

## EAR IRRIGATION GUIDELINE

			PROCEDURAL
Document Category	Clinical		
Document Type	Guideline		
Keywords	Ear care procedure/ Wax removal		
Version	Issue Date:	Review Date:	
4	1 <sup>st</sup> December 2025	1 <sup>st</sup> December 2028	
Supersedes	Guideline for Irrigations November 2022 –November 2025		
Approved by (Committee / group)	Rotherham NHS Foundation Trust  Ear Care and Audiology clinical governance  Heads of Service Clinical governance	Date Approved:	1 <sup>st</sup> December 2025
Scope of target audience	This procedure is only to be carried out by an experienced healthcare worker who has received recognised training in ear care and the use of ear care equipment. This training is available UK-wide from Rotherham Ear Care Centre trainers. The healthcare worker should also access a two yearly update.		
Evidence base/ References	NICE Clinical Knowledge Summaries for management of ear wax (2025)  The Royal Marsden Manual of Clinical and Cancer Nursing Procedures (2020)		
Lead Care Group	Care Group 4		

<b>Lead Specialty/ Department: (Or Care Group if 'care group' owned)</b>	Ear Care and Audiology Specialist Nursing team.
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<b>Sponsor (position/ role):</b>	N/A
<i>Name the documents here or record not applicable (these are documents which are usually developed or reviewed/ amended at the same time – ie a family of documents)</i>	
Associated Policy	N/A
Associated Procedure(s)	N/A
Associated Pathway(s)	N/A
Associated Standard Operating Procedure(s)	N/A
<b>Consultation Undertaken:</b>	Consultation undertaken with Ear Care Specialist Nursing Team and Clinical Lead for Ear Care and Audiology.
<b>Template control:</b>	

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## Amendments from previous version(s)

Version	Issue Date	Section(s) involved (author to record section number/page)	Amendment (author to summarise)
3	November 2025	Section 7. Related documents and guidance	Guidance outdated therefore removed and replaced with information obtained from current NICE Clinical Knowledge Summary.

## CONTENTS

Item	Title	Page
1.0	INTRODUCTION	4
2.0	AIMS/ OBJECTIVES/ PURPOSE (including Related Trust Documents)	4
3.0	ABBREVIATIONS AND DEFINITIONS	5
4.0	ROLES AND RESPONSIBILITIES	5
5.0	GUIDELINE DETAILS (including flowcharts)	5-10
6.0	EDUCATION AND TRAINING	10
7.0	MONITORING COMPLIANCE AND EFFECTIVENESS	10

## **1.0 INTRODUCTION/BACKGROUND**

Cerumen, or wax as it is commonly known, is a normal secretion of the ceruminous glands in the outer meatus. It is slightly acidic, giving bactericidal qualities in both its wet, sticky form (as secreted by Caucasians and African-Caribbeans) or dry, flaky form (as, for example, secreted by S.E. Asian people). In addition to epithelial migration, jaw movement assists the movement of wax to the entrance of the External Auditory Meatus (EAM) where it emerges onto the skin. A small amount of wax is normally found to the entrance of the EAM and its absence may be a sign that dry skin conditions, infection or excessive cleaning have interfered with the normal production of wax. It is only when there is an accumulation of wax that removal may be necessary. Cerumen impaction occurs in up to 6% of the general population, affecting 10% of children and over 30% of the elderly and cognitively impaired populations. Excessive wax should be removed if it is causing the patient a problem such as tinnitus, hearing loss, vertigo, pain/discomfort; or if examination of the tympanic membrane or audiological assessment/ intervention is required.

The experienced practitioner can use his or her clinical judgement on the best method for wax management and removal. Olive oil may be advised in favour of other cerumenolytics. The practitioner should advise patients to instil olive oil 3-5 days prior to their wax removal appointment. The practitioner may decide that extended use of olive oil is required.

These recommendations have been developed to assist practitioners in gaining experience and knowledge in the provision of ear care. They do not replace the need for education, recognised training and supervision in order to perform these procedures.

## **2.0 AIMS/ OBJECTIVES/ PURPOSE (including Related Trust Documents)**

The aim of the document is to promote and provide clear guidance for health care workers trained in the ear irrigation procedure to ensure safe and effective ear care is provided to patients, which will intern reduce litigation in ear irrigation. This document was originally produced by the 'Action On ENT' Steering Board (2002) and endorsed by the Royal College of General Practitioners, The Royal College of Nursing, The Ear Care Centre and the Medical Devices Agency. It has subsequently been revised by the Ear Care Centre (2025).

