**Publications number: GOV-19068**

**Patient Group Direction (PGD) for the supply of oseltamivir for treatment of seasonal influenza**

For the supply of oseltamivir for treatment of seasonal influenza for residents, users and staff of care facilities (with or without nursing), by registered healthcare practitioners identified in [Section 3,](#section3) subject to any limitations to authorisation detailed in [Section 2](#section2).

Reference: 20250808OseltamivirTreatment\_PGD

Version number: 6.0

Valid from: 8 August 2025

Review date: 8 August 2027

Expiry date: 7 August 2028

**The UK Health Security Agency (UKHSA) has developed this PGD for local authorisation in line with national recommendations.**

Those using this PGD must ensure it is organisationally authorised and signed in Section 2 by an appropriate authorising person, relating to the class of person by whom the product is to be supplied, in accordance with Human Medicines Regulations 2012 (HMR2012)[[1]](#footnote-2). **The PGD is not legal or valid without signed authorisation in accordance with** [**HMR2012 Schedule 16 Part 2**](http://www.legislation.gov.uk/uksi/2012/1916/schedule/16/part/2/made)**.**

Authorising organisations must not alter, amend or add to the clinical content of this document ([sections 4,](#Section4) [5](#Section5) [and 6](#Section6)); such action will invalidate the clinical sign-off with which it is provided.

As operation of this PGD is the responsibility of commissioners and service providers, the authorising organisation can decide which staff groups, in keeping with relevant legislation, can work to the PGD. Sections 2, 3 and 7 must be completed and amended within the designated editable fields provided, but only for the purposes for which these sections are provided, that is the responsibilities and governance arrangements of the NHS organisation using the PGD. The fields in Section 2 and 7 cannot be used to alter, amend or add to the clinical content. Such action will invalidate the UKHSA clinical content authorisation which is provided in accordance with the regulations.

The final authorised copy of this PGD should be kept by the authorising organisation completing Section 2 for 25 years after the PGD expires. Provider organisations adopting authorised versions of this PGD should also retain copies for 25 years after the PGD expires.

**Individual practitioners must be authorised by name, under the current version of this PGD before working according to it.**

Practitioners and organisations must check they are using the current version of the PGD. Amendments may become necessary prior to the published expiry date. Current versions of UKHSA seasonal influenza PGDs for authorisation can be found from: [Influenza post exposure prophylaxis and treatment: PGD templates](https://www.gov.uk/government/publications/influenza-post-exposure-prophylaxis-and-treatment-pgd-templates)

Any queries regarding the content of this PGD should be addressed to: [immunisation.resp\_viruses@ukhsa.gov.uk](mailto:immunisation.resp_viruses@ukhsa.gov.uk)

Enquiries relating to the availability of organisationally authorised PGDs and subsequent versions of this PGD should be directed to: insert local contact details

Change history

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| **Version number** | **Change details** | **Date** |
| 01.00 | Original PGD template developed | 11 December 2014 |
| 02.00 | Template reviewed; put into new PHE format and changes to clinical and organisational content made | 7 January 2016 |
| 03.00 | Review:   * updates to title and criteria for inclusion: wording changed to ‘residents/users and staff of care facilities’ * amendments to inclusion and exclusion criteria * additions to actions to be taken if the patient is excluded * renal impairment definitions added * information regarding not splitting packs * additions to patient advice * updated references * minor typographical changes for consistency with other PGDs | 5 June 2018 |
| 04.00 | * amendment to No. 5 criteria for inclusion * addition of haemodialysis to criteria for exclusion * additions to off label use * addition of quantity and duration of treatment of 10 days in immunocompromised individuals * updated references | 7 February 2019 |
| 05.00 | * criteria for inclusion: risk groups updated to align with the Green Book [Chapter 19](https://www.gov.uk/government/publications/influenza-the-green-book-chapter-19) * criteria for exclusion: removed unstable medical conditions, severely unwell, new or worsening breathing difficulties or chest pain and added note under additional information * criterial for exclusion: removed clinically significant drug interactions and added note under drug interactions * criteria for exclusion: added individuals who are immunocompromised and have chronic kidney disease (CKD) with creatinine clearance CrCl ≤10mL/min or who are on peritoneal dialysis; those taking zanamivir * additional information: added information for oseltamivir resistance and information for CKD * route and method of administration: additional information regarding taking with food * dose and frequency of administration: additional information for renal doses, CrCl and eGFR levels, immunocompromised individuals and obesity; updated dose and frequency table * quantity: amended for immunocompromised individuals * drug interactions: information regarding clinically significant drug interactions; information on influenza vaccinations * minor wording changes in line with standard UKHSA PGD text; change from PHE to UKHSA, updated references | 8 August 2022 |
| 06.00 | * updated standard wording for consistency with UKHSA PGD templates * characteristics of staff updated to include allied health care professionals and pharmacy technicians * clinical condition where PGD applies: circumstances updated, definition of ILI aligned with UKHSA management of acute respiratory infection outbreaks in care homes guidance, age defined as per NHSE annual flu letter * action to be taken when excluded: signposting to specialist in those immunocompromised with severe renal impairment * additional information: pregnancy and breastfeeding information updated * quantity to be supplied: addition of alternative supply during periods of shortages * adverse reactions: common and less common side effects updated in line with SPC * referral to non-medical prescribers added where referral to medical practitioner mentioned * references updated | 8 August 2025 |

