Efficacy of dental local anesthetics: A review

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The objective of this review was to investigate the efficacy of dental local anesthetics, as it is well known among clinicians that local anesthesia may be challenging in some circumstances. Therefore, the focus of this review was on the efficacy of the products used in dental local anesthesia.

In a PubMed database literature search conducted, a total of 8646 articles were found to be related to dental local anesthetics. After having applied the inclusion criteria (human research, performed in the last 10 years, written in English language, and focus on dental local anesthetics) and having assessed the quality of the papers, 30 were deemed eligible for inclusion in this review.

The conclusion of this review is that none of the dental local anesthetic amides provide 100% anesthesia. The problem appears to be more pronounced when mandibular teeth are attempted to be anaesthetized and especially if there is irreversible pulpitis involved. The authors conclude that this finding suggest exploration of more efficient techniques to administer dental local anesthesia, especially in the mandible, to establish a 100% efficacy, is needed.

Keywords: Amides; Local Anesthesia.

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INTRODUCTION

Local anesthesia in dentistry provides comfort for the patient, but also as much comfort for the clinician as the planned procedures can be carried out under the best possible conditions. From clinical experience and from the literature, it is clear that dental local anesthesia is not always as successful as anticipated [1-8]. Especially mandibular block anesthesia can be difficult to achieve or challenging in some patients, even in the absence of a tooth with an acute pulpitis. Mandibular block failure rates differ from study to study and teach us that there is no 100% success [1-3,7]. The efficacy of local anesthesia in the maxilla is much higher, based on clinical experience and publications. The main reason is probably the cortical plates of the mandible being thicker and denser and having less porosities that allow for a volume of local anesthetic to be diffused into the cancellous bone in case one attempts a buccal infiltration for instance, explains most of the difference with the maxilla. Another reason is the techniques that are used to achieve local anesthesia. In the maxilla, the most appropriate technique would be a buccal infiltration anesthesia close to the level of the apices of the teeth, while in the mandible, because of the reason mentioned above, local anesthesia is achieved mainly by attempting to deposit a volume of local anesthetic close to the mandibular nerve before it

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enters the mandible [6,7,9,10]. Because of anatomical variations in localisation of the second branch of the trigeminal nerve with regard to the ramus of the mandible, the efficacy of local anesthesia is not 100% [1-11].

The aim of this current review was to assess the peer-reviewed literature on the topic of dental local anesthetics to see if the efficacy of dental local anesthetics depends on the amide or a combination of amides used.

**MATERIAL AND METHODS**

Fig. 1 shows which search terms were used in the PubMed® database to identify relevant publications, how many publications were found per search term and how publications were finally triaged to be reviewed by both researchers. The inclusion criteria used, were that studies had to be related to human research, performed in the last 10 years, written in English language, and with a focus on dental local anesthetics. After having identified the search terms, the search was conducted between January 2017 and January 2018 by one researcher (NB). As can be derived from Fig. 1, an initial total of 8646 manuscripts were identified. After applying the selection criteria, 79 papers were subsequently read by both researchers, and then categorised in consensus in an Excel spreadsheet (Microsoft®, Redmond, Washington, USA) to identify the type of study (e.g. randomised clinical trial), the amides investigated (e.g. articaine), the number of patients included in the study if it was a clinical trial, the country of origin and the year of publication and finally, the conclusion of the study. Both investigators read all 79 publications, and, in consensus, deemed 30 manuscripts eligible for the study’s aim.
Table 1. Review papers included in this current review, indicating reference, year of publication, type of amides involved, type of anesthesia involved and conclusion of the study

