

NATIONAL AMBULANCE SERVICE  
INFECTIOUS PREVENTION & CONTROL GROUP (NASIPCG)

**Monkeypox Guidance**

**POSITION STATEMENT – 09/08/22 (V1.1)**

**Please note this is a moving situation and further updates may be required.**

**Latest Update**

As of 09/08/22 the UK Health Security Agency (UKHSA) has been notified of over 2,914 confirmed cases of monkeypox within the UK. The latest number of cases can be found [here](#).

Monkeypox is a viral zoonotic disease that until May 2022, was primarily identified in Central and West Africa. There are 2 historical clades of monkeypox – a Central African clade with a reported mortality of 10% and a West African clade with a reported mortality of 1% from epidemiological cluster and outbreak reports from Africa. Prior to 2022, it was occasionally identified in other countries related to travel from endemic areas in Central and West Africa.

Within the UK, monkeypox has previously been classified as a High Consequence Infectious Disease (HCID). This is not a legal classification but agreed by UKHSA and the NHS to enable a consistent approach to public health and clinical or NHS management.

Since 13 May 2022, cases of monkeypox have been reported in multiple countries that do not have endemic monkeypox virus in animal or human populations, including countries in Europe, North America and Australasia. Epidemiological investigations are ongoing; however reported cases thus far have no established travel links to an endemic area. This suggests significant community transmission in multiple non-endemic countries in recent weeks. In the UK, all reported cases have been identified as the West African clade through rapid molecular testing.

The situation in the UK has now changed. There are now many cases and it is clear that community transmission is occurring with multiple generations of spread. Illness appears to be generally mild, though some individuals will require hospital admissions to manage secondary infections or complications from the illness. [Pre and post exposure prophylaxis](#) using Imvanex is now available. Due to this changed context, the clade of monkeypox currently circulating in the UK is no longer classified as an HCID.

However, future importations of monkeypox from West Africa and/or monkeypox caused by the Central African clade should remain classified as an HCID as the severity of the original clades remains unknown.

These principles are 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.

However, any Possible, Probable or Confirmed cases of MPX with a travel history to West or Central Africa, **OR** a link to a traveller from those regions. **OR** with a link to an outbreak which is known to be outside Lineage B.1 or sequenced and known to be outside Lineage B.1 should still to be classed and managed as a HCID.

Transmission in this 2022 MPX outbreak is consistent with close direct contact. There is currently no evidence that individuals are infectious before the onset of the prodromal illness.

## Background

The purpose of this document is to provide infection prevention and control (IPC) measures to prevent transmission of monkeypox (MPX) in health and care settings in England. This document should be read in conjunction with [the National Infection Prevention and Control Manual \(NIPCM\) for England](#) specifically appendix 11b, and [UKHSA principles for monkeypox control in the UK: 4 nations consensus statement](#).

## Presentation

The incubation period of monkeypox is between 5 and 21 days after exposure, usually 6 to 16 days, with an initial clinical presentation of:

- Fever.
- Headache.
- Muscle aches.
- Backache.
- Swollen lymph nodes.
- Chills.
- Exhaustion.

Within 1 to 5 days after the appearance of fever, a rash develops, often beginning on the face or genital area, then spreading to other parts of the body. The rash changes and goes through different stages before finally forming a scab which later falls off. An individual is contagious until all the scabs have fallen off and there is intact skin underneath. The scabs may also contain infectious virus material.

*Figure 1 – Images of Monkeypox Rash*



In some of the new cases, the rash has presented on genitals and occurred before the fever prodrome.

The rash is sometimes confused with [chickenpox](#). It starts as raised spots, which turn into small blisters filled with fluid. These blisters eventually form scabs which later fall off. The symptoms usually resolve in 2 to 4 weeks.

### **Transmission**

Monkeypox does not spread easily between people. It may occur when a person comes into close contact with an animal (rodents are believed to be the primary animal reservoir for transmission to humans but monkeypox is not found in UK rodents at present), human, or materials contaminated with the virus.

