# Original Research Article

#### **TITLE**

Efficacy of Piroxicam and Tramadol as a Pre-Emptive Analgesic Agent for Mandibular Third Molar Surgery

#### **ABSTRACT**

**Objectives:** The aim of the present clinical trial study is to compare the pre-emptive analysesic effectiveness of 20 mg of piroxicam and 50 mg of tramadol for mandibular third molar surgery.

Methods: This prospective study included 30 patients who had been referred to the Department of Oral and Maxillofacial Surgery, Chennai for surgical removal of impacted mandibular third molars of similar difficulty index under local anesthesia. The patients were randomly assigned to 2 groups: Group A was administered 20 mg of piroxicam intramuscularly (IM) 50 minutes before the surgery and Group B was given 50 mg of tramadol IM 50 minutes before the surgery. The time to analgesic re-medication, Pain intensity (VAS Scores) at 1st, 2nd, 12th, 24th hour, total analgesic consumption was evaluated.

**Results:** The group receiving 20 mg of piroxicam IM showed differences in pain intensity evaluated by the visual analogue scale and total analgesic consumption [Lesser values] when compared with the group receiving 50 mg of tramadol IM and the results were statistically significant (p<0.05). However, the time to first rescue analgesic medication, number of patients requiring the rescue analgesic procedure (10 mg of oral ketorolac), and number of patients without the need for analgesic during the period of evaluation did not show statistically significant differences between the two groups (P > 0.05).

**Conclusion:** Within the limitations of the present study, the patients receiving 20 mg of preoperative piroxicam had less pain intensity and total analgesic consumption than those receiving 50 mg of preoperative tramadol. Therefore, piroxicam given preoperatively showed

superior analgesic properties for intermediate surgical procedures in comparison to preemptively administered tramadol.

**Keywords:** third molar; pre-emptive analgesia; impacted, mandibular molar; postsurgical pain; piroxicam; tramadol

#### INTRODUCTION

Removal of the impacted third molar teeth is one of the most common oral surgical procedures performed in dentistry and invariably gives rise to a number of postoperative sequelae, among which pain is almost always present [1]. The onset of pain usually begins as the effects of the local anesthetic agent subside. The concept of pre-emptive analgesia involves the administration of analgesic before a painful stimulus is initiated. It consists of antinociceptive treatment that prevents central neural sensitization which amplifies postoperative pain [2]. Analgesics given before surgical trauma are thought to have a pre-emptive effect; implying that analgesia will start before the surgical stimulus, leading to a reduction of CNS input and, hence, reducing pain [2].

Tramadol is an opioid analgesic clinically effective in treating moderate to moderately severe pain, and it has a low addiction potential. In acute therapeutic use, it produces analgesia against multiple pain conditions, including postsurgical pain, obstetric pain, terminal cancer pain, and pain of coronary origin. The analgesic acts at the opioid receptors and appears to modify the transmission of pain impulses by inhibiting monoamine reuptake [3]. Several nonsteroidal anti-inflammatory drugs (NSAIDs) have been used for pain, swelling, and trismus control after mandibular third molar surgery [4]. These medications achieve their therapeutic effect through the inhibition of cyclooxygenase (COX), which determines the inhibition of prostaglandin (PG) production [5].

Piroxicam is a NSAID of the acidic enolic class that preferentially inhibits the inducible Cox-2 enzyme and shows a weaker influence on the constitutive Cox-1 enzyme [6, 7]. Thus, it is largely used for the treatment of acute and chronic pain and inflammatory and degenerative disorders [7]. There also has shown to be an antinociceptive synergism between intraperitoneal piroxicam and morphine [8]. Postoperative analgesia comparable with that of opioids has been demonstrated with the non-steroidal anti-inflammatory drugs (NSAIDS) [9, 10]. An opioid sparing effect has also been observed with NSAIDS, as well as a reduction in opioid induced nausea, vomiting and respiratory depression. This reduction in opioid requirement and side-effects may benefit the patient by producing increased postoperative analgesia and, even, reduce hospital stay [11].