| Authors, Country, Year | Type of study | Amid and Vasoconstrictor Concentrations | Anesthesia Techniques Used | Conclusion of Study |
|------------------------|---------------|-----------------------------------------|---------------------------|---------------------|
| Su N et al., China, 2014 [12] | meta-analysis of randomised controlled trials | • 2% lidocaine and 1:100k epinephrine  
• 3% mepivacaine plain  
• 2% mepivacaine and 1:100k epinephrine  
• 2% mepivacaine and 1:20k levonordefrin  
• 2% lidocaine and 1:50k epinephrine | • Inferior alveolar nerve block  
• Infiltration  
• Maxillary block  
• Intracutaneous  
• Intraligamentary | Given the efficacy and safety of the two solutions, 2% mepivacaine with vasoconstrictors is better than 2% lidocaine with vasoconstrictors in dental treatment. Meanwhile, 3% plain mepivacaine is better for patients with cardiac diseases. |
| Su N et al., China, 2014 [14] | meta-analysis of randomised controlled trials | • 2% lidocaine and 1:100k epinephrine  
• 0.5% bupivacaine and 1:200k epinephrine  
• 2% lidocaine and 1:80k epinephrine  
• 0.75% bupivacaine and 1:200k epinephrine  
• 0.75% levobupivacaine plain  
• 0.5% levobupivacaine plain | • Inferior alveolar nerve block  
• Infiltration | There was no statistical significance in adverse events between two groups. Given the efficacy and safety, the bupivacaine group is better than the lidocaine group in dental operations that take a relatively long time, especially in endodontic treatments or where there is a need for postoperative pain management. |
| Katyal V, Australia, 2010 [42] | meta-analysis | • 4% articaine and 1:100k epinephrine  
• 2% lignocaine and 1:100k epinephrine | • Inferior alveolar nerve block  
• Infiltrations in maxilla and mandible | Articaine is more likely than lignocaine to achieve an anesthetic success in the posterior first molar area with a relative risk for success at 1.31. The results of this systematic review provide support for the argument that articaine is more effective than lignocaine in providing anesthetic success in the first molar region for routine dental procedures. In addition, both drugs appear to have similar adverse effect profiles. The clinical impact of articaine’s higher post-injection pain scores than lignocaine is negligible. Hence, articaine is a superior anesthetic to lignocaine for use in routine dental procedures. Use in children under 4 years of age is not recommended, since no data exists to support such usage. |
| Kung J et al., USA, 2015 [43] | systematic review and meta-analysis | unclear | • Inferior alveolar nerve block  
• Infiltration | For combined studies, articaine was more likely than lidocaine to achieve successful anesthesia. Maxillary infiltration subgroup analysis showed no significant difference between articaine and lidocaine. For combined mandibular anesthesia studies articaine was superior to lidocaine, with further subgroup analysis showing no difference for mandibular block anesthesia. When used for supplemental infiltration after successful mandibular block anesthesia, articaine was significantly more effective than lidocaine. There were no reports of adverse events. In conclusion, the present meta-analysis showed that in patients with symptomatic irreversible pulps, articaine is as effective as lidocaine when used for mandibular block or maxillary infiltration anesthesia. In cases of persistent pulpal pain despite successful mandibular block anesthesia, supplementary infiltration with articaine instead of lidocaine has 3.55 times greater likelihood of achieving successful anesthesia. |


RESULTS

A total of 30 publications (Fig. 1) were considered relevant for the study: 7 reviews, and 23 clinical trials. The details of the different studies and their conclusions can be found in Tables 1 and 2. These tables show data regarding the country of origin, the number of subjects involved, which amides were investigated and the last column the final conclusions as stated in the respective papers. Table 3 tabulates the number of manuscripts that investigated which type of amide and which amide was found to be more or equally in efficacy to achieve local anesthesia. The numbers in the right hand side column refer to the manuscripts reference list.

Of the six amide products used in dental local anesthesia, 13 in vivo studies and 4 reviews found that articaine was the amide with the highest efficacy when compared to either lidocaine, mepivacaine, prilocaine or bupivacaine (see Table 1). Mepivacaine [12,13] and bupivacaine [14,15] as opposed to lidocaine, appeared to have a higher efficacy in one review [12,14] and one in vivo study [13,15], each.

Prilocaine, as opposed to lidocaine and bupivacaine, was found to have a higher efficacy in only one in vivo study [16]. Two in vivo studies claimed that ropivacaine had a high efficacy under different concentrations [17], without comparing it to another amide or when comparing it to lignocaine [18].

