

Who can administer

POTASSIUM phosphate

- Infusions prepared at ward level using ampoules
 - May be administered by registered competent doctor or nurse/midwife, PROVIDED the guidelines below (in Methods of Administration) have been adhered to

Important information

- There are two separate monographs for IV phosphate- sodium phosphate or potassium phosphate- please ensure you are using the correct monograph
- Caution with rate of administration (due to potassium content)
- Suggest: **Senior doctor review** before administration of intravenous phosphate, as it's use **can be dangerous**
 - Caution: the response to any given dose cannot be predicted, and IV use can cause hypocalcaemia (tetany), calcium-phosphate precipitation in the kidneys, and fatal arrhythmias ^(ref 1)
- Patients with HYPOcalcaemia should have their calcium corrected before replacing phosphate (ref 5)
- Patients with severe HYPERcalcaemia who require phosphate replacement: seek specialist advice ^(ref 4)
- **Renal impairment**:Â Requires dose adjustment- see below
- Give in a dedicated line as it may precipitate with other drugsÂ

Available preparations

| Phosphate salt | Volume | Phosphate content per vial/ampoule/bag | Sodium content per vial/ampoule/bag | Potassium content per vial/ampoule/bag |
|--|--------|--|---|--|
| Potassium phosphate ampouleÂ (Braun) | 20ml | 12mmol | nil | 20mmol |
| Phosphate polyfusor pre-mixed bag - very severe hypophosphataemia. Supplied only on request. | 500ml | 50mmol | 81mmol | 9.5mmol |

Reconstitution

Already in solution

Ampoules should be diluted further prior to administration

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Addition of potassium phosphate concentrate to infusion bags

- Preparation must be done jointly by a doctor and a nurse in the clinic room.
- Both the Controlled Drug register, and the Additive label **must be signed** Â by the SAME doctor and

nurse

- UNUSED ampoules must immediately be returned to the CD press and signed back into the CD register by the SAME doctor and nurse
- Clearly over-label the infusion bag to reflect the TOTAL amount of mmol of potassium phosphate
- After adding potassium phosphate concentrate to an infusion bag, squeeze and invert bag a MINIMUMÂ of ten times to avoid inadvertent administration of a toxic bolusÂ

Infusion fluids

Sodium chloride 0.9% (preferred)Â

Glucose 5% may also be used if clinically appropriate

Methods of intravenous administration

Intermittent intravenous infusion (using an electronically controlled infusion device)

• Administer as per guidelines below

Dose in adults

| Table 1: Guidance on rout | e given below but clin | ical judgement is always | s required ^(ref 1) |
|---------------------------|------------------------|--------------------------|-------------------------------|
|---------------------------|------------------------|--------------------------|-------------------------------|

| Route of administration | Phosphate level |
|-----------------------------|--|
| Oral/enteral replacement | PREFERRED if >0.32mmol/L and asymptomaticÂ or if level >0.48mmol/LÂ and symptomaticÂ |
| Intravenous route preferred | <0.32mmol/L or <0.48mmol and symptomatic or if unable to tolerate oral supplementation |

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Table 2: POTASSIUM PHOSPHATEÂ via peripheral line

Consider ONLY IF co-existing hypokalaemiaÂ

• Preferable to treat hypophosphataemia and hypokalaemia separately using two individual infusion bags - rather than using Potasssium phosphate vial at all. Â This allows for the greatest amount of flexibility in the doses of both electrolytes

• It is difficult to provide concrete guidelines for the treatment of severe hypophosphataemia as regimens vary greatly across hospitals in the UK and Ireland - **we have tried to provide guidelines below but clinical judgment is always required**Â

• Use caution when interpreting phosphate levels. Changes in phosphate levels may be transient - treating **underlying causes** may be sufficient to correct level. **Review medications** which may contribute e.g. sevelamar, antacids, diuretics ^(ref 5)

• The response to any given dose cannot be predicted, and IV use can cause hypocalcaemia (tetany), calcium-phosphate precipitation in the kidneys, and fatal arrhythmias ^(ref 1)

Prescribe dose in terms of phosphate dose required and then the phosphate salt required
 eg Â '12mmol phosphate as potassium phosphate'

| Gentle replacement | Dose: 9mmol phosphate over 12 hours, and repeat as necessary $\hat{A}^{(ref2,3)}$ | | | | |
|--|---|----------------|----------------------|--|--|
| More individualised dosing ^(ref 1) | Level (mmol/L) | Phosphate dose | Maximum initial dose | | |
| | less than 0.32 | 0.4mmol/kg | 48mmolÂ phosphate | | |
| | 0.33 to 0.44 | 0.3mmol/kg | 30mmolÂ phosphate | | |
| | >0.45 | 0.2mmol/kg | 20mmolÂ phosphate | | |
| Preparation | Doses up to 24mmol phosphate (40mmol potassium)Â Â Add to 500mL infusion fluid Doses 25 to 48mmol phosphate (40 (approx) to 80mmol potassium)Â Â Add to 1000mL infusion fluid For fluid restricted patients May be added to less volume provided the final concentration does not exceed 40mmol POTASSIUM per 500mL | | | | |
| Administration | Administer the required dose over 12 hours May administer more quickly - however cannot exceed a rate of administration of the POTASSIUM element of 10mmol/hour Doses of 36mmol phosphate or less may be administered over minimum 6 hours if clinically appropriate Doses greater than 36mmol phosphate MUST be administered over | | | | |
| Renal impairment | Use with great caution, consider specialist advice Generally avoid in severe renal impairment ^(ref 6) | | | | |
| Critical care/Fluid restriction | Higher doses and rates may apply in the Critical Care setting | | | | |
| Polyfusor | Available in Critical care areas- note however- only contains 9.5mmol potassium per polyfusorÂ | | | | |
| Repeated doses | May require repeat infusions over subsequent days. Usual maximum is 50mmol phosphate per 24 hours | | | | |
| Switch to oral route | Consider switch to oral route once level >0.48mmol/L | | | | |

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Renal impairment

• Use reduced doses with caution- see tables above

Monitoring

- Monitor the following electrolytes every 6 to 12 hours: phosphate, calcium, potassium, sodium, magnesium
- Monitor fluid balance and blood pressure
- Monitor ECG

Storage

• Potassium phosphate ampoules are treated as a **controlled drug in GUHs** (as it is a potasisum concentrate as well as containing phosphate). The routine supply of potassium phosphate is restricted

to designated wards which are likely to be caring for critically ill patients

- Phosphate Polyfusor is NOT treated as a controlled drug
- Store below 25°C

References

- 1. Uptodate. Hypophosphataemia: Evaluation and Treatment March 2024. Accessed online 23/01/2025
- 2. Martindale- accessed online 23/01/2025
- 3. BNF- accessed online 23/01/2025
- 4. UpToDate Potassium Phosphate monograph accessed March 2025
- 5. Maidstone and Tunbridge Wells NHS Trust 'Treatment of acute hypophosphataemia in adults. Review date August 2027
- 6. Local specialist opinion email on file25/06/2025

These local guidelines were also consulted in the preparation of guide (to try and create a consensus from different sources)

- Grampian staff guideline for the management of hypophosphataemia in adults July 2024
- Worcestershire acute hospitals NHS Trust 'guideline for the treatment of hypophosphataemia in adults, March 2023
- Liverpool University Hospitals NHS TrustÂ
- UKMI Leeds hospital 'How is acute hypophosphataemia treated in adults
- Adults Therapeutic Handbook (NHS Greater Glasgow and Clyde), May 2023 Management of hypophosphataemia