OVERLAPPING MYASTHENIA GRAVIS & GRAVE'S DISEASE Challenging Case Reports

NGUYEN DUY DUAN, MD



A 24-year-old female

3 years	Grave's disease G		Recent Grave's Ophthal.		
	On Tx of Methimazole → euthyroid state		Ptosis and ophthamoplegia follow-up w/o Ta	X	

1. How to recognize Grave's pt has MG or not?

A 24-year-old female



3 years	2 Grave's disease	years	Grave's Ophthal.	Rece	nt GD & MG concomitantly
	On Tx of Methimazole → euthyroid state		Ptosis and ophthamoplegia → follow-up w/o Tx	6	Response well to Pyridostigmin & Corticosteroid

1. How to recognize Grave's pt has MG or not?

A 24-year-old female



3 years	2 Grave's disease	years	Grave's Ophthal.	Rece	nt GD & MG concomitantly
	On Tx of Methimazole → euthyroid state		Ptosis and ophthamoplegia → follow-up w/o Tx	6	Response well to Pyridostigmin & Corticosteroid

Questions was posed are:

• Is it common enough to be noticed?

•Why does it make physicianS confused?

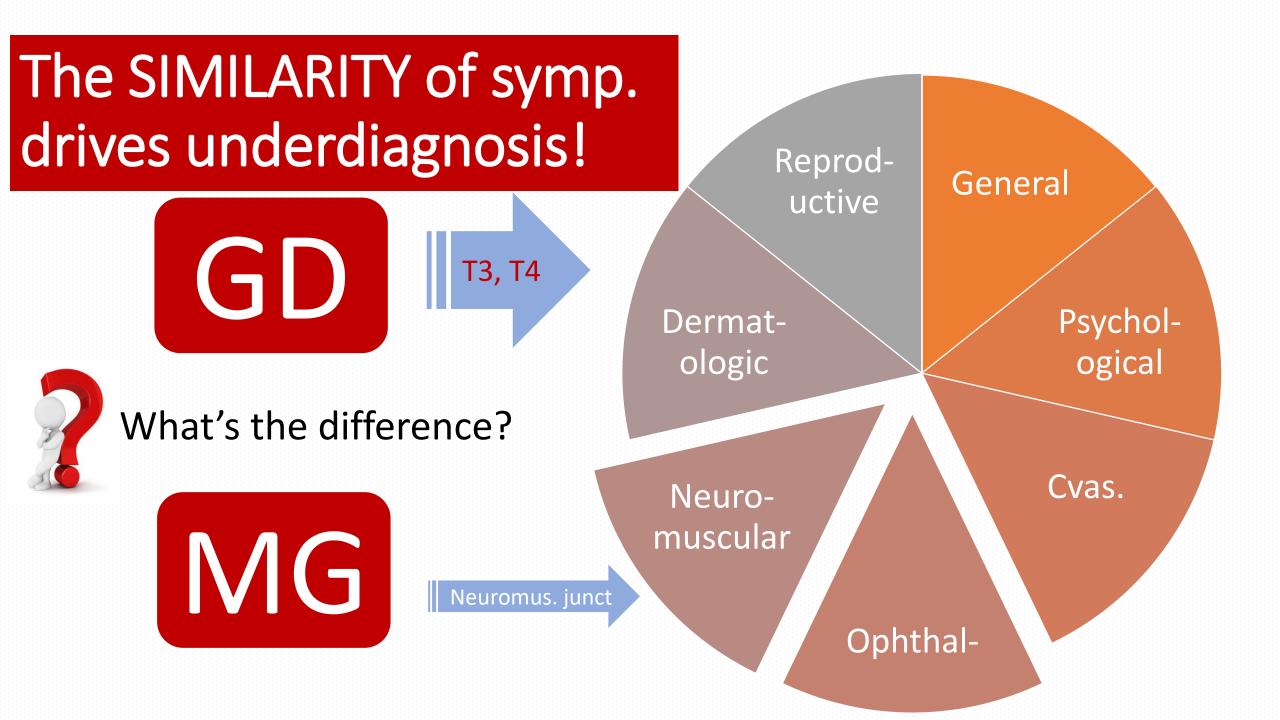
•How to recognize Grave pt has MG or not?

