Newborn use only

Alert	Most often given in conjunction with calcium for the prevention and treatment of metabolic bone		
Aleit	disease in preterm infants.		
	1 mmol phosphorus/phosphate (P) = 31 mg elemental phosphorus.		
	1 mmol elemental calcium (Ca) = 40 mg elemental calcium.		
	Separate oral doses from calcium supplements by at least 2 hours.		
	When using IV preparation, always check plasma sodium and potassium concentrations to assist in		
Indication	choosing the right phosphate preparation (e.g. sodium or potassium phosphate preparation). Treatment of metabolic bone disease.		
indication			
	Treatment of hypophosphataemia.		
Action	Supplementation to meet the recommended daily intakes.		
Action	Phosphorus is a major intracellular mineral and is important in bone mineralisation and energy		
	production.		
Drug Type	Mineral		
Trade Name	IV		
	Sodium dihydrogen phosphate Phebra IV (Preferred inorganic preparation)		
	Each 1 mL vial corresponds to 1 mmol phosphate, 1 mmol sodium and 2 mmol hydrogen.		
	Potassium dihydrogen phosphate concentrated injection DBL IV		
	Potassium dihydrogen phosphate concentrated injection Phebra IV		
	Each 1 mL ampoule corresponds to 1 mmol phosphate, 1 mmol potassium and 2 mmol hydrogen.		
	ORAL		
	Phosphate-Phebra® oral effervescent tablets		
	Each tablet contains: 16.1 mmol phosphate (equivalent to 500 mg elemental phosphorus); 20.4 mmol		
	sodium; 3.1 mmol potassium		
	Sodium dihydrogen phosphate Phebra IV (preferred IV preparation)		
	Each 10 mL vial (sodium dihydrogen phosphate 1.56 g) contains: 10 mmol phosphate; 10 mmol sodium;		
	20 mmol hydrogen		
	Potassium dihydrogen phosphate concentrated injection DBL IV		
	Potassium dihydrogen phosphate concentrated injection Phebra IV		
	Each 10 mL ampoule (potassium dihydrogen phosphate 1.361 g) contains: 10 mmol phosphate; 10 mmol		
	potassium; 20 mmol hydrogen		
Presentation	IV: Sodium dihydrogen phosphate 10 mL vial; Potassium dihydrogen phosphate concentrated injection		
	10 mL ampoule.		
	Oral: 500 mg effervescent tablets; IV preparation (e.g. sodium or potassium dihydrogen phosphate) can		
	be given orally.		
Dose	Treatment of metabolic bone disease (MBD)		
	ORAL: 1 to 3 mmol/kg/day in 2-4 divided doses as an addition to intake from milk and other		
	sources to a maximum intake of 4.5 mmol/kg/day.		
	Use either Sodium dihydrogen phosphate Phebra IV preparation or Phosphate-effervescent		
	tablets.		
	General principles of treatment of MBD:		
7	A. Commence at low dose (e.g. 1 mmol/kg/day) and titrate the dose up as tolerated.		
	B. Given in conjunction with calcium supplementation (but not together - example: Calcium 8		
	AM, 2 PM, 8 PM and Phosphorus 6 AM, 12 MD, 6 PM)		
	C. Aim to reach the upper end of the recommended intake: calcium 5 mmol/kg/day and		
	phosphorus 4.5 mmol/kg/day. ⁸		
	D. Dose can be adjusted with a goal of slight excess supply aiming for urinary calcium		
	≥1.2mmol/L and phosphate ≥0.4 mmol/L.		