However, when duration of the local anesthetic effect was taken into account, mepivacaine was shown to work for a shorter duration compared to lidocaine [19]. Eleven studies [20-30] either found no significant differences between two types of amides or studied the volume of anesthetic and found its efficacy was significantly greater if higher volumes (more than one cartridge) were administered.

With regard to addition of vasoconstrictors epinephrine and clonidine, one in vivo study concluded that clonidine increased efficacy better than epinephrine, combined with
| Authors, Country, Year | Type of study | Amid and Vasoconstrictor Concentrations | Technique Used: Block or Infiltration? Mandibular or Maxillary? | Number of Subjects in Study | Conclusion of Study |
|------------------------|--------------|----------------------------------------|---------------------------------------------------------------|----------------------------|-------------------|
| Colombini BL et al., Brazil, 2006 [45] | randomised double blind cross-over study | • 4% articaine and 1:100k epinephrine  
• 2% mepivacaine and 1:100k epinephrine | Inferior alveolar nerve block (third molar removal) | 20 | Articaine provides a longer period of analgesic effect and a tendency for a longer period of anesthesia as compared to mepivacaine. The presence of a vasoconstrictor agent in local anesthetic solutions does not seem to influence hemodynamic parameters during lower third molar removal in healthy subjects. |
| Jain NK et al. India, 2016 [31] | comparative prospective | • 4% articaine and 1:100k epinephrine  
• 2% lignocaine and 1:80k epinephrine | Inferior alveolar nerve block (third molar surgical removal) | 70 | Results showed that 4% articaine had a significant faster onset and longer duration of action when compared to 2% lignocaine. We concluded that 4% articaine is a safe alternative to 2% lignocaine, which is potent and effective in minor surgical procedures such as removal of mandibular third molars. |
| Pellicer-Chover H et al. Spain, 2013 [46] | comparative split mouth | • 4% articaine and 1:100k epinephrine  
• 0.5% bupivacaine and 1:200k epinephrine | Inferior alveolar nerve block (third molar surgical removal) | 36 | Articaine showed greater clinical efficacy than bupivacaine, reducing latency time, bleeding, anesthetic duration in the soft tissues and achieving higher anesthetic quality, requiring less reinforcement during surgery than bupivacaine. |
| Budharapu A et al. India, 2015 [18] | randomised single blind trial | • 0.5% ropivacaine  
• 2% lignocaine | Inferior alveolar nerve block (third molar surgical removal) | 78 | We had no adverse effects from 0.5% ropivacaine, which provided efficient anesthesia and excellent residual analgesia with no cardiovascular or central nervous system complications. The addition of adrenaline to ropivacaine has been shown not to improve the anesthetic effect and may cause transient increases in arterial pressure and heart rate. Evaluation of the plasma concentration would give more details about its effects on the cardiovascular system, as we confined ourselves to studying only the haemodynamic changes. |
| Brkovic BM et al. Serbia, 2010 [17] | double blind randomised controlled design | • 0.5% ropivacaine plain  
• 0.75% ropivacaine plain  
• 1% ropivacaine plain | Infiltration (third molars maxilla) | 66 | In conclusion, the current data suggest that maxillary infiltration of 0.75 and 1% of ropivacaine offered adequate and safe intraoperative analgesia but not successful postoperative pain control for the surgical removal of upper third molars. |
| Christensen J et al. Denmark, 2013 [15] | randomised double blind cross over design | • 2% lidocaine with or without methylprednisolone or placebo  
• 0.5% bupivacaine with or without methylprednisolone or placebo | Inferior alveolar nerve block (third molar surgical removal) | 126 | Bupivacaine combined with methylprednisolone reduced the postoperative pain and swelling compared with the use of lidocaine and placebo, lidocaine and methylprednisolone, or bupivacaine and placebo. (two to four supplemental injections were administered sometimes) |
| Krzeminski TF et al. Poland, 2011 [25] | randomised parallel group trial | • 0.5% ropivacaine plain  
• 4% articaine and 1:100k epinephrine | Infiltration maxillary incisors and canines | 60 | The efficacy of anesthesia of lateral and central incisors was 100% for both anesthetics. There were insignificant differences in effectiveness of canine pulp anesthesia. Ropivacaine (0.5%) achieved effective and long duration of uninfamed pulp and soft tissue anesthesia. Ropivacaine could be useful for long-lasting operative procedures without the need for a vasoconstrictor. |
Table 2. Continued

