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A gaze palsy is a limitation of extraocular motility that has two specific characteristics—what are they?

- --?
- --?



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--The limitation must involve only one eye, or both

--?





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It is a gaze palsy

A gaze palsy is a limitation of extraocular motility that has two specific characteristics—what are they?

- --The limitation must involve both eyes
- --The limitation is in the same direction of gaze bilaterally (eg, both eyes can't look up; both eyes can't look to the left; etc)



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Before we get into Parinaud syndrome specifically, let's first create a frame for thinking about extraocular motility limitation generally.

't look up;



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It is a gaze palsy

A gaze palsy is a limitation of extraocular motility that has two specific characteristics—what are they?

Before we get into Parinaud syndrome specifically, let's first create a frame for thinking about extraocular motility limitation generally. Once we have this general understanding we will see where Parinaud syndrome fits within it.

't look up;

Q

# **Parinaud Syndrome**



CN... CN...

Which cranial nerves innervate the extraocular muscles (EOMs)?





CN3 CN6 CN4

Which cranial nerves innervate the extraocular muscles (EOMs)?

Q

#### **Parinaud Syndrome**



CN3	CN6	CN4
?	?	?

What is the name for the collections of neurons that give rise to each of these cranial nerves? (This is not a trick question--the answer is as obvious as it seems.)



CN3 Nucleus CN6 Nucleus CN4 Nucleus

What is the name for the collections of neurons that give rise to each of these cranial nerves? (This is not a trick question--the answer is as obvious as it seems.)



Nuclear

CN3 Nucleus CN6 Nucleus CN4 Nucleus

As will be apparent shortly, the 'nuclear level' is the locus around which we organize our understanding of extraocular motility control (and limitations thereof)

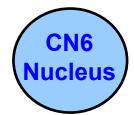


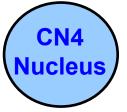
?

?

Nuclear

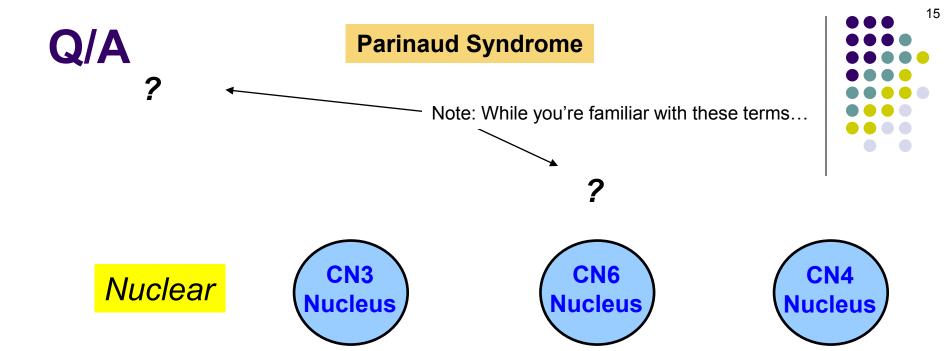
CN3 Nucleus





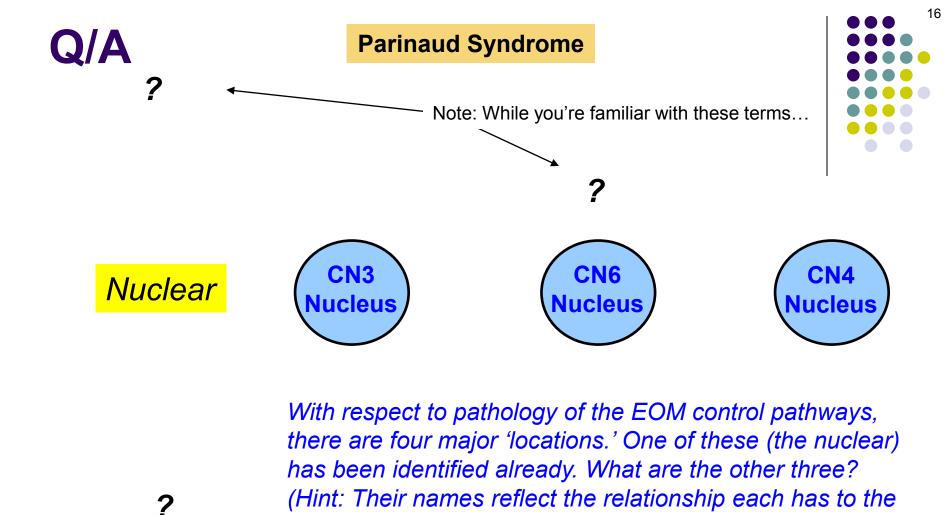
?

