



# Incidence, severity and management of cancer chemotherapy related oral mucositis in Eastern Cape and Western Cape

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This study explored the incidence, duration and severity of oral mucositis in patients receiving chemotherapy in the Eastern and Western Cape, how this symptom was managed and whether the patients considered the management to be effective. An exploratory, contextual, quantitative survey was conducted. The sampling method was convenience. One hundred and sixty patients were recruited, with 106, (66.3%) participating. Data were collected by means of self-reports, using a self-administered questionnaire. Oral mucositis was a common problem, with 71.7% ( $n = 76$ ) reporting to have had mucositis. Pain was not effectively managed, as 69.8% ( $n = 53$ ) of respondents experienced pain whilst only 17.1% ( $n = 13$ ) reported to have used analgesics. More than half of the respondents used prescribed mouth and throat preparations, whilst 28.9% ( $n = 22$ ) used non-prescribed self-care measures including potentially harmful products. A significant difference was found between using non-prescribed self-care measures and the duration of oral mucositis ( $\chi^2 = 0.81$ ;  $p = 0.01$ ). The reported grade of mucositis did not influence the use of non-prescribed self-care measures, whilst the more pain patients experienced the less inclined they were to use these measures. The management of oral mucositis remains a challenge. Failure to palliate this distressing symptom can lead to the use of potentially harmful self-care measures.

Die studie het die insidensie, tydsduur en intensiteit van orale mukositis in Oos en Wes Kaapse pasiënte wat kankerchemoterapie ontvang verken asook hoe hierdie simptome hanteer is en die sukses hiervan volgens die pasiënte. 'n Kwantitatiewe, eksploratiewe, kontekstuele opname is onderneem. 'n Gerieflikheidssteekproef is gebruik om die deelnemers te verkry. Een honderd en sestig persone is genader en 66.3% ( $n = 106$ ) het aan die studie deelgeneem. Die datainsamelingsmetode was self-rapportering met behulp van 'n vraelys en beskrywende statistiek is gebruik om die data te verwerk. Orale mukositis was 'n algemene probleem, aangesien 71.7% ( $n = 76$ ) van die respondente die simptome ondervind het. Pyn was nie goed beheer nie, aangesien 69.8% ( $n = 53$ ) pyn ondervind het terwyl slegs 17.1% ( $n = 13$ ) aangedui het dat hulle analgetika gebruik het. Meer as die helfte van die respondente het voorgeskrewe mond en keel preparate gebruik, terwyl 28.9% ( $n = 22$ ) self gemedikeer het wat potensieel nadelige preparate ingesluit het. 'n Beduidende verskil is tussen die gebruik van nie voorgeskrewe medikasie en die duur van mukositis bevind ( $\chi^2 = 0.81$ ;  $p = 0.01$ ). Die graad van die mukositis het nie die gebruik van die self-medikasie beïnvloed nie. Hoe meer pyn ondervind is, hoe minder was die neiging om die mukositis self te behandel. Die hantering van orale mukositis bly 'n uitdaging. Die onsuksesvolle palliasie van hierdie simptome kan tot die gebruik van potensieel nadelige self-sorg maatreëls lei.

## Introduction

Oral mucositis is a debilitating symptom which can have a profound effect on the quality of life of a person diagnosed with cancer (Witt 2007). Oral mucositis leads to pain which can become so severe that the patient is unable to eat or drink and can undermine the willingness to continue with chemotherapy (Dougherty & Bailey 2008). Oral mucositis can also lead to chemotherapy dose reductions, cessation of chemotherapy, hospitalisation, reliance on parenteral nutrition and even death (Sonis 2007). South African studies focussing on cancer chemotherapy-related mucositis are limited and no evidence is available of how this symptom is managed. This study therefore focused on oral mucositis, not in terms of the chemotherapy regime the patient received, but as a complication of cancer and the treatment thereof – the incidence, severity and management according to the patient.