| Study | Design | Anesthesia Type | Procedure | N | Notes |
|-------|--------|-----------------|-----------|---|-------|
| Thakare A et al. India, 2014 [47] | Randomised cross-over study | • 4% articaine  
• 0.5% bupivacaine  
• [epinephrine concentration unclear] | Infiltration in maxilla (premolar extractions) | 40 | The results showed that 4% articaine had significantly faster onset of action and lower VAS scores when compared with bupivacaine. However, the duration of analgesia and time to first rescue analgesic medication was longer in the bupivacaine group. Articaine seemed to have better potency and efficacy in terms of onset of action and lower pain scores compared to the bupivacaine group. Further studies are required to confirm these results. Moreover, the volume of LA required seemed to be lower in the articaine group, demonstrating better anesthetic potency and efficacy. |
| Ashraf H et al. Iran, 2013 [48] | Randomised double blind study | • 2% lidocaine and 1:100k epinephrine  
• 4% articaine and 1:100k epinephrine | Inferior alveolar nerve block  
Infiltration | 125 | Supplementing an incomplete articaine inferior alveolar nerve block with articaine infiltration raises the anesthetic success more effectively compared with lidocaine in mandibular molars with irreversible pulpitis. The success rate after the administration of the infiltration injections after an incomplete inferior alveolar nerve block by using lidocaine was 29%, whereas by using articaine it was 71%. No statistical differences were detected in the success rates between the 2 anesthetics after the block injections. |
| Brkovic B et al. Serbia, 2008 [49] | Randomised double blind study | • 2% lidocaine and clonidine (15ug/mL)  
• 2% lidocaine and epinephrine (12.5 ug/mL) | Infiltration (maxilla) | 40 | The results of this study indicate for the first time in dental anesthesia that the lidocaine + clonidine combination could be a useful and safe alternative to lidocaine + epinephrine for intraoral infiltration anesthesia. |
| Piccinni C et al. Italy, 2015 [16] | Case-non-case study | Unclear about concentrations of:  
• lidocaine  
• bupivacaine  
• articaine  
• prilocaine | NA | 17246 | In conclusion, among local anesthetics, only articaine and prilocaine generated a signal of paresthesia, especially when used in dentistry. The highest number of reports was found for lidocaine (247 reports), followed by bupivacaine (99 reports), articaine (85 reports), combination of different local anesthetics (45 reports) and prilocaine (30 reports). A significant disproportionality of 'paresthesias and dysaesthesias' was found for articaine and prilocaine. Other local anesthetics did not show disproportionality signals according to the defined thresholds. |
| Lammers E, et al. USA, 2014 [26] | Prospective, randomised, double blind study | • 2% lidocaine with 1:100k epinephrine  
• 2% lidocaine with 1:100k epinephrine | Inferior alveolar nerve block | 100 | The combination of 3% mepivacaine plus 2% lidocaine with 1:100,000 epinephrine was equivalent to the combination of 2 cartridges of 2% lidocaine with 1:100,000 epinephrine in terms of injection pain, onset time, and pulpal anesthetic success for the IAN block. |
| Visconti RP et al. Brasil, 2016 [13] | Double blind randomised clinical trial | • 2% mepivacaine and 1:100k epinephrine  
• 2% lidocaine and 1:100k epinephrine | Inferior alveolar nerve block (irreversible pulpitis) | 42 | All patients tested reported lip anesthesia after application of either type of inferior alveolar nerve block. Pulpal anesthesia success rates measured by using the pulp tester were satisfactory for both solutions (86% for mepivacaine and 67% for lidocaine). Success rates according to patient report of no pain or mild pain during pulpectomy were higher for mepivacaine solution (55%) than for lidocaine solution (14%). The differences between mepivacaine and lidocaine were statistically significant. Mepivacaine resulted in effective pain control during irreversible pulpitis treatments. |
| Study Details | Design Details | Anesthetic Formulations | Anesthesia Type | Success Rate |
|---------------|---------------|-------------------------|----------------|--------------|
| Whitcomb M et al. USA, 2010 [32] | Prospective, randomised, double blind | 2% lidocaine and 1:100k epinephrine, 2% lidocaine buffered with sodium bicarbonate | Inferior alveolar nerve block | 40 |
| Kanaa MD et al. UK, 2012 [28] | Randomised, double blind study | 4% articaine and 1:1000k epinephrine, 2% lidocaine and 1:80k epinephrine | Infiltration (maxilla and irreversible pulpitis) | 100 |
| Mohajeri L et al. Iran, 2015 [36] | Randomised, double blind, clinical trial | 2% lidocaine and 1:100k epinephrine, 2% lidocaine and 1:100k epinephrine with 5% meperidine | Intraligamentary injections | 60 |
| Gazal G, Saudi Arabia, 2017 [50] | Randomised, double blind cross over study | 4% articaine and 1:100k epinephrine, 2% mepivacaine and 1:100k epinephrine | Inferior alveolar nerve block, Buccal infiltration | 23 |
| Srinivasan N et al. India, 2009 [51] | Prospective, randomised, double blind study | 4% articaine and 1:100k epinephrine, 2% lidocaine and 1:100k epinephrine | Infiltration (maxilla and irreversible pulpitis) | 40 |
| Glenn B et al. USA, 2016 [38] | Prospective, randomised, double blind trial | Liposomal bupivacaine, 0.5% bupivacaine and 1:200k epinephrine, 2% lidocaine and 1:100k epinephrine | Buccal infiltrations, inferior alveolar nerve block, infiltration (maxilla) | 100 |
| Schellenberger J et al. USA, 2015 [33] | Prospective, randomised, double blind trial | 4% articaine buffered with sodium bicarbonate and 1:100k epinephrine, 4% articaine and 1:100k epinephrine | Inferior alveolar nerve block | 100 |
| Shurtz R et al. USA, 2015 [34] | Prospective, randomised, double blind trial | 4% articaine buffered with sodium bicarbonate and 1:100k epinephrine, 4% articaine and 1:100k epinephrine | Infiltration (mandible) | 80 |
| Nydegger B et al. USA, 2014 [52] | Prospective, randomised, double blind trial | 4% articaine and 1:100k epinephrine, 4% lidocaine and 1:100k epinephrine, 4% prilocaine 1:200k epinephine | Infiltration (mandible) | 60 |