Previously our team has a rich experience in working on various research projects across multiple disciplines [12–26]. Now the growing trend in this area motivated us to pursue this project. Based on this inspiration, we aim to compare the pre-emptive analysesic effectiveness of 20 mg of piroxicam and 50 mg of tramadol for mandibular third molar surgery.

#### MATERIALS AND METHODS

# **Study Setup:**

This randomized prospective controlled clinical study was conducted among patients reporting to the outpatient dental department of oral surgery clinic during the period between June 2020- March 2021. The study population included 30 adult patients who were randomly selected and had been referred to the department of oral and maxillofacial surgery for surgical removal of impacted mandibular molar. The sample size was divided mainly into two groups, each with 15 patients, namely: Group A was administered 20 mg of piroxicam intramuscularly 50 minutes before the surgery and Group B was administered 50 mg of tramadol intramuscularly 50 minutes before the surgery.

#### **Inclusion Criteria:**

- Patients between 18 years-50 years of age
- Both genders
- Clinical and radiographic diagnosis of a partially bony impacted mandibular third molar
- No pain associated with the subject third molar up to the day of surgery

# **Exclusion Criteria:**

- Patients with incomplete clinical and radiological records.
- Patients with severe systemic conditions like diabetes and hypertension.
- Use of analgesics 3 days before the procedure, history of seizure disorder, pregnancy or lactation, oral contraceptive use, and known hypersensitivity to the study medications.

#### **Procedure:**

All the surgical procedures were carried out at the Department of Oral and Maxillofacial Surgery by the same surgeon, and evaluations were made by a single independent investigator. Anesthesia was achieved through a block of the lingual, buccal, and inferior alveolar nerves by use of two 1.8-mL capsules of 2% lidocaine—containing 1:100,000 epinephrine. Once anesthesia was given, surgery was started. A mucoperiosteal flap was prepared by making an incision distal to the mandibular second molar along the anterior edge of the ascending ramus of the mandible. This flap was used to close the surgical wound. Suturing was done with No. 4-0 silk. In each patient, a partial bony impacted mandibular third molar was extracted. The time to analgesic re-medication was registered. The patients were given four 10-mg oral ketorolac pills and were instructed to take 1 pill as rescue medication at least 6 hours apart, according to their requirements. At the end of the evaluation period (24 hours), the patients returned the unused ketorolac. The pills were counted to determine the number of consumed pills, as well as the number of patients in each group who did not need any pills. The total analgesic consumption was also evaluated.

# Diagnostic Criteria

# Post Operative Pain Evaluation by Visual Analogue Scale:

A 100-mm visual analogue scale (VAS) was used to assess the pain. The VAS consisted of an interval scale ranging from 0, representing no pain or discomfort, to 100, representing maximum pain or discomfort. The VAS report was recorded each at 1st, 2nd, 12 hours after completion of surgery, and the last evaluation was done at 24 hours.

# **Study Parameters:**

The following data were extracted for the purpose of the study:

- Age of the patient
- Gender of the patient
- Post operative VAS pain Scores
- Time to analgesic re-medication (ie, the time from the end of the surgery until the intake of the first rescue analgesic medication became necessary for the patient)
- Number of patients in each group who did not need any pill
- Number of patients requiring the rescue analgesic procedure (10 mg of oral ketorolac)
- Total analgesic consumption

The subjects were divided into four age groups- Group 1: 11-20 years, Group 2: 21-30 years, Group 3: 31-40 years, Group 4: 41-50 years.

#### **Data Collection:**

The data related to the study parameters were obtained from among patients who reported to the Outpatient Department from June 2020- March 2021. Approval for the study was obtained from the Institutional Ethical Committee. All assessments were done by a single examiner and the findings were reviewed and recorded by two investigators. All the subjects were informed of the possible risks of oral surgery and experimental treatments, and they signed an institutionally approved written consent form.

#### **Statistical Analysis:**

The data was tabulated and analysed using IBM SPSS version 23.0 software. Descriptive statistics was expressed by frequency and percentage. Student's t-test was used to compare variables (Time to analgesic re-medication, number of patients in each group who did not need any pill, number of patients requiring the rescue analgesic procedure, total analgesic consumption) between Piroxicam and Tramadol Groups. The effects over time of the pre-emptive analgesics on pain intensity were evaluated by Mann-Whitney U-Test. The significance level was set at P<0.05 with a confidence interval of 95%.

