Evaluation of the Efficacy of Low-Dose Naloxone for the Prevention of Acute Remifentanil-Induced Hyperalgesia in Patients Undergoing General Anesthesia for Laparotomy

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Abstract

Background: Hyperalgesia is a major complication of continuous or intermittent opioid administration. The evidence suggests that concomitant administration of low-dose naloxone could prevent the development of acute opioid-induced hyperalgesia, with no effect on pain control.

Objectives: The current study aimed to assess the effects of intraoperative low-dose naloxone, adding to remifentanil infusion on preventing acute postoperative hyperalgesia in patients undergoing general anesthesia for laparotomy.

Methods: In this randomized clinical trial, patients undergoing general anesthesia for laparotomic hysterectomy in a tertiary referral teaching hospital from February to December 2019 were randomly assigned to one of three groups of remifentanil-naloxone (remifentanil 0.3 µg/kg/min with low-dose naloxone 0.25 µg/kg/h prepared in 50 mL of normal saline), remifentanil (0.3 µg/kg/min), and control (receiving 50 mL saline infusion), intraoperatively. Patients and researchers were blinded to the type of intervention. The severity of hyperalgesia, as the main outcome, was evaluated by the static Tactile test. The severity of pain was assessed by visual analogous scale 0.5, 2, 6, 12, and 24 hours after surgery.

Results: In total, 75 patients were evaluated. The results showed no difference concerning the independent variables (age, body mass index, hypertension, surgery duration, anesthesia duration, and American Society of Anesthesiologists (ASA) class) between the three groups. Heart rate was significantly different in all study time points between the three groups (P < 0.001), but mean arterial pressure and systolic and diastolic blood pressure showed no significant difference (P > 0.05) throughout the study. Assessment of hyperalgesia using the tactile test revealed a higher incidence of hyperalgesia in the remifentanil group in 0.5, 2, 6, 12, and 24 hours after surgery compared to the other two groups, which was statistically significant between the groups at 0.5, 2, and 6 hours after surgery (P < 0.05). Shivering incidence, Morphine dose in 24 hours post-surgery, morphine dose in the recovery room, and VAS for pain were significantly different during the study between the three groups (P < 0.05).

Conclusions: This study demonstrated the efficacy of intraoperative low-dose naloxone (0.25 µg/kg/h) added to remifentanil infusion on reducing the frequency and severity of acute postoperative hyperalgesia in patients undergoing general anesthesia for laparotomy hysterectomy.

Keywords: Naloxone, Acute Hyperalgesia, Remifentanil, General Anesthesia, Hysterectomy

1. Background

Remifentanil is a µ-opioid receptor agonist, which is recognized as a cardinal drug widely using in the total intravenous anesthesia (TIVA) method (1). It also can enhance the quality of anesthesia in those without a cancer diagnosis. Opioids are commonly using for maintaining general anesthesia in the United States (2-4). However, widespread using of opioids and their non-medical misuse have raised concerns about the harms and drawbacks of opioids, including remifentanil (4). Opioid-induced hyperalgesia (OIH) is a major consequence that has attracted the attention of many researchers (5). OIH is a major complication of continuous or intermittent opioid administration, which in turn increases the sensitivity to pain. This condition paradoxically enhances the sensitivity to pain,
which is alleviated by lowering opioid doses. The pain pattern may be identical to the pain due to the underlying disease or appears different in quality. The patient may feel the pain in a different location(s) or use neuropathic pain descriptors (6). Although many animal or clinical studies have reported hyperalgesia following opioid administration, especially remifentanil, there is little known about the measures to prevent or decrease the severity or prevalence of acute OIH (7, 8). The evidence suggests that concomitant administration of low-dose naloxone can prevent the development of acute OIH, with no effect on pain management (8, 9).

2. Objectives

The current study aimed to assess the effects of intraoperative low-dose naloxone, adding to remifentanil infusion on preventing acute postoperative hyperalgesia in patients undergoing general anesthesia for laparotomy.