For the buffered 2% lidocaine with 1:100,000 epinephrine/sodium bicarbonate formulation, successful pulpal anesthesia ranged from 10 to 71%. For the unbuffered 2% lidocaine with 1:100,000 epinephrine formulation, successful pulpal anesthesia ranged from 10 to 72%. We concluded that buffering a 2% lidocaine with 1:100,000 epinephrine with sodium bicarbonate, as was formulated in the current study, did not statistically increase anesthetic success, provide faster onset, or result in less pain of injection when compared with unbuffered 2% lidocaine with 1:100,000 epinephrine for an IAN block.

There was no significant difference in efficacy between 4% articaine with 1:100,000 epinephrine and 2% lidocaine with 1:80,000 epinephrine in achieving anesthesia in maxillary teeth with irreversible pulpsitis after buccal infiltration.

Within the limitations of the present study, it appears that the addition of meperidine as an opioid to 2% lidocaine with 1:100,000 epinephrine will not improve the anesthetic efficacy of a periodontal ligament injection in patients with irreversible pulpsitis for whom an inferior alveolar nerve block was ineffective despite lip numbness.

Articaine has better potency, rapid onset of action, earlier lip and teeth numbness compared to the mepivacaine group. Articaine/mepivacaine buccal injection was significantly more comfortable than mepivacaine inferior alveolar nerve block.

The success rate for maxillary buccal infiltration to produce pulpal anesthesia using articaine was 100% in first premolar and first molar, and for the lidocaine solution, success rate was 80% in first premolar and 30% in first molar.

The success rate was 29% for the liposomal group and 22% for the bupivacaine group, with no significant difference between the groups.