#### **RESULTS**

A total of 30 patients participated in this study, with an overall 100% participation.

# **Age Distribution:**

The youngest and oldest patients were aged 18 and 50 years, respectively. The distribution of study subjects based on age revealed that most patients belonged to 31-40 years of age group (67.50%).

#### **Gender Distribution:**

The distribution of study subjects based on gender, over a ten-month period, revealed that 20 patients (75%) women and 10 patients (25%) men participated in this study.

# Post Operative Pain Evaluation by Visual Analogue Scale:

Pain score at 1st and 2nd hours after surgery was different between the two analysis groups; the mean VAS scores recorded after injection of piroxicam at 1 and 2 hours respectively were significantly lower than after tramadol at 1st and 2 hours, respectively. The pain intensity was also highest at the end of 2nd hour for the tramadol group (Figure 1). No significant

differences in pain score were observed between the two analysesics at 12 and 24 hour post-surgery (P>0.05) [Mann-Whitney U test].



Figure 1: Bar diagram depicting VAS scores of the piroxicam group (blue) and the tramadol group (orange) at the 1st, 2nd, 12th and 24th hour post-surgery. The X-Axis depicts the Post extraction hour and Y-Axis represents the VAS Scores. The VAS scores of the tramadol group were higher than the piroxicam group at the 2<sup>nd</sup> hour post-surgery.

# Time to first rescue analgesic medication, number of patients requiring rescue analgesic, number of patients without need of analgesic, total analgesic consumption:

The parameters: Time to first rescue analgesic medication, number of patients requiring the rescue analgesic procedure (10 mg of oral ketorolac), number of patients without the need for analgesic during the period of evaluation did not show significant statistical differences (P >.05). However, the difference between total analgesic consumption between the 2 groups was statistically significant (p=0.019) [Table 1].

Table 1 depicts the distribution of variables (time to first rescue analgesic medication, number of patients requiring the rescue analgesic procedure (10 mg of oral ketorolac), number of patients without the need for analgesic during the period of evaluation and total analgesic consumption) between Piroxicam Group and Tramadol Group.

Parameters	Piroxicam Group (Mean)	Tramadol Group (Mean)	Test Value	P Value
Time to first rescue analgesic (hr)	1.05	0.95	1.23	0.42
No. of patients (%) requiring rescue analgesic during period of evaluation (24hr)	6	3	5	0.12
No. of patients (%) not requiring analgesic during period of evaluation (24 hr)	1	6	-4	0.15
Total analgesic consumption (mg)	12.6	24.2	-34.3	0.019*

<sup>\*</sup>Statistically significant; Independent sample t test

On comparison of the parameters between the two groups, there was no statistically significant difference in time to first rescue analgesic medication (p=0.42), number of patients requiring the rescue analgesic procedure (10 mg of oral ketorolac) (p=0.12), number of patients without the need for analgesic during the period of evaluation (0.15). However, the difference between total analgesic consumption between the 2 groups was statistically significant (p=0.019).

#### DISCUSSION

Analgesia given immediately before surgical stimulus has been described as "pre-emptive analgesia". It prevents or reduces central hyperexcitability, leading to improved postoperative analgesia and reduced postoperative analgesic requirement [27]. Pre-emptive analgesia usage

is controversial in oral surgery, with reports in favour of it as well as against it [28]. Therefore, some guidelines and protocols have been developed to assess the quality of reports of randomized clinical trials in pain research. It has been reported that blind assessments produce significantly lower and more consistent scores than open assessments [29].

A meta-analysis by Ong et al [30] assessing the ability of pre-emptive analgesic interventions to attenuate and alleviate postoperative pain scores, decrease postoperative analgesic requirements, and prolong the time to first rescue analgesia showed an overall beneficial effect in selected analgesic regimens that was most pronounced after epidural analgesia, local wound infiltrations, and systemic NSAID administration. Recent studies by Richmond et al [31] using opioids have shown that preoperative morphine reduced pain scores and postoperative analgesic requirement in patients undergoing abdominal hysterectomy. Another study showed that administration of 30 mg of ketorolac IV produces better pre-emptive analgesic efficacy in comparison to 50 mg of tramadol IV preoperatively in third molar surgery [32].