3. Methods

In this prospective double-blinded trial, 75 ASA I - II patients aged 18 to 75 years scheduled for an elective hysterectomy in a tertiary referral teaching hospital in Tehran from February to December 2019 were recruited. Informed written consent was obtained from all participants. Those who didn’t meet inclusion criteria were excluded. Exclusion criteria were, having uncontrolled diabetes mellitus, neurologic disorders, psychologic disorder(s) which needs treatment, inflammatory and renal diseases, neuropsychiatric disorders, drug abuse, routine use or taking non-steroidal anti-inflammatory drugs (NSAIDs), opioids, or other analgesics 48 hours prior to surgery, allergy or contraindication to anesthetic agents or pain medications, including NSAIDs, acetaminophen and opioids, and history of any type of chronic pain which needed medical or interventional treatments. Patients who develop any surgical or anesthetic complication, those who needed reoperation or needed more than 2-unit transfusion of packed red blood cells, or received ketamine, antipsychotic, or gabapentinois considered to be excluded latter.

Patients were randomized via a computer-based randomization process. All medications were prepared by investigators who were blinded to the intervention group and a written instruction; however, both patients and investigators were blinded to the intervention group and randomization process. All medications were prepared by an anesthesia nurse who was not engaged in the study.

After administering pre-medication (midazolam 0.05 mg/kg, fentanyl 2 µg/kg, and lidocaine 1 mg/kg, based on Ideal Body Weight for overweight patients) under standard monitoring, including electrocardiography (ECG), oxygen saturation (SpO2), end-tidal CO2 (ETCO2), noninvasive blood pressure monitoring (NIBP), respiratory rate (RR), and heart rate (HR), anesthesia was induced by propofol 2 mg/kg and atracurium 0.5 mg/kg. After tracheal intubation, 1.5 MAC isoflurane and muscle relaxants were used for maintaining anesthesia, if necessary. Fentanyl dose was repeated each hour to adjust the hemodynamic parameters based on anesthesia judgment.

In the remifentanil group, on the occurrence of bradycardia (HR < 50) and blood pressure drop more than 20% of baseline, interventions were provided, including administration of atropine or ephedrine and a bolus injection of crystalloids. If interventions were not efficient, the remifentanil dose was reduced. The appropriate dose of opioids was continuously adjusting based on the hemodynamic parameters during the surgery (HR and BP were increased to higher than 20% of baseline). In the postsurgery period, morphine sulfate PCA (patient-controlled analgesia) was used at 1 mg/ml; Bolus: 1 ml; lock-out interval: 7 min; basal infusion: 0 mL/h., for 24 hours after surgery to control the patient’s pain and precisely assess morphine dose consumption. All patients received IV paracetamol 1 g Q6H and IV ketorolac 30 mg Q8H.

Demographic parameters and baseline characteristics (age and surgery duration) of all participants were recorded. Total fentanyl dose used, morphine dose used in post-anesthesia care unit (PACU), morphine dose used within 24 hours after surgery, the first time-point of opioid administration after surgery and pain severity after surgery during movement, and rest according to visual analogous scale (VAS) in 0.5, 2, 6, 12, and 24 hours after surgery were documented. In PACU, shivering, the need for opioids, nausea, and vomiting (0.5, 2, 6, 12, and 24 hours after surgery) were recorded. Heart rate and blood pressure were analyzed before induction, after intubation, 30 min-
utes post-induction, and 30 minutes post-extubation.

The need for increasing remifentanil dose, based on hemodynamic parameters, was also documented. The severity of hyperalgesia and allodynia was assessed by static tactile tests. In this test, a soft brush that is only a sensory stimulus is contacted to the edge around the patient’s wound, and in the case of allodynia, the patient drastically moves the body away from this stimulus. The severity of pain was reported using the VAS. Patients were asked to grade the highest experienced pain from one to 10 in 0.5, 2, 6, 12, and 24 hours after the surgery. A pilot study was performed on five patients from all three groups, and the results were used for calculating the sample size, by considering $\alpha = 0.5$ and $\beta = 0.2$. Statistical analyses were performed using SPSS v.21.0 (IBM Corp., Armonk, NY, USA). The results are described using frequencies and mean scores. ANOVA test and Bonferroni’s post hoc test were used for comparing the groups.

