THE USE OF DENOSUMAB IN GIANT CELL TUMORS IN THE SACRUM

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Abstract

Giant Cell Tumor (GCT), a benign tumor with local aggression, corresponds to 5% of primary tumors. Fifteen percent of these are located in the sacrum. En bloc resection is an effective treatment, but when it cannot be performed, Denosumab may be indicated as an alternative treatment. The objectives of this work are: to justify the indication; determine the best dose and time of use; and recognize the need for post-treatment surgery. Methods: A systematic search of clinical trials. Twenty-five articles were selected, ten of which met the inclusion criteria. The use of Denosumab is justified in advanced stages, with a dose of 120 mg administered subcutaneously, every 7 days in the first month and then maintained every 4 weeks, for 2.5 to 13 months. Adverse events are mild and can be observed in 84% of patients. With Denosumab, surgery may be less aggressive or even unnecessary. The bibliography justifies the indication of Denosumab in advanced stages of GCT, with a dose of 120 mg administered subcutaneously; there is no consensus as to the maintenance dose, which is a weekly dose in the first month and then every four weeks for 2.5 to 13 months. Complications are frequent but mild. Level of evidence III; Systematic review.

Keywords: Giant Cell Tumor of Bone; Sacral Region; Denosumab; Complications.
metastasis, and in less than 1% of cases, they can affect different locations simultaneously.\(^1\)

In some cases, subchondral fractures or frank cortical destruction and soft tissue extension can be observed. It rarely occurs in skeletally immature patients. The risk of local recurrence varies depending on the treatment modality. Ten percent of patients with recurrent GCT develop pulmonary metastasis. Malignant transformation has been reported spontaneously in the context of previous irradiation, or less frequently, after surgical resection of the tumor.

In GCTs, as in osteoporosis, multiple myeloma and metastatic carcinomas, bone homeostasis is altered.\(^1,2\) In the case of GCT, giant cell-mediated osteolysis is generated.

Denosumab, a human monoclonal antibody, within the antiresorptive group, is reserved for use in cases of advanced stage GCTs, preventing the typical osteolysis observed in this type of tumor and thus, creating the necessary surgical conditions.

A systematic review of the current bibliography on the use of Denosumab in GCTs was carried out, in order to: justify the indication of the drug; determine the best dose and treatment time to prevent complications; and recognize the need for post-treatment surgery.

**METHODS**

A search of the Pubmed and Lilacs databases was performed using the search terms: giant cell tumor, Denosumab, complications and osteonecrosis of the jaw. The search was restricted to clinical trials and articles published in English. Twenty-five articles were selected, ten of which met the inclusion criteria (justification of the indication, dose, treatment time and complications of Denosumab). (Figure 1)

**RESULTS**

In relation to the indication of antiresorptive Denosumab for giant cell tumor, Lizz van der Heijden et al.\(^2\) and Sant Chawla et al.\(^3\) justify its use in advanced tumors where en bloc resection or even amputation are the only therapeutic options. There is consensus on the dose (120 mg) and the mode of administration, which should be subcutaneous, but not on the interval and time, which vary depending on whether it is a primary or recurrent GCT. (Table 1)\(^1,4\)

Chawla et al.\(^3\) found that 84% of 282 patients included in their trial developed some type of adverse effect; 20% developed arthralgias, 18% headaches, 17% nausea, 5% hypocalcemia, 9% severe side effects that did not cause death, and 1% back pain. Raskin et al.\(^5\) mentioned skeletal pain as the most frequent adverse effect. Osteonecrosis of the jaw is a rare event in patients who use Denosumab, with an incidence of 1-1.7% of patients.\(^2,3,6,7\)

Denosumab provides an option in the treatment of advanced GCTs for which surgery is contraindicated, or that would result in high morbidity (hemipelvectomy, amputations, neurological disorders), or in cases where it is not possible to resect the entire tumor due to the increased risk of recurrence this entails. Some GCTs do not show progression with the use of Denosumab, sometimes eliminating the need for surgery or achieving an intervention with lower morbidity.\(^2,6\)