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	Treatment of acute hypophosphataemia
	IV infusion: 0.2 mmol/kg/dose [range 0.15–0.33 mmol/kg/dose]. Repeat as necessary. Aim to
	maintain normophosphataemia of 1.8–2.6 mmol/L (5.6–8.1 mg/dl).
	Daily enteral supplementation to meet the recommended daily intakes (RDI)
	2–4.5 mmol/kg/day (62–140 mg/kg/day of phosphorus) ^{7,8}
	Calculate intake from parenteral and enteral sources
	2. Supplement the difference via IV or oral route.
Dose adjustment	
Maximum dose	4.5 mmol/kg/day
Total cumulative dose	
Route	ORAL IV
Preparation	IV infusion for treatment of acute hypophosphataemia:
Preparation	IV infusion (sodium dihydrogen phosphate): Draw up 1 mL (1 mmol phosphate) and add 19 mL sodium chloride 0.9% or glucose 5% to make a final volume of 20 mL with a concentration of 0.05 mmol/mL. Draw up 4 mL/kg (0.2 mmol/kg).
	IV infusion (potassium dihydrogen phosphate): Draw up 1 mL (1 mmol phosphate) and add 24 mL sodium chloride 0.9% or glucose 5% to make a final volume of 25 mL with a concentration of 0.04 mmol/mL. Draw up 5 mL/kg (0.2 mmol/kg).
	Oral
	Option 1 (preferred option for infants going home or when a long storage time is required in the NICU): Disperse 500 mg (16.1 mmol) phosphate effervescent tablet in water for injection, make up to total volume of 16 mL with an approximate concentration of 1 mmol/mL. Prepare fresh each dose and discard any remaining liquid.
	Option 2 (can be used where preparation with low osmolality is preferred e.g. infants with history of feed intolerance): IV sodium dihydrogen phosphate decanted into a bottle and given orally undiluted (expiry time: 7 days).
Administration	Oral
	Can be administered with feeds (refer to evidence summary section).
	Separate calcium supplements by at least 2 hours.
	IV .
	As part of parenteral nutrition fluid – refer to individual parenteral nutrition formulations.
	IV infusion for treatment of acute hypophosphataemia:
	IV sodium dihydrogen phosphate or IV potassium dihydrogen phosphate: Infuse over at least 6 hours.
	For severe hypophosphataemia infuse over 8–12 hours. Maximum infusion rate of 0.2 mmol/kg/h.
Monitoring	Phosphate, calcium, magnesium, and alkaline phosphatase concentrations are required at least fortnightly or more often if required. Once these concentrations normalise, serum analysis may be
	performed once monthly for 6 months or at the discretion of the clinician. 10
	Urinary calcium and phosphate, and Tubular Reabsorption Phosphate (TRP)%, parathormone, and vitamin D concentrations may be useful under certain circumstances.
Contraindications	Hyperphosphataemia, dehydration, severe renal insufficiency, and shock.
Precautions	Hypernatraemia (avoid sodium dihydrogen phosphate)
	Hyperkalaemia (avoid potassium dihydrogen phosphate)
Drug Interactions	Calcium and magnesium antacids (e.g. acetate, carbonate, citrate, hydroxide etc.) reduce phosphate
-	absorption — separate doses by at least 2 hours.
	Additive effects with other drugs that may prolong QT interval.
	Potassium dihydrogen phosphate preparation may increase the risk of hyperkalaemia when used in
	conjunction with potassium sparing diuretics (e.g. spironolactone).
Adverse Reactions	Diarrhoea (oral use only), hypocalcaemia, nephrotoxicity, prolonged QT interval, hypotension, and
	hypomagnesaemia.
	Hyperphosphataemia – carpopedal spasm, seizures. ²

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Compatibility	Potassium dihydrogen phosphate Fluids: Glucose 5%, glucose 10%, glucose in sodium chloride solutions, sodium chloride 0.45%, sodium chloride 0.9%, sodium chloride 3%. Y-site: No information.		, sodium chloride 0.45%, sodium	
	Y-SITE: NO INFORMATION.			
	Sodium dihydrogen phosphate Fluids: Glucose 5%, sodium chloride 0.9	9%.		
	Y-site: No information			
Incompatibility	Potassium dihydrogen phosphate	ntonal nutrition. Hartmann and	d Diagram Luta (dial) of	
	Fluids: Fluids containing calcium - parenteral nutrition, Hartmanns, and Plasma-Lyte (risk of			
	precipitation). Note: PN solutions contain phosphate; running extra phosphate with PN may cause precipitation.			
	Y-site: Aciclovir, amiodarone, calcium salts, ketamine, lorazepam, magnesium salts, rocuronium Solutions that contain other cations such as calcium, magnesium, iron, and aluminium may also precipitate.			
	Sodium dihydrogen phosphate	ntoral nutrition Hartmanns on	of Diagna Luta (rick of	
	<u>Fluids:</u> Fluids containing calcium - pare precipitation). Note: PN solutions containing calcium - pare			
	precipitation.	tain phosphate, running extra p	mosphate with the may cause	
	Y-site: Aciclovir, amiodarone, calcium s	alts, calcium, aluminium or ma	gnesium, iron, and magnesium	
	containing solutions.		green, weny and magnetical	
Stability	Preparation from oral effervescent tab	lets: It is to be used immediate	ly after preparation and discard	
	unused portion.			
	Oral preparation from IV sodium dihyd	rogen phosphate: 7 days		
Storage	Store below 25°C.			
Excipients	Phosphate-Phebra® oral effervescent t		_	
	4000, citric acid, sucrose, orange 5257	O TP0551 and saccharin sodium		
Special Comments				
Evidence	Recommended daily intakes (RDI) Phosphorus absorption is typically 80% to 90% of dietary intake. ³			
	Filospilorus absorption is typically 80%	to 90% of dietary littake.		
	Parenteral intake: Previously, the reco	mmended doses of parenteral	calcium and phosphate in preterm	
	infants varied from calcium 1.3–3 mmo			
	phosphate ratio in the range of 1.3–1.7	7. ^{1,4-6} ESPGHAN 2018 updated g	uidelines on parenteral nutrition	
	recommends the following calcium and phosphate:12			
		Parenteral calcium	Parenteral phosphate	
		mmol (mg)/kg/day	mmol (mg)/kg/day	
	Preterm during the first days of life	0.8-2.0 (32-80)	1.0-2.0 (31-62)	
	Growing preterm	1.6-3.5 (100-140)	1.6-3.5 (77-108)	
	Term neonate	0.8-1.5 (30-60)	0.7-1.3 (20-40)	
	/			
	Enteral intake: ESPGHAN 2010 Guideli			
	absorbable phosphate source in a ratio with calcium (calcium: phosphate) of 1.5–2.0.7 American Academy of Pediatrics Committee on Nutrition 2013 Guidelines recommend calcium 150-200 mg/kg/c (3.8-5 mmol/kg/day) and phosphate 75-140 mg/kg/day (2.4-4.5 mmol/kg/day) and 200-400 IU/day of vitamin D for enteral nutrition in preterm neonates.8			
	Vitariii 2 for effectal flatition in prete	The flates.		
	The exact serum phosphorus concentration at which to commence supplementation of phosphate is not known and recommendations vary from 1.3 mmol/L ⁸ to 1.8 mmol/L. ⁹			
	Metabolic bone disease			
	Goal: Aim for the upper end of the recommended range to prevent fractures and clinical symptoms of			
	osteopenia: calcium and phosphate of around 4-4.5 mmol/kg/day. Adjust the mineral intake with a			
	of achieving a slight excess of urinary r		= = =	
	mmol/L. ¹⁴	<i>,</i>	· ·	
	_ ······ <i>3·y</i> = ·			

