

Whether Does Acupressure (p6) Prevent Nausea and Vomiting in Patients Undergoing Laparoscopic Surgery

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ABSTRACT

Objective: To investigate whether acupressure (p6) prevents nausea and vomiting in patients undergoing laparoscopic surgery and to compare effect and placebo effect of acupressure (P6) in preventing nausea and vomiting in patients who have undergone laparoscopic surgery.

Methods: This was double blind randomized study in which sixty patients were enrolled. They were divided into three equal groups. First group received acupressure with bilateral stimulation of P6 (A), Second group received bilateral placebo stimulation (P) and a third group received no acupressure stimulation and served as a reference group (R). We observed how many patients developed nausea vomiting and needed rescue antiemetics in each group.

Results: It is clear from the results that 9,7 and 6 patients in group A,P and R respectively develop nausea only while 1, 1 and 8 patients had nausea (8 Vs 1 with $P < 0.05$) 24 h after surgery in A,P and R groups respectively. When compared to placebo acupressure (2 patients vomited and 5 needed rescue medication) significantly ($P < 0.05$) fewer needed rescue antiemetic medication after acupressure at P6 (no vomiting and rescue medication) When compared to the reference group (5 vomited and 4 needed rescue medication) significantly fewer vomited after acupressure ($P < 0.05$).

Conclusion: It has been concluded from our study that acupressure (p6) has significantly reduced vomiting and need of rescue medication in patients undergoing laparoscopic surgeries. While placebo effect of acupressure decreased nausea 24 h after surgery.

Keywords: Nausea, vomiting, acupressure, antiemetics, laparoscopic surgery

INTRODUCTION

Postoperative nausea and vomiting (PONV) are common in patients undergoing laparoscopic surgeries and the main symptoms which delay discharge from recovery. Even though the most efficient pharmacological treatments decrease the incidence of PONV by about 50%^{1,2} but are not without their adverse effects and their cost benefit can also be questioned³. Acupuncture towards the P6 (Neiguan) point, which is located three fingers breadth proximal to the proximal flexor palmar crease between the tendons of flexor carpi radialis and palmaris longus⁴, has been shown to be effective in decreasing the incidence of nausea and vomiting to about the same extent as pharmacological treatments⁵. Acupressure is a method where pressure is applied to acupuncture points. A finger, a round stick or a pea sized pearl is used. Acupressure towards P6 has been shown to be effective in several recent studies, ⁶⁻¹⁰ including studies from Great Britain in the 1990s^{11,12}. No adverse effects have been reported. Acupressure has not reached widespread use and is not even mentioned in recent

review of the subject. Because the placebo effect of a treatment like acupressure is probably considerable¹³. We use a design in which true effect and placebo effect could be measured. For placebo stimulation we used a sensory stimulation of a non-acupressure point of similar intensity as the active acupressure stimulation¹⁰ and a control group to measure the placebo effect. Our aim was to investigate the effect and placebo effect of acupressure on the incidence of nausea and vomiting after laparoscopic surgeries.

MATERIAL & METHODS

The study was conducted at Bhatti International Hospital, Kasur which is attached to Central park medical college. After obtaining approval from the institution from Nov. 2011 to July 2012, sixty patients aged (40-60 years) ASA I and II, who were scheduled for laparoscopic surgery were enrolled in the study.

The study design was randomized double blind. Patients with history of severe adverse reactions to NSAIDs, bronchial asthma, kidney or liver dysfunction, bleeding disorders or history of steroids intake within 24h of surgery were excluded. All patients were divided in to three equal groups. First group (A) received active treatment (n=20), second group (P) received placebo treatment (n=20) and a

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third group (R) was used as a reference group (n=20). Anesthesia protocol was similar for all patients. They were given midazolam 0.1 mg/kg as oral premedication two hours prior to induction. After arrival in operating theatre an intravenous catheter was placed, infusion of crystalloid D5 +0.9% saline started and standard monitoring was established.

Anesthesia was induced with following drugs.

1. Inj. Nalbuphin 0.1mg/kg
2. Inj. Propofol 2.5mg/kg
3. Inj. Atracurium 0.5mg/kg followed by 0.1mg/kg increments as necessary.
4. Oxygen, Nitrous oxide and sevoflurane to maintain anesthesia via tracheal tube and mechanical ventilation.

