Rabies

The disease

Rabies is an acute viral encephalomyelitis caused by members of the lyssavirus genus. The disease may be caused by rabies virus genotype 1 (classical rabies) or less commonly by rabies-related lyssaviruses. The presentations are clinically indistinguishable. Rabies-related lyssaviruses implicated in human disease include European bat lyssaviruses (EBLVs) and Australian bat lyssavirus (ABLV).

Infection is usually via the bite or scratch of a rabid animal, most frequently a dog. In some parts of the world, other animals are important sources of exposure. In parts of Europe (including the UK) EBLV-1 and EBLV-2 are found in insectivorous bats and have occasionally caused human disease.

On rare occasions, transmission of the virus has occurred through body fluids from an infectious animal coming into contact with an individual's mucous membranes. Exposure through mucous membranes has a low probability of infection but must be managed as a significant event. Infection does not occur through intact skin. Virus is present in some tissues and fluids of humans with rabies, but person-to-person spread of the disease has not been documented other than in exceptional circumstances. Cases have occurred rarely outside the UK through corneal grafts and other transplanted tissues taken from individuals with rabies.

The incubation period is generally between three and 12 weeks, but may range from four days to 19 years. In more than 93% of patients, the onset is within one year of exposure. The onset of illness is insidious. Early symptoms may include paraesthesiae around the site of the wound, fever, headache and malaise. The disease may then present with hydrophobia, hallucinations and maniacal behaviour progressing to paralysis and coma, or as an ascending flaccid paralysis and sensory disturbance. Rabies is almost always fatal, death resulting from respiratory paralysis. There is no specific treatment other than supportive care once clinical symptoms develop.

History and epidemiology of the disease

Rabies in animals occurs in all continents except Antarctica, although individual countries and islands are reported to be rabies-free. In the US, classical rabies virus in animals has become more prevalent since the 1950s; skunks, raccoons and bats account for 85% of animal cases. In Asia, Africa, Central and South America, classical rabies virus (genotype 1) is endemic in feral dogs and is also present in domestic dogs. In Mexico and Central and South America, vampire bats carry the classical rabies virus. Most countries that are declared rabies-free probably have rabies-related viruses in their bat populations. In the UK, EBLV1 has been detected in serotine bats, and EBLV 2 has been detected in Daubenton's bats, but no lyssaviruses in the most common bat species in the UK, the pipistrelle (Folly et al, 2021). In other parts of Europe and in Australia, other bat species have been affected.
During the twentieth century, rabies in wildlife spread through parts of Central and Western Europe. Foxes have been the main host, but many other animals have also been infected, particularly dogs. The incidence of endemic, fox-adapted rabies in Western Europe fell dramatically in the last years of the twentieth century. This has been largely due to the immunisation of wild and domestic animals. Rabies continues to be reported in domestic animals imported from rabies endemic countries. Rabies remains prevalent in certain countries in Eastern Europe.

Worldwide an estimated 59,000 people die of rabies each year, with the majority of deaths occurring in Asia (59.6%) and Africa (36.4%) (WHO, 2018a). In the UK, deaths from classical rabies continue to occur in people infected abroad. Such instances are, however, rare, with 26 deaths having been reported since 1946, six of which have occurred since 2000 and the most recent was in 2018. None had received timely and appropriate post-exposure treatment. An estimated 20 million people receive post-exposure treatment across the world each year (WHO, 2018a). Pre-COVID-19, approximately 2000 people each year required post-exposure treatment in England, of which 12% were potentially exposed to bats in the UK and 88% were potentially exposed to an animal overseas (PHE data, 2018).

No case of indigenous human rabies from animals other than bats has been reported in the UK since 1902. In 2002, a man died from rabies caused by EBLV-2 acquired in the UK from a bat (Fooks et al., 2003). Only three other human cases of EBLV infection (all fatal) have been reported in the past 30 years in Europe (Nathwani et al., 2003).

**The rabies immunisation**

There is currently one rabies vaccine licensed for intramuscular use in the UK – purified chick embryo cell rabies vaccine (PCEC, 2.5 IU/ml) (Rabipur®).

