

Eye movement problems in adults

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- Horizontal misalignment
- (Cyclo)vertical misalignment
- Special forms of strabismus
- Pathological nystagmus and other ocular oscillations

OVERVIEW





- Comitant
- 1. Esotropia
- 2. Exotropia

- Non-comitant
- 1. Cranial nerve problem: causes/location
- 2. Neuromuscular junction
- 3. Extraocular muscle disease
- 4. Orbital disease

Horizontal misalignment

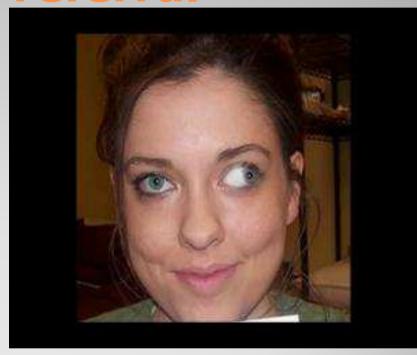
Comitant

Esotropia	Exotropia
Infantile	Infantile
Fully/partially accommodative	Intermittent distance exotropia
Sensory	Sensory
Sudden onset	Sudden onset
Residual	Residual
Consecutive	Consecutive

Horizontal misalignment

Longstanding? Routine referral





What can we do?
Assess, advise, operate

Non-comitant

- 1. Cranial nerve problem
- 2. Neuromuscular junction
- 3. Extraocular muscle disease
- 4. Orbital disease

Horizontal misalignment

Cranial nerve anomaly/disease: causes

Palsy/paresis

Congenital/developmental

Tumour

Vascular

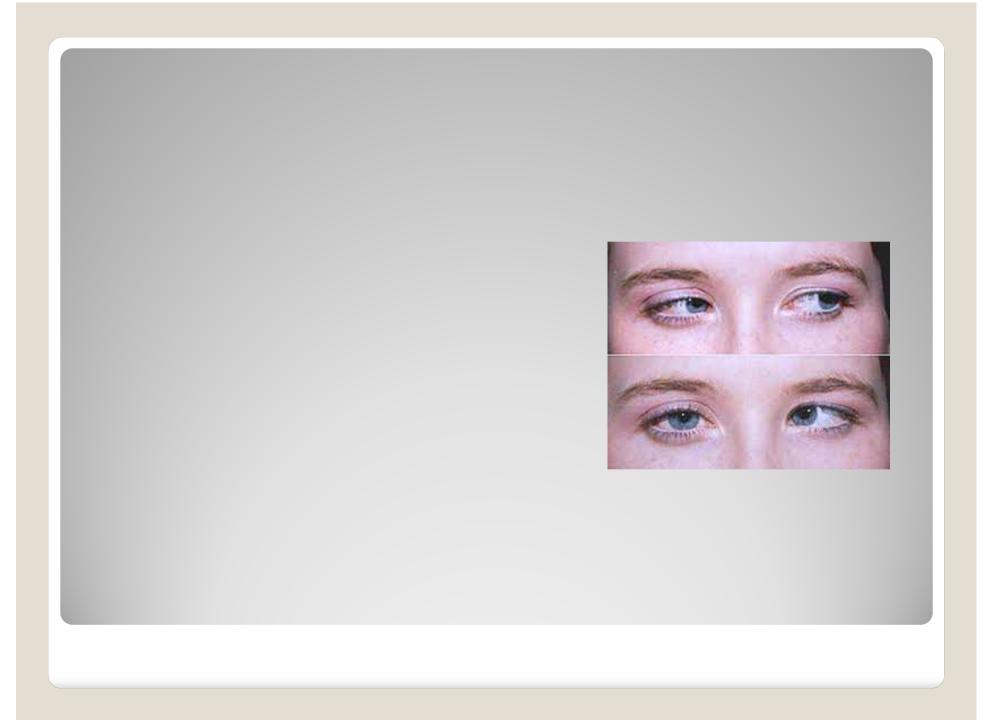
Inflammatory

Trauma

Immune

Other

1. CN problem: Causes



Causes of sixth nerve palsy

Most common: vascular (diabetes, hypertension, atherosclerosis)

Trauma

Watch out for:

Raised intracranial pressure

Giant cell arteritis

Cavernous sinus mass

Brain stem tumour, aneurysm

Multiple sclerosis, sarcoidosis, vasculitis

Urgent referral



Sixth nerve palsy

Investigations for sixth nerve palsy

Check blood pressure, blood sugar

Check full blood count, inflammatory markers, renal function, cholesterol

Fundoscopy, Ishihara, neuro exam

Decide whether imaging is needed, and how urgently

Involve other hospital specialists if underlying condition such as MS identified

Urgent referral



Management of sixth nerve palsy

Treatment for any underlying condition

Ophthalmic:

prisms

occlusion

After 6-12 months of stable measurements: surgery

Botulinum toxin

Urgent referral



Horizontal gaze palsy

Limitation of abduction on one side (abducens nerve)

Limitation of adduction of the other eye (contralateral oculomotor nerve)

Cause: lesion affecting interneurons from abducens nucleus to oculomotor nucleus in the pons

Sixth plus: horizontal gaze palsy



New case

Down and out

Ptosis

Dilated pupil (or not)







New case

Causes of third nerve palsy

ANEURYSM (rare, posterior communicating artery)

Most common: vascular (diabetes, hypertension, atherosclerosis)

Trauma

Watch out for:

Giant cell arteritis

Cavernous sinus mass

Tumour

Multiple sclerosis, sarcoidosis, vasculitis







Investigations for third nerve palsy

Check blood pressure, blood sugar

Check full blood count, inflammatory markers, renal function, cholesterol

Fundoscopy, Ishihara, neuro exam

Decide whether imaging is needed, and how urgently

Involve other hospital specialists if underlying condition identified





Management for third nerve palsy

Treatment for any underlying condition Urge

Ophthalmic:

prisms

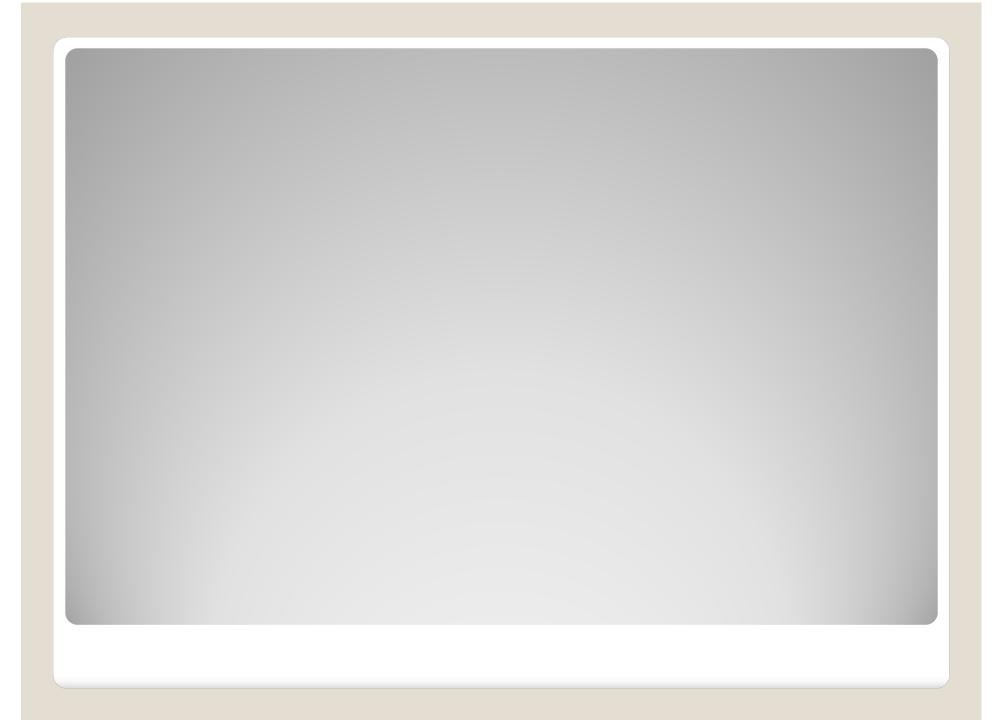
Occlusion if diplopia (usually ptosis)

After 6-12 months of stable measurements: surgery

Botulinum toxin



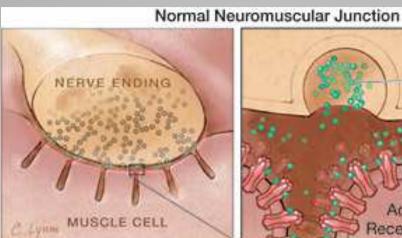
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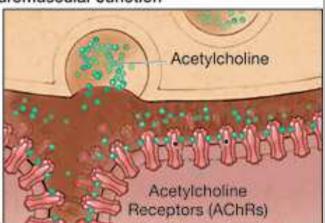






