

## THE EFFECT AND SAFETY OF TRANSDERMAL FENTANYL FOR CONCURRENT CHEMORADIATION THERAPY- INDUCED MUCOSITIS PAIN IN HEAD AND NECK CANCER PATIENTS

Shih-Hua Liu<sup>1</sup>, Yu-Jen Chen<sup>1,6</sup>, Kou-Hua Chang<sup>1,7</sup>, Yi-Shing Leu<sup>2</sup>, Jehn-Chuan Lee<sup>2</sup>,  
Ming-Jer Huang<sup>3</sup>, Hong-Wen Chen<sup>1,4,7</sup>, Yuen-Liang Lai<sup>1,4,5,6\*</sup>

*Department of Radiation Oncology<sup>1</sup>, Department of Otorhinolaryngology<sup>2</sup>, Department of Medical Hematology<sup>3</sup>,  
Hospice and Palliative Care Center<sup>4</sup>, Mackay Memorial Hospital, Taipei, Taiwan  
Center of General Education<sup>5</sup>, National Yang-Min University  
School of Medicine<sup>6</sup>, Taipei Medical University  
Mackay Medicine<sup>7</sup>, Nursing and Management College*

**Purpose** : This prospective, longitudinal study was to evaluate the efficacy and the safety of using TTS-fentanyl for severe oral mucositis pain induced by concurrent chemoradiotherapy (CCRT) in head and neck cancer (HNC) patients.

**Materials and Methods** : Patients diagnosed as HNC and scheduled for concurrent chemoradiotherapy were eligible for this study. TTS-fentanyl was given with the initial dose of 25 µg/h when severe mucositis pain (numeric rating scale [NRS] ≥ 7) occurred in spite of nonsteroid anti-inflammatory drugs (NSAID) and topical steroidal ointment use. The pain intensity was assessed every week and if severe pain persisted, another dose of 25 µg/h TTS-fentanyl was given. The adverse effects were recorded every week according to Common Toxicity Criteria (CTC) Version 2.0.

**Results** : From October 2002 to March 2003, 21 male and 1 female receiving CCRT with HNC were recruited to this study. The mean NRS for pain significantly decreased from 7.65 ± 0.99 (baseline) to 3.9 ± 1.29 within the first week. In the fifth week, the NRS raised to 4.79 ± 1.18 but was still lower than the baseline (p = 0.003). The sufficient analgesia was achieved in 15 cases with a dose of 25 µg/h, in 5 cases with a dose of 50 µg/h, and in 1 case with a dose of 100 µg/h. One patient withdrew the TTS-fentanyl due to intolerable skin rash and vomiting in the second week. Besides, no one developed respiratory depression, mental clouding, or grade III/IV gastrointestinal toxicities.

**Conclusion** : TTS-fentanyl is effective and safe in the treatment of mucositis-induced pain which is resistant to NSAID in HNC patients receiving CCRT. TTS-fentanyl also spares the need to swallow pain killers in these patients suffering from odynophagia and/or dysphagia, as well as the time for titration of conventional analgesic drugs. Further larger series with randomized setting is warranted to evaluate the efficacy and safety of TTS-fentanyl for oral mucositis pain.

[ Therapeut Radiol Oncol 2005; 12(2): 103-111 ]

Key Words: Transdermal fentanyl, Mucositis, Head and neck cancer, Concurrent chemoradiation therapy (CCRT)