### **Related Trust Documents**

To consent to examination or treatment (trust policy available on HUB)  
Heath records policy (trust policy available on HUB)  
Standard infection prevention and control precautions (trust

policy available on HUB)  
Otoscopy guideline (trust policy available on the HUB)  
Electronic Irrigator Cleaning guideline (trust policy available of the HUB)

### 3.0 ABBREVIATIONS AND DEFINITIONS

#### Definitions

Definitions Noots – receiver used to collect water

#### Abbreviations

EAM – External Auditory Meatus

### 4.0 ROLES AND RESPONSIBILITIES

All staff involved in the aural care of patients must follow the guidance within this document or record any justifications for not doing so.

### 5.0 GUIDELINE DETAILS

#### **PURPOSE**

- Correctly treat chronic or fungal otitis externa where the meatus is obscured by debris
- Improve conduction of sound to the tympanic membrane when it is blocked by wax
- Remove keratin or debris to allow examination of the EAM and the tympanic membrane
- Remove wax in order to facilitate hearing aid mould impressions or other audiology assessment
- Facilitate the removal of wax and foreign bodies, which are **not** hygroscopic, from the EAM. Hygroscopic matter (such as peas and lentils) will absorb the water and expand, making removal more difficult

#### **SCOPE**

Irrigation should **NOT** be carried out if:

- the patient has previously experienced complications following this procedure in the past
- there is a history of a middle ear infection in the last six weeks

- the patient has undergone ANY form of ear surgery (apart from grommets that have extruded at least 18 months previously and it is documented subsequently that the tympanic membrane is intact)
- the patient has a perforation
- the patient has a healed perforation
- there is a history of a mucous discharge in the past twelve months
- there is evidence of acute otitis externa with pain and tenderness of the pinna
- there is a history of cleft palate, repaired or not
- the patient has abnormalities of the meatus such as exostosis
- the patient has hearing in only one ear if it is the ear to be treated

**This list is not exhaustive and the practitioner must use his or her own judgement for each individual**

Precautions: (Ear irrigation should be carried out on a low setting)

- the patient has tinnitus
- the patient suffers from dizziness
- the patient is taking anti-coagulants or high dose steroids
- the patient is immunocompromised
- the patient has had radiotherapy of the head or neck

## Children

Irrigation can be carried out on children as long as the child has no contraindications and is happy to co-operate with the procedure. The practitioner must ensure irrigation is appropriate and necessary. It may be advisable to instil olive oil for a longer period of time in children to avoid the need for irrigation. When carrying out otoscopy, gently pull the pinna down and backwards to straighten the EAM.

## EQUIPMENT

The metal syringe is obsolescent for use in the EAM. The syringe design is inherently dangerous. Combined with the danger of the syringe itself and the pressure of water it creates within the EAM, there is the difficulty of disinfecting the

syringe after each use. The Medical Devices Agency (MDA) also has reservations about the use of the metal syringe for wax removal. There are issues around the poor manufacture of some syringes, allowing them to break and cause injury during use, and the pressure of water that can be exerted manually on the tympanic membrane.

Electronic irrigators such as the "Propulse" or "Projet" allow irrigation of the EAM rather than wax removal under pressure. The MDA issued Safety Notice SN 9807 in February 1998 which advised users that the original Propulse electronic irrigator required an isolation transformer for electrical safety. Subsequently, the manufacturer designed and marketed the Propulse II and later the Propulse III to replace the original Propulse. Propulse NG and G5 are now available which is both mains and battery operated.

This guidance document does not recommend the use of manual syringes or the Propulse 1, even with an isolation transformer, but recommends that practitioners should use the Projet or Propulse II, III, NG or G5 irrigators and refer to the procedure as ear irrigation.

The Projet and Propulse II III NG and G5 irrigators have a pressure-variable control, allowing the flow of water to be easily controlled by commencing irrigation on the minimum setting. For patient safety, the manufacturers have limited the maximum pressure available: this limit is stated in the user instructions. The Projet, Propulse III, NG and G5 irrigators have specific disinfecting guidelines issued.

#### Equipment Requirements:

- Otoscope
- Headlight or other suitable light source and spare batteries
- Electronic irrigator
- Tap water at 38°C - 40°C or temperature comfortable for the patient, avoiding cool water
- Noots trough/receiver
- Jobson Horne probe /carbon curette or an appropriate cotton wool carrier and good quality cotton wool or ear mop/ear canal wick
- Tissues
- Disposable waterproof cape and paper towels
- Disposable apron and gloves

## PROCEDURE

This procedure should be carried out with both participants seated and under direct vision, using a headlight or other light source which allows a stream of light to illuminate the external auditory meatus.