Rabies Vaccine BP (human diploid cell vaccine) was discontinued in August 2022.

Other WHO approved cell culture-derived vaccines are available in other countries and may contain different concentrations of rabies antigen. The UK licensed products (Rabipur®) contains 2.5IU rabies antigen in a 1ml dose; other products should be used as recommended by the manufacturer. Rabies Vaccine BP manufactured by Sanofi was discontinued in 2022.

Rabipur® is thiomersal-free. This vaccine is inactivated, does not contain live organism and cannot cause the disease against which it protects.

The PCEC rabies vaccine is a freeze-dried suspension of the Flury LEP-25 rabies virus strain cultured in chick embryo cells and inactivated with betapropiolactone. The potency of the reconstituted vaccine is not less than 2.5 IU per 1.0ml dose. It contains traces of amphotericin B, chlortetracycline and neomycin.

The cell-culture derived rabies vaccines may be used interchangeably to provide protection pre- or post-exposure (Cabasso et al., 1974; Turner et al., 1982; Fekadu et al., 1988; Briggs et al., 1992; Strady et al., 1998; Strady et al., 2000). Intramuscular immunisation with tissue-culture vaccines reliably produces rabies virus neutralising titres in approximately 95% of recipients (Nicholson et al., 1987; Fishbein et al., 1989; Strady et al., 1998). In 95% of individuals, rabies titres are long-lived, particularly if the vaccine has been administered intramuscularly compared with intradermal immunisation (Fishbein et al.,
1989; Briggs et al., 1992; Strady et al., 1998; Suwansrinon et al., 2006). Immunologically competent persons who have received a primary course of rabies vaccine have a primed immune response, and will respond promptly once they receive a booster dose of vaccine (Rosanoff et al., 1979; Turner et al., 1982; Fishbein et al., 1986; Naraporn et al., 1999).

Storage

Presentation
Rabipur
The vaccine is supplied as freeze-dried powder and solvent for suspension and for injection. The powder is white. The solvent is a clear, colourless solution. Following reconstitution with the solvent supplied, the suspension will be a clear-colourless solution and free from particles.

The vaccine should be used immediately and no later than one hour after reconstitution with the solvent supplied.

Dosage, schedule and administration
For primary pre-exposure immunisation, three doses of rabies vaccine (2.5 IU; one vial) should be given intramuscularly on days 0, 7 and 28. The third dose can be given from day 21 if there is insufficient time before travel.

Alternatively, an accelerated course of primary pre-exposure immunisation may be given if there is insufficient time before travel to complete the 21-28 day course. Three doses of rabies vaccine (2.5IU) should be given intramuscularly on days 0, 3 and 7, with an additional dose at one year if they will continue to travel to high risk (enzootic) areas. Where there is sufficient time to complete the 21-28 day course, this is the preferred schedule for those receiving pre-exposure prophylaxis.

For post-exposure treatment schedules, see Tables 27.3 – 27.5, as the doses required will depend on a risk assessment and the calculation of a Composite Rabies Risk (CRR).

Administration
Vaccines are routinely given intramuscularly into the upper arm or anterolateral thigh (Zuckerman, 2000). However, for individuals with a bleeding disorder, vaccines should be given by deep subcutaneous injection to reduce the risk of bleeding.

The Joint Committee on Vaccination and Immunisation recommends the intramuscular rather than the intradermal route for pre-exposure prophylaxis use of rabies vaccine. The committee also recommends that only the intramuscular route (or deep subcutaneous route for those with bleeding disorders) is used for post-exposure treatment.

