Vitamin A

Newborn use only

| Alert | Vitamin A is expressed as microgram retinol activity equivalents (RAE) or international units (IU) or units. |
|-------------------|---|
| | 1 microgram RAE = 1 microgram retinol = 3.3 units of retinol. (1) |
| | Pentavite Infant, a commonly used multi-vitamin supplement in Australia, contains 390 microgram RAE or 1287 units of retinol. |
| | Vitamin A for chronic lung disease is beyond the scope of this formulary. Refer to evidence summary. |
| Indication | Prevention of vitamin A deficiency |
| | Neonatal cholestasis |
| | Cystic fibrosis |
| Action | Fat soluble vitamin required for vision, growth and bone development, immune function and maintenance |
| 71011011 | of epithelial cells particularly in the retina and respiratory tract tissues. |
| Drug type | Fat soluble vitamin |
| Trade name | Bio-Logical Vitamin A oral solution |
| Presentation | Bio-Logical Vitamin A oral solution (50 mL bottle): Contains retinol palmitate 1.375 mg per 0.1 mL (750 |
| | microgram RAE or 2500 units vitamin A/0.1 mL) |
| Dose | Suggested starting dose |
| | Prophylaxis in preterm infants <1800 g birthweight |
| | Bio-Logical Vitamin A oral solution: 0.1 mL/day (2500 units/day) |
| | Range: 1320-3300 units/kg/day |
| | |
| | Neonatal cholestasis: Refer to vitamins in cholestasis formulary |
| | |
| | Supplementation for cystic fibrosis* |
| | Bio-Logical Vitamin A oral solution: 0.1 mL/day (2500 units/day) ⁽⁸⁻¹¹⁾ (ANMF consensus) |
| | *Pentavite 0.45 mL twice daily provides 2574 units/day. |
| Dose adjustment | Therapeutic hypothermia – No information. |
| | ECMO – Not applicable. |
| | Renal impairment - No information. |
| | Hepatic impairment – No information. |
| Maximum dose | |
| Total cumulative | |
| dose | |
| Route | Oral |
| Preparation | No preparation is required |
| Administration | Oral: Administer undiluted with a feed |
| Monitoring | An 'adequate' concentration of plasma vitamin A in VLBW infants is not known. Concentrations below 0.70 |
| | μmol/L have been considered deficient in premature infants and concentrations below 0.35 μmol/L indicate |
| | severe deficiency and depleted liver stores. (4) |
| Contraindications | Hypersensitivity to vitamin A or any component of the formulation, hypervitaminosis A |
| Precautions | |
| Drug interactions | May increase effects of anticoagulant and antiplatelet agents |
| Adverse | Hypervitaminosis A: Irritability, lethargy, vomiting, bulging fontanelle. |
| reactions | |
| Compatibility | Not applicable |
| Incompatibility | Not applicable |
| Stability | |
| Storage | Protect from light |
| Excipients | Sodium benzoate. Avoid exposure to sodium benzoate of >99 mg/kg/day in neonates. |
| Special | Social Senzoate. Avoid exposure to social senzoate of 233 mg/ kg/ day in neonates. |
| comments | |
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| Evidence | Efficacy |
|-----------------|---|
| | Preterm infants <1800 g |
| | Recommended enteral intake in preterm infants <1800 g birthweight (ESPGHAN 2010): 400-1000 |
| | microgram/kg/day (1320-3300 units/kg/day). (4) Recommended parenteral intake (ESPGHAN 2018): Preterm |
| | neonates – 227-455 microgram/kg/day (700-1500 units/kg/day); Term neonates – 150-300 |
| | micrograms/kg/day (495-990 units/kg/day). Alternatively 2300 units (697 micrograms)/day. (5) |
| | <u>Chronic liver disease</u> |
| | Supplementation with 5000–10,000 IU/day may be needed in children with chronic liver disease. (2) The dose |
| | in neonates is unclear. It is important to monitor levels in children receiving supplementation, as |
| | hypervitaminosis A can lead to potentially fatal hepatotoxicity. (2) |
| | Prevention of chronic lung disease and neurodevelopmental impairment |
| | Cochrane review by Darlow et al 2016 ⁽³⁾ evaluated vitamin A supplementation on the incidence of death or |
| | chronic lung disease and long-term neurodevelopmental disability in very low birth weight (VLBW) infants |
| | compared with a control (placebo or no supplementation). Eleven randomised clinical trials (RCTs) were |
| | analysed that included over 1500 VLBW infants, defined as birth weight ≤ 1500 grams or less than 32 weeks' |
| | gestation. All except one RCT in this review used intramuscular vitamin A as an intervention. Doses varied |
| | among the studies. One of the dosing regimens used in RCT by Tyson et al. was IM vitamin A 5,000 |
| | units/dose 3 times weekly initiated within the first 96 hours of life and continued for 4 weeks. (3,7) Meta- |
| | analysis (3) found that vitamin A was associated with a small benefit in reducing death or oxygen use at one |
| | month of age and a marginal reduction in oxygen use at 36 weeks' postmenstrual age. However, |
| | neurodevelopmental assessment in the largest trial showed no difference at 18 to 22 months corrected age. |
| | No adverse effects of vitamin A supplementation were reported, but it was noted that intramuscular |
| | injections of vitamin A were painful. |
| | ANMF consensus: Clinicians need to balance the clinical benefits against painful intramuscular injections and |
| | the decision may depend upon the local incidence of the outcomes. It is not a standard practice in Australia |
| | to administer intramuscular vitamin A for prevention of chronic lung disease. |
| | Supplementation in cystic fibrosis The LIC Cystic Fibrosic Foundation (CFF) recommended deith complementation of 1500 III of vitagein A 100 |
| | The US Cystic Fibrosis Foundation (CFF) recommends daily supplementation of 1500 IU of vitamin A, 400– |
| | 500 IU of vitamin D, and 40–50 IU of vitamin E for infants with CF. These dosages increase to 5000 IU of |
| | vitamin A, 800–1000 IU of vitamin D, and 80–150 IU of vitamin E for children 1–10 years of age. (8-10) A |
| | multicentre prospective longitudinal study known as FIRST determined the prevalence of suboptimal vitamins A, D, and E status in infants supplemented with CF foundation recommended vitamin dosages. The |
| | prevalence of vitamin A, D and E insufficiency were 3%, 22% and 5% on these dosages. The study found |
| | normalisation of serum retinol and α -tocopherol in almost all infants by age 3 years. (11) |
| | ANMF consensus: Bio-Logical Vitamin A oral solution at a dose of 0.1 mL (2500 units/day) is recommended |
| | for ease of administration and to prevent any potential insufficiency in accordance with the US Cystic |
| | Fibrosis Foundation's recommended dosage. |
| Practice points | Recommendations for daily supplementation of vitamin A: |
| | Recommended enteral intake in preterm infants <1800 g birthweight (ESPGHAN 2010): 400-1000 |
| | microgram/kg/day (1320-3300 units/kg/day). (4) |
| | Recommended parenteral intake (ESPGHAN 2018): Preterm neonates – 227-455 microgram/kg/day (700- |
| | 1500 units/kg/day); Term neonates – 150-300 micrograms/kg/day (495-990 units/kg/day). Alternatively |
| | 2300 units (697 micrograms)/day. (5) |
| | Preterm human milk contains 50-400 units of vitamin A/dL. Term human milk contains 60-200 units/dL. (6) |
| | Evaluate vitamin A intake from other sources prior to prescribing e.g. feeds and other concomitant |
| | medications or supplements. |
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