1. The patient's presenting complaints and the result of the initial examination should be documented. Valid consent should be obtained and documented prior to proceeding
2. Examine both ears by first inspecting the pinna and adjacent scalp using direct light. Check for previous surgery incision scars or skin defects, and then inspect the EAM with the otoscope.
3. Check whether the patient has had his/her ears irrigated previously, or if there are any contraindications why irrigation should not be performed.
4. Explain the procedure to the patient and ask the patient to sit in an examination chair (a child could sit on an adult's knee with the child's head held steady).
5. Check that the headlight/light source is in place and is working correctly.
6. Place the protective cape and paper towel on the patient's shoulder and under the ear to be irrigated. Ask the patient to hold the receiver under the same ear.
7. Fill the reservoir of the irrigator; check that the temperature of the water in the tank is approximately 38°C - 40°C. Set the pressure at minimum.
8. Connect a new tip applicator to the tubing of the machine with a firm 'push/twist' action. Push until a "click" is felt.
9. Direct the irrigator tip into the Noots receiver and switch on the machine for 10-20 seconds in order to circulate the water through the system and eliminate any trapped air or cold water. This offers the opportunity for the patient to become accustomed to the noise of the machine. The initial flow of water is discarded, thus removing any static water remaining in the tube. Check the temperature of the water again.



10. Twist the tip so that the water can be aimed along the posterior wall of the EAM (towards the back of the patient's head).

11. Gently pull the pinna upwards and outwards to straighten the EAM (directly backwards in children).

12. Warn the patient that you are about to start irrigating and that the procedure will be stopped if he/she feels dizzy and/or experiences any pain. Ensure that the light is directed down the EAM. Place the tip of the nozzle into the EAM entrance and, using the foot control, direct a stream of water along the roof of the EAM and towards the posterior wall (direct towards the back of the patient's head). Increase the pressure control gradually if there is difficulty removing the wax. It is advisable that a maximum of one reservoir of water (500ml) per ear is used in any one irrigation procedure.

13. There is evidence to suggest that leaving water in the canal for 15 minutes will increase the chance of success. You may find it beneficial to instil water into both ears (if both require irrigation with water) and return to the procedure after a rest of 15 minutes. (Eekhof J et al 2001)

14. Periodically inspect the EAM with the otoscope and inspect the solution running into the receiver.

15. After removal of wax or debris, dry mop excess water from the meatus under direct vision using the Jobson Horne probe/carbon curette/ear canal wick or an appropriate cotton wool carrier and good quality cotton wool. Stagnation of water and any abrasion of skin during the procedure predispose to infection. Removing the water with the cotton wool tipped probe reduces the risk of infection.

16. Examine the ear, both meatus and tympanic membrane, and treat as required following specific guidelines, or refer to a doctor if necessary.

17. Give advice regarding ear care and any relevant information. Advise the patient to return if the ear starts to discharge or become painful. If the presenting symptom was hearing loss and the hearing doesn't improve following wax removal advise patient to seek further advice as per local policy.

18. Document what was observed in both ears, the procedure carried out, the condition of the tympanic membrane and external auditory meatus and treatment given. Findings should be documented, nurses following the NMC guidelines on record keeping and accountability. If any abnormality is found a referral should be made to the ENT Outpatient Department following local policy.

19. All contaminated equipment and PPE should be disposed of in clinical waste, with sharp instruments to be disposed of in appropriate sharps disposal.

**NB. IRRIGATION SHOULD NEVER CAUSE PAIN. IF THE PATIENT COMPLAINS OF PAIN, STOP IMMEDIATELY.**

It is recommended that you follow the manufacturer's guidelines and local policy for cleaning and disinfecting irrigator and its components.

## **RISK FACTORS**

Potential complications following procedure:

- Trauma
- Infection
- Dizziness
- Tinnitus

## **6.0 EDUCATION AND TRAINING**

This procedure is only to be carried out by an experienced healthcare worker who has received recognised training in ear care and the use of ear care equipment. This training is available UK-wide from the Ear Care Centre trainers. The healthcare worker should also access a two yearly update.

An individual assessment should be made of every patient to ensure that it is appropriate for ear irrigation to be carried out.

## **7.0 MONITORING COMPLIANCE AND EFFECTIVENESS**

Compliance with this procedural guideline will be monitored by undertaking yearly peer led clinical supervision.