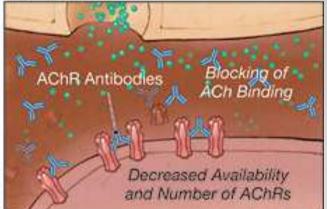
New case





Myasthenia Gravis





2. Neuromuscular Junction

Neuromuscular junction: Myasthenia

Varying symptoms and signs

Fatiguability

Generalised symptoms of weakness, breathing problems, chewing/swallowing difficulties, change of voice

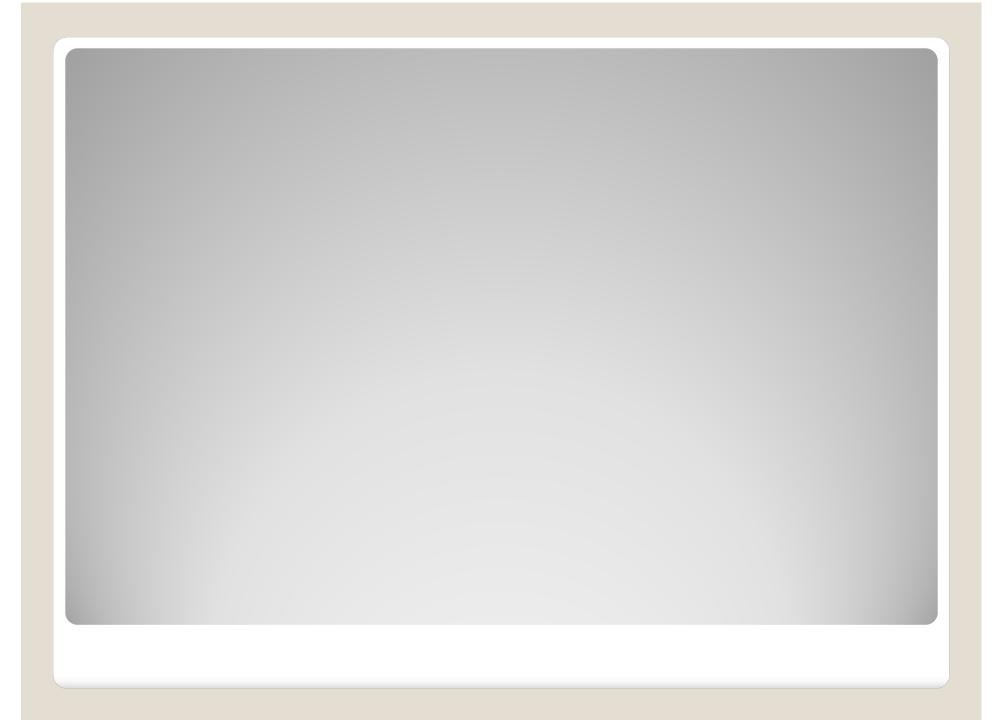
Tests: anti-acetylcholine receptor antibodies (negative in up to 50% of ocular only myasthenia), MuSK (musclespecific kinase)

Chest MRI/CT: thymoma, lung tumour (Lambert Eaton)

Electromyography

Treatment: steroids, pyridostigmine, neostigmine

2. Neuromuscular Junction







3. Extraocular muscle disease

Extraocular muscle disease

Congenital/developmental

Tumour

Vascular

Inflammatory

Trauma

Immune

Other





3. Extraocular muscle disease

Management of TED

Assessment of severity and activity

Thyroid function test, antithyroglobulin antibodies, antithyroid peroxidase antibodies

Treatment: general, then orbital, then strabismus, then lids

Steroids, thyroxine, smoking cessation, ocular artificial tears, orbital decompression, strabismus surgery to increase field of binocular vision, lid surgery to restore appearance





