

ROLE OF MUCOSAMIN® IN THE PREVENTION OF MUCOSITIS IN PATIENTS UNDERGOING HAEMATOPOIETIC STEM CELL TRANSPLANT: CONTROLLED CASE STUDY

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INTRODUCTION:

Oral mucositis, also called stomatitis, is a multifactorial disease defined as "an epithelial thinning associated with intense erythema, ulceration, pain, bleeding, and increased risk of infection". The cytotoxic effects of antineoplastic drugs on high turnover tissues such as the oral epithelium, and the local effects of radiation on the oral mucosa, are responsible for this manifestation, which significantly compromises the patient's quality of life. It is particularly manifested as a complication in patients who have undergone haematopoietic stem cell transplant (HSCT). The most affected sites are the non-keratinised mucosa (floor of the mouth, buccal mucosa, labial mucosa and the tongue). There is currently no effective protocol for its prevention.

PURPOSE:

This aim of this study is to assess the clinical effects of Mucosamin® in the prevention and management of pain caused by oral mucositis following haematopoietic stem cell transplant. Mucosamin® is a sodium hyaluronate preparation combined with a pool of amino acid collagen precursors, including L-Proline, L-Leucine, L-Lysine and Glycine. The importance of professional oral hygiene in reducing the severity of mucositis, as a single therapy or in addition to the Mucosamin® treatment, was also assessed.

MATERIALS AND METHODS:

A case-control type of study was carried out on a sample of 101 patients. They were all recruited from the bone-marrow transplant list at the Oral Surgery Department of the Turin School of Dentistry.

They were then divided into 3 randomised groups

- GROUP 1: (33 patients): professional oral hygiene session and instructed to use Mucosamin® from the first day of admission;
- GROUP 2 (32 patients): professional oral hygiene session and standard treatment with 0.20% Chlorhexidine mouthwash
- GROUP 3 (34 patients): prescribed 0.20% Chlorhexidine only

EVALUATION SYSTEMS

• WHO MUCOSITIS SCALE: 0 = no symptoms; 1 = pain, erythema; 2 = erythema, ulceration but able to eat solids; 3 = ulceration and the need for a liquid diet; 4 = oral feeding not possible

• OMAS Mucositis: (Ulceration 0 = no lesion; 1 = lesion <1 cm²; 2 = lesion from 1 to 3 cm²; 3 = lesion > 3 cm²; Erythema 0 = none; 1 = not severe; 2 = severe)

• PERIODONTAL PLATE: Plaque index. PSR bleeding index

• DURATION OF MUCOSITIS IN DAYS



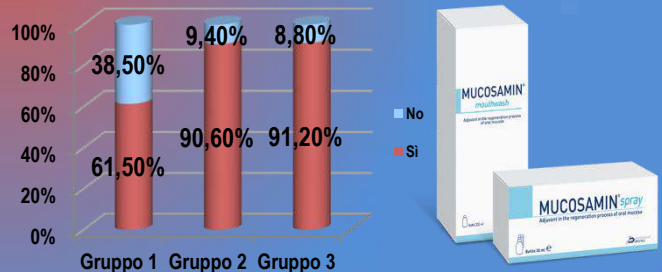
RESULTS:

The data showed that 81% of the entire sample developed oral mucositis; in particular there is a statistically significant difference with regard to the protective action of Mucosamin® as 38.50% of patients belonging to Group 1 have benefited. It has been shown that only 61.5% of the subjects belonging to Group 1 developed mucositis, against 91% of the remaining two groups. In particular, the results that derive from the WHO Scale are very interesting, as Group 1 showed a statistically significant prevalence of mild mucositis (39% WHO 1) compared to the other two groups that showed more cases of severe mucositis (52% WHO3 and 35% WHO 4) (Group 1 vs Group 2 p-value 0.005; Group 1 vs Group 3 p-value 0.003). These results are in agreement with those deriving from the OMAS Scale, in which Group 1 has the highest percentage of mild mucositis (54% OMAS 1) while Group 3 is the one that developed mucositis in the most severe form (51.6% OMAS 3). Analyzing the healing times most of the lesions were resolved altogether between 7 and 30 days, we highlight also in this case the adjuvant action of Mucosamin® as only Group 1 has as a maximum follow-up the value of 21 days.

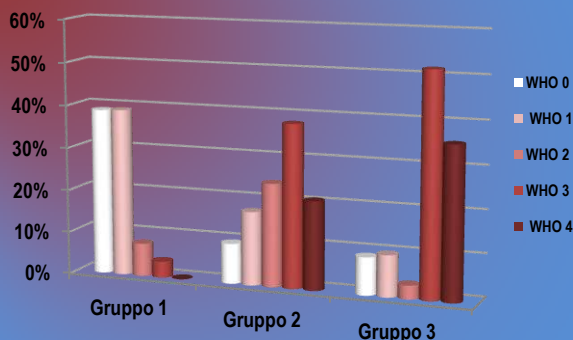
Group	WHO	Patients
Group 1	WHO 1	14 patients out of 33 (41.3%)
Group 2	WHO 3	11 patients out of 34 (38%)
Group 3	WHO 3	16 patients out of 34 (51.6%)

Pvalue	Severe Degree (3+4 on the WHO scale)
Group 1 vs Group 2	0,0004
Group 1 vs Group 3	3,3 x 10 ⁻⁸

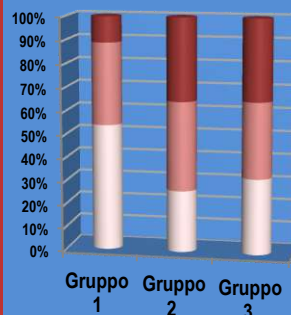
Development of mucositis



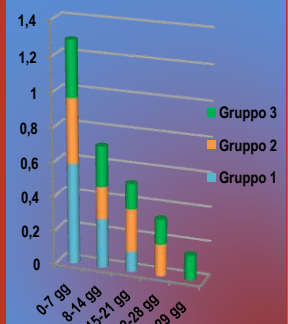
WHO Degree Scale



Omas Degree Scale



Healing time



CONCLUSIONS:

This study allows us to conclude that the combination of proper oral hygiene during hospitalization and the correct use of Mucosamin® reduce both the onset of oral mucositis and, it appears, the severity of the latter, influencing consequently on the patient's discomfort, in a statistically significant way. The mouthwash in prevention and the spray in treatment, both based on hyaluronic acid and amino acids (Mucosamin®), can therefore be a valuable therapeutic aid for oral mucositis in patients undergoing HSCT.

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