A study by Isiordia et al [28] showed that patients receiving 15 mg of preoperative meloxicam had less pain intensity and total analgesic consumption than those receiving 50 mg of preoperative tramadol. All of these results were in accordance with the results of our study. However, a study performed by Nekoofar et al [7] showed no significant differences in the analgesic efficacy of meloxicam, piroxicam, and placebo but showed a significant effect of the time factor on reducing postoperative pain after endodontic treatment. In this study 50 mg of tramadol was administered because this dose has been used widely in the treatment of postoperative pain after third molar surgery and has been shown to be effective and safe [32–34]. Tramadol is an effective postoperative analgesic and can be used for a much longer time than morphine. Unfortunately, the widespread use of tramadol is hindered by its major adverse effects of nausea and vomiting [35]. In this study the major side effects were not evident because it was a single-dose study.

The main action mechanism of piroxicam is the inhibition of COX, which determines the inhibition of PG. The PGs are released from the damaged tissues and directly sensitize the peripheral nociceptors, and they also play a role in primary and secondary hyperalgesia and these are important in the modulation of pain [36]. The inhibition of peroxidase enzyme by piroxicam provides the advantage of a better gastrointestinal tolerance in comparison to other

NSAIDs. Moreover, piroxicam with its long half-life and when given preoperatively may provide a longer effect that is clinically relevant [37].

This is the first study comparing piroxicam with an opioid analgesic both as pre-emptive analgesics for pain control after third molar surgery, and few studies have reported on its analgesic efficacy in this acute pain clinical model in comparison to other NSAIDs [38–40]. These studies have shown that piroxicam can be a good alternative in pain treatment after the extraction of a mandibular third molar. It is possible that higher doses of tramadol (100 or 200 mg) may have a better analgesic effect in comparison to meloxicam. However, the incidence of side effects, particularly nausea and vomiting, may be high. Dental pain is largely inflammatory, and evidence-based medicine has shown that NSAIDs are the best analgesic for dental pain [41, 42]. Our institution is passionate about high quality evidence based-research and has excelled in various fields [16, 43–62].

#### Limitations

As the VAS Scores were based on patients' perception, a subjective opinion regarding the results was obtained, hence it would be a limitation of our study. Also, the pain threshold for different patients would not be similar.

# **Future Scope**

Although the literature provides a number of studies on pre-emptive analysis efficacy of piroxicam and tramadol, there are limited studies related to comparing piroxicam with an opioid analysis both as pre-emptive analysis for pain control after third molar surgery.

# **CONCLUSION**

Within the limits of this study, it can be concluded that the patients receiving 20 mg of preoperative piroxicam had less pain intensity and total analgesic consumption than those receiving 50 mg of preoperative tramadol. Therefore, piroxicam given preoperatively showed superior analgesic properties for intermediate surgical procedures in comparison to preemptively administered tramadol.

**COMPETING INTERESTS DISCLAIMER:** 

Authors have declared that no competing interests exist. The products used for this research are commonly and predominantly use products in our area of research and country. There is absolutely no conflict of interest between the authors and producers of the products because we do not intend to use these products as an avenue for any litigation but for the advancement of knowledge. Also, the research was not funded by the producing company rather it was funded by personal efforts of the authors.