The success rate for the inferior alveolar nerve block was 32% for the buffered group and 40% for the non buffered group, with no significant difference between the groups.

Buffered articaine did not provide any advantage over non buffered articaine for anesthetic success, anesthesia onset, or pain of injection for a primary buccal infiltration of the mandibular first molar.

The success rate for the 4% articaine formulation was 55%, 33% for the 4% lidocaine formulation, and 32% for the 4% prilocaine formulation. There was a significant difference between articaine and both lidocaine and prilocaine formulations.
| Study | Design Details | Anesthetic Details | Results/Conclusions |
|-------|----------------|-------------------|---------------------|
| Rogers BS et al. USA, 2014 [53] | Prospective, randomised, double blind trial | 4% lidocaine and 1:100k epinephrine, 2% lidocaine and 1:100k epinephrine | Improved anesthesia success rate, reduced postoperative discomfort. |
| Cohen H et al. USA, 2013 [37] | Prospective, randomised, single blind trial | 1.72 mL 4% lidocaine and 50 ug/mL epinephrine, 5 mL 68.8 mg lidocaine with 50 ug epinephrine plus 0.9 M mannitol | Improved anesthesia success rate, reduced postoperative discomfort. |
| Martin M et al. USA, 2011 [29] | Prospective, randomised, cross-over design | 4% lidocaine and 1:100k epinephrine | Improved anesthesia success rate, reduced postoperative discomfort. |
| McEntire M et al. USA, 2011 [54] | Prospective, randomised, double blind crossed-over trial | 4% lidocaine and 1:200k epinephrine, 4% lidocaine and 1:100k epinephrine | Improved anesthesia success rate, reduced postoperative discomfort. |
| Wall M et al. USA, 2010 [55] | Prospective, randomised, single blind trial | 2% lidocaine and 1:50k epinephrine, 2% lidocaine and 1:100k epinephrine | Improved anesthesia success rate, reduced postoperative discomfort. |
| Goodman A et al. USA, 2006 [35] | Prospective, randomised, single blind cross-over trial | 4% lidocaine and 18 ug/1.8 mL epinephrine, 4% lidocaine and 36 ug/3.6 mL epinephrine and 36 mg meperidine | Improved anesthesia success rate, reduced postoperative discomfort. |
| Brunetto PC et al. Brasil, 2008 [30] | Randomised, double blind cross-over study | 2% lidocaine and 1:100k epinephrine | Improved anesthesia success rate, reduced postoperative discomfort. |
| Poomi S et al. India, 2011 [20] | Prospective randomised double blind clinical trial | 4% articaine and 1:100k epinephrine, 2% lidocaine and 1:100k epinephrine | Improved anesthesia success rate, reduced postoperative discomfort. |
| Evans G et al. USA, 2008 [56] | Prospective, randomised double blind study | 4% articaine and 1:100k epinephrine, 2% lidocaine and 1:100k epinephrine | Improved anesthesia success rate, reduced postoperative discomfort. |
Table 2. Continued

