



Keywords: TAP blocks; Postoperative analgesia; Regional Anesthesia; Literature to evaluate the combinations of approaches. O'Hara et al. demonstrated that when morphine was added to bupivacaine, when compared to bupivacaine alone, there was reduced total morphine consumption as well as prolonged time to request for analgesics. Also, there was a decrease in VAS for pain in the abdominal wall. By utilizing anatomic landmarks to determine the insertion site within the lumbar triangle of Petit, a single "pop" sensation served as an endpoint for appropriate needle depth to provide analgesia to the parietal peritoneum as well as the skin and muscles of the anterior abdominal wall [1]. Later, O'Donnell described a "double pop" technique, which resulted from the blunt needle passing through the fascial plane and transverse abdominis muscle where the nerves are located.

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satisfaction score, requirement of analgesics in the 24 h, and the side effects of clonidine (sedation, dryness of mouth, hypotension, and bradycardia) were observed. Duration of analgesia was longer in the second group receiving bupivacaine and clonidine compared to the group receiving just bupivacaine (P<0.001). Additionally the combination with clonidine prolonged analgesia by 10-12 h and reduced overall postoperative analgesic requirements. For those who received the combination with clonidine, none of the patients experienced hypotension or bradycardia [8].

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Y of bupivacaine and dexmedetomidine added to bupivacaine used in TAP block by US guidance on postoperative pain was evaluated by Aksu et al. Sixty three patient enrolled, divided in 3 groups. Group C (Control) had TAP block with 21 mL 0.9% saline, Group B had 20 mL 0.5% bupivacaine+ 1 mL saline, Group BD and 20 mL 0.5% bupivacaine+1 mL 100 µg dexmedetomidine. Results demonstrated that with the addition of dexmedetomidine, the treatment of postoperative pain compared to control group and group B was VAS was lower at 10-24 hours, morphine consumption was lower and higher patient satisfaction for group BD. No was found on nausea and vomiting score, neither in the requirement of antiemetics [9].

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In a double blind study by Manjaree Mishra et al. the of dexmedetomidina with ropivacaine was studied using 40 patients. Patients were divided into two groups. Group R (n=20) where patients received bilateral TAP block using 20ml of ropivacaine 0.2% and 2 ml of normal saline, and Group RD (n=20) who received dexmedetomidine 0.5 mcg/kg dissolved in 2 ml of normal saline and added to 20 ml of ropivacaine 0.2%. Blocks were performed in the operating room intubation. Age and gender were taken in consideration in this study, showing no between those variables, as well as nausea and vomiting results of this study showed that the group receiving combination of ropivacaine and dexmedetomidine has lower pain scores postoperatively

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