With respect to pathology of the EOM control pathways, there are four major 'locations.' One of these (the nuclear) has been identified already. What are the other three? (Hint: Their names reflect the relationship each has to the nuclear level.)



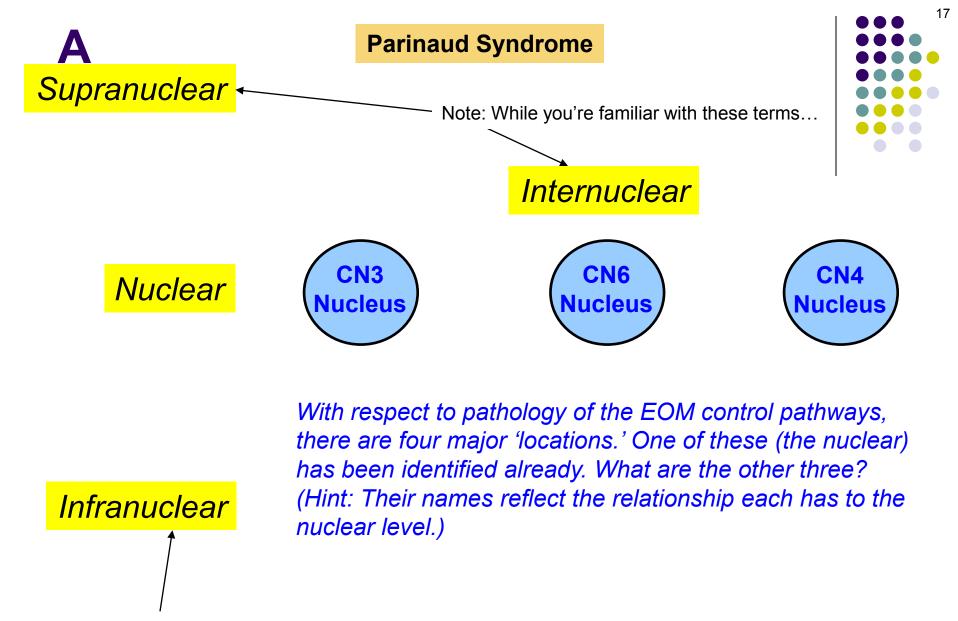
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...you may not be with this one, although you'll agree it makes sense in context. (Further, and importantly, it is used in the BCSC *Neuro* book.)

nuclear level.)



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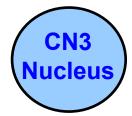
# Supranuclear

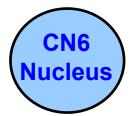
The *supranuclear pathways* consists of inputs to the nuclei from centers in the cortex, cerebellum, vestibular system, etc.



Internuclear







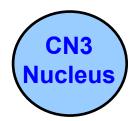


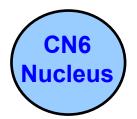
# Supranuclear

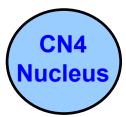
The *supranuclear pathways* consists of inputs to the nuclei from centers in the cortex, cerebellum, vestibular system, etc. These locations are 'supra' in that they carry signals *to* the nuclei.









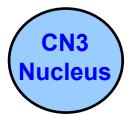






Internuclear

Nuclear







Are lesions of the CN3 nucleus commonly encountered in clinical practice?





Internuclear

Nuclear







Are lesions of the CN3 nucleus commonly encountered in clinical practice?

No, they are rare



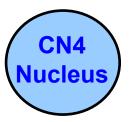


Internuclear

Nuclear







Are lesions of the CN4 nucleus commonly encountered in clinical practice?



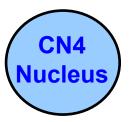


Internuclear

Nuclear







Are lesions of the CN4 nucleus commonly encountered in clinical practice?

No, these are even rarer

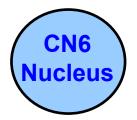




Internuclear

Nuclear







Are lesions of the CN6 nucleus commonly encountered in clinical practice?

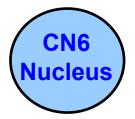




Internuclear

Nuclear







Are lesions of the CN6 nucleus commonly encountered in clinical practice?