Whilst the intramuscular route is preferred, suitably qualified and experienced healthcare professionals may give the vaccine via the intradermal route for pre-exposure prophylaxis. This ‘off label’ use of the intradermal route is on the prescriber’s own responsibility as this is not covered by the manufacturer’s Product Licence. For pre-exposure intradermal immunisation, 0.1 ml (0.25 IU) of the vaccine can be used according to the schedule above. Although approved by the World Health Organization, a two-site two-dose
intradermal vaccine course has not been recommended for use in the UK by the Joint Committee on Vaccination and Immunisation. Intradermal immunisation is reliable only if the whole of the 0.1 ml dose is given properly into the dermis and should only be given by those experienced in the intradermal technique. It should not be used in those taking chloroquine for malaria prophylaxis as this drug suppresses the antibody response if the vaccine is given by the intradermal route (chloroquine does not suppress the antibody response if the vaccine is given by the intramuscular route). Whilst the use of the intradermal route potentially allows the contents of a vial of rabies vaccine to be shared amongst more than one individual, this practice is not recommended and carries the risks of contamination (see chapter 4).

Rabies vaccines can be given at the same time as other vaccines, including other travel vaccines. The vaccines should be given at separate sites, preferably in different limbs. If given in the same limb, they should be given at least 2.5cm apart (American Academy of Pediatrics, 2006). The site at which each vaccine was given should be noted in the individual’s records. The vaccinee must keep a record of the vaccine and regimen received as it will influence future post-exposure treatment (see table 27.5).

Disposal

Rabies-specific immunoglobulin

Human rabies immunoglobulin (HRIG) is obtained from the plasma of immunised and screened human donors. Because of a theoretical risk of transmission of vCJD from plasma products, HRIG used in the UK is prepared from plasma sourced from outside the UK. All donors are screened for HIV and hepatitis B and C, and all plasma pools are tested for the presence of nucleic acid from these viruses. A solvent detergent inactivation step for envelope viruses is included in the intramuscular/sub-cutaneous products.

HRIG is used after high risk exposure to rabies to give rapid protection by neutralising the rabies virus at the wound site until rabies vaccine, which should be given at a separate site at the same time, becomes effective.

Storage
Human rabies immunoglobulin (HRIG) should be stored in a refrigerator between +2°C and +8°C. This product is tolerant to ambient temperatures for up to one week, and can be distributed in sturdy packaging outside the cold chain if needed.

Administration
When indicated for post-exposure treatment (see Table 27.5), HRIG at a dose of 20 IU/kg body weight should be infiltrated in and around the cleansed wound. HRIG is of greatest value when infiltrated at the wound site as it neutralises rabies virus at the wound site before the immune system can respond to the vaccine by producing antibodies. Where HRIG is recommended, every effort should be made to administer HRIG at the wound site rather than intramuscularly, as the benefit of intramuscular administration away from the site of the wound is likely to be negligible (WHO, 2018b).
HRIG dosage must be calculated using the weight of the patient and potency of the HRIG batch. The quantity of HRIG on the packaging is the minimum content of the vial, and must NOT be used for calculating the dose.

If infiltration of the whole volume is not possible, any remaining HRIG should be given intramuscularly in the anterolateral thigh, remote from the immunisation site although the additional benefit is likely to be limited. If more than 2 ml is to be given intramuscularly to children, or more than 5 ml to adults, the HRIG should be divided into smaller amounts and given into different sites.

If vaccine is given but HRIG treatment is delayed, HRIG can still be given up to seven days after starting the course of vaccine. HRIG is no longer required once an active antibody response to the rabies vaccine has started to develop. Therefore, HRIG is not indicated more than seven days after the first dose of vaccine, or more than one day after the second dose of vaccine.

Equine immunoglobulin (eRIG) or monoclonal antibody (mAb) products may have been given as part of rabies post-exposure treatment in other countries where access to HRIG is limited. If eRIG or mAb have been administered overseas, HRIG is not required.

**Disposal**

HRIG is for single use and any unused solution should be disposed - see chapter 3.

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**Recommendations for the use of the vaccine**

**Pre-exposure (prophylactic) immunisation and reinforcing immunisations**

To determine the need for pre-exposure immunisation, an individual risk assessment of potential exposures should be carried out. Individuals considered to be at risk of exposure to rabies virus are listed in Table 27.1 and Table 27.2 and should be offered pre-exposure rabies immunisation according to the schedule above.