1. **PGD development**

This PGD has been developed by the following on behalf of the UKHSA:

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| **Developed by:** | **Name** | **Signature** | **Date** |
| **Pharmacist**  (Lead author) | Shilan Ghafoor  Lead Pharmacist - Medicines Governance, UKHSA |  | 8 August 2025 |
| **Doctor** | Dr Matthew Donati  Consultant Medical Virologist, UKHSA SW Regional Clinical Network Laboratory and Severn Infection Sciences |  | 8 August 2025 |
| **Registered nurse** | Lesley McFarlane  Lead Immunisation Nurse Specialist, Immunisation Programmes Division, UKHSA |  | 8 August 2025 |

This PGD has been peer reviewed by the Seasonal influenza PGD Expert panel in accordance with the UKHSA PGD Policy. It has been agreed by the UKHSA Medicines Governance Group and ratified by the UKHSA Clinical Quality and Oversight Board.

**Expert panel**

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| **Name** | **Designation** |
| Dr Jamie Lopez Bernal | Chair, Consultant Epidemiologist, UKHSA |
| Dr Conall Watson | Consultant Epidemiologist – influenza and seasonal respiratory viruses, Immunisation and Vaccine-Preventable Diseases Division, UKHSA. Registered pharmacist |
| Dr Jamie Lopez Bernal | Consultant Epidemiologist, UKHSA |
| Mark Borthwick | Consultant Pharmacist, Oxford University Hospitals NHS Foundation Trust |
| Gemma Hudspeth | Senior Health Protection Practitioner, North East and Yorkshire Region, UKHSA. Registered nurse |
| Jo Jenkins | Associate Director Medicines Governance, Medicines Use and Safety, NHS Specialist Pharmacy Service |
| Michelle Jones | Principal Medicines Optimisation Pharmacist, NHS Bristol North Somerset and South Gloucestershire Integrated Care Board |

**2.** **Organisational authorisations**

The PGD is not legally valid until it has had the relevant organisational authorisation.

It is the responsibility of the organisation that has legal authority to authorise the PGD, to ensure all legal and governance requirements are met. The authorising body accepts governance responsibility for the appropriate use of the PGD.

**Insert authorising body name** authorises this PGD for use by the services or providers listed below:

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| Authorised for use by the following organisations and/or services |
| For instance, NHSE services |
| Limitations to authorisation |
| For instance, any local limitations the authorising organisation feels they need to apply in line with the way services are commissioned locally. This organisation does not authorise the use of this PGD by …. |

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| **Organisational approval (legal requirement)** | | | |
| **Role** | **Name** | **Sign** | **Date** |
| For instance, NHSE Governance Lead, Medical Director |  |  |  |

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| **Additional signatories according to locally agreed policy** | | | |
| **Role** | **Name** | **Sign** | **Date** |
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Section 7 provides a practitioner authorisation sheet. Individual practitioners must be authorised by name to work to this PGD. Alternative practitioner authorisation sheets may be used where appropriate in accordance with local policy, but this should be an individual agreement, or a multiple practitioner authorisation sheet as included at the end of this PGD