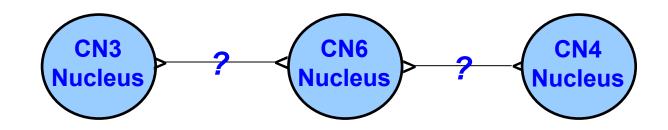
While not common, they are a well-known clinical entity





# Internuclear

Nuclear

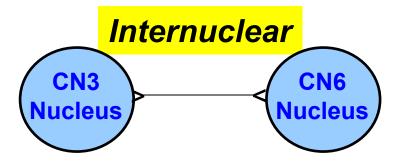


Which two nuclei share an internuclear connection that is of well-established clinical importance?





Nuclear





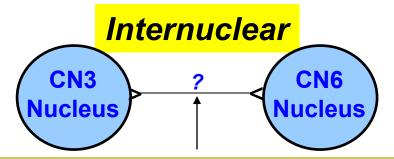
Which two nuclei share an internuclear connection that is of well-established clinical importance?

3 and 6





Nuclear



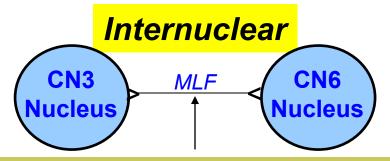
CN4 Nucleus

What is the name of the internuclear pathway connecting the CN3 and CN6 nuclei?





Nuclear



CN4 Nucleus

What is the name of the internuclear pathway connecting the CN3 and CN6 nuclei?
The medial longitudinal fasciculus (MLF)



Nuclear

Internuclear

CN3
Nucleus

MLF
CN6
Nucleus

CN4 Nucleus

*Infranuclear* 

The *infranuclear pathway* consists of everything below the nuclei: the axons as they run from the nuclei to the neuromuscular junction; the junction itself; and finally the EOMs themselves.

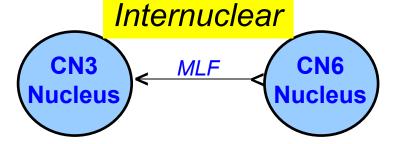
Extraocular muscle

(No question—proceed when ready)

# Supranuclear









*Infranuclear* 



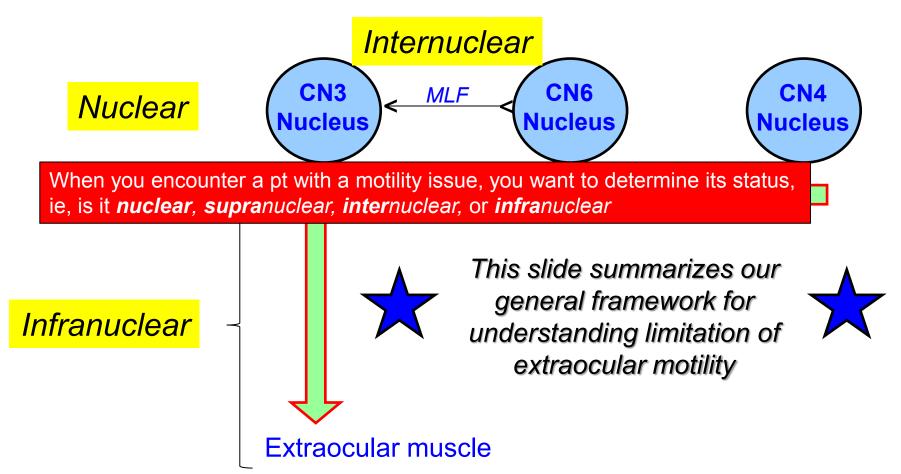
This slide summarizes our general framework for understanding limitation of extraocular motility



Extraocular muscle

(No question—proceed when ready)