Mucositis, or inflammation of the mucous membranes, is a common side effect for cancer patients receiving chemotherapy and/or radiotherapy and can occur anywhere along the digestive



tract – from the mouth to the anus (Newton, Hicky & Mars 2009). Oral mucositis is defined as mucositis of the oral and oropharyngeal mucous membranes and includes mucositis of the lips, tongue, gingiva, buccal mucosa, palate and floor of the mouth (Dougherty & Bailey 2008). Oral mucositis, a condition characterised by inflammation and ulceration of the mouth with pseudomembrane formation, affects more than 40.0% of patients receiving chemotherapy and/or radiotherapy (Naidu, Ramana, Rani, Mohan, Suman & Roy 2004; Volpato, Silva, Oliveira, Sakai & Machado 2007). According to Brown, McGuire, Peterson, Beck, Dudley and Mooney (2009), 51.0% of cancer patients receiving outpatient chemotherapy experience a sore mouth and all those receiving radiotherapy of the head and neck, including the oral cavity, will experience oral mucositis (Fulton & Treon 2007; Cawley & Benson 2009).

Although the incidence and severity of mucositis differs between patients and the treatment received, the risk of developing this condition increases with the number of chemotherapy cycles as well as previous episodes experienced. Chemotherapy which is cell cycle specific for the S-phase of cell division, like fluorouracil, methotrexate and Gemcitabine mercaptopurine, leads to more severe oral mucositis. It is also suspected that cancer chemotherapy administered as bolus and continuous infusion poses a greater risk for oral mucositis compared to prolonged and repetitive administration of lower doses (Naidu *et al.* 2004).

The pathophysiology of mucositis is not fully understood (Naidu *et al.* 2004) but it is thought that chemotherapy and radiotherapy-induced mucositis has two mechanisms, direct and indirect. Direct mucositis refers to the influence that chemotherapy and radiotherapy have on the maturity and cellular growth of the mucosa. The epithelial cells of the oral mucosa undergo rapid turnover – usually every seven to 14 days – and are therefore susceptible to the effects of chemotherapy. Indirect mucositis of the oral cavity refers to an invasion by Gram-negative bacteria and fungal species. Neutropenic patients are especially at risk for oral infections and when the oral cavity is infected, indirect mucositis appears. The onset of mucositis secondary to myelosuppression varies, depending on the timing of the neutrophil nadir associated with the specific chemotherapy drugs the patient receives. However, mucositis will typically develop at any point between the 10th and 21st day after receiving chemotherapy (Naidu *et al.* 2004) and presents as a shallow ulcer, probably caused by depletion of the basal epithelial layer leading to denudation. The healing response consists of the formation of a pseudomembrane formed by an inflammatory cell infiltrate, an interstitial exudate and the remains of cells and fibrin. This membrane is analogous to an eschar in a superficial skin lesion (Bensadoun, Magné, Marcy & Demard 2001, in Volpato, Silva, Oliveira, Sakai & Machado 2007).

## Problem statement

According to Volpato *et al.* (2007), there is no consensus on the effectiveness of a variety of measures used to prevent or treat

oral mucositis, therefore standard treatment does not exist. It is unclear how many South African patients experience oral mucositis as it is suspected that patients under-report their cancer and cancer treatment related symptoms. It was also not known which self-care measures patients used to manage their mucositis and how effective this was. The research question for the study was therefore: What is the incidence, duration and severity of oral mucositis in patients receiving chemotherapy in the Eastern and Western Cape, how was this symptom managed and how effective did the patients consider the management to be?