4. Results

In the present study, 75 patients were separated into three groups. All participants were eligible for participation. The three groups were not significantly different concerning the baseline characteristics (age, BMI, ASA class, duration of anesthesia and surgery, and presence of underlying diseases (hypertension, asthma, and other systemic diseases) (Table 1). Hemodynamic status of patients, including HR and mean arterial pressure (MAP), were assessed at different time points. Hemodynamic parameters are described in Table 2. Except for HR in 30 minutes post-intubation and 30 minutes post-extubation, there was no significant difference between the three groups regarding the HR and MAP.

Comparing EtCO$_2$ and SpO$_2$ in PACU revealed no significant difference in various time points between the three groups. The three groups were compared concerning intraoperative fentanyl dose, time of the first dose of rescue analgesia in recovery, rescue morphine dose in the recovery room, and total 24-hours postoperative morphine dose (Table 3). Cumulative intraoperative fentanyl dose was significantly higher in the control group than the other two groups (P-value = 0.001). The time of the first dose of rescue analgesia in recovery was significantly earlier in the remifentanil group compared to the control and remifentanil-naloxone groups. Also, this time was significantly earlier in the control group than the remifentanil-naloxone group (P-value = 0.001). The dose of rescue morphine sulfate in the remifentanil group was significantly higher than the two other groups (P-value = 0.002). Also, 24-hours postoperative morphine sulfate consumption was significantly higher in the remifentanil group compared to other groups (P-value = 0.001). The need for atropine was not significantly different between the three groups (P-value = 0.810), but the incidence of shivering in PACU was significantly lower in the remifentanil-naloxone group (8%) compared to the remifentanil (40%) and control (20%) groups (P-value = 0.02). The incidence of nausea and vomiting after surgery in the remifentanil-naloxone, remifentanil, and control groups was 28%, 32%, and 32%, respectively (P-value = 0.922). Pain severity was assessed postoperatively in 5-time points, which was significantly lower in the remifentanil-naloxone group compared to the other two groups. The details of postoperative pain, measured by VAS, are provided in Table 4.

Assessment of hyperalgesia with tactile test revealed a higher incidence of hyperalgesia in the remifentanil group in 0.5, 2, 6, 12, and 24 hours after surgery compared to the other two groups, which was statistically significant at 0.5, 2, and 6 hours after surgery (P < 0.05). The details of the test at different time points are shown in Table 5.

5. Discussion

This study demonstrated that adding ultra-low doses of naloxone can improve analgesia during the postoperative period after intraoperative remifentanil infusion. Patients in the remifentanil group received significantly higher amounts of opioids and had higher scores of hyperalgesia and pain intensity compared to the other two groups. A few mechanisms, including opioid-induced hyperalgesia, tolerance, and withdrawal syndrome, have been proposed for increased pain intensity after opioids infusion. The pain resulting from tolerance to opioids usually happens in the site of tissue injury and relieved by additional opioids. OIH will be perceived even in the distant areas (far from the tissue injury site), almost everywhere in the body. The intensity of the pain caused by the OIH will be increased if additional opioids be administered. OIH can be managed by reducing opioid doses, opioid rotation, or anti-N-methyl-D-aspartate (NMDA) receptor medications such as ketamine (6). However, abrupt cessation of the remifentanil infusion may cause acute withdrawal syndrome. Several mechanisms contribute to the appearance of OIH. These mechanisms include central sensitization due to suppressed reuptake or the increased re-
Makarem J et al.

Table 1. Baseline Characteristics of the Patients

|                     | Remifentanil-Naloxone | Remifentanil | Placebo | P-Value |
|---------------------|------------------------|--------------|---------|---------|
| Age (y)             | 50.56 ± 7.0            | 50.32 ± 6.6  | 50.24 ± 5.8 | 0.7     |
| BMI (kg/m²)         | 25.37 ± 4.1            | 26.04 ± 3.4  | 26.90 ± 4.4 | 0.4     |
| Surgery duration (min) | 122 ± 18              | 130 ± 24    | 128 ± 23  | 0.4     |
| Anesthesia duration (min) | 138 ± 19            | 146 ± 26    | 143 ± 25  | 0.5     |
| ASA II class (%)    | 9 (36)                 | 8 (32)      | 6 (24)   | 0.6     |
| Underlying diseases (%) | 9 (36)               | 8 (32)      | 6 (24)   | 0.6     |

Abbreviations: ASA, American Society of Anesthesiologists; BMI, body mass index.

Values are expressed as mean ± SD unless otherwise indicated.