#### REFERENCES

- Seymour RA, Meechan JG, Blair GS. An investigation into post-operative pain after third molar surgery under local analgesia. Br J Oral Maxillofac Surg 1985; 23: 410– 418.
- 2. Woolf CJ, Chong MS. Preemptive analgesia—treating postoperative pain by preventing the establishment of central sensitization. Anesthesia & Analgesia. 1993 Aug 1;77(2):362-79.
- 3. Scott LJ, Perry CM. Tramadol. Drugs 2000; 60: 139–176.
- 4. Barden J, Edwards JE, McQuay HJ, et al. Relative efficacy of oral analgesics after third molar extraction. Br Dent J 2004; 197: 407–11.
- 5. Dionne RA, Berthold CW. Therapeutic uses of non-steroidal anti-inflammatory drugs in dentistry. Crit Rev Oral Biol Med 2001; 12: 315–330.
- 6. Euller-Ziegler L, Vélicitat P, Bluhmki E, et al. Meloxicam: a review of its pharmacokinetics, efficacy and tolerability following intramuscular administration. Inflamm Res 2001; 50 Suppl 1: S5–9.
- 7. Nekoofar MH, Sadeghipanah M, Dehpour AR. Evaluation of meloxicam (A cox-2 inhibitor) for management of postoperative endodontic pain: a double-blind placebocontrolled study. J Endod 2003; 29: 634–637.
- 8. Miranda HF, Pinardi G. Lack of effect of naltrexone on the spinal synergism between morphine and non steroidal anti-inflammatory drugs. Pharmacol Rep 2009; 61: 268–274.
- 9. Gillies GW, Kenny GN, Bullingham RE, McArdle CS. The morphine sparing effect of ketorolac tromethamine: A study of a new, parenteral non-steroidal anti-inflammatory agent after abdominal surgery. Anaesthesia. 1987 Jul;42(7):727-31.
- 10. Rosenblum M, Weller RS, Conard PL, et al. Ibuprofen provides longer lasting analysesia than fentanyl after laparoscopic surgery. Anesth Analy 1991; 73: 255–259.
- 11. Gold BS, Kitz DS, Lecky JH, et al. Unanticipated admission to the hospital following

- ambulatory surgery. JAMA 1989; 262: 3008–3010.
- 12. Govindaraju L, Gurunathan D. Effectiveness of Chewable Tooth Brush in Children-A Prospective Clinical Study. J Clin Diagn Res 2017; 11: ZC31–ZC34.
- 13. Christabel A, Anantanarayanan P, Subash P, et al. Comparison of pterygomaxillary dysjunction with tuberosity separation in isolated Le Fort I osteotomies: a prospective, multi-centre, triple-blind, randomized controlled trial. Int J Oral Maxillofac Surg 2016; 45: 180–185.
- 14. Soh CL, Narayanan V. Quality of life assessment in patients with dentofacial deformity undergoing orthognathic surgery--a systematic review. Int J Oral Maxillofac Surg 2013; 42: 974–980.
- 15. Mehta M, Deeksha, Tewari D, et al. Oligonucleotide therapy: An emerging focus area for drug delivery in chronic inflammatory respiratory diseases. Chem Biol Interact 2019; 308: 206–215.
- Ezhilarasan D, Apoorva VS, Ashok Vardhan N. Syzygium cumini extract induced reactive oxygen species-mediated apoptosis in human oral squamous carcinoma cells. J Oral Pathol Med 2019; 48: 115–121.
- 17. Campeau PM, Kasperaviciute D, Lu JT, et al. The genetic basis of DOORS syndrome: an exome-sequencing study. Lancet Neurol 2014; 13: 44–58.
- 18. Kumar S, S S. Knowledge and awareness regarding antibiotic prophylaxis for infective endocarditis among undergraduate dental students. Asian J Pharm Clin Res 2016; 154.
- 19. Christabel SL. Prevalence of type of Frenal Attachment and morphology of frenum in children, Chennai, Tamil Nadu. World J Dent 2015; 6: 203–207.
- 20. Kumar S, Rahman R. Knowledge, awareness, and practices regarding biomedical waste management among undergraduate dental students. Asian J Pharm Clin Res 2017; 10: 341.
- 21. Sridharan G, Ramani P, Patankar S. Serum metabolomics in oral leukoplakia and oral squamous cell carcinoma. J Cancer Res Ther 2017; 13: 556–561.
- 22. Ramesh A, Varghese SS, Doraiswamy JN, et al. Herbs as an antioxidant arsenal for periodontal diseases. J Intercult Ethnopharmacol 2016; 5: 92–96.
- 23. Thamaraiselvan M, Elavarasu S, Thangakumaran S, et al. Comparative clinical evaluation of coronally advanced flap with or without platelet rich fibrin membrane in the treatment of isolated gingival recession. J Indian Soc Periodontol 2015; 19: 66–71.
- 24. Thangaraj SV, Shyamsundar V, Krishnamurthy A, et al. Molecular Portrait of Oral