1. **Characteristics of staff**

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| **Qualifications and professional registration** | To be completed by the organisation authorising the PGD for instance: Registered professional with one of the following bodies:   * nurses currently registered with the Nursing and Midwifery Council (NMC) * pharmacists and pharmacy technicians currently registered with the General Pharmaceutical Council (GPhC * allied health care professionals currently registered with the Health and Care Professions Council (HCPC) but must be one of the registered professionals who can legally supply and administer under a PGD * additional registered healthcare professionals to be added by organisation authorising the PGD   The practitioners above must also fulfil the [Additional requirements](#Additionalrequirements) detailed below.  Check [Section 2 Limitations to authorisation](#limitations) to confirm whether all practitioners listed above have organisational authorisation to work under this PGD. |
| **Additional requirements** | Additionally practitioners:   * must be authorised by name as an approved practitioner under the current terms of this PGD before working to it * must have undertaken appropriate training for working under PGDs for supply/administration of medicines for example [Patient Group Directions - elearning for healthcare](https://www.e-lfh.org.uk/programmes/patient-group-directions/) * must be competent in the use of PGDs (see [NICE Competency framework](https://www.nice.org.uk/guidance/mpg2/resources) for health professionals using PGDs) * must be familiar with the product and alert to changes in the Summary of Product Characteristics (SPC) * must be competent to assess the individual and discuss treatment options * must have undertaken training appropriate for working under this PGD * must have access to the PGD and associated online resources * should fulfil any additional requirements defined by local policy * authorising organisation to insert any additional requirements   **The practitioner must be authorised by name, under the current version of the PGD, before working according to it.** |
| **Continued training requirements** | Authorising organisation to insert any continued training requirements |

**Note:** The authorising organisation should ensure staff working with this PGD are trained in addressing issues of consent, including those individuals with dementia. The healthcare professional working under this PGD should follow their existing organisational procedures in relation to consent.

