

# SOP25 Standard Operating Procedure (SOP) The Safe Use of Ultrasound Gel to Reduce Infection Risk

#### **1.0** To ensure the safe use of non-sterile and sterile ultrasound gel

This document provides guidance on the safe use of ultrasound gel to reduce risk of transmission of infection arising from these products.

#### 2.0 Accountabilities

Owner: Karen Hill

Review: November 2024

Ratification:

**Dissemination:** Publication on Intranet. Incorporation into Induction Training

Compliance: All users

#### 3.0 Procedure/Guidelines Details/Action

#### 3.1 Introduction

This Standard Operating Procedure introduces a protocol to ensure the safety of patients and remove the risk transmission of infection when using ultrasound gel.

This is to be achieved in every setting where ultrasound is used as an imaging modality.

#### 3.2 Definitions

To ensure there is no transmission of infection between ultrasound gel and patients

#### 3.3 Regulatory Background

The UK Health Security Agency (UKHSA) published <u>guidance on the use of ultrasound gel</u> on 10 November 2021. This replaced the interim guidance published in January 2021, in response to the increased awareness of infections caused by contaminated non-sterile ultrasound gel products. The SoR and the SoR ultrasound advisory group were involved in the development of this guidance along with a wide range of stakeholders.

On 11 November 2021 the Medicine and Healthcare products Regulatory Agency (MHRA) released a <u>national patient safety alert</u>, which clarifies the actions required to reduce harm and practice safely.

# 3.4 Procedures to Follow For both Non-Sterile Gel and Sterile gel

- Warming of gel is not advised unless there is a clinical benefit that outweighs applying gel
  at room temperature. Where warming of gel is performed, dry heat warmers must be used
  with the bottle stored in the upright position. Gel must not be warmed in water. Gel warmers
  must be cleaned regularly according to the manufacturer's instructions
- Ensure healthcare workers carry out hand hygiene before and after use of ultrasound gel.



- Gel must be stored according to manufacturer's instructions in an area that is dry and away from potential sources of contamination.
- Dispose of container if it appears soiled, is damaged or is out of date

### Non-Sterile gel

- Non-sterile gel **must** not be decanted from larger gel containers.
- Gel bottles <u>must</u> be dated when opened and used within one month, unless the expiry date is earlier.
- The whole of the gel bottle, including the tip, must be wiped with a disinfectant wipe between use.
- The gel must be removed from the individual's skin with suitable tissue. Soft tissue should be provided if wiping sensitive areas.
- Low-risk outpatients must be advised to wash with soap and water to remove any residue when they return home. Assistance should be offered by staff in the department if patients are unable to do this independently in their home setting.

# Sterile gel Indications for Use

- Sterile single use gel sachets <u>must</u> be used when invasive procedures are to be performed during the examination, or likely to be within the next 24 hours, as decontamination of the skin does not fully remove gel.
- For examinations on non-intact skin or where contact with mucous membranes occur,
   e.g. transvaginal or transrectal scans, sterile gel must be used both inside and outside the probe cover, as small perforations may be present in probe covers.
- Any examination taking place on severely immunocompromised individuals or in high dependency settings require sterile gel.
- In labour where there is high likelihood of C-section or invasive instrumentation during delivery
- Where the ultrasound examination is near to an indwelling invasive device, such as an intravenous line or suprapubic catheter
- Where there is contact with or near to non-intact skin (any alteration in skin integrity such as a rash or surgical wound, including umbilicus in neonates)

# **Guidance for Use**

- Ensure that **only** unopened sachets and containers labelled as sterile are used
- Do not reuse the container or sachet once opened, either with the same or other patients. The container or sachet **must** be discarded after use



# 4.0 Equipment Required

Non-sterile gel. Sterile gel

#### 5.0 Training

This document will be made available and accessible for all intended user groups.

It will form part of the Local Induction for new staff members. No formalized training required

#### 6.0 Financial Risk Assessment

1	Does the implementation of this policy require any additional Capital resources				
2	Does the implementation of revenue this policy require additional resources	No			
3	Does the implementation remove of this policy require additional manpower	No			
4	Does the implementation of this policy release any manpower costs through a change in practice				
5	Are there additional staff training costs associated with implementing this policy which cannot be delivered through current training programmes or allocated training times for staff.				
	Other comments				

No financial risks have been identified at the time that this policy was developed.

#### 7.0 Equality Impact Assessment

The initial screening of this policy has not identified any adverse/negative impact and therefore a full equality analysis is not required. The completed general screening proforma has been sent to relevant personnel (from OP73 Undertaking an Equality Analysis (EA)

#### 8.0 Maintenance

This document will be reviewed and kept up to date by the Ultrasound Manager Karen Hill. Prior to review date all intended heads of department will be consulted to consolidate any changes and or amendments

#### 9.0 Communication and Training

No training required

Following approval and ratification, this procedural document will be published in the Trust's formal documents library and all intended staff will be notified through the Radiology Directorates normal notification process.



#### 10.0 Audit Process

This Standard Operating Procedure will be monitored through Radiology Governance Meetings.

Criterion	Lead	Monitoring method	Frequency	Committee
Peer to peer audit will apply to ensure compliance	Karen Hill	Random audit of practice	Quarterly	Modality Leads Meeting - held weekly . Results and any recommendat ions will be discussed quarterly

#### 11. References

The Medicine and Healthcare products Regulatory Agency

(MHRA) National Patient Safety Alert which clarifies the actions required to reduce harm and practice safely. 11 November 2021

Guidelines for Professional Ultrasound Practice, Society and College of Radiographers (SCoR) and British Medical Ultrasound Society (BMUS) 2020



# **Part A Document Control**

Procedure/	Title of	Status:		Author:				
Guidelines	Procedure/Guideline	s						
number and				Karen Hill				
version				Ultrasound Manager				
Version 2.0	safe use of			Sponsor: Chief				
	ultrasound gel			Nursing Officer				
SOP25	Ū			0				
Version /	Version	Date	Author	Reason				
Amendment								
History	1.0	Feb. 2022	Karen Hill I	Introduction of SOP				
	2.0	July 2025	Karen Hill	Review				
Intended Recipie	nts:							
	diologists, intensivists, m	nidwives, vascula	ar access spe	ecialists, emergency				
•	nedical clinicians, physio							
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	her locally approved							
committee (if loc								
document)								
Date of Procedur	Septembe	September 2025						
Review Date and	Frequency (standard	September	September 2028 and every 3 years thereafter					
review frequency	is 3 yearly unless							
otherwise indicate	d)							
Training and Diss								
No training required								
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in an alternative format e.g., larger print please contact Policy Manager Officer for Trust- wide documents or your line manager or Divisional Management office for Localdocuments.								
Contact for Review  Miss Karen Hill								
Monitoring arrangements Radiology Governance								
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Day to Day practice  Document summary/key issues covered: The safe storage and use of ultrasound gel								
Key words for intranet searching Ultrasound Gel								
purposes								
Parhoses								