

CKD-Mineral & Bone Disorder (CKD-MBD)

Biochemical and bone monitoring, vascular calcification assessment, phosphate/calcium and PTH management, bone-targeted therapy, and kidney transplant bone disease.

2017 KDIGO Clinical Practice Guideline Update
Diagnosis, Evaluation, Prevention, and Treatment of CKD-MBD
Ketteler, Block, Evenepoel et al. • Kidney Int Suppl
2017;7(1):1-59

GRADE 1 • We Recommend 2 • We Suggest NG • Not Graded

1 3.1 — Biochemical Monitoring

1C

Monitor serum **calcium, phosphate, PTH, and ALP** beginning CKD G3a (children: G2).

NG

Base monitoring frequency on abnormality severity/progression — Ca/phosphate q6–12mo (G3a–b) to q1–3mo (G5D); PTH q6–12mo to q3–6mo; ALP annually (G4–G5D).

2C

25(OH)D may be measured, repeated per baseline/intervention; correct deficiency as in general population.

1C

Base therapeutic decisions on **trends** rather than a single lab value.

2D

Use individual calcium and phosphate values together rather than the **Ca × P product**.

1B

Labs should report assay method/changes and specimen source to aid interpretation.

2 3.2 — Diagnosis: Bone

2B

Consider **BMD testing** if evidence of CKD-MBD/osteoporosis risk factors AND result will impact treatment.

NG

Bone biopsy reasonable if knowledge of renal osteodystrophy type will impact treatment.

2B

Serum PTH or bone-specific ALP can evaluate bone disease — markedly high/low values predict turnover.

2C

Do **not** routinely measure bone-derived collagen turnover markers (e.g., P1NP, CTX).

1B

Measure **length** quarterly in infants (G2–G5D); assess linear growth annually in children.

3 3.3 — Diagnosis: Vascular Calcification

2C

Lateral abdominal radiograph (vascular) or echocardiogram (valvular) are reasonable alternatives to CT-based imaging.

2A

Known vascular or valvular calcification = **highest cardiovascular risk category** — use to guide CKD-MBD management.

4 4.1 — Phosphate & Calcium Treatment

NG

Base treatment on serial phosphate, calcium, and PTH assessed together; base phosphate-lowering treatment on progressive or persistent elevation.

2C

Lower elevated phosphate toward the normal range.

2C

Avoid hypercalcemia in adults; age-appropriate normal calcium in children. Dialysate calcium 1.25–1.50 mmol/L (2.5–3.0 mEq/L).

2B

Restrict **calcium-based binder** dose in adults; base pediatric choice on serum calcium.

1C

Avoid long-term aluminum-containing binders; avoid dialysate aluminum contamination.

2D

Limit dietary phosphate intake alone or combined with other treatments; consider phosphate source (animal/vegetable/additive).

2C

Increase **dialytic phosphate removal** for persistent hyperphosphatemia.

5 4.2 — PTH Management

2C

Evaluate progressively rising/persistently elevated iPTH for modifiable factors — hyperphosphatemia, hypocalcemia, high phosphate intake, vitamin D deficiency.

2C

Don't routinely use **calcitriol/vitamin D analogs** in non-dialysis G3a–G5 adults; reserve for severe/progressive HPT in G4–G5.

2C

Maintain G5D iPTH ~2–9× ULN; treat marked within-range shifts to avoid drifting outside this range.

2B

G5D PTH-lowering therapy: **calcimimetics, calcitriol, vitamin D analogs**, or a combination.

2B

Parathyroidectomy for severe hyperparathyroidism refractory to medical/pharmacologic therapy.

6 4.3 — Bone Treatment

1A

CKD G1–G2 with osteoporosis/high fracture risk: manage **as general population**.

2B

CKD G3a–G3b with normal PTH + osteoporosis/high fracture risk: treat as general population.

2D

CKD G3a–G5D with CKD-MBD abnormalities + low BMD/fragility fracture: **individualize**, consider bone biopsy.

1A

Children/adolescents, CKD G2–G5D + height deficit: **recombinant GH** after addressing malnutrition and CKD-MBD abnormalities.

7 5 — Kidney Transplant Bone Disease

1B

Measure serum calcium/phosphate **≥weekly** in the immediate post-transplant period until stable.

NG

Base post-transplant monitoring frequency on abnormality severity/CKD progression; ALP annually.

2C

BMD testing if osteoporosis risk factors AND result will alter therapy.

2D

First 12mo post-transplant, eGFR > ~30, low BMD: consider vitamin D/calcitriol/antiresorptive agents; bone biopsy before bisphosphonates.

2C

G4T–G5T with known low BMD: manage as **non-transplant CKD G4–G5**.

2C

25(OH)D may be measured, repeated per baseline values/interventions; correct deficiency as in general population.