| Southern Sud<br>Health   | REGIONAL ADULT PARENTERAL DRUG MONOGRAP  |   |                  |  |
|--|--|---|------------------|--|
| Effective Date: Mar12-2014<br>Revised Date: Mar 13 2019<br>Revised Date:           | CLASSIFICATION<br>Bone Metabolism<br>Regulator   | OTHER NAMES<br>Aclasta, Zometa                                      | PAGE<br>1 of 2   |  |
| IV Bolus: - Not recom   IM: - Not recom   Subcut: - Not recom   RECONSTITUTION/DIL | ministered by a nurse<br>mended<br>mended<br>UTION/ADMINISTRATION:   |   |                  |  |
| IV intermittent: Dilu<br>Infu  | g – 5 mL vial OR 5 mg/100 mL<br>the dose in 100 mL normal saline<br>OR<br>use contents of ready to use IV be<br>th IV line with 10 mL normal sal | e and infuse over 15 minutes<br>ottle over 15 minutes. Use a ventee | d set.           |  |
| Breast Cancer – adjuvant o<br>Breast Cancer –metastatic                            | 4 mg every 6 months (bone metastases):   |   | √ every 12 weeks |  |
| Multiple myeloma:  | 4 mg IV every 3 to 4 weeks   |   |                  |  |
| Hypercalcemia:   | 4 mg every 7 to 28 days  |   |                  |  |
| Osteoporosis:  | 5 mg IV once yearly  |   |                  |  |
| Maximum rate:<br>Maximum concentration:<br>Maximum single dose:                    | 5 mg over 15 minutes<br>5 mg in 100 mL<br>5 mg   |   |                  |  |
| STABILITY/COMPATIB<br>Stability of Final Admixtu                                   |  | Allow solution to reach room temp                                   | perature before  |  |
|  |  |   |                  |  |

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| Southern | Santé |
| Health   | Sud   |

## **REGIONAL ADULT PARENTERAL DRUG MONOGRAPH**

## GENERIC NAME **zoledronic acid**

| Effective Date: Mar12-2014                 | CLASSIFICATION<br>Bone Metabolism | OTHER NAMES<br>Aclasta, Zometa            | PAGE<br>2 of 2     |
|--|-----------------------------------|---|--------------------|
| Revised Date: Mar 13 2019<br>Revised Date: | Regulator                         |   | 2 01 2             |
| PRECAUTIONS, POTENT                        | TIAL ADVERSE REACTIO              | NS:                                       |                    |
| • Mild fever, chills, pain a               | t infusion site                   |   |                    |
| • Hypophosphatemia, hyp                    | omagnesemia and asymptoma         | tic hypocalcemia, hypokalemia             |                    |
| • Peripheral edema, hypot                  | ension, hypertension, chest pa    | in  |                    |
| • Muscle, bone or joint pa                 | in                                |   |                    |
| • GI: nausea, vomiting, al                 | bdominal pain, diarrhea           |   |                    |
| • Neurological: Pain, head                 | lache, fatigue, dizziness, mala   | ise, insomnia                             |                    |
| • Renal: Elevated serum c                  | reatinine                         |   |                    |
| Respiratory: dyspnea, cough                |                                   |   |                    |
|  | D NURSING CONSIDERA               | TIONS:                                    |                    |
| • Patients with hypercalce                 | mia: establish adequate hydra     | tion and urinary output prior to and dur  | ing treatment in   |
| order to increase renal e                  |                                   |   |                    |
|  | ma patients who have broncho      |   |                    |
| • Acetaminophen after ad symptoms, myalgia | ministration may reduce the ir    | cidence of acute reaction (eg. arthralgia | a, fever, flu-like |
|  | ould receive zoledronic acid f    | or osteoporosis should be informed to d   | lrink two glasses  |

- Not recommended for osteoporosis in patients with renal dysfunction (CrCl less than 30 mL/minute)
- Increased risk of renal impairment with rapid infusion rates (must be given over at least 15 minutes)