The requirement for booster doses is dependent on an individual's indication for pre-exposure prophylaxis and the likely frequency of ongoing exposures. In those who may have frequent unrecognised exposures to the virus, e.g. bat handlers, a single reinforcing dose of vaccine should be given one year after the primary course has been completed. Further booster doses should then be given every three to five years or based on serology. Laboratory staff routinely working with the rabies virus should have their vaccine antibody tested at six monthly intervals to determine optimal timing for booster doses. Antibody levels of at least 0.5 IU/ml are considered protective (WHO, 2018b).

**Table 27.1 Pre-exposure (prophylactic) immunisation for those within the UK**

- laboratory staff routinely working with rabies virus
- workers at DEFRA-authorised quarantine premises and carriers
- those who regularly handle bats, including on a voluntary basis, in the UK
- veterinary and technical staff who, by reason of their employment, encounter enhanced risk
Table 27.2. Pre-exposure (prophylactic) immunisation for those travelling outside the UK

- animal control and wildlife workers, veterinary staff or zoologists who regularly work in rabies enzootic areas
- travellers to rabies enzootic areas, especially if:
  - post-exposure medical care and rabies biologics at the destination are lacking or in short supply
  - or they are undertaking higher risk activities such as cycling or running
  - or they are living or staying for more than one month

Routine booster doses are not recommended for most travellers. A single booster dose of vaccine can be considered, following a risk assessment, in those who have completed a primary course over one year ago and are travelling again to a high risk (enzootic) area. A complete pre-exposure primary course is considered to be three doses over 21-28 days or an accelerated three dose course (over 7 days) plus an additional dose of vaccine at one year.

Further information on country-specific rabies travel risk is available from the National Travel Health Network and Centre (http://travelhealthpro.org.uk) and, in Scotland, from Public Health Scotland’s, Travax (www.travax.nhs.uk – health professionals only, login required) or FitForTravel (www.fitfortravel.nhs.uk). All travellers to enzootic areas should also be informed by their medical advisers of the practical steps to be taken if they are bitten by an animal or have some other types of exposure that puts them at risk of rabies (e.g. when saliva from an infected animal comes into contact with broken skin or mucous membranes such as the eyes, nose, or mouth).

**Post-exposure management**
Post-exposure management normally consists of wound treatment and risk assessment for appropriate post-exposure treatment. Treatment and immunisation after a possible rabies exposure will depend on the circumstances of the exposure, including the nature of the exposure, the species involved, the country/area and the immune status of the exposed person.

Detailed guidance on risk assessment and management of potential rabies exposure can be found on the UK Health Security Agency (UKHSA) and Health Protection Scotland websites.


https://www.hps.scot.nhs.uk/a-to-z-of-topics/rabies/

**Wound treatment**
As soon as possible after the incident, the wound should be cleaned by thorough flushing under a running tap for several minutes and washing with soap or detergent and water. A suitable disinfectant should be applied and the wound covered with a simple dressing. Suitable disinfectants include 40 to 70% alcohol, tincture or aqueous solution of povidone-iodine.
Salivary exposures to mucous membranes such as eyes, nose or mouth should be washed thoroughly with clean water as soon as possible.

Primary suture could cause further damage to the wound and may increase the risk of introduction of rabies virus to the nerves. It should be avoided or postponed until post-exposure treatment has commenced. In patients requiring HRIG, sutures (and infiltration of local anaesthetic) should be delayed until HRIG has been infiltrated into the wound.

**Risk assessment**

Each case requires a full risk assessment based on detailed information about the circumstances of the potential exposure. Health care professionals should try to collect as much of this information as possible to inform the risk assessment. Risk assessment should be carried out rapidly, so that post-exposure treatment, if indicated, can be started promptly. Treatment may need to start before full information is available on the ownership and condition of the biting animal.