- Tongue Squamous Cell Carcinoma Shown by Integrative Meta-Analysis of Expression Profiles with Validations. PLoS One 2016; 11: e0156582.
- 25. Ponnulakshmi R, Shyamaladevi B, Vijayalakshmi P, et al. In silico and in vivo analysis to identify the antidiabetic activity of beta sitosterol in adipose tissue of high fat diet and sucrose induced type-2 diabetic experimental rats. Toxicol Mech Methods 2019; 29: 276–290.
- 26. Ramakrishnan M, Shukri M. Fluoride, Fluoridated Toothpaste Efficacy And Its Safety In Children-Review. International Journal of Pharmaceutical Research. 2018 Oct 1;10(04):109-14.
- 27. McQuay HJ. Pre-emptive analgesia: a systematic review of clinical studies. Ann Med 1995; 27: 249–256.
- 28. Isiordia-Espinoza MA, Sánchez-Prieto M, Tobías-Azúa F, et al. Pre-emptive analgesic effectiveness of meloxicam versus tramadol after mandibular third molar surgery: a pilot study. J Oral Maxillofac Surg 2012; 70: 31–36.
- 29. Jadad AR, Moore RA, Carroll D, et al. Assessing the quality of reports of randomized clinical trials: is blinding necessary? Control Clin Trials 1996; 17: 1–12.
- 30. Ong CK-S, Lirk P, Seymour RA, et al. The efficacy of preemptive analysis for acute postoperative pain management: a meta-analysis. Anesth Analg 2005; 100: 757–73.
- 31. Richmond CE, Bromley LM, Woolf CJ. Preoperative morphine pre-empts postoperative pain. Lancet 1993; 342: 73–75.
- 32. Ong KS, Tan JML. Preoperative intravenous tramadol versus ketorolac for preventing postoperative pain after third molar surgery. Int J Oral Maxillofac Surg 2004; 33: 274–278.
- 33. Ong CKS, Lirk P, Tan JMH, et al. The analgesic efficacy of intravenous versus oral tramadol for preventing postoperative pain after third molar surgery. J Oral Maxillofac Surg 2005; 63: 1162–1168.
- 34. Pozos-Guillen A, Martinez-Rider R, Aguirre-Banuelos P, et al. Pre-emptive analgesic effect of tramadol after mandibular third molar extraction: a pilot study. J Oral Maxillofac Surg 2007; 65: 1315–1320.
- 35. Farshchi A, Ghiasi G. Comparison the analgesic effects of single dose administration of tramadol or piroxicam on postoperative pain after cesarean delivery. Acta Med Iran 2010; 48: 148–153.
- 36. Dahl JB, Kehlet H. Non-steroidal anti-inflammatory drugs: rationale for use in severe postoperative pain. Br J Anaesth 1991; 66: 703–712.