1. **Clinical condition or situation to which this PGD applies.**

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| **Clinical condition or situation to which this PGD applies** | Treatment of influenza A and B:   1. When **all** of the following circumstances apply:  * there is indication that influenza virus is circulating in the community**[[2]](#footnote-3)**, such as UKHSA surveillance or advice from the Chief Medical Officer (CMO) or Department of Health and Social Care **and** * the person is in an ‘at-risk’ group, including being aged 65 years and over**[[3]](#footnote-4)** (see [inclusion criteria](#inclusion)) **and** * the person has an ‘influenza-like illness’ (ILI)**[[4]](#footnote-5)** and can start treatment within 48 hours of the onset of symptoms  1. Outside the periods when surveillance indicates that influenza virus is circulating in the community, if there is an outbreak of an ILI in a long-term residential or nursing home (care homes), oseltamivir may be offered to ‘at risk’ residents and ‘at risk’ staff as part of treatment for those who have symptoms of influenza. This is regardless of vaccination status. However, this should only be done if there is a high level of certainty that the causative agent in a localised outbreak is influenza, usually based on virological evidence of infection with influenza in the index case or cases.   UKHSA Health Protection Teams (HPTs) will advise on whether influenza is the likely causative agent. |
| **Criteria for inclusion**  Continued overleaf  **Criteria for inclusion** (continued) | This PGD will come into force only when either there is an indication that influenza virus is circulating in the community or when, in a localised outbreak, there is a high level of certainty the causative agent is influenza, as advised by the local HPT.  Individuals must:   1. Be a resident or user of a care facility or staff working in a care facility **and** 2. Be exhibiting signs or symptoms of an influenza-like illness (ILI) or confirmed to have tested positive for influenza **and** 3. Either be aged 65 years and over (regardless of risk group) **or,** if aged 13 – 64 years, must be in one of the defined risk groups as detailed in the Green Book chapter 19 and summarised below:  * chronic (long-term) respiratory disease such as asthma that requires continuous or repeated use of inhaled or systemic steroids or with previous exacerbations requiring hospital admission, chronic obstructive pulmonary disease (COPD) or chronic bronchitis * chronic heart disease or vascular disease, such as heart failure * chronic kidney disease (CKD) at stage three, four or five**[[5]](#footnote-6)** (see [Additional information](#ckd)) with some exceptions (see [criteria for exclusion](#renalexclusions)) * chronic liver disease * chronic neurological disease, such as Parkinson’s disease or motor neurone disease, or learning disability * diabetes or adrenal insufficiency * immunosuppression due to disease or treatment (refer to [the Green Book Chapter 19)](https://www.gov.uk/government/publications/influenza-the-green-book-chapter-19) * asplenia or dysfunction of the spleen * morbid obesity (defined as a BMI of 40 and above) * any other clinical risk group, as listed in [the Green Book chapter 19,](https://www.gov.uk/government/publications/influenza-the-green-book-chapter-19) that puts the individual at risk of complications of influenza * pregnant women at any stage of pregnancy (first, second or third trimesters) and up to 2 weeks post-partum (see [Additional information](#breastfeeding))  1. Be able to begin therapy within 48 hours of the onset of symptoms. Alternatively supply can be considered after 48 hours of the onset of symptoms, when the local HPT or a specialist in infection such as a medical microbiologist, virologist or specialist in infectious disease advises this could be considered**[[6]](#footnote-7)**. Note such supplies are a recommendation, not a direction to supply (see [footnote](#footnote6) 6 below). This is a clinical decision which rests with the practitioner working under this PGD and this is [off-label use](#offlabel). |
| **Criteria for exclusion[[7]](#footnote-8)**  Continued overleaf  **Criteria for exclusion** (continued) | Individuals will not be considered for treatment with oseltamivir under this PGD if the following criteria apply:   * they are not a resident or user of or working in a care facility * they are less than 13 years of age * they are receiving haemodialysis * they are immunocompromised and have CKD with creatinine clearance CrCl ≤10mL/min * they are immunocompromised and on peritoneal dialysis * they have a known allergy to oseltamivir or to any of the excipients in the capsules * they have been symptomatic with this episode of ILI for more than 48 hours, unless initiation is advised by the local HPT or infection specialist(see [footnote 6](#footnote6)) * they have disturbance of consciousness, delirium or excessive drowsiness * they have significant vomiting or are unable to drink fluids * they are receiving zanamivir or baloxavir * infected with oseltamivir resistant influenza |
| **Action to be taken if the individual or carer declines treatment** | Advise the individual or their carer of the possible consequences of refusing the treatment, the protective effects of the treatment, the risks of infection, the risks of spreading the disease to others in the care facility, disease complications and alternative sources of treatment.  Consider if the individual is suitable for treatment with zanamivir or refer to the local HPT or a specialist in infection such as a medical microbiologist, virologist or infectious diseases specialist for further guidance.  Document the refusal and the advice given in the individual’s patient record.  Inform the care home manager andthe GP or care home doctor without delay.  These individuals should be managed with bed rest, fluids and symptomatic remedies such as analgesics or referred to NHS services if necessary.  All individuals and their carers should be advised to seek medical advice if symptoms worsen or do not improve within a week. |
| **Action to be taken if the individual is excluded**  Continued overleaf  **Action to be taken if the individual is excluded**  (continued) | Some individuals excluded under this PGD may still be suitable for treatment with oseltamivir if clinically assessed and prescribed.  If they have CKD with CrCl ≤10mL/min or on peritoneal dialysis and are immunocompromised, refer to a specialist in infection such as a medical microbiologist, virologist or infectious diseases specialist for advice. If a decision to supply oseltamivir is made, a prescription or a Patient Specific Direction (PSD) will be required.  If more than 48 hours from symptom onset and there is no advice in place from the local HPT or specialist in infection such as a medical microbiologist, virologist or specialist in infectious disease, the HPT should be consulted or advice sought from a medical or non-medical prescriber.  Consider if the individual is suitable for treatment with zanamivir (see PGD for treatment with zanamivir in care facilities).  Any individual excluded under this PGD who is clinically assessed as requiring treatment and who is not suitable for treatment with zanamivir should be referred to local NHS services for advice without delay. |
| **Additional information**  Continued overleaf  **Additional information** (continued) | If an individual is severely unwell, has new or worsening breathing difficulties, chest pain or is otherwise medically unstable and may be at risk of hospitalisation, initiate the antiviral but ensure the individual is referred for assessment by an appropriate clinician, typically a doctor.  Zanamivir inhaler is recommended as first line therapy (see PGD for treatment with zanamivir in care facilities) in the following circumstances:   * if the HPT has advised the confirmed or dominant circulating influenza strain is higher risk for oseltamivir resistance and the individual is immunocompromised[[8]](#footnote-9) or * the individual is known to have oseltamivir resistant influenza whether immunocompromised or not or * the individual is strongly suspected to have oseltamivir resistant influenza whether immunocompromised or not, for example they have been in contact with known oseltamivir resistant influenza   If the individual is unable to use inhalers, seek advice from a specialist in infection such as a medical microbiologist, virologist or specialist in infectious disease.  It is normal practice to administer only one neuraminidase inhibitor to an individual at a time. Therefore supply either oseltamivir or zanamivir but not both (see PGD for treatment with zanamivir in care facilities) and confirm another neuraminidase inhibitor or baloxavir has not been prescribed.  **Chronic kidney disease:** the SPC dose recommendations for renal impairment are based on creatinine clearance (CrCl) which is no longer routinely reported by laboratories; the estimated Glomerular Filtration Rate (eGFR) is usually reported. There may therefore be a misalignment between the laboratory result reported for renal impairment and the result required to ensure the correct dosage and frequency. Where not reported, do not delay treatment but substitute the CrCl value with the eGFR result in the [dosage table](#table) and supply a dose according to eGFR. Some individuals may receive a larger oseltamivir dose as a result, but this is unlikely to be harmful as clinical experience reveals a wide margin of safety. Further information on CKD classifications can be found at [Chronic kidney disease: assessment and management NICE guidance (NG203)](https://www.nice.org.uk/guidance/ng203/chapter/Recommendations#classification-of-ckd-in-adults).  **Breastfeeding**: the UK Drugs in Lactation Advisory Service (UK DILAS) from the Specialist Pharmacy Service advises oseltamivir is acceptable for use in those who are breastfeeding, and the benefits of breastfeeding are considered to outweigh any, albeit unidentified, risks. Use of oseltamivir is not a reason to discontinue or put limitations on breastfeeding. Limited data suggest that oseltamivir passes into breastmilk in negligible amounts but as a precaution, infants should be monitored for vomiting, diarrhoea, irritability and changes in sleep. If an infant is unwell, premature, or the mother is taking multiple medicines, then an individual risk assessment will need to be made (see [SPS: Using oseltamivir and zanamivir during breastfeeding)](https://www.sps.nhs.uk/articles/using-oseltamivir-and-zanamivir-during-breastfeeding/#:~:text=%E2%80%A2Using%20oseltamivir%20and%20zanamivir%20during%20breastfeeding)  **Pregnancy:** Although safety data are limited, oseltamivir can be used during pregnancy when the potential benefit outweighs the risk. |