Information required to complete the risk assessment and initiate post-exposure treatment:

- patient name, date of birth, age, address, and NHS number if possible
- date of exposure
- species and current health status of animal involved if possible
- country of exposure
- category of exposure
- site (on body) of exposure
- whether the patient is immunosuppressed or has any allergies
- any previous rabies vaccinations or immunoglobulin treatment
- weight of the patient if HRIG is being considered

As the incubation period for rabies can be prolonged, treatment should still be considered even if the interval from exposure is lengthy. Risk assessment should always be done, even if the exposure occurred many months or years previously.

The following features are important considerations in the likelihood of transmission associated with a possible exposure:

**Country:**

- countries are classified as high, low and no risk for rabies in different animal species.

**Animal source:**

- rabies can be transmitted by the saliva of any warm blooded animal, including bats, monkeys and rodents as well as cats, dogs and foxes
- animals behaving abnormally represent a higher risk of infection (but normal appearance and behaviour do not exclude rabies)
- unprovoked bites carry greater risk than provoked bites
a regularly vaccinated animal is unlikely to be rabid but, rarely, vaccinated dogs have transmitted rabies

domestic dogs or cats behaving normally at 15 days after an exposure would not have been infectious at the time of the exposure. Therefore, where possible, domestic dogs and cats should be observed for 15 days to see if they begin to behave abnormally

Category and site of exposure

higher risk exposures are those with broken skin, including single or multiple transdermal bites, severe lacerations, or where mucous membranes or an existing skin lesion have been contaminated by the animal’s saliva or other body fluid. Intact skin is a barrier against infection

bites and severe lacerations represent a higher risk than scratches

bat bites from species in the UK are usually felt and not seen

proximal bites (e.g. head and neck) represent a shorter incubation period than distal wounds

Immune status of the individual

Individuals who are primed by pre-exposure rabies immunisation respond more rapidly to post-exposure treatment and in most cases do not require HRIG.

Post-exposure treatment may often have been started elsewhere and further treatment will depend upon the dates and type of vaccine already received.

Immunosuppressed individuals (as defined in Chapter 6) should be managed with a greater degree of caution as they may not respond as effectively to rabies immunisation

Post-exposure treatment

Using a combination of the country where the incident occurred and the type of terrestrial mammal or bat involved, and category of exposure (Table 27.3), a Composite Rabies Risk will be calculated (Table 27.4). This is then used, along with the immunisation status and immune competence of the patient to determine the post-exposure treatment required (see Table 27.5).
### Table 27.3 Categories of exposure (adapted from WHO)

<table>
<thead>
<tr>
<th>Category</th>
<th>Terrestrial mammals</th>
<th>Bats</th>
</tr>
</thead>
</table>
| 1        | No physical contact with saliva  
For example:  
• touching, stroking or feeding animals | No physical contact (i.e. no direct contact with the bat's saliva)  
For example:  
• touching a dead bat  
• touching a bat where the person was protected by a barrier capable of preventing saliva contact, such as a boot, shoe, or appropriate protective clothing  
• a bat in the same room as a person (including a sleeping person) in the UK/Ireland |
| 2        | Minimal contact with saliva and/or unable to infiltrate wound with HRIG if needed  
For example:  
• bruising or abrasions  
• licks to broken skin (i.e. over insect bites or scratches)  
• scratches  
• bites which do not break the skin | Uncertain physical contact (i.e. where there has been no observed direct physical contact (with saliva) but this could have occurred)  
For example:  
• handling a bat without appropriate protective clothing (i.e. gloves)  
• a bat becoming tangled in hair  
• potential contact with a bat in the UK/Ireland in someone who is unable to give an accurate history of an exposure (i.e. intoxicated individual, young child, individual with mental impairment)  
• any bat found in the room of a sleeping person outside of the UK or Ireland |
| 3        | Direct contact with saliva  
For example:  
• severe/deep lacerations (i.e. down to the muscle)  
• bites that break the skin  
• contact of mucous membranes with saliva (e.g. licks) | Direct physical contact with bat's saliva  
For example:  
• all bites or scratches  
• contamination of mucous membrane with saliva or bat droppings/urine |