3. Extraocular muscle disease

Extraocular muscle disease

Congenital/developmental

Tumour

Vascular

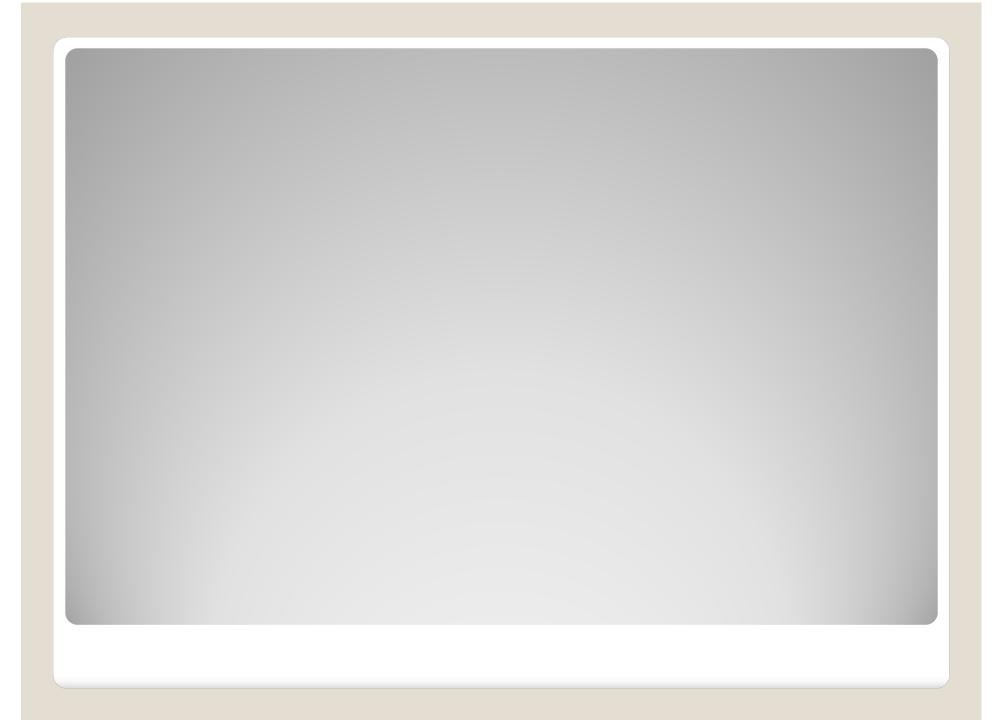
Inflammatory

Trauma

Immune

Other

4. Orbital disease



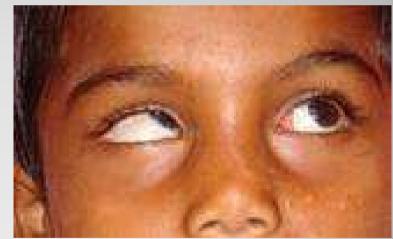
- 1. Apparent oblique muscle dysfunction
- 2. Paresis

Cyclovertical misalignment

 Inferior oblique overaction

 Very common, in particular in association with childhood strabismus

Asymptomatic: others notice



Cyclovertical muscle dysfunction





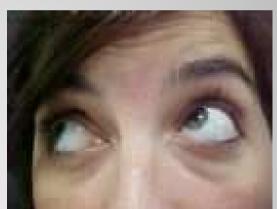
New case

Vertical double vision

Worse when looking to one side

Better when tilting head





New case

Causes of fourth nerve palsy

Most common: Congenital

Most common acquired: head injury

less common: vascular (diabetes, hypertension, atherosclerosis)

Watch out for:

Giant cell arteritis

Cavernous sinus mass

Tumour

Multiple sclerosis, sarcoidosis, vasculitis

Usually longstanding and decompensating -> diplopia

If in doubt referurgently

Fourth nerve palsy

Investigations for fourth nerve palsy

History: old photos head tilt?

Orthoptic assessment: vertical fusion range

Check blood pressure, blood sugar

Check full blood count, inflammatory markers, renal function, cholesterol

Fundoscopy, Ishihara, neuro exam

Decide whether imaging is needed, and how urgently

Involve other hospital specialists if underlying condition identified

Management for fourth nerve palsy

Treatment for any underlying condition

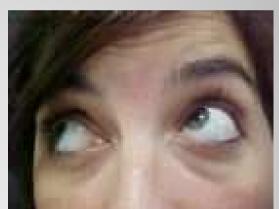
Ophthalmic:

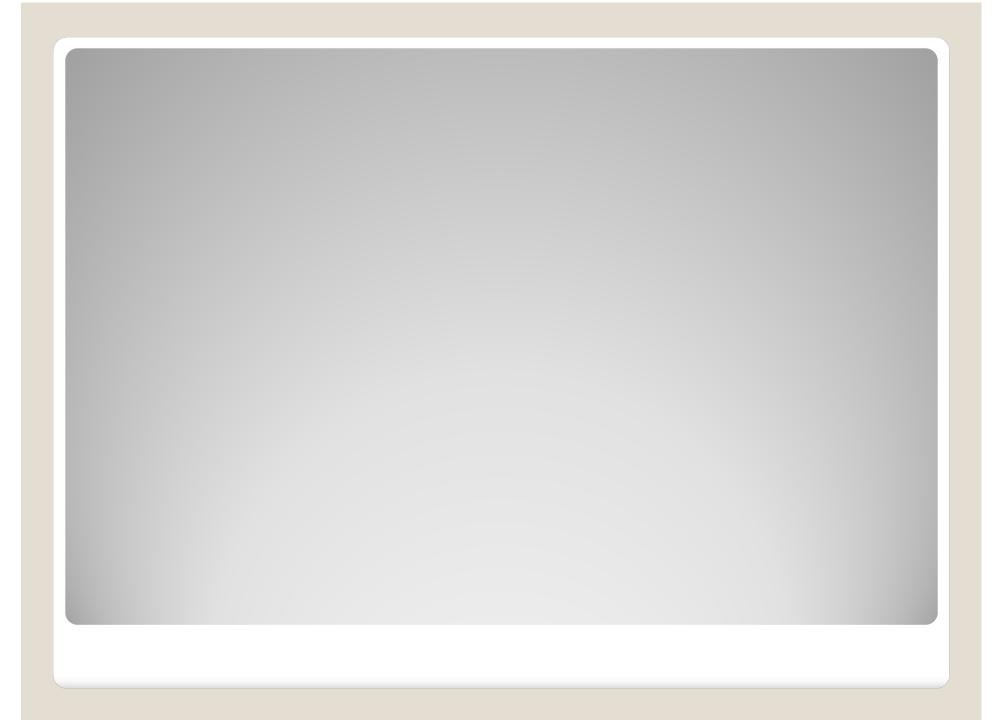
prisms

occlusion

After 6-12 months of stable measurements: surgery







Restrictive/Mechanical Strabismus

Secondary to Muscular Disease

Associated with Orbital Bony Disease

Iatrogenic Cyclovertical Deviations ("Induced Adhesive Syndromes")

- a) internuclear ophthalmoplegia
- b) supranuclear eye movement disorders
- b1) Progressive supranuclear palsy
- b2) Dorsal midbrain syndrome (Parinaud)
- b3) Tonic gaze deviation
- c) Skew deviation
- d) Eye movement changes associated with Parkinson disease

a) internuclear ophthalmoplegia

Lesion in the medial longitudinal fasciculus

Usually multiple sclerosis or microvascular

Limitation of adduction plus contralateral nystagmus on abduction

Usually improve spontaneously

Diplopia: occlusion. Long-term: surgery

b) supranuclear eye movement disordersb1) Progressive supranuclear palsy

Degenerative condition

Blurring of vision, photophobia

Slowing of saccades, then pursuits

Swallowing, speech, cognitive function, tone gait

all progressively impaired

b2) Dorsal midbrain syndrome (Parinaud)

b3) Tonic gaze deviation

- b) supranuclear eye movement disorders
- b1) Progressive supranuclear palsy
- **b2)** Dorsal midbrain syndrome (Parinaud)

Posterior commissure lesion: pineal gland,

hydrocephalus

loss of upgaze, Bell's phenomenon

Convergence-retraction nystagmus on attempted

upgaze

Lid retraction (Collier sign)

Pupils: light-near dissociation

Convergence paralysis

b3) Tonic gaze deviation

- b) supranuclear eye movement disorders
- b1) Progressive supranuclear palsy
- b2) Dorsal midbrain syndrome (Parinaud)
- **b3)** Tonic gaze deviation

Lesion in frontal eye field (infarct) sustained horizontal conjugate deviation

c) Skew deviation

Vertical misalignment
lesion of the vestibular input into nuclei of third,
fourth, sometimes sixth nerve nucleus
One eye hypotropic/excyclotorted
Fellow eye hypertropic/incyclotorted

d) Eye movement changes associated with Parkinson disease

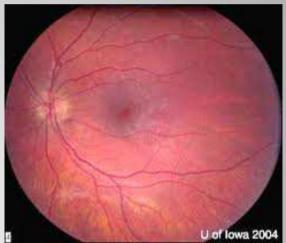
Convergence deficit
Sometimes lid lag on downgaze
Hypometric saccades
Impaired smooth pursuits
Saccadic intrusions