1. **Description of treatment**

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| **Name, strength and formulation of drug** | Oseltamivir 30mg, 45mg and 75mg capsules |
| **Legal category** | POM - Prescription only medicine |
| **Black triangle▼** | No |
| **Off-label use** | Yes   * when used outside the periods when national surveillance indicates that influenza virus is circulating generally in the community - see footnote below**[[9]](#footnote-10)** * when supplied after 48 hours of the onset of symptoms * in renal impairment with CrCl ≤10mL/min, the SPC states ‘not recommended’. [The UKHSA guidance on use of antiviral agents for the treatment and prophylaxis of seasonal influenza](https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment_data/file/1037465/ukhsa-guidance-antivirals-influenza-11v4.pdf) gives the dose in the [table overleaf](#table) * the duration of treatment given for individuals who are severely immunocompromised with renal impairment is outside the SPC but is recommended in [The UKHSA guidance on use of antiviral agents for the treatment and prophylaxis of seasonal influenza](https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment_data/file/1037465/ukhsa-guidance-antivirals-influenza-11v4.pdf)   Where a product is recommended off-label, consider, as part of the consent process, informing the individual/carer the product is being offered in accordance with national guidance but that this is outside the product licence. |
| **Route / method of administration** | Oral  The capsules should be taken in the morning and evening with food to reduce the chance of nausea or vomiting.  The capsules should be swallowed whole with water. For individuals with swallowing difficulties, the capsules can be opened and the contents mixed with a small amount of sweetened food, such as syrup, dessert toppings or sugared water, just before administration (see [Patient Information Leaflet](https://www.medicines.org.uk/emc/)). |
| **Dose and frequency of administration**  Continued overleaf  **Dose and frequency of administration** (continued) | See [table overleaf](#table)  Initiate treatment as soon as possible within the first two days (48 hours) of onset of symptoms, unless otherwise advised by the local HPT.  The doses given in the table below are for individuals with stable CKD. If there is a history of renal impairment supply as per the latest documented creatinine clearance (CrCl) results.  Estimated glomerular filtration rate (eGFR) may be more readily available. If eGFR is the only value available, do not delay treatment and supply a dose according to eGFR, substituting eGFR for the CrCl figure in the table below. Some individuals may receive a larger dose of oseltamivir as a result, but this is unlikely to be harmful as clinical experience reveals a wide margin of safety.  If the individual is definitely known to have chronic renal impairment and CrCl or eGFR results are not available, consider if they are suitable for treatment with zanamivir (see PGD for treatment with zanamivir in care homes) or refer to a medical practitioner or a non-medical prescriber. If a decision to supply oseltamivir is made, a prescription or a Patient Specific Direction (PSD) or a prescription will be required.  No dose adjustment is needed in obese individuals.  For severely immunocompromised individuals**10**, supply a course for **10** days rather than 5 days, except those with CrCl ≤10mL/min or those on peritoneal dialysis, who are excluded from this PGD (see dosage table below).   |  |  |  |  | | --- | --- | --- | --- | | **Characteristics of individual** | **Daily dose** | **Duration for normal immune function** | **Duration for severely immunocompromised individuals[[10]](#footnote-11)** | | Normal renal function;  weight 40kg+ | One 75mg capsule twice daily\* | 5 days | 10 days | | Normal renal function;  weight >23kg to 40kg | Two 30mg capsules twice a day | | CrCl >30 to 60 mL/min | One 30mg capsule twice a day | | CrCl >10 to 30mL/min | One 30mg capsule once a day | | CrCl ≤10mL/min | One 30mg capsule | One dose only | Do not supply under this PGD; refer to a medical practitioner or a non-medical prescriber; | | Haemodialysis | Refer to a medical practitioner or a non-medical prescriber; do not supply under this PGD | | | | Peritoneal dialysis | One 30mg capsule | One dose only | Do not supply under this PGD; refer to a medical practitioner or a non-medical prescriber; |   \*In the event the 75mg capsules are not available due to supply issues, the dose can be made up of the 30mg and 45mg presentation. The individual should be counselled on using the two strengths to make up the required dose. |
| **Duration of treatment** | See dosage schedule above |
| **Quantity to be supplied** | **Not severely immunocompromised**   * no known chronic renal impairment and weight above 40kg: 10 x 75mg capsules (*if the 75mg capsules are not available due to supply disruptions, give 10 x30mg and 10x45mg capsules)* * no known chronic renal impairment and weighing >23kg to 40kg: 20 x 30mg capsules * CrCl >30 to 60 mL/min: 10 x 30 mg capsules * CrCl >10 to 30mL/min: 5 x 30mg capsules * CrCl ≤10mL/min: 1 x 30mg capsule * peritoneal dialysis: 1 x 30mg capsule   **Severely immunocompromised individuals**   * no known renal impairment 20 x 75mg capsules (*if the 75mg capsules are not available due to supply disruptions, give 20 x30mg and 20x45mg capsules)* * no known chronic renal impairment and weighing >23kg to 40kg: 40 x 30mg capsules * CrCl >30 to 60 mL/min: 20 x 30 mg capsules * CrCl >10 to 30mL/min: 10 x 30mg capsules * CrCl ≤10mL/min, on haemodialysis or peritoneal dialysis: excluded from the PGD   When supplying under PGD, this should be from the manufacturer’s original pack or over-labelled pre-packs so the individual’s name, the date and additional instructions can be written on the label at the time of supply. As split packs cannot be supplied, an over-supply might be required. Individuals must be advised to take any remaining capsules to a community pharmacy for destruction. |
