

METABOLIC BONE DISEASE (MBD) IN PRE-TERM INFANTS - Paediatric Full Clinical Guideline

Reference no.: NIC ME 08/ Jan 17/v003

1. Introduction

To ensure accurate diagnosis and management of metabolic bone disease in preterm babies.

2. Aim and Purpose

For medical staff to optimise the management of metabolic bone disease in preterm babies.

3. Implementing the Guideline

Background

In-utero mineral accretion is at its most rapid in the last trimester of pregnancy. Prematurity will interrupt this critical period of growth. Preterm infants are therefore at risk of inadequate bone mineralisation if dietary intake of calcium, phosphate, magnesium and Vitamin D is insufficient. In practice calcium and magnesium deficiency is rare and current vitamin supplements give adequate Vitamin D. At low mineral intakes phosphate deficiency is most likely to arise first.

Babies Who Are At Risk

- < 30 weeks of gestation
- Weight < 1000gms
- Prolonged TPN
- Breast fed babies (without any supplementation)
- Prolonged use of Medications – diuretics, steroids ,bicarbonate
- Cholestatic jaundice
- Short gut syndrome (malabsorption of Vitamin D and calcium)

Biomechanical Features

- ↑ Alkaline phosphatase (>1200 IU)
- Hypophosphatemia (<1.2mmol/L).
- Abnormal Ca: PO₄ ratio in urine. This should be < 1 after 3 weeks of age
- (Both measured in mmols/L)

Clinical Features

Metabolic bone disease presents between 6-12 weeks of age. Babies may remain asymptomatic for weeks. Symptoms include poor weight gain, respiratory difficulty

Some show evidence of severe demineralisation. Features include:

- Radiological features, e.g. fractures, rickets, osteoporosis. Peak time for these changes are 36-40 weeks after conception
- Skeletal deformity, e.g. rib cage softening, craniotables, reduced growth velocity

Investigations

- Weekly Alkaline phosphate, Calcium (serum calcium may remain normal until late) and phosphate on babies at risk
- Urine calcium and phosphate
- Consider measurement of Vit D and Parathyroid hormone (PTH)

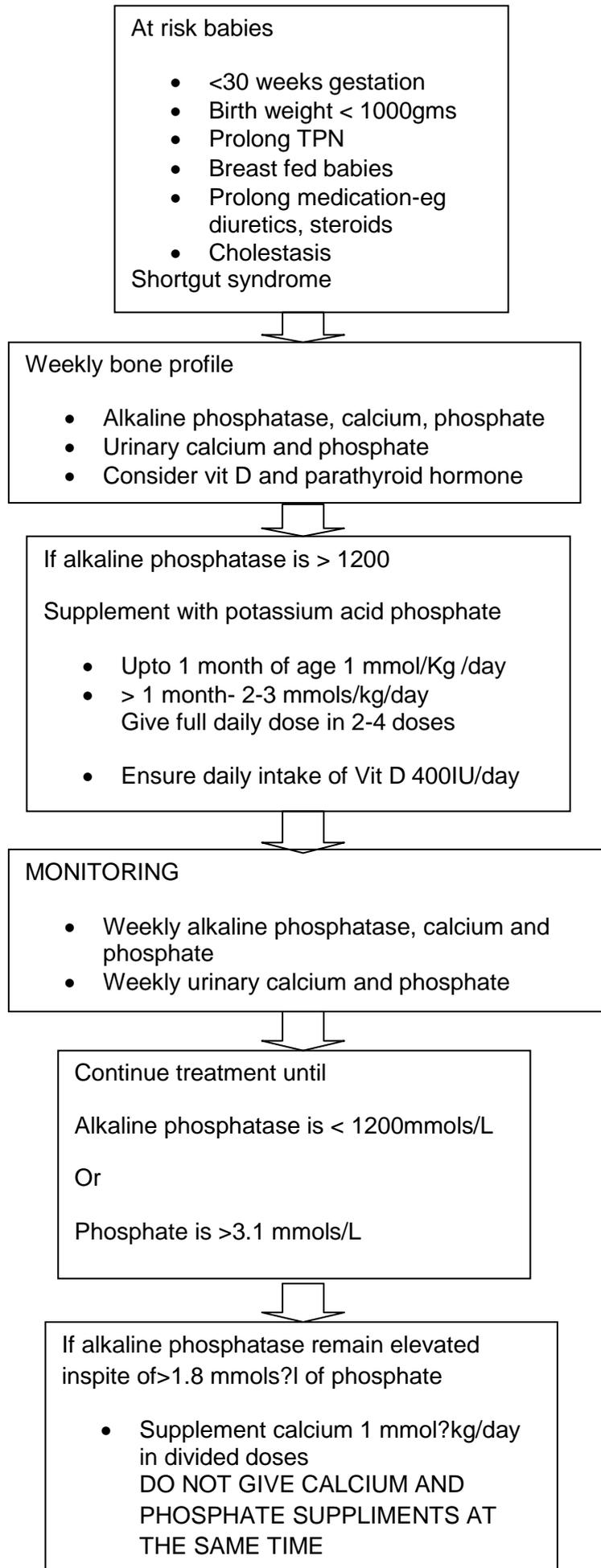
(Low urinary calcium and raised Parathyroid hormone suggest calcium deficiency)

Management

For those at risk of MBD and established on full enteral feeds:

- If alkaline phosphatase >1200 IU, with or without low phosphate, supplement the expressed breast milk (EBM) with potassium acid phosphate 1mmol/kg/day
- From 1 month of age 2-3 mmols /kg/Day in 2-4 divided doses
- Continue surveillance (weekly check of alkaline phosphatase and phosphate) and supplementation while on EBM, until alkaline phosphatase <1200 IU or the feed is changed to formula milk. Phosphate supplements should be stopped if serum phosphate increase above 3.1 mmols/L
- Once phosphate supplementation is adequate , a relative deficiency of calcium may become evident (elevated alkaline phosphatase in spite of PO₄ > 1.8 mmols/L). If there is deficiency of both calcium and phosphate and PO₄ is replaced without replacing calcium, it will provoke more PTH production and PO₄ loss in the urine. These babies need replacement of both calcium and phosphate. Calcium 1mmol/kg should be given with feeds. Calcium and phosphate should not be given at the same time, as intestinal obstruction has been associated with calcium supplementation, incremental advancement of calcium is recommended.
- Ensure Daily intake of 400IU vitamin D/day

X-rays may be indicated on clinical grounds e.g. suspected fracture. Routine X-rays to detect rickets are not necessary.



4. References (including any links to NICE Guidance etc.)

1. Nutritional Needs of the Preterm Infant: Scientific Basis and Practical Guidelines (1993); Williams & Wilkins; Eds: R. Tsang, A. Lucas, R. Uauy, S. Zlotkin.
2. Ryan, S. (1996) Nutritional aspects of metabolic bone disease in the newborn; Arch Dis Child; 74; F145-148
3. Robertson's text book of Neonatology
4. Newborn Services clinical guidelines- Auckland District Health board -

5. Documentation Controls

Development of Guideline:	Dr M. Ratnayaka
Consultation with:	Neonatal Consultants
Approved By:	Dr R. Bowker (Lead Clinician) - 20 th Jan 2017 Integrated Care Division - Feb 2017
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Key Contact:	Dr M. Ratnayaka