### Table 27.4 Composite rabies risk (CRR) table

<table>
<thead>
<tr>
<th>Combined Country/Animal risk</th>
<th>Category 1 exposure</th>
<th>Category 2 Exposure</th>
<th>Category 3 exposure</th>
</tr>
</thead>
<tbody>
<tr>
<td>No risk</td>
<td>Green</td>
<td>Green</td>
<td>Green</td>
</tr>
<tr>
<td>Low risk</td>
<td>Green</td>
<td>Amber</td>
<td>Amber</td>
</tr>
<tr>
<td>High risk</td>
<td>Green</td>
<td>Amber</td>
<td>Red</td>
</tr>
<tr>
<td>Confirmed rabid animal</td>
<td>Green/Amber*</td>
<td>Red</td>
<td>Red</td>
</tr>
</tbody>
</table>

* Advice should be sought from Rabies and Immunoglobulin Service (RIGS) team in the assessment of these contacts. In Scotland, advice should be sought from local teams, see page 11 for contact details
Following a risk assessment, if post-exposure treatment is indicated, this should be given as in Table 27.5. It is particularly important that the first three doses are given as close as possible (within a day or two) of the recommended schedule. Post-exposure rabies vaccine should be given via the intramuscular route (or by deep subcutaneous injection for people with bleeding disorders). Treatment for immunosuppressed individuals should include the most immunogenic immunisation course available and HRIG, irrespective of previous rabies immunisation status, with antibody testing to ensure response to post-exposure treatment (see Table 27.5).

Table 27.5 Post-exposure treatment based on Composite rabies risk (CRR)

<table>
<thead>
<tr>
<th>Composite rabies risk</th>
<th>Post-exposure treatment</th>
<th>Fully immunised</th>
<th>Immunosuppressed</th>
</tr>
</thead>
<tbody>
<tr>
<td>Green</td>
<td>None</td>
<td>None</td>
<td>None</td>
</tr>
<tr>
<td>Amber</td>
<td>Four doses of vaccine on days 0, 3, 7, and 21</td>
<td>Two doses of vaccine on days 0 and 3-7</td>
<td>HRIG plus five doses of vaccine on days 0, 3, 7, 14 and 30***</td>
</tr>
<tr>
<td>Red</td>
<td>HRIG** plus four doses of vaccine on days 0, 3, 7, and 21</td>
<td>Two doses of vaccine on days 0 and 3-7</td>
<td>HRIG plus five doses of vaccine on days 0, 3, 7, 14 and 30***</td>
</tr>
</tbody>
</table>

* Persons who have incomplete pre- or post-exposure course of rabies vaccine seek advice (as below).

** HRIG is not required more than seven days after the first dose of vaccine, or more than one day after the second dose of vaccine (unless immunosuppressed).

*** Send sample for antibody testing.

If an individual arrives in the UK having started post-exposure treatment via the intradermal route, they should receive the remaining doses via the intramuscular route. Where a regime has been started that is different to that used in the UK, specialist advice should be sought.
Further details can be found in the detailed guidance on risk assessment and management of potential rabies exposures. Specialist advice on the assessment of the risk and appropriate management can be obtained from one of the following:

<table>
<thead>
<tr>
<th>Country</th>
<th>Contact</th>
<th>Telephone Number</th>
</tr>
</thead>
<tbody>
<tr>
<td>England</td>
<td>Rabies and Immunoglobulin Service (RigS), UK Health Security</td>
<td>0330 128 1020</td>
</tr>
<tr>
<td></td>
<td>Or local health protection team (HPT)</td>
<td>See <a href="https://www.gov.uk/health-protection-team">https://www.gov.uk/health-protection-team</a> for contact details</td>
</tr>
<tr>
<td>Wales</td>
<td>Public Health Wales All Wales Acute Response Service (AWARe)</td>
<td>0300 003 0032</td>
</tr>
<tr>
<td>Scotland</td>
<td>Local on-call infectious diseases:</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Aberdeen Royal Infirmary On-call infectious diseases consultant</td>
<td>0345 456 6000</td>
</tr>
<tr>
<td></td>
<td>University Hospital Crosshouse, Ayr On- call infection specialist</td>
<td>01563 521 133</td>
</tr>
<tr>
<td></td>
<td>Dumfries &amp; Galloway Royal Infirmary On- call infection specialist</td>
<td>01387 246 246</td>
</tr>
<tr>
<td></td>
<td>Forth Valley Royal Hospital On-call infectious diseases consultant</td>
<td>01324 566000</td>
</tr>
<tr>
<td></td>
<td>Queen Elizabeth University Hospital, Glasgow On-call infectious diseases consultant</td>
<td>0141 201 1100</td>
</tr>
<tr>
<td></td>
<td>Raigmore Hospital, Inverness On- call infection specialist</td>
<td>01463 704 000</td>
</tr>
<tr>
<td></td>
<td>University Hospital Monklands, Lanarkshire On- call infection specialist</td>
<td>01236 748 748</td>
</tr>
<tr>
<td></td>
<td>Ninewells Hospital, Dundee On-call infectious diseases consultant</td>
<td>01382 660 111</td>
</tr>
<tr>
<td></td>
<td>Victoria Hospital, Fife On-call public health team</td>
<td>01592 643 355</td>
</tr>
<tr>
<td></td>
<td>Western General Hospital, Edinburgh On-call infectious diseases consultant</td>
<td>0131 537 1000</td>
</tr>
<tr>
<td>Northern Ireland</td>
<td>Public Health Agency Duty Room</td>
<td>030 0555 0119</td>
</tr>
</tbody>
</table>

**Contraindications**

Pre-exposure rabies vaccine should not be given to those who have had:

- a confirmed anaphylactic reaction to a previous dose of rabies vaccine, or
- a confirmed anaphylactic reaction to any component of the vaccine
The PCEC rabies vaccine (Rabipur®) contains residues of chicken proteins (e.g. ovalbumin) so an alternative rabies vaccine may be considered for pre-exposure immunisation in those with severe egg allergy.

The intradermal 0.1ml pre-exposure vaccine regimens should not be used in those with immunosuppression or those taking chloroquine for malaria prophylaxis, as this can suppress the antibody response.

There are no contraindications to post-exposure treatment with rabies vaccine (including Rabipur® for those with severe egg allergies). In the event of a hypersensitivity reaction to a dose of a pre-exposure course, such individuals should still receive post-exposure immunisation if indicated, because the risks of rabies outweigh the risks of hypersensitivity. When there is a history of a hypersensitivity reaction to rabies immunisation, specialist advice should be sought and further doses given under close medical supervision.

### Precautions

Minor illnesses without fever or systemic upset are not valid reasons to postpone pre-exposure immunisation.

If an individual is acutely unwell, pre-exposure immunisation should be postponed until they have recovered. This is to avoid confusing the differential diagnosis of any acute illness by wrongly attributing any signs or symptoms to the adverse effects of the vaccine.

Post-exposure treatment should always be started without delay, even in the presence of other illness.

**Pregnant women and breast-feeding**

Pregnant women and breast-feeding mothers should be given pre-exposure prophylaxis if the risk of exposure to rabies is high following a risk assessment by a health professional and rapid access to post-exposure treatment would be limited.

Post-exposure treatment should be given to pregnant women and breast-feeding mothers when indicated.

**Immunosuppression and HIV infection**

Individuals with immunosuppression and HIV infection (regardless of CD4 count) may be given pre-exposure rabies vaccines in accordance with the recommendations above. These individuals may not make a full antibody response. Re-immunisation should be considered after treatment is finished and recovery has occurred or, in the case of those with HIV, when there has been immune recovery following commencing antiretroviral treatment (e.g. CD4 count is greater than 200 cells/μl).