## Trends

The incidence of cancer chemotherapy-related oral mucositis has been described in various studies (Chan, Chang, Molassiotis, Lee & Lee 2003; Sonis, Elting, Keefe, Peterson, Schubert, Hauer-Jensen, Bekele, Raber-Durlacher, Donnelly & Rubenstein 2004; Avritscher, Cooksley & Elting 2004; Nishimura, Nakano, Ueda, Kodaira, Yamada, Mishima, Yokoyama, Terui, Takahashi & Hatake 2010). The prevention, management, treatment and treatment guidelines for oral mucositis have also been explored, described and evaluated in Europe and countries including the United States, United Kingdom, Japan and Europe (Eilers & Million 2007; Harris, Eilers, Harriman, Cashavelly & Maxwell 2008; Clarkson, Worthington, Furness, McCabe, Khalid & Meyer 2010; Feller, Essop, Wood, Khammasisa, Chikte, Meyerov & Lemmer 2010; Wu, Beale & Ma 2010; Eilers & Million 2011; Worthington, Clarkson, Bryan, Furness, Glenney, Littlewood, McCabe, Meyer & Khalid 2011). How patients experienced oral mucositis have been explored and described by Chen (2008) and Brown, Beck, Peterson, McGuire, Dudley & Mooney (2009) and the impact of severe oral mucositis on the function and quality of life by Cheng (2010). Sieracki, Johannik, Kopaczewski & Hubert (2009) described the development and implementation of a patient-centred oral care protocol whilst Kearney, Miller, Maguire, Dolan, MacDonald, McLeod, Maher, Sinclair, Norrie & Wengström (2008) assessed a nursing intervention aimed at the reduction of chemotherapy-related symptoms including oral problems. The South African literature describes the pathobiology, epidemiology and management of chemotherapy and radiotherapy-related oral mucositis (Feller, Essop, Wood, Khammasisa, Chikte, Meyerov & Lemmer 2010) and a nursing management approach relating to oral mucositis (Robinson 2008).

## Objectives of the study

Little is known about oral mucositis as side effect of South African patients receiving cancer chemotherapy as no literature investigating this side effect in the specific setting could be found. The objectives of the study were to explore the incidence, duration and severity of oral mucositis in cancer patients receiving chemotherapy in the Eastern and Western Cape; how this symptom was managed and how effective the patients considered the management to have been.



## Contribution to field

The current study provides baseline data allowing reflective practice in terms of how this side effect is currently prevented and managed. The study furthermore provides a baseline allowing comparative studies in terms of the application of evidence based practice and interventions to improve disease and treatment outcomes of patients suffering from cancer.

## Research methods and design

### Strategy and setting

An exploratory strategy was used for this study. The setting was a private oncology practice based in Cape Town, South Africa. This practice forms part of the private health care system and has various satellite practices in Western and Eastern Cape. Cancer patients with a wide variety of diagnoses are treated at this practice, most commonly on an outpatient basis. Modern equipment and technology are used and patients are managed in a holistic manner by a multi-professional team.

### Design and recruitment strategy

A quantitative survey was conducted and 160 patients were recruited but 106 ( $n = 106$ ), 66.3%, participated. The inclusion criteria were older than 18 years, treated with chemotherapy and willing to participate. Sampling was convenience as all patients 18 years and older, receiving chemotherapy and willing to participate were included.

### Survey instrument

To enable data collection a self-administered questionnaire based on literature and expert opinion was developed. The questionnaire consisted of two sections and contained both open-ended and closed-ended questions. Section A allowed the gathering of general information whilst Section B focused on oral mucositis. Mucositis related questions concentrated on the number of times the respondent experienced this symptom, the severity, the pain experienced, how this symptom was managed and how effective the respondent considered the management to have been. The oral toxicity scale of the World Health Organization was used as the grading scale. The questionnaire was pre-tested using the first 10 respondents ( $n = 10$ ), with no changes made as respondents raised no queries. The data gathered during the pre-test forms part of the final data.

### Data collection and analyses

The data were collected during respondents' scheduled appointments in August and September 2010. The field worker explained the purpose of the study and informed consent; handed the questionnaire to the respondent and requested the respondent to hand it back once completed. No time limit applied. Completed questionnaires were numbered and sealed in an envelope. The data were entered onto an Excel spreadsheet and analysed by means of the SPSS 14 computer programme. The data are presented as descriptive statistics. The chi-square was used for secondary data analyses.

## Results

The median age of respondents was 60 years with the average age 55.8 years. Females were in the majority (59.4%) with the most common diagnosis being breast cancer. The general characteristics of the respondents are outlined in Table 1.