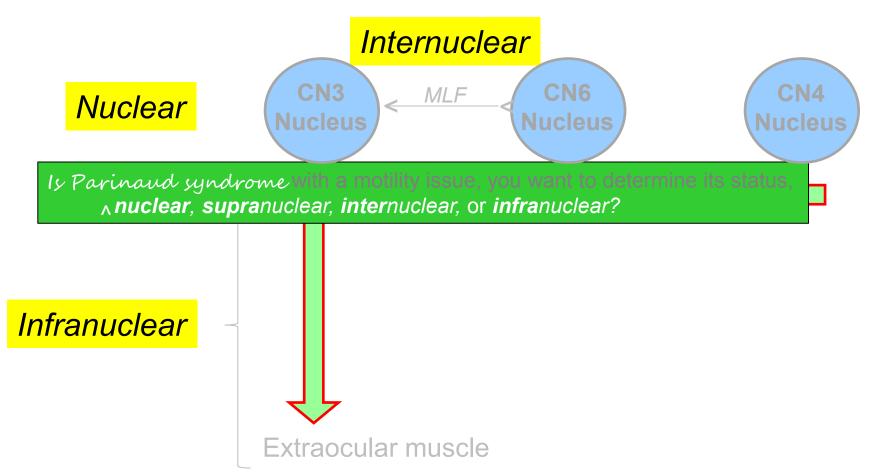




(No question—proceed when ready)

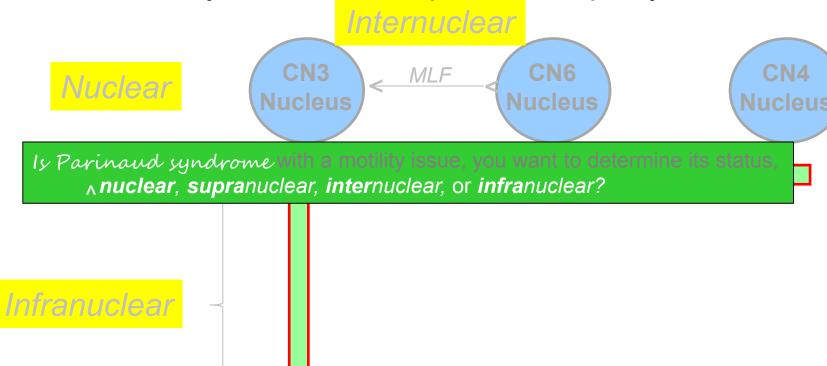








# Parinaud syndrome is a supranuclear palsy



Extraocular muscle



#### Daringud Syndromo

Internuclear

Recall we said Parinaud syndrome is a gaze palsy. Are all gaze palsies supranuclear?



Parinaud syndrome is a supranuclear palsy

Nuclear

CN3 Nucleus

MLF CN6
Nucleus

CN4 Nucleus

ly Parinaud syndrome with a motility issue, you want to determine its status ^nuclear, supranuclear, internuclear, or infranuclear?

Infranuclear

Extraocular muscle



#### Daringud Cundromo

Recall we said Parinaud syndrome is a gaze palsy. Are all gaze palsies supranuclear? No (although many are)



Parinaud syndrome is a supranuclear palsy

Nuclear

CN3 Nucleus

MLF CN6
Nucleus

CN4 Nucleus

Is Parinaud syndrome with a motility issue, you want to determine its status Anuclear, supranuclear, internuclear, or infranuclear?

Infranuclear

Extraocular muscle



- Parinaud syndrome has four main features:
  - ?
  - ?
  - ?
  - ?



- Parinaud syndrome has four main features:
  - Impaired upgaze
  - Lid retraction
  - Convergence-retraction nystagmus
  - Light-near dissociation



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Do all Parinaud pts manifest all four features?



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Is it common, or rare, for a pt to present with just a subset of them?





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Is it common, or rare, for a pt to present with just a subset of them?
It is fairly common



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Of the four, which is the one most likely to be present?





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Impaired upgaze





Impaired upgaze in Parinaud's



- Parinaud syndrome has four main features:
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  - Parinaud syndrome is characterized by tonic downward
  - displacement of the eyes, with impaired upgaze.

(No question yet—keep going)

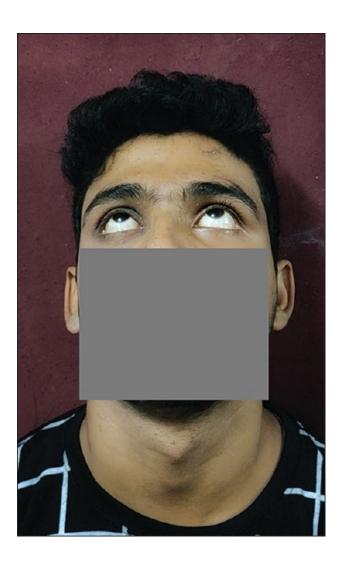


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Which two classes of drugs are most commonly implicated and which is number one?





