

CLINICAL GUIDELINE

Myasthenia Gravis or Lambert-Eaton Myasthenia Syndrome – Medicines that may affect patients

A guideline is intended to assist healthcare professionals in the choice of disease-specific treatments.

Clinical judgement should be exercised on the applicability of any guideline, influenced by individual patient characteristics. Clinicians should be mindful of the potential for harmful polypharmacy and increased susceptibility to adverse drug reactions in patients with multiple morbidities or frailty.

If, after discussion with the patient or carer, there are good reasons for not following a guideline, it is good practice to record these and communicate them to others involved in the care of the patient.

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Important Note:

The Intranet version of this document is the only version that is maintained.

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Information for healthcare professionals

Medicines that may affect patients with Myasthenia Gravis or Lambert-Eaton Myasthenic Syndrome

There are certain medicines that have been reported to worsen or induce myasthenia gravis (MG), often by increasing muscular weakness, and should be used with caution in patients with this condition. The list of medicines in table 1 has been compiled to assist prescribers in the decision making process when prescribing medicines for patients with myasthenia gravis.

The medicines in this list have been classed according to those which should be:

- ▲ Absolutely contraindicated
- Avoided
- Used with caution
- Probably safe with patient monitoring.

This list is not exhaustive, best attempts have been made to identify problematic medicines but it is acknowledged that information may not be complete. If a medicine is not on the list it is not guaranteed that it will be safe in myasthenia gravis, please be vigilant with new medicines where safety in myasthenia gravis may be lacking. It is important that patients and doctors be alert to the early signs of an exacerbation of myasthenia gravis (MG) when a new medicine is commenced, even if it is not on the list or is listed as probably safe.

If further advice is required this can be obtained from the neurology pharmacy team on 61978 or the on call neurology registrar who can be contacted via switchboard.

Table 1: Medicines that may affect patients with Myasthenia Gravis or Lambert Eaton Myasthenic Syndrome

Avoid 🔺	Caution	Probably Safe
Infections		
▲ ▲ Telit	thromycin is Absolutely Contr	aindicated
Aminoglycosides ¹ ▲ Gentamicin, amikacin, tobramycin, streptomycin,	Macrolides ☐ Clarithromycin, erythromycin, azithromycin	Penicillins (except ampicillin) ●
neomycin	Penicillins	Aztreonam ● Meropenem ●
Antimalarials ▲ Chloroquine, mefloquine,	Ampicillin	Trimethoprim ● Ceftriaxone ●
quinine	Quinolones Ciprofloxacin, ofloxacin,	Metronidazole •
Polymyxins A Colistimethate sodium	levofloxacin, moxifloxacin, norfloxacin	Aciclovir ●
Clindamycin ▲ (caution with topical)	Tetracyclines Doxycycline, lymecycline, minocycline, oxytetracycline, tetracycline, demeclocycline	
Ribavirin A	Vancomycin	
	Co-trimoxazole (sulphonamides) Nitrofurantoin	
	Imipenem-cilastin ■ Rifampicin ■	
	Ritonavir =	
Cardiovascular	Anti-platalata	
Anti-arrhythmics¹ ▲ Procainamide, lidocaine, disopyramide, propafenone	Anti-platelets Dipyridamole (may counteract the effect of pyridostigmine)	
	Beta-blockers ² ■ (class effect including ophthalmic preparations)	
	Statins ³ ■ (class effect)	
	Diuretics ☐ (class effect, monitor electrolytes)	
	Calcium Channel Blockers Felodipine, verapamil, nifedipine, diltiazem	Calcium Channel Blockers Amlodipine
	Methyldopa	
Pain & Musculoskeletal		
Strong opioids ▲ (class effect)	Weak opioids ☐ (class effect, includes tramadol)	Paracetamol •
Rheumatology Penicillamine	Etanercept -	Ibuprofen ● Naproxen ●
Hydroxychloroquine A	Riluzole	
Methocarbamol 🛦		
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Quinine A		

- 1. If this medicine is clinically indicated and no alternative exists, seek specialist advice.
- 2. Do not discontinue if patient on existing therapy.
- 3. Consider withdrawal if clinical condition worsens.

Table 1: Medicines that may affect patients with Myasthenia Gravis or Lambert Eaton Myasthenic Syndrome cont'd.

Avoid A	Caution <a> 	Probably Safe
Nervous System		
Anxiolytics & Hypnotics Benzodiazepines ▲ (class effect) Zopiclone, Zolpidem ▲ Diphenhydramine (Nytol®) ▲ Movement Disorders Orphenadrine ▲ Procyclidine ▲ Trihexyphenidyl ▲	Anti-epileptics Carbamazepine, Ethosuximide, Phenobarbitone, Phenytoin, Gabapentin Antipsychotics (class effect, includes first and second generation) Tricyclic antidepressants (class effect) Buspirone Lithium Monoamine-Oxidase Inhibitors	Anti-epileptics Levetiracetam, Lamotrigine, Sodium Valproate
	Isocarboxacid, phenelzine, tranylcypromine Amantadine	
Gastrointestinal		
Co-phenotrope ▲ Magnesium containing antacids/laxatives ▲	Antimuscarinics Hyoscine (hydrobromide and butylbromide), dicycloverine H2-receptor antagonists	Antimuscarinics Propantheline ● (this is routinely used to counteract the side effects of pyridostigmine)
Endocrine	Anti-emetics Prochlorperazine, droperidol	
Levothyroxine Avoid over replacement	Corticosteroids (commonly used to treat MG. Sudden increases in dose could exacerbate symptoms – please consult neurology)	
Genito-urinary	Bisphosphonates -	
	Apha-adrenoceptor blockers Alfuzosin, doxazosin, indoramin, prazosin, tamsulosin, terazosin Antimuscarinics Darifenacin, fesoterodine, flavoxate, oxybutynin, solifenacin, trospium, propiverine, tolterodine Oestrogens/Progestogens	
	Hormonal changes may affect MG symptoms	
Skin/Ophthalmology Polymyxin or bacitracin containing preparations	Acetazolamide Antimuscarinic eye drops atropine, cyclopentolate, tropicamide Beta-blocker eye drops Imiquimod	

Table 1: Medicines that may affect patients with Myasthenia Gravis or Lambert Eaton Myasthenic Syndrome cont'd.

Syndrome cont'd.	T	T		
Avoid 🔺	Caution <a> 	Probably Safe		
Anaesthesia				
Discussion with neurology ad	vised prior to myasthenic patient un	dergoing anaesthesia		
	Inhalational agents ■ myasthenic patients require smaller amounts of these agents Non-depolarising drugs ■ myasthenic patients are particularly sensitive to these agents Depolarising drugs ■ variable response in myasthenia gravis Local anaesthetics ■ myasthenic patients require smaller amounts of these agents Antimuscarincs ■ Atropine, glycopyrronium	Propofol •		
Chamathayanautia Aganta				
Chemotherapeutic Agents	vised prior to commencing chemoth	orany in myasthania nationt		
Zissassion manifest sology ad	Doxorubicin, etoposide, cisplatin, fludarabine lipilimumab, nivolumab, pembrolizumab, atezolizumab	S. S. J. M. M. S.		
Miscellaneous				
Sedating antihistamines A Alimemazine, clemastine, chlorphenamine, cyproheptadine, hydroxyzine, ketotifen, promethazine Avoid live vaccines if on immunosupression	Interferon (alpha and beta) Trientene Iodinated Contrast Media Glatiramer			
Botulinum Toxin Magnesium supplements (unless treating hypomagnesaemia)				

References

The following references have been used to compile this list in conjunction with local expert opinion:

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