1. Is it common enough to be noticed?

CONCOMITANCE is NOT RARE

MG has been reported to be discovered simultaneously with, or prior to, the diagnosis of Graves' disease, but is most commonly subsequent to it.

Genetic predisposition for autoimmune disease

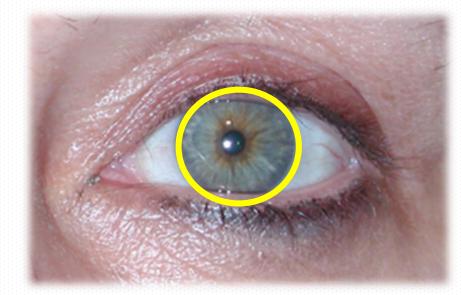


3. How to recognize Grave's pt has MG or not?

• Ptosis:

- GO: Eyelid retraction due to increased sympathetic stimulation of Muller's muscle by thyroid hormone.
- MG: Ptosis because of weakness of superioris levator.





Normal



Dalrymple's sign



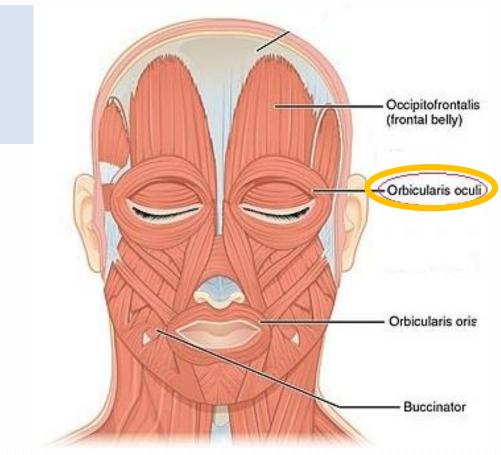
Ptosis

3. How to recognize Grave's pt has MG or not?

Orbicularis Oculi weakness:

- GO doesn't affect this muscle
- Occular MG is usually involved in

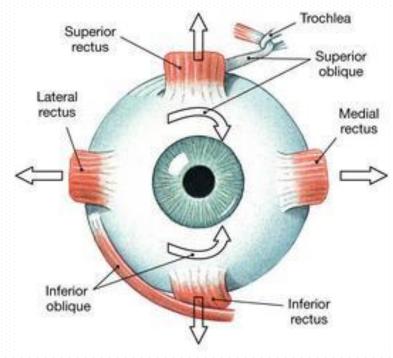




3. How to recognize Grave's pt has MG or not?

• Exotropia (deviate outward):

- GO: usually affects the Medial and inferior recti → Restrictive Myopathy of MR and IR→ eye move inward (esotropia) and downward (hypotropia) whileas exotropia RARELY occur.
- MG: doesn't have any predilection for particular extraocular muscles.



British Journal of Ophthalmology 1993; 77: 822-823

Exotropia as a sign of myasthenia gravis in dysthyroid ophthalmopathy

3. How to recognize Grave's pt has MG or not?

- Weakness of the voluntary muscles of the head and neck:
 - MG: Common
 - G-Myopathy: Rare \rightarrow usually induces proximal limbs weakness.

• Respiratory muscle involvement:

- MG: could happen in Myasthenic Crisis
- G-Myopathy: Rarely even in Acute thyrotoxic myopathy
- Diurnal fluctuation of weakness:
 - MG: more severe
 - GD: vary the degree of wkn during the long course in Grave's Myopathy and thyrotoxic periodic paralysis .

Questions was posed are:

- Is it common enough to be noticed?
- •Why does it make physicians confused? SIMILARITY
- •How to recognize Grave pt has MG or not?
 - Ptosis
 - Obicularis Occuli wk.
 - Exotropia

• Head & Neck musl. wk.