Table 2. Comparison of Heart Rate and Mean Arterial Pressure Between Three Groups

|                     | Remifentanil-Naloxone | Remifentanil | Placebo | P-Value |
|---------------------|------------------------|--------------|---------|---------|
| HR at baseline      | 85.1 ± 9.4             | 82.1 ± 13.1  | 81.4 ± 13.1 | 0.89    |
| HR after intubation | 91.9 ± 9.4             | 95.1 ± 14.5  | 98.6 ± 14.8 | 0.25    |
| HR at 30 mins after intubation | 60.6 ± 11.1          | 59.1 ± 9.5   | 87.2 ± 12.3 | 0.001   |
| HR after extubation | 75.2 ± 9.4             | 72.5 ± 9     | 94.6 ± 11.9 | 0.001   |
| MAP at baseline (mmHg) | 79.5 ± 6.7           | 79.1 ± 6.5   | 78.1 ± 6.2  | 0.83    |
| MAP after intubation (mmHg) | 94.3 ± 6.1           | 93.5 ± 6.9   | 92.8 ± 5.7  | 0.74    |
| MAP at 30 mins after intubation (mmHg) | 73.9 ± 6.2         | 72.7 ± 5.5   | 71.3 ± 5.6  | 0.65    |
| MAP after extubation (mmHg) | 84.6 ± 6.6           | 83.1 ± 6.2   | 84.7 ± 6.8  | 0.55    |

Abbreviations: HR, heart rate; MAP, mean arterial pressure.

Values are expressed as mean ± SD.

One-way ANOVA.

Table 3. Comparison of Opioids Administered Intra- and Post-Operatively and Time of the First Dose of Rescue Analgesia in Recovery Between Three Groups

|                     | Remifentanil-Naloxone | Remifentanil | Placebo | P-Value |
|---------------------|------------------------|--------------|---------|---------|
| Cumulative intraoperative fentanyl (µg) | 76 ± 38                | 90 ± 35      | 160 ± 47  | 0.001   |
| Time of first dose of rescue analgesia in recovery (min) | 41 ± 6                 | 26 ± 4       | 33 ± 4    | 0.001   |
| Rescue morphine sulphate (mg) | 8.2 ± 3.5              | 12.1 ± 4.7   | 9.1 ± 3.3  | 0.002   |
| 24-hours postoperative morphine sulphate (mg) | 27.7 ± 5.1             | 39.1 ± 7.8   | 36.1 ± 2.8 | 0.001   |

Values are expressed as Mean ± SD unless otherwise indicated.

One-way ANOVA.

A review by Kim et al. (2014) reported that clinical studies support the development of OIH in healthy subjects and patients after administration of short-acting opioids, such as remifentanil, with an infusion rate of > 0.1 µg/kg/min. The findings also revealed that remifentanil infusion with a dose of 0.3 µg/kg/min leads to OIH.

In the present study, hyperalgesia was confirmed by the presence of the following signs: development of allodynia (assessed by brush test with more frequent positive results), the need for rescue analgesia in PACU (both required dose and the time of need to the first dose), total 24-hours postoperative morphine dose used, and the severity of pain (measured using the VAS). In the present study, patients in the remifentanil-naloxone group showed less severe pain and had lower analgesic requirements in PACU and 24 hours after surgery compared to the remifentanil and placebo groups. Intraoperative fentanyl requirement...
The present study also showed that post-surgery shivering that naloxone contains anti-shivering properties (24, 25). The main limitation of the present study was not using appropriate devices to assess allodynia and hyperalgesia (e.g., von Frey hair kit), which may have affected the precision of hyperalgesia and allodynia evaluation. Moreover, we could not differentiate OIH from acute tolerance. Therefore, the improved analgesic effect of naloxone in this study can be attributed to two issues: low dose naloxone has an analgesic effect, or it can reduce the OIH.

5.1. Conclusion

This study demonstrated the efficacy of intraoperative low-dose naloxone (0.25 µg/kg/h) added to remifentanil infusion on reducing the frequency and severity of acute postoperative hyperalgesia in patients undergoing general anesthesia for laparotomic hysterectomy.