### Incidence, duration and severity

When asking respondents whether they had ever had oral mucositis 71.7% ( $n = 76$ ) indicated they had this complication, whilst 28.3% ( $n = 30$ ) denied ever having it. When asked how many times they had had oral mucositis, 46.1% indicated once or twice, whilst 19.7% indicated three to five times, 9.2% did not know whilst the rest (25%) indicated more than five times. When respondents were asked how long their mouths were sore, the number of days ranged from one to 14 (Table 2); one respondent (0.9%) indicated her mouth was

**TABLE 1:** General characteristics ( $n = 106$ ).

Variable	<i>n</i>	%
<b>Age group</b>		
18–19	1	0.9
20–29	4	3.8
30–39	7	6.6
40–49	20	18.9
50–59	18	17.0
60–69	44	41.5
70 and older	12	11.3
<b>Gender</b>		
Male	43	40.6
Female	63	59.4
<b>Marital status</b>		
Married civil	77	72.6
Married traditional	4	3.8
Single	12	11.3
Widowed	4	3.8
Divorced	9	8.5
<b>Diagnosis</b>		
Breast cancer	28	26.4
Cervical cancer	4	3.8
Colorectal cancer	21	19.8
Liver cancer	5	4.7
Lung cancer	9	8.5
Lymphoma	6	5.7
Ovarian cancer	10	9.4
Prostate cancer	4	3.8
Other	19	17.9

**TABLE 2:** Duration of a sore mouth ( $n = 76$ ).

Days	<i>n</i>	%
1	3	3.9
2	11	14.5
3	16	21.1
4	10	13.2
5	7	9.2
6	1	1.3
7	9	11.8
10	6	7.9
14	5	6.6
Could not tell	7	9.2
Permanently	1	1.3
<b>Total</b>	<b>76</b>	<b>100</b>



'permanently' sore, whilst seven (9.2%) could not tell. The average number of days respondents experienced a sore mouth was 5.1 days and the median 4 days.

The grading scale of the WHO was used when asking respondents ( $n = 76$ ) to describe their mouths when at their worst. Two (2.9%) did not know, whilst 29.9% graded their mucositis as Grade 1, 40.8% as Grade 2, 25% as Grade 3 and 2.9% as Grade 4. Respondents were also asked to quantify the pain they experienced when their mouths were most painful and a numerical scale was provided for this purpose. Most (69.8%;  $n = 53$ ) indicated they experienced pain and the rating varied from 0, having no pain, to 10, worst possible pain. A comparison between the reported grade of mucositis and pain level is presented (Table 3).

## Management

To determine how the mucositis was managed, three themes were explored – disclosure, prescribed medicine and self medication. Respondents were asked whether they reported the oral mucositis, to whom they reported it to and whether they were open and honest about their experience. The majority (73.7%) indicated they told a health care professional about their oral mucositis, whilst 21.5% told no one, and 5.3% did not answer the question. A greater percentage of females compared to males (25.5% vs 13.8%) did not tell either the nurse or the oncologist of their mucositis. More than half (56.3%) of the total number of respondents who did not disclose their mucositis ( $n = 16$ ) had moderate pain, whilst the rest (43.7%) experienced mild pain.

Not all respondents were prepared to disclose to whom they reported the mucositis as 25.0% did not answer the question; however, 17.1% informed a nurse, 40.8% the doctor and the rest (17.1%) told both the nurse and the doctor. When exploring whether respondents were open and honest about what they disclosed, the majority (55.2%) indicated they told the doctor and/or nurse exactly how they experienced the mucositis, 6.6% indicated they told the doctor and/or nurse but did not tell them how bad it really was, whilst 7.9% indicated to have only mentioned it but did not discuss it. The rest (30.3%) did not answer the question. No respondent selected the option 'I told them, but it made it worse than it was for me'.

More than half of the respondents who had had a sore mouth (56.6%;  $n = 43$ ) indicated they used prescribed mouth and

throat preparations. Approximately 20.6% were unable to name the medication. However, Benzidamine oral rinse was used by 47.1% alone or in combination with nystatin (38.2%) and miconazole (20.6%). Triamcinolone acetonide ointment was used by 8.8%. Respondents were asked to indicate on a numerical scale how effective they found the prescribed medication. Answers varied from 0, not effective at all to 10 extremely effective could not be better, with the average being 7.