| **Storage** | Medicines must be stored securely according to national guidelines and in accordance with the product’s SPC. Do not store above 25oC |
| **Disposal** | Any unused stock should be disposed of in accordance with local arrangements.  Individuals receiving an over-supply should be advised to return any remaining capsules to a community pharmacy for destruction. |
| **Drug interactions** | Clinically important drug interactions involving competition for renal tubular secretion are unlikely, due to the known safety margin for most of these products, the elimination characteristics of the active metabolite (glomerular filtration and anionic tubular secretion) and the excretion capacity of these pathways. However, care should be taken when prescribing oseltamivir in subjects when taking co-excreted agents with a narrow therapeutic margin (e.g. chlorpropamide, methotrexate, phenylbutazone).  The [Green Book](https://www.gov.uk/government/publications/influenza-the-green-book-chapter-19) states administration of influenza antiviral agents within two weeks of administration of a live attenuated influenza nasal spray vaccine, as used in the school-age vaccination programme, may adversely affect the effectiveness of the vaccine. Adult influenza vaccinations are inactivated and are not affected by antiviral administration. A detailed list of drug interactions is available in the [SPC](https://www.medicines.org.uk/emc/) |
| **Identification and management of adverse reactions** | Frequently reported adverse reactions include nausea and headache.  These reactions may only occur on a single occasion, on either the first or second treatment day, and resolve spontaneously within one to two days. However, if symptoms persist, individuals should consult a healthcare professional.  Individuals should be advised not to discontinue treatment without consulting a doctor or pharmacist.  Other commonly reported adverse reactions include bronchitis, dizziness (including vertigo), fatigue, vomiting, insomnia, herpes simplex, nasopharyngitis, upper respiratory tract infections, sinusitis, cough, sore throat, pyrexia, rhinorrhoea, pain including limb pain, abdominal pain and dyspepsia.  A detailed list of adverse reactions is available in the [SPC](https://www.medicines.org.uk/emc/) |
| **Reporting procedure of adverse reactions** | Document any reported adverse reaction to the product in the individual’s medical records.  Alert an appropriate clinician in the event of a serious adverse reaction.  Report any suspected severe adverse reactions to the Medicines and Healthcare products Regulatory Agency (MHRA) using the [Yellow Card](http://yellowcard.mhra.gov.uk) reporting scheme or search for MHRA Yellow Card in the Google Play or Apple App Store. |
| **Written information to be given** | Supply the marketing authorisation holder's patient information leaflet (PIL)**[[11]](#footnote-12)**. Where applicable, inform the individual or their carer that the PIL with large print, Braille or audio CD may be ordered from the manufacturer (see [electronic medicines compendium](https://www.medicines.org.uk/emc)) |
| **Advice /follow up** | Inform the individual or their carer:   * to read the PIL before taking the medication * taking the medication with food can reduce nausea or vomiting * if they have difficulty swallowing, the capsules can be opened and taken with a small amount of sweetened food (see PIL) * of any possible side effects and their management * to seek advice if common side effects do not spontaneously resolve 48 hours after presentation * to seek medical advice in the event of a severe adverse reaction, if breathing difficulties develop or if health declines quickly * to complete the course * if an over-supply has been, individuals must be advised to take any remaining capsules to a community pharmacy for destruction   Promote bed rest, fluids and symptomatic remedies such as analgesics  Advise to isolate or stay away from work to prevent transmission |
| **Special considerations** | Use of oseltamivir is not a substitute for influenza vaccination. The protection against influenza lasts only as long as oseltamivir is taken. |
| **Records** | Record:   * whether valid informed consent was given or a decision to supply was made in the individual’s best interests in accordance with the [Mental Capacity Act 2005](https://www.legislation.gov.uk/ukpga/2005/9/contents) * name of the individual, address, date of birth and their GP * name of the member of staff who supplied the product * name and brand of product * date of supply * dose, form and route of administration of product * quantity supplied * batch number and expiry date * advice given; including advice given if the individual is excluded or declines treatment * details of any adverse drug reactions and actions taken * the medicine was supplied via PGD * if an over-supply is required and advice to return the remaining product to a community pharmacy for destruction has been given   All records should be signed and dated (or password-controlled e-records), contemporaneous, clear and legible.  A record of all individuals receiving treatment under this PGD should also be kept for audit purposes in accordance with local policy  Inform the individual’s GP that oseltamivir has been supplied under this PGD. |

