



## **ARHAI Scotland**

Antimicrobial Resistance and Healthcare Associated Infection

**Disclaimer:** When an organisation, for example a health and care setting, uses products or adopts practices that differ from those stated in the National Infection Prevention and Control Manual, that individual organisation is responsible for ensuring safe systems of work including the completion of a risk assessment approved through local governance procedures.

This Guidance has been developed as part of an emergency response to the current mpox outbreak and therefore does not follow normal NIPCM methodology and process.

# **Version history**

Version	Date	Summary of changes
V1.0	01/06/22	First publication
V1.1	06/06/22	Greater clarity on symptomology associated with PPE use – section 2.2
		Clarity on cleaning and PPE requirements – <u>section 2.3</u>
		Inclusion of PPE requirements for vaccinators when vaccinating contacts who do not meet possible/probable/confirmed case definition – <a href="mailto:section2.7">section 2.7</a>
V1.2	08/07/22	Updated to reflect changes to HCID categorisation of MPX outbreak clade.
		Changes throughout the document to refer to MPX outbreak clade.
		Updated to include Department of Transport Multilateral Agreement M347 on the re-categorisation of waste for MPX outbreak clade to Category B.
V1.3	18/08/22	Update to reflect agreed name changes to Monkeypox variants with current outbreak variant identified as Monkeypox (Clade IIb)
V1.4	26/01/23	Update to reflect UKHSA derogation from HCID for MPXV (organism that causes mpox)
		Change of name from Monkeypox to mpox.
V1.5	29/06/2023	Review of extant guidance following update to UKHSA IPC guidance. No changes to ARHAI Scotland guidance, condensed general information and updated electronic links to UKHSA guidance as appropriate.

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#### 1. General information

General information on mpox including case definitions, clinical pathways and contact tracing can be found in the <a href="PHS guidance: Mpox guidance for health protection teams.">PHS guidance: Mpox guidance for health protection teams.</a>

This document outlines the infection prevention and control advice for healthcare workers who may be involved in the care of possible, probable or confirmed cases of mpox. The guidance is based on the published <a href="Moox Principles for control of non-HCID mpox in the UK: 4 nations">Moox Principles for control of non-HCID mpox in the UK: 4 nations</a> <a href="Consensus statement">Consensus statement</a> from the UK Health Security Agency (UKHSA) and associated <a href="UKHSA">UKHSA</a> <a href="Moox guidance collection">Mpox guidance collection</a>. The guidance should be read in conjunction with the <a href="MIPCM">NIPCM</a> and links to the appropriate sections of the NIPCM will be provided within this guidance.

Transmission of non-HCID mpox is consistent with close direct contact. There is currently no evidence that individuals are infectious before the onset of the prodromal illness. For individuals with infection who have evidence of lower respiratory tract involvement or severe systemic illness requiring hospitalisation, the possibility of airborne transmission has not been excluded: See <a href="https://doi.org/10.1007/jhei.com/hcic.new/">The Approved List of Biological Agents, Advisory Committee on Dangerous Pathogens (ACDP)</a>.

#### 2. Patient management

The derogation of <u>HCID</u> status therefore means a change in how mpox is defined. The operational definition going forward is therefore:

# Confirmed or highly probable cases of mpox are considered an HCID if the lineage responsible is:

confirmed Clade I mpox virus (MPXV),

or

- not known and:
  - there is a travel history to Central Africa, or a link to a traveller from those regions and/or
  - o there is an epidemiological link to a case of Clade I mpox and/or
  - o the case results from a new zoonotic jump in any country or setting

#### Mpox is not considered an HCID where:

confirmed as Clade II MPXV

or

- not known, and all the following conditions apply:
  - o there is no history of travel to Central Africa
  - o there is no link to a traveller from Central Africa
  - o the case has not resulted from a new zoonotic jump

When assessing a patient for possible mpox, ensure you assess the travel and contact history as above (see also the <u>case definitions</u>).

This derogation of HCID status for mpox is to ensure a proportionate response to deliver on achievable strategic outcomes. These principles do not replace the need for local dynamic risk assessments which remain key.

In preparation, healthcare settings that may receive and care for probable, possible, or confirmed non-HCID mpox cases should ensure that staff are:

- aware of what actions to take if a possible, probable or confirmed case presents
- familiar with all IPC controls required as per this guidance

#### 2.1 Patient placement

In line with the <u>hierarchy of controls</u>, efforts should be made to perform telephone triage/assessment to help establish symptoms present and risk associated with potential mpox in advance of any face-to-face contact where possible.

Possible, probable, or confirmed non-HCID mpox cases who require to be seen by healthcare staff within a healthcare facility should be placed in a negative pressure room (or a single neutral pressure room where negative pressure room is unavailable). Where there are minimal numbers of negatively pressurised rooms, these should be prioritised for confirmed cases.

All possible, probable, or confirmed patients should be provided with a Fluid Resistant Surgical Mask (FRSM) to wear where this can be tolerated and does not compromise their clinical care, for example when receiving oxygen therapy.

