

Using Patient Reported Outcome Measure (PROM) to Manage Pain During Radiation Therapy

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Introduction

Pain severity is an important symptom to assess for in head and neck cancer patients undergoing radiotherapy (RT) due to its effects on the quality of life [1]. Although radiation itself is painless, it induces an acute postradiation reaction that can cause many side effects, with the most debilitating being oral mucositis [2]. Studies have shown a correlation between severity of pain and severity of oral mucositis [3]. To help manage the pain, the World Health Organization (WHO) has a recommended escalation chain of analgesics guideline, starting with NSAIDS, then weak opioids, followed by potent opioids [4]. Due to the subjective nature of pain, which can be difficult for health systems to measure, patient reported outcome measures (PROMs) is a tool that can be used to better gauge and improve patient's health experience [5]. Significant pain signals that modifications in pain management are needed to better monitor the pain. In this study, we used an institution-based PROM, The MD Anderson Symptom Inventory (MDASI), for pain assessment. The objective of our project is to determine the efficacy of pain management during RT using PROM.

Pain Management During RT



Results

MDASI pain scores reported during weekly see visits revealed a significant increase through the course of RT (p <0.0001), with the highest severity of pain is experienced at the end of RT. Pain scores were tested across age groups. There were no significant differences in pain scores between age cohorts, with a similar trend in each week across the cohorts.

Methods

Retrospective data was obtained for 500 oral cavity and oropharyngeal (OC/OPC) cancer patients, treated with RT at the Radiation Oncology department, MD Anderson Cancer Center in Houston, TX, USA. Pain scores were collected during the weekly see visits using MDASI, which is a multi-symptom PROM used to assess the severity of symptoms and how much of that disrupts daily life [6], using a scale from 0 to 10 (0 = no pain, 10=the worst pain). Clinical data was collected using EPIC and Brocade software. Pain scores were compared to mucositis intensity (0 =none, 1 =mild, 2 =moderate, 3 =severe), assessed weekly during radiation therapy visits. Patients were divided into cohorts based on age (19-33 =1, 34-48 =2, 49-64 =3, 65-78 =4, 79-98 =5). T-test and ANOVA test were used for statistical analysis.



mucositis severity displayed a significant difference from week 1 to week 7 of RT. (p <0.0001).



Future Steps

Future steps to take can include analyzing which types of analgesics are given and at what point during radiation therapy. Stronger pain medications may be needed earlier during the course to prevent high levels of pain seen towards the end of radiation therapy. By analyzing pain scores alongside the type and when analgesics are administered, this opens the possibility for modification of the current pain management plan, improving the patient's overall quality of life and experience.

Conclusions

The aim of this study is to determine the efficacy of pain management during radiation therapy using PROM. Our data suggests that pain levels consistently increased during RT. This suggests the pain management currently implemented may not adequately control pain during the weeks of RT and may be in need for improvement. This is important to address due to the negative effect pain can have on the quality of life [1].



=5).

Pain Scores Across Age Cohorts

Cohort 1

Cohort 2

weekly visits

Week 7-

Types of Analgesics Administered



Fig. 4 Collected the types of analgesics administered (based on the WHO analgesics ladder) to 224 out of the 500 patients during the course of RT. 0=no and 1=yes. The mean values for each analgesic is displayed.

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