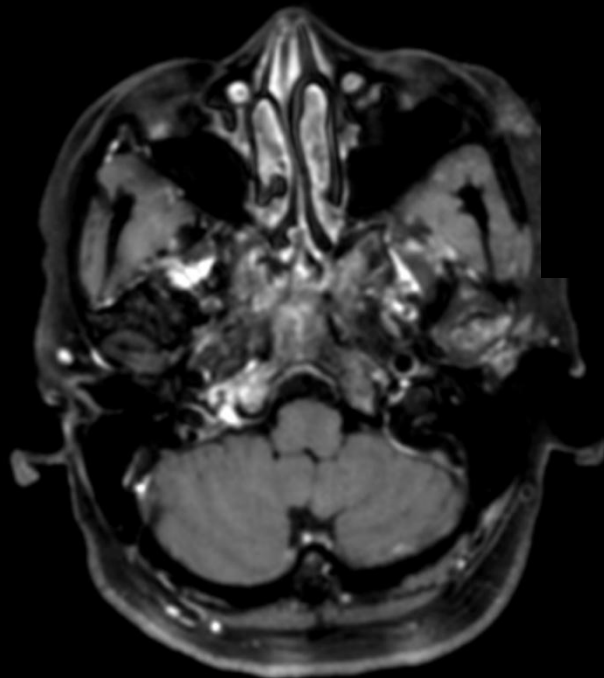
A 25 year old lady came with blurring of vision of 7 days duration followed by weakness of all 4 limbs of three days duration.



Anti-MOG (Myelin oligodendrocyte glycoprotein) IgG antibodies

STRONGLY POSITIVE

A 41 year old lady with recurrent neurological illness since 2019, came with slurring of speech, difficulty swallowing liquids with nasal regurgitation.



RHP

5

LFA

cm

An 18 year old girl with history of swaying while walking since 10 years of age and frequent falls. History of similar illness in father.

She has scanning dysarthria and cerebellar signs on exam.

Genetics s/o SCA-1