#### 2.2 Personal Protective Equipment

Table 1: Minimum PPE requirements for possible, probable, and confirmed non-HCID mpox

Definition categories	Minimum PPE required
Where symptoms are mild (this may include a localised rash) and there is no evidence of respiratory symptoms.	<ul> <li>Gloves – single pair. Ensure hand hygiene is performed appropriately prior to and after removal of gloves</li> <li>Fluid Resistant Surgical Facemask (FRSM)</li> <li>Apron</li> </ul>
Possible, Probable and Confirmed – Where symptomology includes respiratory symptoms, widespread rash AND/OR	<ul> <li>Gloves – single pair. Ensure hand hygiene is performed appropriately prior to and after removal of gloves.</li> <li>FFP3 respirator<sup>1</sup></li> </ul>

Definition categories	Minimum PPE required
clinically deteriorating as a direct result	Fluid resistant gown/coveralls
of mpox	Eye/face protection
AND/OR	
prolonged close contact with a patient	
and their environment for example an	
overnight inpatient admission stay	

A full face visor is required in addition to a FFP3 respirator where the respirator is not fluid resistant. HCWs must be fit tested prior to donning a respirator and perform a fit check each time it is donned.

Donning and doffing step-by-step instructions can be found in Appendix 6 of the NIPCM.

ARHAI Scotland recognise that some Infectious Disease Units will have established processes and procedures for HCID PPE which will be clearly defined as 'standard practice' for that area, which may include the application of a high-level unified ensemble for all HCIDs. Should they choose to adopt these to prevent confusion they may do so.

#### 2.3 Decontamination

Equipment in the room where a possible, probable or confirmed case is being managed should be kept to a minimum. Reusable patient care equipment should be dedicated to the patient as far as practicable. The non-HCID mpox virus will be destroyed by hospital detergents and disinfectants.

Decontamination of reusable patient care equipment after use on the possible, probable or confirmed case should be in line with Appendix 7 of the NIPCM.

Cleaning and decontamination of the patient room within healthcare settings should be undertaken using:

- a combined detergent/disinfectant solution at a dilution of 1,000 parts per million available chlorine (ppm available chlorine (av.cl.)); or
- a general-purpose neutral detergent in a solution of warm water followed by disinfection solution of 1,000ppm av.cl.

Manufacturers' guidance and recommended product "contact time" must be followed for all cleaning/disinfection solutions.

Increased frequency of decontamination/cleaning schedules should be incorporated into the environmental decontamination schedules for rooms occupied by possible, probable, or confirmed non-HCID mpox cases where there may be higher environmental contamination rates for example inpatient rooms. Staff cleaning the room should wear PPE in line with <a href="section 2.2">section 2.2</a> whist the patient is still present.

Inpatient rooms must be terminally cleaned following resolution of symptoms, discharge and/or transfer of the possible, probable, or confirmed non-HCID mpox case. Equipment should be cleaned, linen bagged, and waste removed by the clinical staff who have been caring for the patient wearing the PPE outline in <a href="mailto:section 2.2">section 2.2</a>. Curtains and bed screens should also be removed for laundering. Where domestic staff are required to remove curtains/screens they should do so wearing PPE outlined in <a href="mailto:section 2.2">section 2.2</a>. Subsequent cleaning of the empty room to complete the terminal clean thereafter may be undertaken whilst wearing an apron, gloves and FRSM.

In primary care settings where a consultation with a possible, probable, or confirmed non-HCID mpox case has taken place, ensure thorough cleaning using the same products stated above once the patient leaves the room and wearing the PPE outlined in <u>section 2.2</u>. This must include all equipment and surfaces and as a minimum, a full floor clean at the end of the day.

#### 2.4 Waste

By international agreement, samples and waste from **all** mpox cases are classified as Category B for transport and waste management. Infectious waste from these individuals can be treated as healthcare (clinical) category B waste as per <u>Multilateral Agreement M347 under section</u>

1.5.1 of ADR on the carriage of monkeypox virus and can be disposed of in an orange bag for alternative treatment and does not have to be sent for incineration. The waste will be assigned to UN3291, clinical waste.

If there is any chemical or pharmaceutical contamination, the waste must be consigned in a yellow container (or purple if cytotoxic/cytostatic) and incinerated or sent to a permitted site for disposal as per national regulation.

Laboratory cultures of MPXV remain classified as Category A.

#### Linen management

Contaminated clothing and linen are a potential source of transmission. Care must be taken not to shake the linen and prevent dispersal of skin scales. All linen generated during the care of a possible, probable, or confirmed case of non-HCID mpox must be managed as infectious linen in line with Appendix 8 of the NIPCM.

#### 2.5 Safe management of blood and body fluids

Spillages of blood and/or other body fluids associated with a possible, probable or confirmed case of non-HCID mpox must be treated immediately in line with <u>Appendix 9</u> of the NIPCM.

#### 2.6 IPC advice for MPX vaccination teams

If vaccinating a possible, probable, or confirmed non-HCID mpox case, the content of this guidance document should be followed.

If vaccinating asymptomatic contacts who **do not** fit the possible, probable, or confirmed case definition or HCWs as part of a routine response where that HCW has not had any exposure to a case of non-HCID mpox, IPC requirements are those in line with standard IPC vaccination protocols and there is considered no additional risk. Gloves are not routinely required for vaccination. An <u>SBAR detailing the evidence around this</u> is available.

# 2.7 Visitors to an inpatient who is a possible, probable, or confirmed case of non-HCID mpox

Visitors to possible, probable, or confirmed non-HCID mpox inpatients should be restricted. If essential, for example carer/parents/guardians, individual advice should be sought from IPCT/HPT regarding the safest way to arrange a visit.