- a) Myasthenia Gravis
- b) Lambert Eaton myasthenic syndrome
- c) Miller Fisher syndrome
- d) Chronic progressive external ophthalmoplegia
- e) Myotonic and oculopharyngeal dystrophy

- a) Myasthenia Gravis
- b) Lambert Eaton myasthenic syndrome
- c) Miller Fisher syndrome
 Ophthalmoplegia, ataxia, areflexia
 Auto-antibodies against ganglioside QG1b

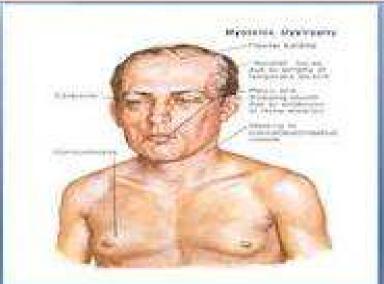
d) Chronic progressive
external ophthalmoplegia
Mutation in mitochondrial DNA:
not enough ATP produced
Ptosis, progressive paresis of eye
muscles,
retinal salt-and-pepper
appearance,
cardiac conduction defect
(Kearns-Sayre)





e) Myotonic and oculopharyngeal dystrophy Expansion of unstable DNA trinucleotide repeats







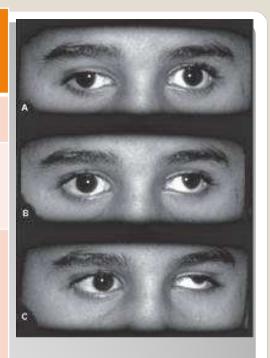
- 1. Duane syndrome
- 2. Lower cranial nerve maldevelopment: Moebius sequence
- 3. Marcus Gunn Jaw-Eyelid Synkinesis Syndrome
- 4. Congenital Fibrosis of the Extraocular Muscles (CFEOM)
- 5. Restrictive Hypotropia in Adduction: Brown Syndrome

1. Duane syndrome

Missing development of abducens nucleus

Lateral rectus either not innervated, or innervated by branch from oculomotor nerve, usually inferior division

Result: deficit of abduction, sometimes deficit of adduction, narrowing of palpebral fissure on horizontal gaze; abnormal head posture



2. Lower cranial nerve maldevelopment: Moebius sequence

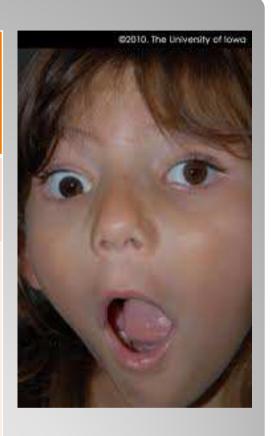
Sixth and seventh nucleus maldeveloped

Plus additional nuclei affected (swallowing, speech etc)



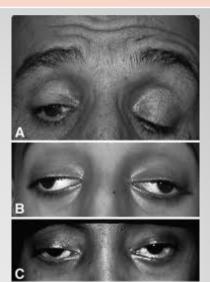
3. Marcus Gunn Jaw-Eyelid Synkinesis Syndrome

Maldevelopment of lid innervation Levator palpebrae instead supplied by nerve providing input to muscles of mastication Result: eyelid lifts when chewing or sucking