NOT RARE

- Respiratory musl. Wk
- Diurnal fluctuation of sev.

SUSPECTED!!

FURTHER TESTs!

- Ptosis
- Obicularis Occuli wk.
- Exotropia
- Head & Neck musl. wk.
- Respiratory musl. Wk
- Diurnal fluctuation of sev.



- Ice-pack test
- Tensilon/Prostig. test
- Autoantibodies

• EMG

BEDSIDE TESTs

- Ice-pack test: ptosis improve within one minute applying.
- Edrophonium (or Prostigmin) test: the weakness is reversed



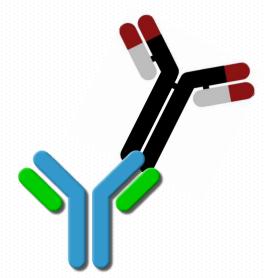
Ice-pack test



Edrophonium test: before and after test

OTHER TESTs

- Acetylcholin receptor antibodies (AChR-Ab) and Muscular Specific receptor tyrosin kinase (MuSK)
 - MG: 6-12% pts have Negative both test called Seronegative MG.
 - GD: Negative
- Electromyography (EMG):
 - MG: repetitive nerve stimulation (+)ive w/ decremental response in 75-80%, while that single-fiber EMG w/ jitter (+) in 95%.
 - G-Myopathy: myopathic findings of increased polyphasic, low amplitude motor unit potentials







A 18-year-old male

5 years	s 2 y	ears Several d	ay latter	
	GD	Grave's Myopathy		
	On Tx of	Muscular Weakness	Breathless &	
	Methimazole	and Fatigue esp. in	dysphagia	
		the end of the day		
		MTZ + Propanolol		

CASE 2
2. What must be
noticed in mag.?

A 18-year-old male



5 yea	rs 2 y	ears Several c	lay latter	GD & MG
	GD	Grave's Myopathy	Myasthenic Crisis	concomitantly
	On Tx of	Muscular Weakness	Breathless &	
	Methimazole	and Fatigue esp. in	dysphagia	Response well to Pyridostig.
		the end of the day	\rightarrow Prostig. test (+)	& CS
		MTZ + Propanolol		



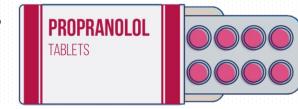
2. What must be noticed in mag.?

• Beta-Blocker (BB):

- GD: Propranolol typically rapidly reverse the paralysis in patients with thyrotoxic periodic paralysis. decrease the adrenergic imbalance of thyrotoxicosis.
- MG: it makes the weakness more severe, leads to Myasthenic Crisis

• Benzodiazepine:

- GD: relief the frequent thyrotoxicosis-associated anxiety
- MG: cause Myasthenic Crisis







2. What must be noticed in mag.?

• Glucocorticoid:

- GD: G-ophthalmopathy, after Radioidodine therapy
- MG: usually well-tolerated but occasionally ass/w an exacerbation.







2. What must be noticed in mag.?

- Routine Pre-operative management:
 - nondepolarising neuromuscular blocking drugs (used to facilitate the mandatory tracheal intubation in thyroidectomy surgery)
 - Antibiotics





TAKE HOME MESSAGE

Routine checking: Extraocular muscle invol. Atypical weakness of GD

- Ptosis
- Obicularis Occuli wk.
- Exotropia
- Head & Neck musl. wk.
- Respiratory musl. Wk
- Diurnal fluctuation of sev.

The Grave-treating drugs make the symptom worse

- Beta Blocker
- Benzodiazepine
- Corticosteroids
- Antibiotics (AG, Quinolone..)
- Neuromuscular blocking drugs

TAKE HOME MESSAGE

Bedside test

- Ice-pack test
- Tensilon/Prostig. test

Laboratory test

- Autoantibodies
- EMG