Individuals who are immunosuppressed (and fulfil criteria in Chapter 6) who have a significant rabies exposure (i.e. red or amber Composite Rabies Risk) should receive five intramuscular doses (2.5IU) of rabies vaccine on days 0, 3, 7, 14 and 30, plus HRIG. Antibody tests are recommended to confirm response to the post-exposure treatment, for example at the same time as the fifth dose (i.e. day 30) and following completion of the course if needed.
Adverse reactions

All suspected adverse reactions should be reported to the Medicines and Healthcare products Regulatory Agency (MHRA). Anyone can report a suspected adverse reaction to the MHRA using the Yellow Card reporting scheme (https://yellowcard.mhra.gov.uk/).

Rabies vaccine may cause local reactions such as redness, swelling or pain at the site of injection within 24 to 48 hours of administration. Systemic reactions such as headache, fever, muscle aches, vomiting and urticarial rashes are rare. Delayed hypersensitivity reactions have been reported from the US. Reactions may become more severe with repeated doses. Neurological conditions, such as Guillain-Barré syndrome, have been reported extremely rarely; a causal association with immunisation is not established.

HRIG may cause local pain and low-grade fever, but no serious adverse reactions have been reported.

Management of cases

Human rabies is a notifiable disease. In the event of a case, or suspected case, of human rabies, the local health protection team should be informed.


Supplies

Rabipur is available from Valneva UK (Tel: 01252 762208) or MASTA (Tel: 0113 238 7500).

Under the Health and Safety at Work Act (HSWA) 1974 and the control of Substances Hazardous to Health (COSHH) Regulations 2002 employers have a responsibility to assess the risks for exposure to hazardous agents, including rabies virus, and to protect employees from those risks as is reasonably practicable (See also Chapter 12). This includes the provision of pre-exposure rabies immunisation if indicated.

In England, rabies vaccine for pre-exposure immunisation will only be provided by UK Health Security Agency for bat handlers where no formal employer can be identified. This can be obtained from the Rabies and Immunoglobulin Service (rigs.ig.clerks@nhs.net) using the dedicated form (https://www.gov.uk/government/publications/rabies-pre-exposure-request-form). For other pre-exposure indications, it can be obtained through local pharmacies by private prescription. UKHSA does not supply pre-exposure rabies immunisation for travellers. Rabies vaccine and HRIG for use in post-exposure treatment are available free of charge to patients and providers from UKHSA. To facilitate prompt access for patients, NHS services are expected to arrange local seven-day access to a small number of doses of rabies vaccine to initiate a course. Any vaccine held locally that is used for post-exposure treatment will be replaced free of charge by UKHSA. Information may be obtained from the local health protection team (for contact details see https://www.gov.uk/health-protection-team) or UKHSA Rabies and Immunoglobulin Service (Tel 0330 128 1020) or out of hours via the UKHSA Colindale Duty Doctor (Tel: 020 8200 4400).
In Northern Ireland, rabies vaccine for licensed bat handlers where no formal employer can be identified may be obtained on HS21 prescription for administration by the individual’s GP; HS21 must not be used for pre-exposure rabies immunisation for travel purposes. Rabies vaccine for post-exposure treatment is available from the Royal Victoria Hospital Pharmacy Department, BHSCT (Tel: 028 9024 0503) and HRIG from the Northern Ireland Blood Transfusion Service (Tel: 077 7461 9337, Duty Biomedical Scientist), advice on risk assessment is available from the PHA Duty Room (Tel: 030 0555 0119), if required.

In Scotland, pre-exposure immunisation is available for bat handlers, where no formal employer can be identified, through normal Health Board channels. Post-exposure treatment is available through local on-call infectious disease consultant/infection specialist or public health team: check Table 27.6 for local contact numbers.

In Wales, rabies vaccine for pre-exposure immunisation will only be for bat handlers where no formal employer can be identified. This can be obtained from the UKHSA Rabies and Immunoglobulin Service (see above for contact details and form). Post-exposure treatment is provided through the Public Health Wales All Wales Acute Response Service (AWARe) (Tel: 030 0003 0032).
References


Naraporn N, Khawplod P, Limsuwan K et al. (1999) Immune response to rabies booster vaccination in subjects who had postexposure treatment more than 5 years previously. J Travel Med 6(2): 134-6.