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Step 1: Calculate the mineral intake from enteral feed:

Example: 150 mL/kg/day of mature preterm EBM contains: calcium 1 mmol/kg/day and phosphate 0.6 mmol/kg/day. 150 mL/kg/day preterm EBM+24kcal HMF contains: calcium 4.5 mmol/kg/day and phosphate 2.7 mmol/kg/day.

Preterm milk	Calcium, mmol (mg)/100 mL	Phosphate, mmol (mg)/100 mL
1 st week	0.7 (26)	0.4 (11)
2 nd week	0.6 (25)	0.5 (15)
Week 3/4	0.6 (25)	0.5 (14)
Week 10/12	0.7 (29)	0.4 (12)
Term milk		
1 st week	0.7 (26)	0.4 (12)
2 nd week	0.7 (28)	0.6 (17)
Week 3/4	0.7 (27)	0.5 (16)
Week 10/12	0.7 (26)	0.5 (16)

Elemental Ca, 1 mmol = 40 mg. Elemental Phosphorus, 1 mmol = 31 mg. Adapted from Gidrewicz and Fenton BMC Pediatrics 2014, 14:216. 15

Step 2: Calculate the gap in calcium and phosphate intake/requirement: This will be the dose required.

Step 3: Prescribe 50% of the required dose of calcium and phosphate in 2-3 divided doses alternatively but not together. (example: calcium 8 AM, 2 PM, 8 PM and phosphate 6 AM, 12 MD, 6 PM).

Step 4: Once 50% dose is tolerated for 1 week, increase to 100% required dose.

ORAL preparation during NICU stay: Sodium dihydrogen phosphate Phebra IV is the preferred preparation for oral administration due to its low osmolality.

ORAL preparation at discharge or stable neonates: Phosphate effervescent tablets can be used.

American Academy of Pediatrics Committee on nutrition 2013 Guidelines on management for Enterally Fed Preterm Infants With Radiologic Evidence of Rickets: 1. Maximize nutrient intake. 2. If no further increases in these can be made, add elemental calcium and phosphorus as tolerated. Usually beginning at 20 mg/kg per day of elemental calcium and 10–20 mg/kg per day elemental phosphorus and increasing, as tolerated, usually to a maximum of 70–80 mg/kg per day of elemental calcium and 40–50 mg/kg per day elemental phosphorus. May consider targeting 25-OH-D concentration of >20 ng/mL (50 nmol/L).8 However, breast milk content of phosphorus is variable and harder to estimate the intakes accurately. A more pragmatic approach suggested by our consensus group: start with phosphate 0.5-1.0 mmol/kg/day in divided doses and increase as tolerated to a maximum of phsophate 3 mmol/kg/day.

Efficacy and safety

An ideal oral form of phosphate for use in preterm infants does not exist. Administering the intravenous preparations orally can be considered, because they are lower in osmolarity than are commercially available phosphorus-containing liquids. For example, potassium dihydrogen phosphate provides 31 mg of elemental phosphorus per millimole. A dose of 10 to 20 mg/kg per day of elemental phosphorus is reasonable and will likely resolve hypophosphataemia in most preterm infants.⁸

Oral phosphorus and feeds

It is recommended to separate oral doses from calcium and antacids containing agents such as aluminium hydroxide, calcium or magnesium salts, as these may reduce the bioavailability of phosphate. Oral phosphate preparation has high osmolality and administration with feeds may have theoretical benefit of reducing the osmolality (consensus opinion).

Practice points

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VERSION/NUMBER	DATE
Original 1.0	15/11/2016
Version 1.1	22/11/2018
Version 2.0	20/05/2021
Version 3.0	15/07/2021
Version 4.0	21/10/2021
Version 4.0(minor errata)	23/06/2023
Version 4.0(minor errata)	1/08/2024
Version 5.0	23/10/2025
REVIEW	23/10/2030

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