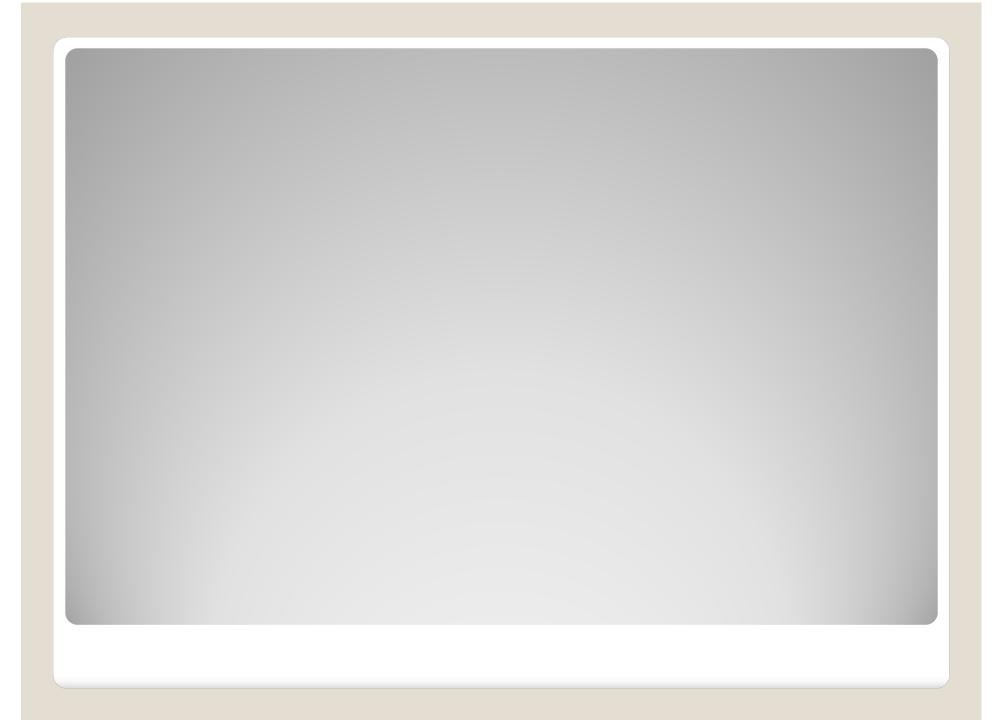
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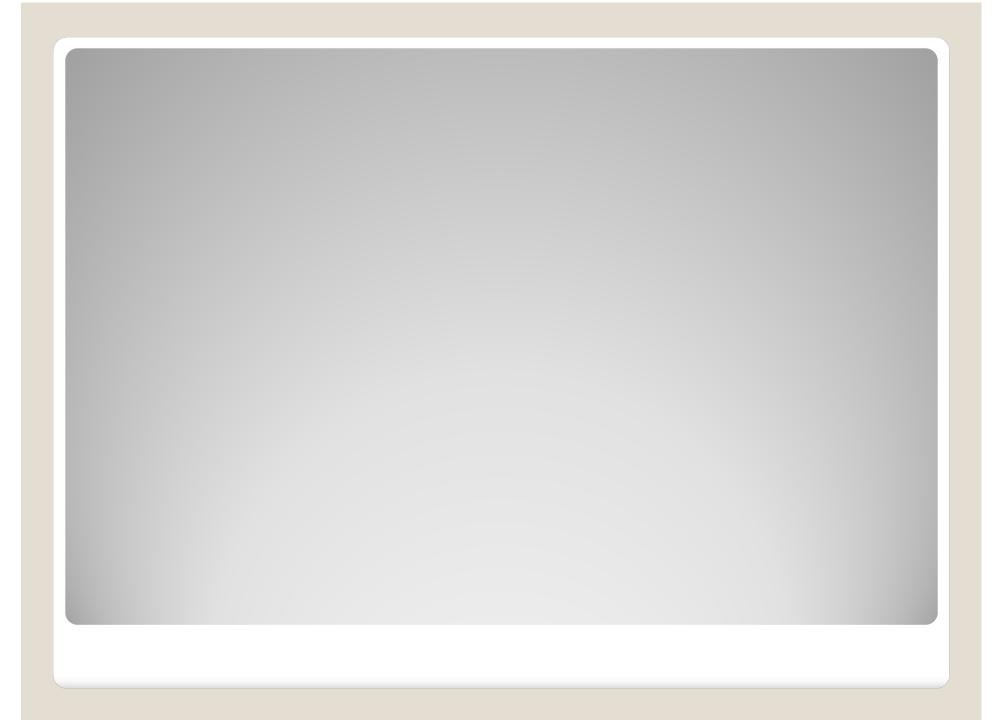




5. Restrictive Hypotropia in Adduction: Brown Syndrome







- Acquired
- Oscillopsia
- Central/Vestibular causes
- Drugs, alcohol, medication, thiamine deficiency
- Downbeat: posterior fossa malformation, cerebellar ectopia (Arnold Chiari)
- Seesaw (chiasmal, posterior fossa, trauma)
- Periodic alternating: cerebellar, visual, idiopathic

Pathologic nystagmus and other oscillations