The virus enters the body through broken skin (even if not visible), the respiratory tract, or the mucous membranes (eyes, nose, or mouth). Person-to-person spread is uncommon, but may occur through:

- Contact with clothing or linen (such as bedding or towels) used by an infected person.
- Direct contact with monkeypox skin lesions or scabs.
- Coughing or sneezing of an individual with a monkeypox rash.

It is not a sexually transmitted disease but may be spread by close skin contact during sexual activity.

### **Infection Prevention and Control**

Monkeypox is a hazard group 3 organism ([ACDP/HSE](#)). Other organisms in this category include Salmonella typhi, HIV, Hepatitis B and C, and Mycobacterium tuberculosis that are managed routinely in the community.

**Frontline and other operational ambulance resources may transport these patients. HART (Hazardous Area Response Team) should not be used for these transfers, but must be considered for any Monkeypox cases that fall outside the current outbreak clade.**

These principles do not replace the need for local dynamic risk assessments which remain key. Based on the level of exposure and risk assessment consideration of the hierarchy of controls will help determine the level of personal protective equipment (PPE) to use.

For any individual presenting in person for advice/treatment, an infection risk assessment requires clinical judgement to identify the risk of cross transmission based on the presenting case/operational definition of MPX as defined UKHSA 9<sup>th</sup> August (link below), and the PPE required: [Monkeypox: case definitions - GOV.UK \(www.gov.uk\)](#)

### **Monkeypox:**

- i. Without travel to West or Central Africa and without a link to a traveller from those regions  
  
AND/OR
- ii. Confirmed by sequencing to be within the current outbreak clade (Lineage B.1)

**is NOT considered a high consequence infectious disease.**

### **Monkeypox:**

- i. With a travel history to West or Central Africa, a link to a traveller from those regions

OR

- ii. With a link to an outbreak which is known to be outside Lineage B.1 OR iii. sequenced and known to be outside Lineage B.1 OR iv. which results from a new zoonotic jump in any country or setting.

**is considered a high consequence infectious disease.**

PPE requirements will differ based on clinical judgement to include:

- the operational/ case definition and whether the case is considered a HCID
- the individual's presenting symptoms if presenting with an unexplained rash or other symptoms such as disseminated lesions or deteriorating condition resulting in a clinical suspicion of MPX
- presence of respiratory symptoms / disseminated lesions or if the patients condition is deteriorating
- the clinical procedures being undertaken e.g. prolonged clinical contact within 1 metre.

Table 1: Minimum PPE requirements for possible, probable and confirmed MPX cases

Case groups	MINIMUM PPE required
<p><b>Cases NOT considered as high consequence infectious disease. Possible, Probable and Confirmed MPX.</b></p> <p>Without travel to West or Central Africa and without a link to a traveller from those regions and/or confirmed by sequencing to be within the current outbreak clade (Lineage B.1)</p> <p><b>AND</b></p> <p>Where symptomatology is limited to a rash and patient is generally well – <b>NO respiratory symptoms.</b></p>	<ul style="list-style-type: none"> <li>• Disposable gloves – single pair</li> <li>• Fluid Resistant surgical facemask (FRSM – Type IIR):</li> <li>• An Apron</li> <li>• Face/eye protection (if there is a risk of spraying/splashing)</li> </ul>
<p><b>Possible, Probable and Confirmed MPX</b></p> <p>Where symptomatology <b>includes respiratory symptoms</b>, widespread rash and/or clinically deteriorating as a direct result of MPX</p> <p><b>AND/OR</b></p> <p>Prolonged* close contact with a patient and their environment, for example inpatient care or repeated assessment of an individual who is clinically unwell or deteriorating</p>	<ul style="list-style-type: none"> <li>• An FFP3 respirator<sup>1</sup> (fit-tested and fit-checked) or equivalent e.g. powered air purifying respirator (PAPR)<sup>1</sup> rather than FRSM</li> <li>• Disposable gloves</li> <li>• A disposable, fluid-resistant gown (coveralls may be worn in some settings e.g. ambulance)</li> <li>• A full face visor<sup>1</sup></li> </ul>
<p><b>Cases considered as high consequence infectious disease</b></p> <p>Possible, Probable or Confirmed cases of MPX with a travel history to West or Central Africa, <b>OR</b> a link to a traveller from those regions. <b>OR</b> with a link to an outbreak which is known to be outside Lineage B.1 or sequenced and known to be</p>	<p><b>TREAT AS HCID – FULL HCID PPE ENSEMBLE</b></p>