1. **Key references**

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| **Key references** | * [Summary of Product Characteristics](https://www.medicines.org.uk/emc/search?q=oseltamivir) accessed 12 June 2025 * [British National Formulary](https://bnf.nice.org.uk/drug/oseltamivir.html) accessed 10 June 2025 * [NICE guidelines on the use of amantadine, oseltamivir and zanamivir for the treatment of influenza](https://www.nice.org.uk/guidance/TA168) TA168 reviewed November 2014 * [UKHSA Guidance on Influenza-like illness (ILI): managing outbreaks in care homes](https://www.gov.uk/government/publications/acute-respiratory-disease-managing-outbreaks-in-care-homes) updated July 2024  * [Specialist Pharmacy Service: Using oseltamivir and zanamivir during breastfeeding](https://www.sps.nhs.uk/articles/using-oseltamivir-and-zanamivir-during-breastfeeding/) updated 12 October 2023 * [UKHSA guidance on use of antiviral agents for the treatment and prophylaxis of seasonal influenza](https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment_data/file/1037465/ukhsa-guidance-antivirals-influenza-11v4.pdf), version 11, updated November 2021 * [Green Book Chapter 19 Influenza](https://www.gov.uk/government/publications/influenza-the-green-book-chapter-19) updated 28 May 2025  * [Chronic kidney disease: assessment and management NICE guidance NG203)](https://www.nice.org.uk/guidance/ng203) updated 24 November 2021 * [NICE Medicines Practice Guideline 2 (MPG2): Patient Group Directions](https://www.nice.org.uk/guidance/mpg2) updated 27 March 2017 * [NICE MPG2 Patient group directions: competency framework for health professionals using patient group directions](https://www.nice.org.uk/guidance/mpg2/resources) updated 27 March 2017 * [Health Technical Memorandum 07-01: Safe and Sustainable Management of Healthcare Waste](https://www.england.nhs.uk/publication/management-and-disposal-of-healthcare-waste-htm-07-01/) NHS England updated 7 March 2023 |

