INTRODUCTION

According to international association for study of pain, it is defined as unpleasant emotional and sensory experience due to actual or potential tissue damage.[1] In postoperative period following abdominal surgeries majority of the patients complain of varying degrees of pain as documented in various studies conducted in postoperative period if pain is not managed well it may lead to various complications such as deep vein thrombosis, hypertension, myocardial ischemia, cardiac arrhythmias, respiratory distress & poor wound healing.[2,3,6,7,8] Combination of simple NSAIDS and opioids remain suitable postoperative analgesics alternative to epidural and USG guided blocks in a minimal resource set up.[9,10,11] The multimodal therapy for analgesia is recommended in the management of postoperative pain as per recent guidelines which improves the recovery process.[12,13,14,15]

Pre-emptive analgesia is a method of administering analgesics prior to surgery which acts by blocking central nervous system hyper excitability and thereby reducing the postoperative pain. Paracetamol, Diclofenac and Tramadol were assessed in combination in our study as they are easily accessible in the market.

Hence we took this study to compare postoperative analgesia using combination of paracetamol-diclofenac and paracetamol-tramadol as a pre-emptive analgesia.

MATERIALS AND METHODS

This study was undertaken at Raichur Institute of Medical Sciences teaching Hospital and RGSSH Hospital, Raichur from 1st July 2021 to 28th February 2022. A randomised controlled clinical trial was conducted at Raichur Institute of Medical Sciences.
0.5 mg was given orally to both the groups. Prior to surgery (60 mins before surgery) 1gmiv (intravenous infusion) Paracetamol and 75mg inj. Diclofenac IM (intramuscular) were administered to the patients belonging to group A whereas for Group B patients, inj. Diclofenac 75 mg intramuscular (IM) and tramadol 100mg intravenous (IV) was given to the patients allocated randomly.

In preoperative room baseline vital signs were recorded & pre medication drugs of Inj. Ondansetron 4 mg iv, Inj. Midazolam 1mg iv and Inj. Glycopyrolate 0.2 mg iv was given. In the operative room standard monitors were connected which included, pulse-oximeter, electrocardiography, non-invasive blood pressure. Following 3 minutes of pre oxygenation general anaesthesia was induced by Inj. propofol 2.5mg /kg, Inj. Fentanyl (2mg/kg) and inj. succinylcholine (2mg/kg) followed by endotracheal intubation. Isoflurane with oxygen and nitrous oxide mixture was used to maintain the anaesthesia. Inj. Vecuronium was given for neuromuscular blockage after completion of surgery & it was reversed with inj. myopyrolate (inj. glycopyrolate 0.01mg/kg and neostigmine 0.05mg/kg) after surgery. Patients were monitored in surgical ICU post operatively for postoperative pain. The rate of pain intensity according to standard numeric rating scale ranging from 0 to 10 was recorded & it was further graded as mild, moderate, severe depending on NRS scale 0-3(mild), 4-6(moderate) and NRS: 7-10 (Severe)respectively.

Till 12th hour every 2hrly, and later at 24th Hour NRS Score were recorded at recovery. Consumption of analgesia in 24hrs postoperatively, request by the patient for first analgesia and analgesia duration was documented.

Results

The statistics were analysed using SPSS V 23 software. Analysis of variants (ANOVA) was used for normally distributed continues data. Chi-squared Test was used to analyse categorical data. Statistical analysis showing p value of <0.05 is significant.

Median NRS score were not significant at 2nd ,12th & 24 hrs (p>0.05) between 2 groups during coughing as per Kruskal walls test. At the 4,6th and 8th hour between groups statistically significant difference was noted (p=0.009 and 0.006). During 24th (p>0.06) all the time between the two groups statistical difference was not found. However, paracetamol – diclofenac NRS scores were higher than paracetamol – tramadol at all time.

First analgesia requirement time in the paracetamol – diclofenac group A (M ± SD:110.01 ±15.02min) was significantly shorter than in the paracetamol-tramadol group B (150.01 ± 10.53min p=0.030), similarly analgesia requirement for the first time in the paracetamol – diclofenac was significantly higher than paracetamol – tramadol group (p<0.001) [Table 2].

Within 24hrs the mean total tramadol consumption between the groups was statistically significant as shown in the [Table 2]. It was observed that total tramadol consumed in 24h shown significantly higher in group A (p -0.007) compared to group B (p-0.02) that is paracetamol & tramadol. Group B showed less significant statistically compared to group A.
Nausea and vomiting were lower (15%) in Group A compared to group B (20%) which was not statistically significant (p=0.6) as well as no serious complications observed in either group.

DISCUSSION

Over 24 hr the total tramadol consumption was less in paracetamol – tramadol compared to paracetamol – diclofenac combination in our study. The mean total tramadol consumption was 160.76 ± 30.53mg in paracetamol – tramadol group as compared to 190.81 ± 80.49mg in paracetamol diclofenac group with p value of 0.001.

<table>
<thead>
<tr>
<th>First dose analgesic requirement time (mins)</th>
<th>Group A</th>
<th>Group B</th>
<th>P value</th>
</tr>
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<tbody>
<tr>
<td>110.01 +/- 15.40</td>
<td>150.05 +/- 10.52</td>
<td>&lt;0.001</td>
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<table>
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<tr>
<th>Total analgesic consumption of Tramadol in mg(IV)</th>
<th>Group A</th>
<th>Group B</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>190.21 +/- 80.29</td>
<td>160.76 +/- 30.54</td>
<td>&lt;0.001</td>
<td></td>
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</tbody>
</table>

Data: mean +/- SD

In our study first dose of analgesia mean time was shorter in paracetamol - diclofenac group (M ± SD:110.62±15.95min) compared to paracetamol – tramadol group (150.01±10.53 min, p=0.001), Similar observation were made in the study conducted by Samimi et al (p<0.05).

In patients who received the paracetamol-diclofenac combination in the first 24 hrs had markedly less mean total morphine consumed compared to diclofenac group (20.1 ± 3.6 mg) with p<0.05 was noted in a study done by Samimi et al.

In the present study the weakest opioid (tramadol) was used for post-operative pain management in order to compare our result with the study done by Samimi et al using opioid conversion factor tramadol to morphine (0.1). An estimated mean of 17.4 mg morphine was consumed in paracetamol-diclofenac group in our study.

In Monotgomery et al study, it was shown that total morphine consumed postoperatively in paracetamol-diclofenac group (18.5–35.8) (mean, 95% CI) was less compared to paracetamol group (36.1–53.6) with p<0.01. Monotgomery et al showed that the use of paracetamol-diclofenac combination is known to reduce the consumption of morphine by 1/3rd in comparison to paracetamol group alone.

Similar results were observed in a study done by Moussa and Riad, which showed total morphine consumed by patients in paracetamol-diclofenac group was significantly lower than paracetamol group alone

To limit the nervous system sensitisation by noxious stimuli multi modal analgesia holds a greatest promise. Multimodal analgesia is combining many modalities of analgesia into a single analgesic regimen. Multilevel interruption to the nociceptive inputs as well as to enhance the pain threshold & to decrease the activation of nociceptor can be done by using effective pre-emptive analgesic techniques. Through blockade of nociceptive stimuli at different site before tissue damage is explained by decreasing the excitability in central nervous system which further leads to decrease in pain score especially in combination groups.

CONCLUSION

In elective abdominal surgeries the first analgesic dose request for paracetamol – tramadol combination has increased time interval compared to paracetamol-diclofenac, hence paracetamol - tramadol combination has better postoperative analgesia in comparison to combination of paracetamol – diclofenac.

REFERENCES


