

Buprenorphine for the Treatment of Pain in Cancer Patients

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ABSTRACT

Background: Opioids remain the cornerstone for the treatment of moderate to severe cancer pain. Due to benefits over full agonist opioids (FAO), buprenorphine has emerged as an alternative treatment.

Purpose/hypothesis: Buprenorphine is only approved for the treatment of pain that is chronic non-cancer. Cancer-related pain is often progressive with breakthrough pain. There is limited evidence for using short-acting FAO in combination with buprenorphine. There are concerns about withdrawal and the efficacy of pain control using buprenorphine. We hypothesize buprenorphine, in combination with short-acting FAOs, can adequately control cancer-related pain without causing withdrawal symptoms.

Procedures/data/observations: Our prospective, single-arm, open-label study enrolls patients with cancer-related pain who are on buprenorphine in combination with an FAO at > 30 mg OME/day, either requiring long-acting pain relief or their pain is not controlled with an FAO alone. Our study is ongoing, with 15 patients enrolled and a target of 50. The patient's pain is self-assessed daily using a mobile application. Withdrawal is assessed regularly using a modified Clinical Opioid Withdrawal Scale (COWS) score.

Conclusions/applications: Buprenorphine appears to be effective for the treatment of cancer pain without causing withdrawal in combination with short-acting FAO >30 mg/day.

Presentation link:
<https://bupe2021.com/10-5055-bupe-24-rp-1015/>

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