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What is the classic cause of oculogyric crisis? A drug reaction

Which two classes of drugs are most commonly implicated and which is number one?
The neuroleptics (#1), and the antiemetics





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Which neuroleptic is most commonly implicated?

nonly implicated

and which is number one:





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Which neuroleptic is most commonly implicated? **Haloperidol** 

nonly implicated

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Which antiemetic is most commonly implicated?

Metoclopramide

nonly implicated



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• Impaired upgaze What is the eponymous name for lid retraction in Parinaud syndrome?

- Convergence-retraction mystaginus
- Light-near dissociation





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Lid retraction in Parinaud's





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Speaking of...The phrase *lid retraction* should bring to mind several conditions...

First, what is the most common cause of lid retraction?





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TED: Lid retraction





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Unilateral lid lag in TED. Note how the normal right upper lid has 'followed' the globe into downgaze



Bilateral lid lag in TED

TED: Lid lag





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What is the eponymous name for lid lag 2ndry to TED?





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There is a form of ptosis that has a strong association with lid lag—which one?





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There is a form of **ptosis** that has a strong association with lid lag—which one? Congenital myogenic ptosis





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In a nutshell, what is the pathogenesis of congenital myogenic ptosis?





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The levator fails to develop properly, with some or all of its muscle fibers replaced by fibrofatty tissue





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OK, I can see how a lack of functioning levator leads to ptosis, but why do these pts have lid lag?





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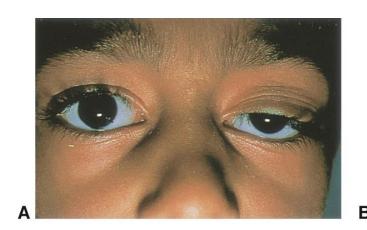
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Congenital myogenic ptosis. A, ptosis in primary. B, failed elevation in upgaze. C, lid lag in downgaze.

What well-known phenomenon associated with a cranial-nerve palsy frequently manifests with lid lag?



Thursid and da (TED). When you have 'lid retreation' thinks

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Aberrant regeneration of the right third nerve. A, In primary gaze, there is mild ptosis, pupillary mydriasis, and exotropia, all on the right.







Aberrant regeneration of the right third nerve. A, In primary gaze, there is mild ptosis, pupillary mydriasis, and exotropia, all on the right. B, With attempted downward gaze, the right eyelid retracts as fibers of the right third nerve supplying the inferior rectus now also innervate the levator muscle

What well-known phenomenon associated with a cranial-nerve palsy frequently manifests with lid lag?

#### **Aberrant regeneration after CN3 palsy**

More info: Slide-set N13

What the heck is aberrant regeneration?

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# These are some of the other conditions that must be considered when faced with a case of lid retraction

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--Compressive

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More info: Slide-set O5

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- Parinaud syndrome has four main features:
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The combination of downward-deviated eyes + lid retraction produces an appearance that has resulted in this being known as the two-words sign

- Convergence-retraction nystagmus
- Light-near dissociation



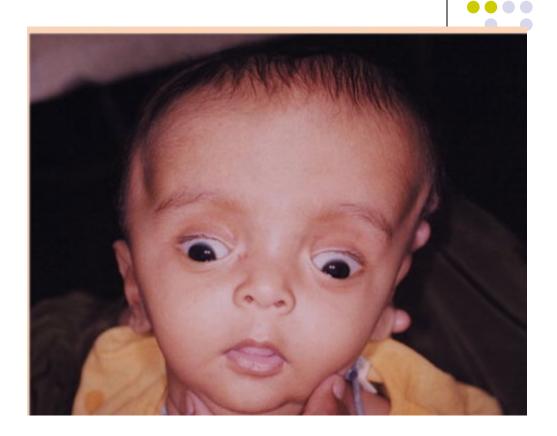


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Parinaud syndrome. The combination of lid retraction + impaired upgaze gives rise to a characteristic appearance known as *setting sun sign* 



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A phenomena in which attempted upgaze causes the globes to retract (ie, sink deeper into the orbit), converge, and 'shimmy' (for lack of a better word)



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Why is the name convergence-retraction nystagmus a misnomer?





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Duane syndrome: Globe retraction



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Because the medial recti contract as well, unopposed by the lateral recti



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What is light-near dissociation?





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Light-near dissociation in Parinaud's



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(Hint: It's iatrogenic)

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These nerves run fairly close to the inner wall of the eye, and thus are frequently impacted by thermal laser procedures that cover extensive portions of the retinal periphery

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--Affe
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Damage to which type/location is implicated in the light-near dissociation associated with **Parinaud's?** 

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Damage to what dorsal midbrain structure is causative in Parinaud syndrome?