**DISCUSSION**

In the diagnosis of GCT, radiographs and tomography can be used, to identify a lytic, well-circumscribed, non-scleral, eccentric lesion.\(^1,2\) Magnetic resonance imaging (MRI) is useful for staging and predicting clinical behavior.\(^2\)

Histopathologically, the GCT contains three cell populations: multinucleated giant cells that express the receptor activator of nuclear factor kappa (RANK), monocytes and neoplastic stromal cells that express the ligand RANK (RANKL). It is believed that the pathophysiology is driven by the proliferation of mesenchymal stromal cells that exhibit an immature osteoblastic phenotype. These cells secrete high levels of chemokines that attract circulating monocytes to the site of the injury. Once there, they are initiated to form positive osteoclastic cells, resulting in giant cell-mediated osteolysis.\(^1,2,4,8,9\)

The standard treatment of GCT consists of surgical removal of

| Table 1. Use of denosumab. Dose/interval/duration. |
|-----------------------------------------------|
| Author (year) | Dose (mg) | Interval (days) | Duration (months) |
| Chawla, S (2013)\(^3\) | 120 | 7 | 2.5 – 13 |
| Ngl, VY (2014)\(^1\) | 120 | 7 (1st month) and later 30 |
| Rotman, PD (2017)\(^4\) | 120 | 7 | 3.5 – 13 |
| van del Heijden (2017)\(^2\) | 120 | 7 | 3.5 – 13 |

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Figure 1. Flowchart of search.
the tumor, or extensive curettage with or without adjuvants.1,2 Surgical treatment in the sacrum is subject to multiple barriers associated with the anatomy and the difficulty of using local coadjuvance.1 En bloc resection and radiotherapy should be reserved for patients in whom intralesional surgery and systemic therapy are impossible, as they result in an increased risk of complications such as the development of a sarcoma (3-11%) and poorer functional outcome in the case of En bloc resections.1,2 Serial embolization is used as neoadjuvant therapy to facilitate surgical curettage or resection. Patients with GCT in the sacrum exhibit a favorable initial response; 50% experience local control.1

Most recurrences occur within 2 years of treatment, rates vary from 27 to 65% for those treated with isolated curettage, 12-27% if adjuvants such as phenol, PMMC, liquid nitrogen are added to the curettage, and 0 -12% for those which are resected en bloc. Intralesional resection in the sacrum or pelvis has a recurrence rate of 50%.1 Individual parameters that strongly increase local recurrence include: soft tissue extension and spinal or sacral localization.

Denosumab is a human monoclonal antibody inhibitor of RANKL recently registered by the Food and Drug Administration of the United States (FDA) and the European Medicines Agency as a neoadjuvant pharmacological treatment for GCT.2,4,6 It is indicated in patients with advanced GCT. The objectives are to achieve local control and create a surgical situation, inhibiting the recruitment of multinucleated giant cells by the neoplastic stromal cells, thus preventing the osteolysis typically observed in the GCT. If the tumor responds to Denosumab, a partial maturation of the neoplastic stromal cells with formation of fibrous and osteoid tissue is observed.

Radiographically, a calcified border surrounding the soft tissue and/or a decrease in the size of the lesion can be seen after an average of 3-4 months of drug use, facilitating curettage with local adjuvants in a previously non-curetable GCT.1,2,4 With longer treatment, a gummy substance is likely to be observed during surgery, which alters the typical macroscopic appearance of the GCT and complicates the complete excision of the tumor by intralesional curettage.2,3 After the use of Denosumab, surgery should involve the extension of the entire original tumor in order to reduce the risk of recurrence.2,4 In an open phase II study (NCT00396279), 86% of patients had an objective response to Denosumab therapy, defined as more than 90% elimination of giant cells on histological evaluation or without radiographic progression of the lesion.1 A second, open trial, also phase II, evaluated 282 patients, restoring the high efficiency of Denosumab in GCT. Stable disease was reported in 96% of GCTs cases without indication of previous curettage after a mean follow-up of 13 months. Most patients had planned surgery with severe morbidity, and with a mean follow-up of 9.2 months, 74% did not require surgery and 16% underwent surgery resulting in low morbidity.3 The complications and side effects of Denosumab, include pain, fatigue, headache, nausea (18-25%), hypocalcemia and hypophosphatemia (3%), and osteonecrosis of the jaw (1-2%), with a malignant transformation in 1.8%.2,4,10 The effect on the fetus is unknown, therefore, the use of adequate contraceptives during treatment is recommended with Denosumab. There is no published clinical experience suggesting recurrence after 6-12 months of interruption of treatment.2,3

