Therapeutics Advisory Group



MEDICINES TO PRESCRIBE BY BRAND NAME IN PRIMARY CARE

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Background

Prescribing medicines by generic name is generally preferred but there are some circumstances where brand-name prescribing is necessary. Reasons for brand prescribing include differences in bioavailability, where patient training differs between products and for biological medicines.

Norfolk and Waveney ICB recommend the following list of medicines are prescribed by brand to ensure supply of the same product. Medicines have been grouped by therapeutic area. This list is not exhaustive, for further information refer to the BNF or contact the medicines optimisation team. Please refer to the netformulary for formulary preferred brands across Norfolk & Waveney.

Colour code			
	Strongly recommend pre	escribing by brand due to clinical reason	
		dation for consistency of supply and cost	
	Good practice to aid pro-	duct identification	
Therapeutic area	Drug / Drug class	Reason for brand-name prescribing &	
		additional information	
Allergy and	Adrenaline auto-injectors	Prescribe by brand name to ensure patients receive an	
immunology		auto-injector device they have been trained to	
		use.Instructions for use vary between brands.	
		If switching between brands, patients should receive full	
		training in use of the new device.	
Anaesthesia and	Buprenorphine patches	Buprenorphine transdermal patches are available as 72-	
pain			
		prescribing is recommended to reduce the risk of confusion	
		and error in dispensing and administration.	
		Transtec® 96-hourly patches are formulary choice.	
		Formulary choice 7-day patches are Reletrans® and Sevodyne ® patches.	
	Fentanyl patches Fentanyl patches Fentanyl patches Fentanyl transdermal patches are available as ma		
	remanyi patches	reservoir formulations. Reservoir patches must not be cut	
		because damage to the rate-limiting membrane can lead to	
		a rapid release of fentanyl resulting in overdose. If the	
		prescriber intends the patch to be cut (NB: unlicensed and	
		not recommended) then the prescription must specify a	
		brand of matrix formulation patch.	
	Morphine Modified	These medicines are available as 12-hourly and 24-hourly	
	Release (MR)	oral formulations. Brand-name prescribing is recommended	
	preparations	to reduce the risk of confusion in dispensing and	
		administration.	
	Oxycodone MR	These medicines are available as 12-hourly and 24-hourly	
	preparations	oral formulations. Brand-name prescribing is recommended	
		to reduce the risk of confusion in dispensing and	
		administration.	
	Tramadol MR	These medicines are available as 12-hourly and 24-hourly	
	preparations	oral formulations. Brand-name prescribing is recommended	

Blood and nutrition Oral rehydration salts To aid identification. Products contain multiple ingredients.			
Calcium salts			to reduce the risk of confusion in dispensing and
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		glycols)	Laxido® and Cosmocol® are first line formulary choices

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	Pancreatin supplements	To aid identification. Products contain multiple ingredients. Enzyme activity can vary between brands	
Genito-urinary system	nito-urinary Combined oral Different brands of the same formulation are a		
	Progestogen only oral	Different brands of the same formulation are available.	
	contraceptive Levonorgesterel-	Patient familiarity with one brand is important. Products have different indications, durations of use and	
	releasing intrauterine systems	introducers.	
	Alprostadil injection	Patient familiarity with one brand is important; instructions	
Malignant disease	Ciclosporin – when used	for use vary between preparations. Patients should be stabilised on a particular brand of oral	
	for transplant rejection	ciclosporin because switching between formulations without close monitoring may lead to clinically important changes in blood ciclosporin concentration. Ciclosporin has a narrow therapeutic index. Switching between a brand and generic formulation, or between generic formulations, should be initiated only by a transplant specialist. If switching is necessary, the patient should be monitored closely for changes in blood-ciclosporin concentration, serum creatinine, blood pressure, and transplant function.	
	Tacrolimus – when used	Inadvertent switching between oral tacrolimus products has	
	for transplant rejection	been associated with reports of toxicity and graft rejection. Oral tacrolimus products should be prescribed and dispensed by brand name only. Tacrolimus has a narrow therapeutic index. Switching between a brand and generic formulation, or between generic formulations, should be initiated only by a transplant specialist.	
	Mycophenolate	Mycophenolate mofetil and mycophenolic acid preparations are not interchangeable. Please ensure correct salt is prescribed. For patients prescribed mycophenolic acid, they should receive the Myfortic® brand which is not interchangeable with any other form of mycophenolate.	
Mental Health	Lithium	Lithium has a narrow therapeutic index and preparations vary widely in bioavailability. Changing the preparation requires the same precautions as initiation of treatment.	
	Methylphenidate MR	Methylphenidate modified-release MR preparations contain both immediate-release (IR) and MR methylphenidate. The proportion of IR and MR methylphenidate differs between brands; different preparations may not have the same clinical effect.	
Neurology	Antiseizure medications for seizure disorders	NICE epilepsy guidelines recommends consistent supply of the same preparation for patients with seizure disorders,	
	Category 1	unless the prescriber, in consultation with the patient and their family or carers, considers this not to be a concern. MHRA guidance groups antiseizure medications into three categories of risk to help healthcare professionals decide whether it is necessary to maintain continuity of a specific manufacturer's product. These groups are: Category 1: Specific measures are necessary to ensure consistent supply of a particular product (which could be either a branded product or specified manufacturer's generic product) for medicines in this category. Medicines: carbamazepine, phenobarbital, phenytoin, primidone For further information see The Use of Generic Anti Epileptics Drugs in Patients with Epilepsy — SPS -	
		Specialist Pharmacy Service – The first stop for professional medicines advice	