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--Affe

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  - Convergence-retraction nystagmus
  - Light-near dissociation

```
What is light-near dissociation?

A phe part of the dorsal midbrain

Damage to what dorsal midbrain structure is causative in Parinaud syndrome?

The pretectum (specifically, the pretectal nuclei)

--Affe
--Cen
--Peri
```

Damage to which type/location is implicated in the light-near dissociation associated with **Parinaud's?**Central



- Parinaud syndrome has four main features:
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  - Lid retraction
  - Convergence-retraction nystagmus
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--Cen What are the two noneponymous names for Parinaud syndrome?

1)
2)
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**Central** 





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  - Light-near dissociation

#### What is light-near dissociation?

More precisely, where does Parinaud syndrome localize to?
The dorsal midbrain

as

Damage to what dorsal midbrain structure is causative in Parinaud syndrome? The pretectum (specifically, the **pretectal** nuclei)

What are the two noneponymous names for Parinaud syndrome?

- 1) Dorsal midbrain syndrome
- 2) Pretectal syndrome

Damage to which type/location is implicated in the light-near dissociation associated with **Parinaud's?** 

Central



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The etiology of a Parinaud syndrome is often a function of who the at is.

What other signs will be present in an infant with hydrocephalus?

--?

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Hydrocephalus: Enlarged head





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The etiology of a Parinaud s	unduance is often a function of tube the at in
	- What other signs will be present in an infant with hydrocephalus?
For each of these pts with P	Enlarged head
A child: <b>Hydrocephalus</b>	
7 torma. Hydrocopilalas	Bulging
	<del>-?</del>





- Parinaud syndrome has four main features:
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  - Light-near dissociation

The etiology of a Parinaud service For each of these pts with Parinaud service.

--A child: **Hydrocephalus** 

What other signs will be present in an infant with hydrocephalus?

- --Enlarged head
- --Bulging fontanelle

--?





Hydrocephalus: Bulging fontanelle





- Parinaud syndrome has four main features:
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  - Lid retraction
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  - Light-near dissociation

The etiology of a Parinaud syndrome is often a function of who the ret is What other signs will be present in an infant with hydrocephalus?

--Enlarged head
--Bulging fontanelle
--Dilated





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  - Impaired upgaze
  - Lid retraction
  - Convergence-retraction nystagmus
  - Light-near dissociation

The etiology of a Parinaud specification For each of these pts with P

--A child: **Hydrocephalus** 

What other signs will be present in an infant with hydrocephalus?

- --Enlarged head
- --Bulging fontanelle
- --Dilated scalp vessels





Hydrocephalus: Dilated scalp vessels



- Parinaud syndrome has four main features:
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  - Lid retraction
  - Convergence-retraction nystagmus
  - Light-near dissociation

- --A child: Hydrocephalus
- --A young man:





- Parinaud syndrome has four main features:
  - Impaired upgaze
  - Lid retraction
  - Convergence-retraction nystagmus
  - Light-near dissociation

- --A child: Hydrocephalus
- --A young man: A pineal tumor



- Parinaud syndrome has four main features:
  - Impaired upgaze
  - Lid retraction
  - Convergence-retraction nystagmus
  - Light-near dissociation

- --A child: Hydrocephalus
- --A young man: A pineal tumor
- --A young woman:





- Parinaud syndrome has four main features:
  - Impaired upgaze
  - Lid retraction
  - Convergence-retraction nystagmus
  - Light-near dissociation

- --A child: Hydrocephalus
- --A young man: A pineal tumor
- --A young woman: MS



- Parinaud syndrome has four main features:
  - Impaired upgaze
  - Lid retraction
  - Convergence-retraction nystagmus
  - Light-near dissociation

- --A child: Hydrocephalus
- --A young man: A pineal tumor
- --A young woman: **MS**
- --An **older** man:





- Parinaud syndrome has four main features:
  - Impaired upgaze
  - Lid retraction
  - Convergence-retraction nystagmus
  - Light-near dissociation

The etiology of a Parinaud syndrome is often a function of who the pt is. For each of these pts with Parinaud's, state the most likely cause:

--A child: Hydrocephalus

--A young man: A pineal tumor

--A young woman: **MS** 

--An older man: CVA