1. **Practitioner authorisation sheet**

**Oseltamivir Treatment PGD v06.00 Valid from: 08/08/2025 Expiry: 07/08/2028**

**Before signing this PGD, check that the document has had the necessary authorisations in section two. Without these, this PGD is not lawfully valid.**

**Practitioner**

By signing this PGD you are indicating that you agree to its contents and that you will work within it.

PGDs do not remove inherent professional obligations or accountability.

It is the responsibility of each professional to practise only within the bounds of their own competence and professional code of conduct.

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| I confirm that I have read and understood the content of this PGD and that I am willing and competent to work to it within my professional code of conduct. | | | |
| Name | Designation | Signature | Date |
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**Authorising manager**

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| I confirm that the practitioners named above have declared themselves suitably trained and competent to work under this PGD. I give authorisation on behalf of **insert name of organisation** for the above-named health care professionals who have signed the PGD to work under it. | | | |
| Name | Designation | Signature | Date |
|  |  |  |  |

**Note to authorising manager**

Score through unused rows in the list of practitioners to prevent practitioner additions post managerial authorisation.

This authorisation sheet should be retained to serve as a record of those practitioners authorised to work under this PGD

1. This includes any relevant amendments to legislation [↑](#footnote-ref-2)
2. The UKHSA uses information from a range of clinical, virological and epidemiological influenza surveillance schemes to identify periods when there is a substantial likelihood that people presenting with an influenza-like illness are infected with influenza virus [↑](#footnote-ref-3)
3. For definition, please see the annual flu letter for the coming/current season, which is also linked in the Green Book chapter 19 [↑](#footnote-ref-4)
4. [UKHSA ILI](https://www.gov.uk/government/publications/acute-respiratory-disease-managing-outbreaks-in-care-homes/management-of-acute-respiratory-infection-outbreaks-in-care-homes-guidance) case definition is temperature of ≥37.8°C and acute onset of one or more of: cough (with or without sputum), sore throat, coryza (nasal discharge or congestion), shortness of breath, hoarseness, sneezing, wheezing or alternatively an acute deterioration in physical or mental ability without other known cause. Note: >40% of older persons with influenza will not develop a fever of this magnitude. [↑](#footnote-ref-5)
5. [Chronic kidney disease: assessment and management NICE Guidance (NG203)](https://www.nice.org.uk/guidance/ng203/chapter/Recommendations" \l "classification-of-ckd-in-adults) [↑](#footnote-ref-6)
6. The practitioner making the supply under this PGD remains professionally accountable and clinically responsible for ensuring a supply is appropriate for an individual as assessed under this PGD. Where the HPT advise a course of treatment can be considered, they are not directing the supply must be made – this is a clinical decision which rests with the practitioner working under this PGD [↑](#footnote-ref-7)
7. Exclusion under this PGD does not necessarily mean the medication is contraindicated, but it would be outside the remit of the PGD and another form of authorisation will be required [↑](#footnote-ref-8)
8. For definition of immunocompromised see [Green Book Chapter 19](https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment_data/file/931139/Green_book_chapter_19_influenza_V7_OCT_2020.pdf) [↑](#footnote-ref-9)
9. The product licence covers treatment of influenza *when influenza virus is circulating in the community.* However [NICE guidelines](https://www.nice.org.uk/Guidance/ta158) recommend oseltamivir can be used during localised outbreaks of ILI *outside the periods when national surveillance indicates that influenza virus is circulating generally in the community,* in ‘at-risk’ people living in long-term residential or nursing homes (care homes). [↑](#footnote-ref-10)
10. For definition of severely immunocompromised refer to [UKHSA guidance on use of antiviral agents for the treatment and prophylaxis of seasonal influenza](https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment_data/file/1058443/ukhsa-guidance-antivirals-influenza-11v4.pdf) [↑](#footnote-ref-11)
11. Pre-packs will contain a copy of the PIL [↑](#footnote-ref-12)