Respondents were also asked whether they used any self-care measures not prescribed by the oncologist. More than one quarter of respondents (28.9%;  $n = 22$ ) confirmed this. The duration of mucositis played a role in using non-prescribed self-care measures as a significant difference was found between the group who reported to have had mucositis for longer than 7 days and those whose mucositis lasted 7 days and less ( $\chi^2 = 0.81$ ;  $p = 0.01$ ). The reported grade of mucositis did not influence the use of non-prescribed self-care measures, as less than 50.0% of respondents with Grade 2 and 3 mucositis and more than 50.0% of respondents with Grade 1 and 4 mucositis used these measures. A greater percentage of respondents with mild pain (56%;  $n = 14$ ) than those with moderate pain (51.2%;  $n = 18$ ) reported the use of non-prescribed self-care measures, whilst only 20.0% of those with severe pain ( $n = 10$ ) reported using these measures.

Non-prescribed self-care measures consisted mainly of oral rinses with bicarbonate of soda (19.1%) followed by oral rinses with salt water (14.3%). Other self-care measures included oral rinses with a combination of salt and bicarbonate of soda, oral rinses with Epsom salts, mouthwash containing calendula and tea-tree oil, Zam-Buk, glycerine, cold pressed sunflower oil, baby teething oral gels, commercial mouthwash, and probiotics. The use of these products were mostly recommended by a family member or friend (28.5%), pharmacist (23.8%) or nurse (23.8%); the rest used their own initiative or did not disclose the source of advice. Once again, a numerical scale was used for respondents to indicate the efficacy of their self-care measures, with answers ranging from three to 10, with the average being 7.4.

Finally, respondents were asked whether they took analgesics to alleviate the pain caused by the oral mucositis. Only 17.1% of the total number of respondents ( $n = 76$ ) indicated that they did. When comparing the use of analgesics with the pain level, it was found that none of the respondents who experienced mild pain ( $n = 25$ ) took an analgesic; 23.1% who had moderate pain ( $n = 39$ ) took analgesics; whilst 40.0% who

**TABLE 3:** Severity of oral mucositis and pain experienced ( $n = 74$ ).

Grade	Description	N	Mild pain		Moderate pain		Severe pain		Average pain score
			n	%	n	%	n	%	
1	Red and painful but no sores (ulcers)	22	22	100	0	0	0	0	2.3
2	Sores (ulcers) with or without redness. Could eat solid food	31	3	9.7	28	90.3	0	0	5.3
3	Sores (ulcers) with or without redness. Could not eat solid food but could swallow liquids	19	0	0	11	57.9	8	42.1	7.4
4	Could neither eat or drink	2	0	0	0	0	2	100	8.8
	<b>Total</b>	<b>74</b>	<b>25</b>	<b>-</b>	<b>39</b>	<b>-</b>	<b>10</b>	<b>-</b>	<b>5</b>





experienced severe pain ( $n = 10$ ) used analgesics. Over-the-counter as well as prescribed analgesics were used (Table 4).

When the respondents were asked how effective the analgesics were in relieving their pain, they could choose from four options: 'The medication worked: (1) well, it took the pain away completely; (2) well, I only had slight pain; (3) not so well, I still had pain; (4) not well at all, it made no difference to the pain'. Slightly more than one fifth (22.2%) of respondents who had moderate pain found the analgesic they used to be very effective, 66.7% reported it as effective and 11.1% as not so effective, as they still experienced pain. Of the respondents who had severe pain ( $n = 4$ ), the majority (75.0%) indicated that the pain medication was not so effective, as they still experienced pain, whilst one respondent (25.0%) indicated the medication worked well, as she only had slight pain.