The cause of osteonecrosis of the jaw (ONJ) development related to Denosumab suggests that by blocking the RANK-RANKL interaction, the function and survival of monocytes and macrophages would be affected, inhibiting normal bone repair at local microdamage. Dentalalveolar trauma (dental extractions) is the most prevalent and consistent risk to develop it. However, there is limited data on the possible mechanism of ONJ related to therapy directed to the bones. More studies focused on this subject are required. It is recommended that patients who are going to receive Denosumab have a dental evaluation before starting treatment.2,7 A meta-analysis was conducted that included a total of 7 prospective clinical trials, with a total of 8963 patients, comparing the role of osteosclerosis of the jaw of Denosumab and bisphosphonates, with an average treatment time of 7-20 months. The use of this drug significantly increased the risk of developing the disease (RR 1.613), with an incidence of 1.7%.6,8 The recommended dose of Denosumab is 120 mg sc on day 1, 8, 15, 29 and then 4 weekly.1 Another of the recommended schemes is the administration of 120 mg sc, every 4 weeks, with an additional 120 mg on days 8 and 15 the first month of treatment for a period of 2.5 to 13 months. In patients presenting primary GCT, treatment should be shorter, 3.5 months, and in patients with recurrent GCT, 6-13 months.4

CONCLUSION

In patients with unsecetable advanced GCT, radiotherapy and serial embolization provide palliation of symptoms, but long-term response is infrequent and malignant transformation may occur after radiation. This is where systemic therapy with Denosumab becomes important. The objective of this neoadjuvant therapy is to facilitate surgery at a later stage, making intralesional resection technically easier and therefore improving immediate local tumor control. Further prospective studies are needed on the use of Denosumab, to reach a consensus regarding its dosage, estimated time of therapy and safety profiles. The risk of recurrence presented by the GCT treated with Denosumab and then surgery has yet to be defined. For GCT with surgical contraindication, Denosumab therapy may even be required for life.

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REFERENCES

1. Ng VY, Davidson DJ, Kim EY, Pollock SM, Conrad IEu, Jones RL. The multidisciplinary management of giant cell tumor of bone. Expert Rev Anticancer Ther. 2014;14(7):783-90.
2. Van der Heijden L, Dijkstra S, Blay JY, Gelderblom H. Giant cell tumour of bone in the sacrum: interim analysis of an open-label, parallel-group, phase 2 study. Lancet Oncol. 2013;14(9):901-8.
3. Chawla S, Hershaw R, Seeger L, Choy E, Blay JY, Ferrari S, et al. Safety and efficacy of denosumab for adults and skeletally mature adolescents with giant cell tumour of bone in the denosumab era. Eur J Cancer. 2017;77:75-83.
4. Ruggiero S. Osteonecrosis of the jaw: BRONJ and ARONJ. Faculty Dental Journal. 2014;5(2):90-3.
5. Hanley DA, Adachi JD, Bell A, Brown V. Denosumab: mechanism of action and clinical outcomes- Available Int J Clin Pract. 2012;66(12):1139-46.
6. Turner B, Ali S, Drudge-Coutes L, Pati J, Nargund V, Wells R. Skeletal Health Part 1: Overview Of Bone Health and Management In the Cancer Setting. Urol Nurs. 2016;36 (1):17-21.
7. Thomas D, Hershaw R, Skubitz K, Chawla S, Staddon A, Blay JY, et al. Denosumab in patients with giant-cell tumour of bone: an open-label, phase 2 study. Lancet Oncol. 2010;11(3):275-90.