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The Comparison of Static Tactile Test Results Between Three Groups

|                      | Remifentanil-Naloxone, No. (%) | Remifentanil, No. (%) | Control, No. (%) | P-Value * |
|----------------------|---------------------------------|-----------------------|-----------------|-----------|
| Positive test after 30 minutes | 4 (16)                          | 13 (52)               | 8 (32)          | 0.02      |
| Positive test after 2 hours       | 3 (12)                          | 10 (40)               | 4 (16)          | 0.03      |
| Positive test after 6 hours        | 2 (8)                           | 7 (28)                | 1 (4)           | 0.03      |
| Positive test after 12 hours       | 2 (8)                           | 5 (20)                | 0               | 0.06      |
| Positive test after 24 hours       | 0                               | 2 (8)                 | 0               | 0.32      |

*One way ANOVA.

Footnotes

Authors' Contribution: Study concept and design: J.M., F.B and S.M.M.; analysis and interpretation of data: F.A., A.J. and F.Y; drafting of the manuscript: F.B., F.A; critical revision of the manuscript for important intellectual content: K.K., S.S and S.M.M; statistical analysis: J.M. and H.M.

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References

1. Chapman CR, Lipschitz DL, Angst MS, Chou R, Denisco RC, Donaldson GW, et al. Opioid pharmacotherapy for chronic non-cancer pain in the United States: a research guideline for developing an evidence-base. J Pain. 2010;11(9):807-29. doi: 10.1016/j.jpain.2010.02.018. [PubMed: 20430701].

2. Okie S. A flood of opioids, a rising tide of deaths. N Engl J Med. 2010;363(21):1981-2. doi: 10.1056/NEJMp1011512. [PubMed: 21083382].

3. Trescot AM, Helm S, Hansen H, Benyamin R, Glaser SE, Adlaka R, et al. Opioids in the management of chronic non-cancer pain: an update of American Society of the Interventional Pain Physicians' (ASIPP) Guidelines. Pain Physician. 2008;11(2 Suppl):S5-S62. [PubMed: 18443640].

4. Benyamin R, Trescot AM, Datta S, Buenaventura R, Adlaka R, Sehgal N, et al. Opioid complications and side effects. Pain Physician. 2008;11(2 Suppl):S305-20. [PubMed: 18443635].

5. Pud D, Cohen D, Lawental E, Eisenberg E. Opioids and abnormal pain perception: New evidence from a study of chronic opioid addicts and healthy subjects. Drug Alcohol Depend. 2006;82(3):228-33. doi: 10.1016/j.drugalcdep.2005.09.007. [PubMed: 16229972].

6. Colvin LA, Bull F, Hales TG. Perioperative opioid analgesia-when is enough too much? A review of opioid-induced tolerance and hyperalgesia. Lancet. 2009;373(9980):1558-68. doi: 10.1016/S0140-6736(09)60430-1. [PubMed: 19838591].

7. Velazuddan A, Bellingham G, Morley-Forster P. Opioid-induced hyperalgesia. Curr Opin Pharmacol. 2014;14(3):125-9. doi: 10.1016/j.cophas.2014.01.005.

8. Kim SH, Stoica N, Soghomonyan S, Bergese SD. Intraoperative use of remifentanil and opioid induced hyperalgesias:acute opioid tolerance: systematic review. Front Pharmacol. 2014;5:208. doi: 10.3389/fphar.2014.00208. [PubMed: 24847273]. [PubMed Central: PMC420143].

9. Koo CH, Yoon S, Kim BR, Cho YJ, Kim TK, Jeon Y, et al. Intraoperative naloxone reduces remifentanil-induced postoperative hyperalgesia but not pain: a randomized controlled trial. Br J Anaesth. 2017;119(6):861-8. doi: 10.1016/j.bja.2015.12.025. [PubMed: 29029049].

10. Buchsbaum MS, Davis GC, Bunney WJ. Naloxone alters pain perception and somatosensory evoked potentials in normal subjects. Nature. 1977;270(5638):620-2. doi: 10.1038/270620a0. [PubMed: 31910].

11. Movafegh A, Shoebi G, Ansari M, Sadeghi M, Azimarahgi O, Aghajani Y. Naloxone infusion and post-hysterectomy morphone consumption: a double-blind, placebo-controlled study. Acta Anesthesiol Scand. 2012;56(10):1241-9. doi: 10.1111/j.1399-6576.2012.02764.x. [PubMed: 22946762].