| Study                        | Design                  | Anesthetic formulations | Outcome |
|------------------------------|-------------------------|-------------------------|---------|
| Sreekumar K et al. India, 2011 [27] | Randomised double blind cross over study | 4% articaine and 1:100k epinephrine | Inferior alveolar nerve block | Maxillary infiltration anesthesia with articaine and epinephrine has a faster onset, a greater success rate, and a longer duration when a volume of 1.2 mL is used than when volumes less than 1.0 mL are used. Palatal tissues were anesthetized with the highest concentration (1.2 mL) in our study (30% of cases). |
| Sampaio RM et al. Brasil, 2012 [21] | Randomised double blind study | 0.5% bupivacaine and 1:200k epinephrine | Inferior alveolar nerve block | Neither of the solutions resulted in an effective pain control during irreversible pulpitis treatments of mandibular molars. Before initiation of the pulpectomy procedure, 15 patients (42.9%) in the lidocaine group and 7 patients (20%) in the bupivacaine group exhibited pulpal anesthesia (ie, a negative response to electrical stimuli generated with an electric pulp tester). A significant difference between the 2 experimental groups for the pulpal anesthesia was observed, with more individuals in the lidocaine group presenting a negative response to electrical stimuli. During the pulpectomy, 7 patients in the bupivacaine group (20%) and 13 in the lidocaine group (37.1%) reported pain, however this difference was not statistically significant. |
| Meen R et al. USA, 2009 [22] | Prospective, randomised, double-blind study | 2% lidocaine and 1:100 epinephrine | Infiltrations (maxilla) | Anesthetic success and the onset of pulpal anesthesia were not significantly different between 2% lidocaine with either 1:100,000 or 1:50,000 epinephrine and 3% mepivacaine for the lateral incisor and first molar. Increasing the epinephrine concentration from 1:100,000 to 1:50,000 in a 2% lidocaine formulation significantly decreased pulpal anesthesia of short duration for the lateral incisor but not the first molar. For both the lateral incisor and first molar, 3% mepivacaine significantly increased pulpal anesthesia of short duration compared with 2% lidocaine with either 1:100,000 or 1:50,000 epinephrine. |
| Forloine A et al. USA, 2010 [23] | Prospective, randomised, double-blind study | 2% lidocaine and 1:100 epinephrine | Maxillary block | The high tuberosity approach to the maxillary second division nerve block with both anesthetic formulations resulted in a high success rate (92%-98%) for the first and second molars. Approximately 76%-78% of the second premolars were anesthetized with both anesthetic formulations. Both anesthetic formulations were ineffective for the anterior teeth and first premolars. The use of 3% mepivacaine provided a significantly shorter duration of pulpal anesthesia than 2% lidocaine with 1:100,000 epinephrine in the molars and premolars. |
| Lawaty I et al. USA, 2010 [24] | Prospective, randomised, double-blind study | 2% mepivacaine and 1:20k levonordefrin | Infiltration (maxilla) | Anesthetic success (obtaining 2 consecutive 80 readings with the electric pulp tester within 10 minutes) was not significantly different between 2% mepivacaine with 1:20,000 levonordefrin and 2% lidocaine with 1:100,000 epinephrine for the central incisor and first molar. However, neither anesthetic agent provided an hour of pulpal anesthesia. |
| Berberich G et al. USA, 2009 [19] | Prospective, randomised, double-blind study | 2% lidocaine and 1:100k epinephrine | Intraoral, infraorbital nerve block | The intraoral, infraorbital nerve block was ineffective in providing profound pulpal anesthesia of the maxillary central incisor, lateral incisor, and first molar. Successful pulpal anesthesia of the canine and first and second premolars ranged from 75%-92% by using 2% lidocaine with 1:100,000 and 1:50,000 epinephrine. However, pulpal anesthesia did not last for 60 minutes. The use of 3% mepivacaine provided a shorter duration of anesthesia than the lidocaine formulations with epinephrine in the canines and premolars. |
| Sreevininith K et al. Thailand, 2017 [57] | Randomised single blind comparative split mouth design | 4% articaine and 1:100k epinephrine | Inferior alveolar nerve block | The use of 4% articaine for the inferior alveolar nerve block was clinically more effective in the onset of subjective and objective anesthesias compared with the use of 4% lidocaine. |
Table 3. Efficacy comparisons of amides in the literature (”>” indicating a higher efficacy and “=” indicating an equal efficacy).

| Comparing efficacy                                      | Literature list reference |
|---------------------------------------------------------|---------------------------|
| Articaine > Mepivacaine                                 | [45], [50]                |
| Articaine > Lidocaine                                   | [31], [42], [43], [48], [16], [44], [32], [52], [53], [56], [57] |
| Articaine > Bupivacaine                                 | [46], [47], [16]         |
| Articaine > Prilocaine                                  | [51]                      |
| Articaine = Lidocaine                                   | [20]                      |
| Lidocaine > Mepivacaine                                 | [19]                      |
| Mepivacaine > Lidocaine                                 | [12], [13]                |
| Mepivacaine = Lidocaine                                 | [22], [23], [24]         |
| Bupivacaine > Lidocaine                                 | [14], [15]                |
| Bupivacaine = Lidocaine                                 | [21]                      |
| Ropivacaine no comparison                              | [18], [17]                |
| Ropivacaine = Articaine                                 | [25]                      |
| Prilocaine > Lidocaine                                  | [16]                      |
| Prilocaine > Bupivacaine                                | [16]                      |
| Mepivacaine + Lidocaine = Lidocaine + Lidocaine         | [26]                      |
| Articaine volume comparison                             | [27]                      |