Group A received acupressure at P6 (Neiguan), a point located on the pericardial meridian, which is found three fingers breadth (app 5cm) proximal to the proximal flexor palmar crease, about 1cm deep, between the tendons of flexor carpi radialis and Palmaris longus, is supposed to have an effect on post operative nausea and vomiting⁴. A Sea-Band carries a plastic pearl which is fastened to apply pressure on P6. Both forearms were used. These points were marked with water-resistant ink so that the band could be properly replaced if removed. The areas were draped with a dressing during the stay in the hospital. Group (P) was given placebo acupressure. A point on the dorsal side of both forearms, four finger breadth proximal to the proximal flexor palmar crease was used for placebo stimulation. These points were marked in the same way as with the active acupressure. Sea-Band was used for stimulation, and the same precautions were taken to keep the stimulation blinded. The doctors and nurses giving anesthesia and the nurses on the postoperative ward, although aware that stimulation was being performed were not aware of the location of P6. Group (R) received neither true acupressure nor placebo acupressure. But anesthesia protocol, instructions for post operative care and assessment were the same. Nausea was estimated by the patient on a horizontal visual analogue scale (VAS), 100mm. The end point was assigned "no nausea" to the left and "worst possible nausea" to the right. The patients were asked to assess their degree of nausea 30, 60, and 120 min after arriving at the postoperative ward. Metoclopramide 10 mg was administered i.v at the patient's request. Vomiting was noted by the nurses as was the need for antiemetic. After discharge the patients were asked to assess the degree of nausea at 6 pm, when going to bed, at breakfast time and at noon the day after surgery. They were also asked to note vomiting. A scoring on the VAS over 10 mm was classified as nausea and scoring below 10mm was

classified as no nausea. Nausea score 24h after surgery was used as the patients evaluation of the treatment. Pain was assessed by the patient on a horizontal VAS 100 mm. The endpoints assigned were "no pain" to the left and "worst possible pain" to the right. Morphine was used in the postoperative ward in doses of 2mg i.v. If additional analgesic was needed (Table 1). The patients received Paracetamol 1 gram six hourly orally for 24 hours after discharge from recovery when oral sips allowed. Demographic data are given as median (range). Kruskal-Wallis test was used to test for difference between demographic data. Comparison of treatment effects was performed with Fishers exact test using the Ciba-Geigy table. The outcome was number of patients experiencing nausea (only), vomiting need for rescue medication and nausea after 24 h. A P- value of <0.05 was considered to be significant.

RESULTS

Demographic data and factors prognostic for PONV are given in Table 1 and results as number of patients are given in table 2. No patients (0%) in the acupressure group vomited, but 2 patients (10%) in the placebo acupressure and 5 patient (25%) in reference group (P< 0.05, vs. acupressure) vomited. No patient (0%) in the acupressure group requested rescue medication, but 5 (25%) patients in the placebo group (P< 0.05 vs. acupressure) and 4 patient (20%) in the reference group requested for rescue medication. There was no difference in the postoperative need for morphine between the groups (Table 1). Twenty-four hours after surgery, one patient (5%) in the acupressure group and one patient (5%) in the placebo group reported nausea, while 8 patients (40%) in reference group developed nausea.

DISCUSSION

An effect of acupressure^{6,7,9} as well as of acupuncture⁵ on nausea and vomiting after gynecological surgery has been reported before. But in our study, we observed the effect of acupressure on nausea and vomiting after laparoscopic surgeries (cholecystectomy, appendectomy, herniorrhaphy). The mechanism of action of acupuncture and acupressure on nausea and vomiting has not been established and a placebo-type mechanism has been suggested¹⁰. Autonomic dysfunction seems to have a correlation to nausea. Considering the multifactorial etiology of PONV, it is unlikely that a single drug or treatment could counteract all causative factors. Different types of surgeries could have different profiles with reference to etiological factors. Most

studies on acupuncture and acupressure have investigated gynecological patients. Methodological considerations that have to be evaluated are whether a bilateral stimulation is superior to a unilateral stimulation and whether the timing is important. An effect of timing on acupressure has been suggested by two studies one reporting no effect of acupressure with start of stimulation after opioid medication and one reporting an effect when stimulation was started before opioid medication¹². How to measure nausea and vomiting has been debated during the last decade as nausea and vomiting have been identified as a major quality factor from the patient's point of view. Nausea can be assessed by the nurse or the patient. A poor correlation between the assessments has been demonstrated. The assessment can be performed by using a nausea questionnaire, verbal categorical scales or visual analogue scale. Similar results have been obtained with these three methods. We have used nausea (only), vomiting, need for antiemetic, and nausea after 24h as outcome measures. That is part of the outcome measurement is dependent on nurse assessment, and part on the patients assessment with a visual analogue scale. To further clarify the usefulness of acupressure on PONV we propose a study group with a higher incidence of PONV but similar design as in this study so that the placebo effect could be measured. If the patients were not only randomized but also stratified according to age sex history of postoperative nausea or vomiting or motion sickness, additional strength would be added.

Table 1: Demographic variables of the three groups of patients

	Reference group (n=20)	Acupressure (n=20)	Placebo (n=20)
Weight (Kg)	64(51-91)	60 (53-81)	63(49-85)
Height (cm)	167 (155-182)	168 (150-175)	163 (153-175)
Age (years)	52	50	51
Previous PONV or motion sickness (n)	7	8	7
Patient given morphine in postop ward	8	7	7
Morphine (mg) given in postop ward	0 (0-4)	0 (0-8)	0 (0-5)

Table: 2 Results in nausea (only), vomiting, rescue medication and nausea after 24hr value as no. of patients

	Reference group(n=20)	Acupressure (n=20)	Placebo (n=20)
Nausea (only)	6	9	7
Vomiting	5	0	2
Rescue medication	4	0	5
Nausea after 24 hr of operation	8	1	

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