## Ethical consideration

The ethical principles outlined in the Belmont Report (Polit & Beck 2010), of beneficence, respect for human dignity, justice, confidentiality and debriefing, were followed. The study was explained to all patients who met the entrance requirements and only those who volunteered were entered. Informed consent was discussed with respondents and respondents consented to participate in the study by completing the questionnaire. Anonymity and confidentiality were ensured by numbering the questionnaires sequentially. After completing the questionnaire, time was allowed for them to raise queries. Furthermore, the research proposal was peer-reviewed by the Departmental and Faculty Research and Innovations Committees of the Tshwane University of Technology and approved by the Ethics Committee of the same university as well as the Ethics Committee of the private oncology practice.

## Validity and reliability

The following measures were taken to increase the validity and reliability of the findings:

- The questionnaire was formulated and specifically planned to explore oral mucositis in patients receiving out-patient chemotherapy.

**TABLE 4:** Pain level and analgesics used ( $n = 13$ ).

Pain level	<i>n</i>	%
<b>Moderate pain (<math>n = 9</math>)</b>		
Paracetamol	1	11.1
Combination drug: ibuprofen 200mg, paracetamol 300mg, codeine phosphate 10mg	4	44.4
Combination drug: paracetamol 320mg, codeine phosphate 8mg, caffeine Anhydrous 32mg, meprobamate 150mg	3	33.3
Combination drug: paracetamol 500mg, d-propoxyphene napsylate 50mg, dyphenhydramine HCl 5mg, caffeine 50mg	1	11.1
<b>Severe pain (<math>n = 4</math>)</b>		
Paracetamol	1	25.0
Aspirin	1	25.0
Combination drug: tramadol hydrochloride 37.5mg, paracetamol, 325mg	1	25.0
Tilidine HCl	1	25.0
<b>Total</b>	<b>13</b>	<b>-</b>

- The questionnaire was pre-tested before data gathering commenced.
- Oral mucositis was graded according to the Oral Toxicity Scale of the World Health Organization (Quinn, Stone, Uhlenhopp, McCann & Blijlevens 2007).
- Data were gathered at venues and times specifically planned for data gathering.
- To create trust, registered nurses practicing at the various oncology practices were trained to gather the data.
- The lead researcher is an oncology nurse with more than 20 years experience in oncology nursing, and the co-workers who did the fieldwork are registered nurses engaged in a post-registration learning programme in oncology nursing.

## Discussion

The study provided evidence that oral mucositis is a common complication, with most patients reporting having had mucositis on more than one occasion. This incidence level (71.7%) was slightly lower than the 75.4% reported by Chen (2008), but higher than the 32.0% reported by Goldberg, Chiang, Selina & Hamarman (2004) and the 51.0% reported by Elting, Cooksley, Chambers, Cantor, Manzullo & Rubenstein (2003). The re-occurrence of oral mucositis is not unique, as Elting *et al.* (2003) reported that 53.0% of their sample had previously had this complication. In the current study, most patients (89.4%) reported that their mucositis lasted 14 days or less, a finding supported by Cheng (2006), who reports that more than 90.0% of patients will recover from oral mucositis within one to two weeks. One respondent's statement that her mouth was 'permanently' sore could raise questions, but in their study interpreting patients' experience of oral mucositis, Borbasi, Cameron, Queded, Olver, To & Evans (2002) found that oral symptoms have the potential to persist and become chronic, and that it does not mean that 'all is fine' once the ulcers have healed, allowing the patient to swallow. Although these authors did not describe pain as part of the chronic problems associated with oral mucositis, the current study provides evidence of this.

As illustrated in this study, pain is a problem associated with oral mucositis. Feller *et al.* (2010) support this finding, stating that oral mucositis is 'almost always' painful, whilst Elting *et al.* (2003) state that pain 'frequently' accompanies oral mucositis. Interestingly, patients rated their pain according to the way they graded their mucositis. This finding alleviates the fears of some nurses who, according to Maree (2009), still believe cancer patients over-exaggerate the level of pain they experience.

The finding that some patients would not report their oral mucositis, or how bad it really was for them, reiterates the importance of assessment, as patient problems cannot be addressed without it. Farrell *et al.* (2005) support this statement with their finding that nurses are unable to identify 80.0% of the concerns of cancer patients undergoing chemotherapy. Pain, especially severe pain, was not effectively managed. Being unable to prescribe medication cannot serve as justification for



this situation, as the advocacy role of the nurse requires acting as a 'go-between' for the patient and doctor (Anstay 1997).