lidocaine [31], while increasing the concentration of epinephrine, with articaine and lidocaine, did not significantly impact the anesthetic efficacy [14,15]. Three in vivo studies showed that adding a sodium bicarbonate buffer to lidocaine [32,33] and articaine [34], yielded no improvement in efficacy for either. One study [24] investigated the difference between mepivacaine combined with levonordefrin and lidocaine with epinephrine, and concluded that there were no significant differences between them with regard to efficacy.

Other in vivo studies incorporated drugs such as meperidine [35,36] and mannitol [37] to lidocaine, and neither appeared to have a significant influence on anesthetic efficacy. Also, liposomal bupivacaine was assessed, but did not improve the efficacy either [38].

DISCUSSION

The authors noticed that the terms efficiency and efficacy are sometimes incorrectly interchanged. Efficacy is the correct term to be used when assessing the outcome of a product, for instance, or the ability to produce a desired result (e.g. 100% pulpal anesthesia), while efficiency is to be used to assess a process (e.g. how successful is mandibular block anesthesia?) or the state or the quality of being efficient. The mix up is understandable if English is not the native language of the authors, but it should be avoided at all times as it makes the search for papers for a review more complicated.

Based on this review, which covers the past 10 years, the findings suggest that articaine has been researched the most and that it also has the highest efficacy of the amides used in dental local anesthesia. The fact that articaine received so much attention is probably attributable to the fact that before the year 2000, articaine was not available in the USA, whereas in Europe it was already marketed in 1976. As can be derived from tables 1 and 2, 20 of the 31 papers included in our study were conducted in the USA. Although, it was not within the scope of this review paper, nevertheless, the authors are aware of the dubious reputation of articaine with regard to post-operative paresthesia and the discussion about it being manufactured as a 4% solution instead of 2% like lidocaine for dental local anesthesia [16,39,40]. It deserves to be emphasized, however, that in vitro laboratory studies on cell lines have shown that articaine is not the most neurotoxic amide used in dental local anesthesia. One in vitro study, conducted by Mallet et al., tested the toxicity of 6 local anesthetic products on human neuroblastoma cells and found that articaine is the...
least toxic amide [40], while another in vitro study, by Perez-Castro et al., conducted on a neuronal cell-line, concluded that bupivacaine is the most toxic amide [41]. These findings are in contrast to the reported potential adverse effects, published in two review articles, stating that articaine is harmful in high concentrations, such as 4% [39], and may cause paresthesia [40]. It has to be emphasized that in the latter review, it was found that also prilocaine can potentially cause paresthesia [40]. It is noteworthy that the clinical reports about paresthesia and apparent toxicity almost always involve mandibular block anesthesia. However, it seems strange to us that articaine, for instance, would have a high neurotoxic preference for the second branch of the trigeminal nerve only. Since that was not the aim of the present study, this issue will not be discussed further here, but it definitely deserves further in depth attention.

It is our impression that, after having read the papers regarding efficacy of dental local anesthetics, none of the amides studied and used in dentistry guarantee a 100% success, especially not in the mandible. Therefore, one could conclude that perhaps the technique of administration is inefficient and therefore the efficacy is poor. Intraosseous anesthesia could be the key to increase the efficacy of local anesthetics in the mandible.

CONCLUSIONS

From the consulted literature, it is clear that local anesthetics used in dentistry do not show a 100% efficacy, especially not if administered in the mandible or in cases of inflammation (acute irreversible pulpitis). The authors suggest that this opens perspectives to explore more successful ways of administering local anesthesia, in order for the local anesthetic to be more efficacious.

CONFLICT OF INTEREST

One of the authors (JA) is a key opinion leader for Dental Hi Tec®, the French manufacturer of SleeperOne® and Quicksleeper®.

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