

SPECIALTY GUIDELINE MANAGEMENT

ZOMETA (zoledronic acid) zoledronic acid

POLICY

I. INDICATIONS

The indications below including FDA-approved indications and compendial uses are considered a covered benefit provided that all the approval criteria are met and the member has no exclusions to the prescribed therapy.

A. FDA-Approved Indications^{1,2}

1. Zometa/zoledronic acid is indicated for the treatment of hypercalcemia of malignancy defined as an albumin-corrected calcium (cCa) of greater than or equal to 12mg/dL [3.0 mmol/L] using the formula: $cCa \text{ in mg/dL} = Ca \text{ in mg/dL} + 0.8 (4.0 \text{ g/dL} - \text{patient albumin [g/dL]})$.
2. Zometa/zoledronic acid is indicated for the treatment of patients with multiple myeloma and patients with documented bone metastases from solid tumors, in conjunction with standard antineoplastic therapy. Prostate cancer should have progressed after treatment with at least one hormonal therapy.

Limitation of Use: The safety and efficacy of Zometa/zoledronic acid in the treatment of hypercalcemia associated with hyperparathyroidism or with other non-tumor-related conditions have not been established.

B. Compendial Uses⁵

1. Treatment or prevention of osteoporosis during androgen-deprivation therapy (ADT) in prostate cancer patients with high fracture risk
2. Treatment in postmenopausal patients with breast cancer who are receiving adjuvant therapy to maintain or improve bone mineral density and reduce risk of fractures
3. Treatment for osteopenia or osteoporosis in patients with systemic mastocytosis

All other indications are considered experimental/investigational and not medically necessary.

II. CRITERIA FOR INITIAL APPROVAL

A. Hypercalcemia of Malignancy¹⁻⁴

Authorization of 2 months may be granted for treatment of hypercalcemia of malignancy.

B. Multiple Myeloma¹⁻⁵

Authorization of 12 months may be granted for prevention of skeletal-related events in members with multiple myeloma.

C. Bone Metastases from a Solid Tumor¹⁻⁵

Authorization of 12 months may be granted for prevention of skeletal-related events in member with bone metastases from a solid tumor.

D. Prostate Cancer⁵

Reference number(s)
2382-A

Authorization of 12 months may be granted for members with prostate cancer for treatment or prevention of osteoporosis during androgen deprivation therapy (ADT)

E. Breast Cancer⁵

Authorization of 12 months may be granted for postmenopausal (natural or induced by ovarian suppression) members receiving adjuvant therapy for treatment of breast cancer to maintain or improve bone mineral density and reduce the risk of fractures.

F. Systemic Mastocytosis⁵

Authorization of 12 months may be granted for treatment of osteopenia or osteoporosis in members with systemic mastocytosis.

III. CONTINUATION OF THERAPY

A. Hypercalcemia of malignancy

Authorization of 2 months will be granted for continued treatment in members requesting reauthorization for hypercalcemia of malignancy who are experiencing benefit from therapy as evidenced by disease stability or disease improvement.

B. All Diagnosis (excluding hypercalcemia of malignancy)

Authorization of 12 months will be granted for continued treatment in members requesting reauthorization for an indication listed in Section II (excluding hypercalcemia of malignancy) who are experiencing benefit from therapy as evidenced by disease stability or disease improvement.

IV. REFERENCES

1. Zometa [package insert]. East Hanover, NJ: Novartis Pharmaceuticals Corporation; December 2018.
2. Zoledronic acid [package insert]. Memphis, TN: Northstar Rx LLC; December 2018.
3. IBM Micromedex DRUGDEX (electronic version). IBM Watson Health, Greenwood Village, Colorado, USA. Available at: <https://www.micromedexsolutions.com/> (cited: October 5, 2020).
4. American Society of Health System Pharmacists. AHFS Drug Information (electronic version). Bethesda, MD. Available at: <http://online.lexi.com>. Accessed October 5, 2020.
5. The NCCN Drugs & Biologics Compendium 2019 National Comprehensive Cancer Network, Inc. <http://www.nccn.org>. Accessed October 5, 2020.