

Standard Operating Procedures

Sublingual Immunotherapy

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Approved by	Quality Improvement meeting and Medicines Safety Committee Meeting
Approved by Medicines	8 th May 2024
Safety Committee:	
Where medicines included in	
guideline	
Date of Approval	20 th March 2024
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This is the most current	
document and is to be used	
until a revised version is	
available	

Aim and scope of Standard Operating Procedure

Allergen immunotherapy is recommended for patients who, despite allergen avoidance and a supervised trial of maximum pharmacotherapy, still have uncontrolled symptoms of allergic rhinitis. Subcutaneous immunotherapy (SLIT) has been shown to be effective in reducing the symptoms and medications required in patients with pollen related seasonal allergic rhinitis. There is also evidence suggesting it may prevent the development of asthma. Sublingual immunotherapy is a treatment that should be prescribed in a specialist allergy service and initiated at least eight weeks prior to the pollen season, after which the treatment may be self-administered and continued at home. Patients will receive regular follow up in allergy clinic.

Target Staff Categories

This SOP is for all healthcare professionals in the Paediatric Department involved in the care of patients with allergic rhinitis

Key amendments to this Standard Operating Procedure

Date	Amendment	Approved by:
March 2024	Additional products added	QI and MSC
	Timing of food before/after treatment changed to 5 minutes	

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Product	Dose	Considerations
Grazax® (Alk Abello) (Phlegm Pretense/Timothy Grass)	75,000 SQ-T	First dose needs to be administered in hospital. Only sublingual product licensed for use in the UK for patients aged 6 years and above. All prescriptions should be hospital provided.
Acarizax® (ALK Abello) (Dermatophagoides pteronyssinus & Dermatophagoides farina / House Dust Mite)	12,000 SQ-T	First dose needs to be administered in hospital. Only sublingual product for HDM allergy.Licensed for use in the UK for patients aged 12 years and above. All prescriptions should be hospital provided.
Itulazax (ALK Abello) (Betula verrucosa (birch), Alnus glutinosa (alder), Carpinus betulus (hornbeam), Corylus avellana (hazel), Quercus alba (oak) and Fagus sylvatica (beech))	12,000 SQ-T	First dose needs to be administered in hospital. All prescriptions should be hospital provided. Licensed from 18 years and above

How to administer SLIT

- Sublingual immunotherapy should be initiated in a clinical area that has resuscitation facilities and where clinical staff are equipped to manage anaphylaxis.
- Written informed consent should be obtained by patient and/or parent/legal guardian.
- Patients should be advised that they must be well on the day of administration and if they have asthma this should be well controlled.
- Wash hands prior to administration and adhere to Worcestershire Acute Hospitals NHS Trust Infection Control Policy
- Discharge with 'Your Immunotherapy Treatment' information

Procedure	Rationale
Check patients name and date of birth to ensure the correct product SLIT is being administered to the correct patient.	To ensure the patient receives the correct treatment
Ensure the patient has a supply of antihistamine at home.	In case mild, local symptoms develop.
Obtain baseline observations prior to commencing SLIT, including chest auscultation.	To reduce the risk of a severe or systemic reaction.
Lung function testing and/or peak flow levels for patients with asthma. Ideally, the patient should not have used their reliever	To ensure patient can safely commence on administration of SLIT.



inhaler to treat wheeze in the two weeks	
prior to commencing treatment.	This could worsen any reaction or side
Assessment of skin condition in those with	effects.
eczema.	
Complete a visual inspection of the mouth,	
observing for oral lesions or loose teeth. If	
oral lesions are present, do not proceed with SLIT until the lesions have healed.	
with Schi dritti the lesions have healed.	
Ensure patient is not unwell.	
In the older patient ensure they are not	SLIT should not be initiated in patients who
pregnant.	are pregnant, due to the risk of anaphylaxis
The patient should refrain from eating and	Food or drink in the oral cavity could affect
drinking for 5 minutes prior to and following	the absorption of the SLIT
administration of SLIT. Administer SLIT by placing the treatment in	SLIT is absorbed via the sublingual route
the sublingual pocket under the base of the	SLIT is absorbed via the sublingual route
tongue. Please see individual summary of	
characteristics for product specific advice.	
SLIT should remain under the tongue for	To ensure SLIT is absorbed by the
between 1 – 2 minutes. It must not be	sublingual glands
swallowed Monitor patient for any sign of an allergic	An allergic reaction could occur following
reaction for 60 minutes following	administration of SLIT
administration of SLIT	daministration of OLIT
Reassure patient if they experience	These are common side effects and should
symptoms such as oral tingling pruritus,	resolve 1 – 2 weeks after beginning SLIT.
mild tongue swelling, itchy throat or ears	They may not occur until 10-15 minutes
that these are common side effects in the early phase of treatment	after administration.
Promptly treat any allergic reaction or side	To ease the discomfort and prevent the
effects of SLIT	development of moderate symptoms
Advise patient that if side effects are	Pre-dosing with an oral antihistamine in the
unpleasant that they can take an oral	first 1 – 2 weeks may help reduce
antihistamine 30 – 60 minutes prior to	unpleasant side effects in the early stage of
taking their SLIT	treatment.
Advise patients to stop taking SLIT in the following situations:	To reduce the risk of SLIT being absorbed systemically through an open lesion rather
 For 7 days following oral surgery, 	than through the sublingual mucosa.
including dental extraction	anan amough ano oublingual muoosa.
For 7 days after shedding a	
deciduous tooth	
 If patient has an oral ulcer or open 	
wound in the mouth or oral mucosa	
 to temporarily discontinue until area has healed. 	
area rias riealeu.	
If patient is unwell with a fever, or unwell	
enough to be absent from school or work,	To reduce the risk of patient experiencing
they should temporarily discontinue their	exacerbation of asthma and/or respiratory symptoms.
SLIT until their illness has resolved. If	symptoms.
patient receives a vaccine which causes	



aide effects auch as favor ar jaint a significant	
side effects such as fever or joint pain, they	
should stop their treatment until side effects	
resolve.	
Detients with concentrate at eathers and	
Patients with concomitant asthma and	
experiencing an acute upper respiratory	
tract infection – to temporarily discontinue	
treatment until treatment has resolved.	
Document administration of SLIT, any side	To record the administration and treatments
effects and treatment given.	for governance.
Reassess the patient prior to discharge and	To ensure the patient has not had an
repeat baseline observations.	allergic reaction and is fit for discharge.
Ensure the patient and/or family have the	To ensure the patient and family are
following information on discharge:	supported and aware of how to overcome
 How to manage an allergic reaction 	any problems with their treatment
 Advise the patient to ensure that 	
they have immediate access to	To treat side effects of SLIT
antihistamine	
 Revision and reinforcement of the 	
importance of compliance with	
medication	
 Have a supply of initial treatment 	
and are aware of when next supply	
will be provided	
Written information relating to the	
product they are using	
 Contact details should they require 	
ongoing support and repeat	
prescriptions	
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