- 37. O'Hanlon JJ, Muldoon T, Lowry D, et al. Improved postoperative analgesia with preoperative piroxicam. Canadian Journal of Anaesthesia 1996; 43: 102–105.
- 38. Aoki T, Yamaguchi H, Naito H, et al. Premedication with cyclooxygenase-2 inhibitor meloxicam reduced postoperative pain in patients after oral surgery. Int J Oral Maxillofac Surg 2006; 35: 613–617.
- 39. De Menezes SAF, Cury PR. Efficacy of nimesulide versus meloxicam in the control of pain, swelling and trismus following extraction of impacted lower third molar. Int J Oral Maxillofac Surg 2010; 39: 580–584.
- 40. Calvo AM, Sakai VT, Giglio FPM, et al. Analgesic and anti-inflammatory dose-response relationship of 7.5 and 15 mg meloxicam after lower third molar removal: a double-blind, randomized, crossover study. Int J Oral Maxillofac Surg 2007; 36: 26–31.
- 41. Mehlisch DR. The efficacy of combination analgesic therapy in relieving dental pain. J Am Dent Assoc 2002; 133: 861–871.
- 42. Ong CKS, Seymour RA. Pathogenesis of postoperative oral surgical pain. Anesth Prog 2003; 50: 5–17.
- 43. Vijayashree Priyadharsini J. In silico validation of the non-antibiotic drugs acetaminophen and ibuprofen as antibacterial agents against red complex pathogens. J Periodontol 2019; 90: 1441–1448.
- 44. Pc J, Marimuthu T, Devadoss P, Kumar SM. Prevalence and measurement of anterior loop of the mandibular canal using CBCT: A cross sectional study. Clinical implant dentistry and related research. 2018 Apr 6;20(4):531-4.
- 45. Ramesh A, Varghese S, Jayakumar ND, et al. Comparative estimation of sulfiredoxin levels between chronic periodontitis and healthy patients A case-control study. J Periodontol 2018; 89: 1241–1248.
- 46. Ramadurai N, Gurunathan D, Samuel AV, et al. Effectiveness of 2% Articaine as an anesthetic agent in children: randomized controlled trial. Clin Oral Investig 2019; 23: 3543–3550.
- 47. Sridharan G, Ramani P, Patankar S, et al. Evaluation of salivary metabolomics in oral leukoplakia and oral squamous cell carcinoma. J Oral Pathol Med 2019; 48: 299–306.
- 48. Mathew MG, Samuel SR, Soni AJ, et al. Evaluation of adhesion of Streptococcus mutans, plaque accumulation on zirconia and stainless steel crowns, and surrounding gingival inflammation in primary molars: Randomized controlled trial. Clin Oral Investig 2020; 1–6.

- 49. Samuel SR. Can 5-year-olds sensibly self-report the impact of developmental enamel defects on their quality of life? Int J Paediatr Dent 2021; 31: 285–286.
- 50. R H, Hannah R, Ramani P, et al. CYP2 C9 polymorphism among patients with oral squamous cell carcinoma and its role in altering the metabolism of benzo[a]pyrene. Oral Surgery, Oral Medicine, Oral Pathology and Oral Radiology 2020; 130: 306–312.
- 51. Chandrasekar R, Chandrasekhar S, Sundari KKS, et al. Development and validation of a formula for objective assessment of cervical vertebral bone age. Prog Orthod 2020; 21: 38.
- 52. Vijayashree Priyadharsini J, Smiline Girija AS, Paramasivam A. In silico analysis of virulence genes in an emerging dental pathogen A. baumannii and related species. Arch Oral Biol 2018; 94: 93–98.
- 53. MP SK. Knowledge, Attitude and practices regarding needlestick injuries among dental students. Asian J Pharm Clin Res. 2016;9(4):312-5.
- 54. SK M. Knowledge, attitude, and practices regarding infection control among undergraduate dental students. Asian J Pharm Clin Res. 2016;9(1):220-4.
- 55. Ak H. Knowledge and awareness about oral cancer among undergraduate dental students. Asian Journal of Pharmaceutical and Clinical Research. 2016 Jul 1:165-7.
- 56. Gayathri MM. Knowledge and awareness among patients about dental implants. Journal of Pharmaceutical Sciences and Research. 2016 May 1;8(5):351.
- 57. Vijayalakshmi B, Kumar MS. Knowledge of students about Local anaesthetics used during oral surgical procedures. Journal of Pharmaceutical Sciences and Research. 2015 Nov 1;7(11):1011.
- 58. Gayathri MM. Knowledge, Awareness and Attitude among dental students about hepatitis B infection. Journal of Pharmaceutical Sciences and Research. 2016 Mar 1:8(3):168.
- 59. Ahamed A, Kumar MS. Knowledge, attitude and perceived confidence in handling medical emergencies among dental students. Journal of Pharmaceutical Sciences and Research. 2016 Jul 1;8(7):645.
- 60. Kumar S. Knowledge, attitude and practices of dental students toward dental management of patients on antiplatelet therapy. Asian J Pharm Clin Res. 2016;9(30):270-6.
- 61. MP SK. Local hemostatic agents in the management of bleeding in oral surgery. Asian J Pharm Clin Res. 2016;9(3):35-41.

62. Kumar MP. Newer methods of extraction of teeth. Int J Pharm Bio Sci. 2015;6(3):679-85.