	Category 2	Category 2: By default, this category includes all antiseizure medications not listed in categories 1 or 3. The need for continued supply of a particular manufacturer's product should be based on clinical judgement and consultation with patient and/or carer. Consider clinical factors such as seizure frequency, treatment history and the potential implications for the individual of having a breakthrough seizure. Medicines: clobazam, clonazepam, eslicarbazepine, lamotrigine, oxcarbazepine, perampanel, rufinamide, topiramate, valproate, zonisamide. For further information see The Use of Generic Anti Epileptics Drugs in Patients with Epilepsy – SPS - Specialist Pharmacy Service – The first stop for professional medicines advice	
Respiratory	Beclometasone CFC-	Beclometasone dipropionate CFC-free pressurised	
. toophatory	free metered dose	metered-dose inhalers are not interchangeable; Qvar and	
NICE CKS	inhalers	Kelhale have extra-fine particules and are more potent than	
<u>asthma</u>		Clenil Modulite and Soprobec. MHRA advice to prescribe	
guidelines advise			
generic prescribing of	Tiotropoim	2008. To ensure patients receive an inhaler they have been	
inhalers should	Попоронн	trained to use, tiotropium capsules and administration	
be avoided as it		devices should be prescribed by brand name.	
can lead to		First line formulary choice are	
people with		Tiogiva® dry powder inhaler and Spiriva Respimat® soft	
asthma being		mist inhaler.	
given an	Formoterol dry powder	Patient familiarity with one brand is important; instructions	
unfamiliar device, affecting	inhalers	for use vary between preparations.	
usage and	Beclometasone and	Easyhaler® and Oxis Turbohaler® are formulary choice Fostair has extra-fine particles and is approximately twice	
adherence.	Formoterol CFC-free	as potent as Clenil Modulite and CFC-containing	
	metered dose inhalers	beclomethasone inhalers.	
	Salbutamol dry powder	Patient familiarity with one brand is important; instructions	
	inhalers	for use vary between preparations.	
		Where appropriate for the patient use a dry powder inhaler	
		– low global warming potential. Easyhaler® or Ventolin	
	The ambudling AAD	Accuhaler® preferred formulary choices.	
	Theophylline MR	MR preparations have different release characteristics and	
	preparations	are not interchangeable. Theophylline has a narrow therapeutic index.	
	Aminophylline MR	MR preparations have different release characteristics and	
	preparations	are not interchangeable.	
	'	Aminophylline has a narrow therapeutic index.	
Skin	Preparations for skin	To aid identification. Products contain multiple ingredients,	
	and scalp conditions	including topical corticosteroids, of different potencies.	
	containing multiple		
	ingredients		

Prescribing information

Providers commissioned to provide services on behalf of Norfolk and Waveney ICB are reminded that they are required to follow the local joint formulary and prescribing guidance, as detailed in the medicines management service specification of their contract.

References

Specialist Pharmacy Service. Example medicines to prescribe by brand name in primary care. Available from https://www.sps.nhs.uk/articles/example-medicines-to-prescribe-by-brand-name-in-primary-care/ [Accessed August 2023]

Adcock, J. Items considered unsuitable for generic prescribing. Great Yarmouth and Waveney Clinical Commissioning Group. Version 1, 2016.

*Adapted from Mid & South East Essex document

Title	MEDICINES TO PRESCRIBE BY BRAND NAME IN PRIMARY CARE	
Description of policy	To inform healthcare professionals	
Scope	Norfolk and Waveney Integrated Care System	
Prepared by	Norfolk and Waveney ICB Medicines Optimisation Team	
Impact Assessment (Equalities and Environmental)	Please indicate impact assessment outcome: Positive impact Adverse impact - low - action plan completed as per guidance Adverse impact - medium - action plan completed as per guidance Adverse impact - high - action plan completed as per guidance No impact No policy will be approved without a completed equality impact assessment	
Other relevant approved documents		
Evidence base / Legislation	Level of Evidence: A. based on national research-based evidence and is considered best evidence B. mix of national and local consensus C. based on local good practice and consensus in the absence of national research based information.	
Dissemination Is there any reason why any part of this document should not be avaithe public web site? ☐ Yes / No ☒		
Approved by	Norfolk & Waveney Therapeutics Advisory Group (TAG) (Date)	
Authorised by	Norfolk & Waveney Drug Integrated Care Board on behalf of the ICS (Date)	
Review date and by whom	Medicines Optimisation Team	
Date of issue	August 2023	

Version Number	Author	Purpose / Change	Date
1.0	Jessica Adcock		September 2016
1.1	N.Cunninghan	Reviewed by author and updated. Interferon, botulinum toxin, somatropin, filgrastim, pegfilgrastim, erythropoietin and apomorphine removed as all hospital only drugs and won't be prescribed in primary care.	August 2023