One interesting finding is the fact that the more pain the patients experienced, the less inclined they were to use self-care measures. The reason for this is unclear, especially in the light of pain not being managed effectively, and should be explored. Symptom management is essential for patients with mucositis and nurses should use their skills to intervene before the situation becomes difficult (Borbasi *et al.* 2002).

Not all patients found the prescribed oral and throat preparations helpful, and some reverted to self-care measures, which they found to be more helpful. Köstler, Hejna, Wenzel & Zielinski (2001), as well as Cheng (2006), support this finding by stating that no single uniformly efficacious agent or intervention has been identified to manage chemotherapy-related oral mucositis which could serve as evidence-based standard therapy. However, there is reasonable evidence that using Benzadimine oral rinse, a non-steroid anti-inflammatory drug seems efficacious in reducing ulceration (Sonis, Clairmont, Lockhart & Connolly 1985). Although topical Triamcinolone is used for aphthous ulcers (Flint 2006), evidence is lacking of its efficacy, specifically in chemotherapy-related oral mucositis.

To assume that patients suffering from cancer do not use non-prescribed self-care measures and medication is unrealistic and the study provides evidence that patients do indeed use non-prescribed self-care. The danger is the potential harm of these measures. A saline solution (0.9% sodium chloride) is non-irritant and might be the least harmful mouthwash available (Miller & Kearney 2001); using sodium bicarbonate is also part of recommended practice (The Johanna Briggs Institute 2010) but it is unclear whether the patients in this study used the correct formula to constitute the solution. Commercial mouthwashes may contain oils, antiseptics, alcohol, astringents and aromatic substances which have the potential to irritate the mucosa and lead to hypersensitive mucositis. Using glycerine or glycerine-based products could also be harmful, as they dry the oral mucosa (Kirshnasamy 1995, in Atkinson & Virdee 2001). Evidence is not available as to the efficacy or potentially harmful effects of other products used topically or as a mouthwash.

## Limitations of the study

This study has several limitations. The study was conducted in specific private cancer care settings and the results are therefore only applicable to this specific patient population. Convenience sampling was used for sample selection, which could have led to bias. Using a questionnaire as a data-collection instrument allowed the respondents to not answer all the questions and furthermore they were familiar with the investigators and could have provided socially acceptable answers. Respondents did not necessarily have oral mucositis during the gathering of the data and recall bias might have been possible.

## Recommendations

Unfortunately, research to date has not been able to identify a universal effective intervention for the prevention and treatment of mucositis (Rubenstein, Peterson, Schubert, Keefe, McGuire, Epstein, Elting, Fox, Cooksley & Sonis 2004; Feller *et al.* 2010). Using a standard approach based on the current evidence is therefore recommended. A standard approach could include a baseline assessment consisting of a risk assessment and oral inspection before any patient commences chemotherapy, ongoing assessment, an oral mucositis prevention protocol as well as a protocol for the management of this side-effect (The Johanna Briggs Institute 2010). Once standard care plans are used, individual care plans could be developed to ensure the best outcomes for each patient and allow the development of practice-based evidence. Furthermore, nurses should fulfil their advocacy role to ensure that appropriate analgesics, informed by the patient's reported pain level, are prescribed for every patient suffering from oral mucositis.

## Conclusion

The management of oral mucositis remains a challenge. Failure to palliate this distressing symptom can lead to the use of potentially harmful self-care measures. This may be avoided by thorough assessment, the implementation of standard care and developing individualised care plans to improve patient outcomes.

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## Competing interests

The authors declare that they have no financial or personal relationship(s) which may have inappropriately influenced them in writing this article.

## Authors' contributions

J.E. Maree (University of the Witwatersrand) was the project leader and wrote the manuscript. M.J. Combrink, T. de Lange and A.S. Toerien (Tshwane University of Technology) assisted with the data gathering and reviewed the manuscript.

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