12. Przewlocki R, Przewlocka B. Opioids in chronic pain. Eur J Pharmacol. 2001;429(1-3):79-91. doi: 10.1016/s0014-2999(00)01308-6. [PubMed: 1198029].

13. Bleakman D, Alt A, Nisenbaum ES. Glutamate receptors and pain. Semin Cell Dev Biol. 2006;17(5):592-604. doi: 10.1016/j.semcdb.2006.10.008. [PubMed: 1710195].

14. Kim AH, Kerchner GA, Choi DW. Blocking Excitotoxicity. In: Marcoux Fw, Choi Dw, editors. CNS Neuroprotection. 155. New York: Springer; 2002. p. 3-36. doi: 10.1007/978-3-662-06274-6_1.

15. Griffith OW, Kilbourn RG. Nitric oxide synthase inhibitors: amino acids. Methods Enzymol. 1998;284(1):375-92. doi: 10.1016/S0076-6879(98)60840-9. [PubMed: 8782604].

16. Gang TJ, Ginsberg B, Glass PS, Fortney J, Jhaveri R, Perno R. Opioid-sparing effects of a low-dose infusion of naloxone in patient-administered morphine sulfate. Anesthesiology. 1997;87(5):705-11. doi: 10.1097/00000542-199705000-00008. [PubMed: 9364592].

17. Tilson HA, Rech RH, Stolman S. Hyperalgesia during withdrawal as a means of measuring the degree of dependence in morphine dependent rats. Psychopharmacology. 1973;28(3):287-300. doi: 10.1007/BF00429309. [PubMed: 4734481].

18. Jacquet YF, Latjha A. Morphine action at central nervous system sites in rat: analgesia or hyperalgesia depending on site and dose. Science. 1973;182(4111):490-2. doi: 10.1126/science.182.4111.490. [PubMed: 4582903].

19. Chu LF, Angst MS, Clark D. Opioid-induced hyperalgesia in humans: molecular mechanisms and clinical considerations. Clin J Pain. 2008;24(6):479-96. doi: 10.1097/AJP.0b013e3181bb2f43. [PubMed: 18574534].
20. Compton P, Charuvastra VC, Ling W. Pain intolerance in opioid-maintained former opiate addicts: effect of long-acting maintenance agent. Drug Alcohol Depend. 2001;63(2):139–46. doi: 10.1016/s0376-8716(00)00200-1. [PubMed: 11376918].

21. Vanderah TW, Ossipov MH, Lai J, Malan TJ, Porreca F. Mechanisms of opioid-induced pain and antinociceptive tolerance: descending facilitation and spinal dynorphin. Pain. 2001;92(1-2):5–9. doi: 10.1016/s0304-3959(01)00311-6. [PubMed: 11323121].

22. Angst MS, Clark JD. Opioid-induced hyperalgesia: a qualitative systematic review. Anesthesiology. 2006;104(3):570–87. doi: 10.1097/00000542-200603000-00025. [PubMed: 16508405].

23. Hoshijima H, Takeuchi R, Kuratani N, Nishizawa S, Denawa Y, Shiga T, et al. Incidence of postoperative shivering comparing remifentanil with other opioids: a meta-analysis. J Clin Anesth. 2018;32:300-12. doi: 10.1016/j.jclinane.2015.08.017. [PubMed: 26432635].

24. Alfonsi P. Postanaesthetic shivering: epidemiology, pathophysiology, and approaches to prevention and management. Drugs. 2001;61(15):2193-205. doi: 10.2165/00003495-200161150-00004. [PubMed: 11722130].

25. Kurz M, Belani KG, Sessler DI, Kurz A, Larson MD, Schroeder M, et al. Naloxone, meperidine, and shivering. Anesthesiology. 1993;79(6):1193-201. doi: 10.1097/00000542-199312000-00009. [PubMed: 8267194].

26. May LM, Kosek P, Zeidan F, Berkman ET. Enhancement of meditation analgesia by opioid antagonist in experienced meditators. Psychosom Med. 2018;80(9):807-11. doi: 10.1097/PSY.0000000000000580. [PubMed: 29355707]. [PubMed Central: PMC6462167].