outside Lineage B.1 <b>OR</b> which results from a new zoonotic jump in any country or setting.	
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1. \*Please note many FFP3 respirators are not fully fluid-resistant, therefore a full face visor is recommended.

The PPE requirements in table 1 apply to all patient care activities including triage, testing, direct clinical care, cleaning of the equipment and the environment, management of waste, linen and blood and body fluid spillages.

Guidance for safe donning and doffing of PPE is available in the NIPCM appendix 6.

All possible, probable and confirmed cases of MPX should be provided with a facemask (type II or type IIR) in all healthcare settings to be worn for the duration of the treatment/consultation/transport unless removed for clinical assessment/treatment.

### **Waste including PPE:**

From July 2022, the international agreed approach for managing clinical waste and diagnostic samples is that they are now managed as Category B, similar to the other organisms in Hazard Group 3.

All waste generated in the care of possible, probable, or confirmed MPX cases must be managed as category B waste. Refer to the Department for Transport Multilateral Agreement M347 under section 1.5.1 of ADR on the carriage of monkeypox virus (see UKHSA guidance)– applicable to ALL clades of MPX until 2025.

Advice can be sought from the local waste contractor, a Dangerous Goods Safety Adviser, or in [Health Technical Memorandum 07:01 'Safe Management of Healthcare Waste'](#).

### **Linen:**

All linen generated in the care of possible, probable, or confirmed MPX must be managed as infectious linen and bagged into a water soluble or soluble seam (alginate) bag then placed into a polythene bag or impermeable sack.

### **Decontamination**

- Following convey of a possible case; carefully clean all equipment and vehicle environment with combined detergent/disinfectant wipe wipes and allow to air dry.
- For the convey of a highly probable (displaying extensive symptoms including weeping pustules and respiratory symptoms) or confirmed cases, vehicles must be initially cleaned using a detergent/disinfect wipe followed by a solution of 1,000ppm av.cl. (or alternative locally agreed cleaning product).
- PPE for cleaning should match the level of PPE for clinical patient care.

**Discharge of patients:** Advice outlined in this document on safe working practices apply. Ambulance services should follow ambulance standard operating procedures for managing infectious individuals (including HCIDs).

Patient discharges by an organisations PTS provider will be undertaken in a suitable vehicle and ensuring:

- the patient does not share the transport with other patients
- the patient lesions are covered if visible (face/hands) wherever possible
- the patient wears an FRSM (if tolerated)
- the patient is physically separated from the driver.
- Staff undertaking the transfer are wearing appropriate PPE as per table 1
- the vehicle must be decontaminated after each discharge /transfer following agreed local SOP (terminal clean) using:
  - a combined detergent disinfectant solution at a dilution (1,000ppm av.cl.); or
  - a general-purpose neutral detergent in warm water followed by a solution of 1,000ppm av.cl. (or alternative locally agreed cleaning product)

Further information and guidance can be found on UKHSA and NHSE websites. If you have any questions please speak to your trust IPC lead.

**Ref:**



IPC measures for  
possible probable anc

[Principles for monkeypox control in the UK: 4 nations consensus statement - GOV.UK \(www.gov.uk\)](https://www.gov.uk/guidance/4-nations-consensus-statement-on-monkeypox)

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[https://www.gov.uk/guidance/hcid-status-of-monkeypox,](https://www.gov.uk/guidance/hcid-status-of-monkeypox)

[C1636-national-ipc-manual-for-england-v2.pdf](#) 5/7/22

[Monkeypox: case definitions - GOV.UK \(www.gov.uk\)](https://www.gov.uk/guidance/monkeypox-case-definitions) 9/8/22