Narrative Review

An update on the chemistry, pharmacology and dose calculations of mepivacaine hydrochloride for podiatrists in the United Kingdom

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Abstract: (1) Background: Local anaesthetic agents suppress action potentials in excitable tissues by blocking voltage-gated sodium channels. In doing so they inhibit the depolarisation of nociceptive nerve fibres and so prevent the transmission of pain impulses. UK legislation allows HCPC-registered Podiatrists with POM-A annotation access to six local anaesthetic drugs and two of these with the addition of adrenaline. The use of local anaesthesia has transformed the treatment of nail pathology by Podiatrists. (2) Methods: a narrative review followed a comprehensive literature search. (3) Results: In the UK, the drug of choice in podiatric practice is 3% mepivacaine hydrochloride: it is a good choice of drug for digital anaesthesia. (4) Conclusion: This paper will review the chemistry, pharmacology and dose calculation of mepivacaine, and challenge some of the orthodoxy over the rigid calculation of maximum safe dosages..

Keywords: Mepivacaine, Scandonest, local anaesthesia, pharmacology, podiatry, maximum safe dose.

1. Introduction

Local anaesthesia (LA) is defined as any technique that renders part of the body insensitive to pain or sensation without affecting consciousness (Khan, 2017). LA drugs do this by interrupting neural conduction via the inhibition sodium ion influx through ionophores within neuronal membranes (Malamed, 2019). Their key application is to facilitate pain free treatment (Pope and Brown, 2020). Clinical experience and pharmacological discoveries have allowed the development of safe and efficient practice using a range of different drugs since Halstead's first peripheral nerve block in 1885 (Calatayud, 2003). The history of LA is a fascinating subject in its own right. For those that want to know more about the origins and advancement of this branch of medicine, refer to the chapter by Chuan and Harrop-Griffiths in Hadzic's Textbook of Regional Anesthesia and Acute Pain Management (Chuan and Harrop-Griffiths, 2017), and work by Borthwick (Borthwick, 2001/05) who details the Podiatric history of access to these drugs.

UK legislation allows registered podiatrists with Health and Care Professions Council (HCPC) POM-A annotation (POM-A annotation replaced the 'LA Certificate' in 2016 - see table 1) access to six amino-amide LA drugs, and two of these with the addition of adrenaline (HMRs, 2012). LA use has transformed podiatric practice over the last 60 years, with most infiltrations (outside of the sub-group of UK Podiatric Surgeons) being given as a digital block prior to nail surgery (Maher, 2020). Lidocaine hydrochloride is the most widely used anaesthetic in medicine and dentistry worldwide (Becker, 2012) but mepivacaine hydrochloride (M.HCl) appears to be* the most commonly used agent in UK podiatric practice (*based on the senior author's experience of teaching under- and post-graduate LA theory and practice).

The aim of this paper is to give podiatric colleagues an update on the chemistry and pharmacology of M.HCl, to give an overview of the variability in dose calculations and to challenge some of the orthodoxy over the rigid calculation of maximum safe dosages.   It will be provided as a preprint to obtain wider professional feedback before the next iteration of this work.

Persons exempted	Prescription only medicines to which the exemption applies	Conditions
1. Registered chiropodists or podiatrists against whose names are recorded in the relevant register annotations signifying that they are qualified to use the medicines specified in column 2.	1. Prescription only medicines for parenteral administration that contain— (a) Adrenaline, (b) Bupivacaine hydrochloride, (c) Bupivacaine hydrochloride with adrenaline where the maximum strength of adrenaline does not exceed 1 mg in 200 ml of bupivacaine hydrochloride, (d) Levobupivacaine hydrochloride, (e) Lidocaine hydrochloride, (f) Lidocaine hydrochloride with adrenaline where the maximum strength of adrenaline does not exceed 1 mg in 200 ml of lignocaine hydrochloride, (g) Mepivacaine hydrochloride, (h) Methylprednisolone, (i) Prilocaine hydrochloride, (j) Ropivacaine hydrochloride.	1. The administration shall only be in the course of their professional practice and where the medicine includes a combination of substances in column 2, those substances shall not have been combined by the chiropodist or podiatrist.

Table 1: The Human Medicines Regulations 2012, Schedule 17, Part 3

2. Materials and Methods

For this review, a strategy that involved searching for research evidence via three different sources was adopted:

- Electronic databases - healthcare databases advanced search (HDAS)
- Reference lists
- Google scholar

HDAS allowed access to the following databases, which were considered the give the best chance of identifying appropriate sources:

- AMED (Allied and Complementary Medicine: 1985 - present)
- CINHAL (Cumulative Index to Nursing and Allied Health Lit.: 1981 - present)
- EMBASE (Excerpta Medica Database: 1974 - present)
- MEDLINE (Medical Literature Analysis and Retrieval Online: 1946 - present)

Additionally, NYSORA (New York School of Regional Anaesthesia), the Cochrane Database of Systematic Reviews, the Royal Society of Medicine and hand searches of the Journal of Foot and Ankle Surgery, the Journal of Foot and Ankle Research were searched. Keywords used were: mepivacaine, toxicity, podiatr*, (with searches limited to humans) with Boolean operators AND and OR for: dose, local anaesthesia, anaesthesia, maximum safe dose, and/or history.   Truncation was with n=*[2](+/-1) were additional to the latter keywords as to avoid omitting corresponding articles.   Inclusion criteria or selected articles were:

- Mepivacaine (or Scandonest)
- Maximum safe dose (MSD) or calculations used in local anaesthesia
- Systemic toxicity of mepivacaine in both adults and children

Exclusion criteria were those of studies involving general anaesthesia, studies involving analgesia outside the podiatric field (though some dental articles were of use), studies specific to other LA involvement, and studies transcribed and translated from languages other than English.

3. Review

Histology and Physiology of Action Potentials

The sodium-potassium pump is an electrogenic transmembrane ATPase structure that pumps three sodium ions out of the cell and two potassium ions into the cell, for every single ATP consumed (Pirahanchi et al, 2020). The voltage-gated sodium channel sits across the cell membrane. It is a complex structure composed of a pore-forming protein made up from an alpha subunit and beta subunits. The alpha subunit is comprised of four domains, each containing six segments that wrap round to form a bell-shaped central channel. The channel itself is formed by the S5 and S6 segments and the short loops of amino acids that link them (Taylor and McLeod, 2019).

First described by Hodgkin and Huxley (Hodgkin and Huxley, 1952), three conformational states of the channel exist: resting, open, and inactivated. The membrane potential is approximately -70 mV in the resting state: this charge is generated by the movement of positively charged potassium ions from outside the neuron (to the inside) via the sodium-potassium pump and the negatively charged anions that remain within the cell (Taylor and McLeod, 2019). After a stimulus, the sodium channels open during depolarisation, allowing a rapid influx of sodium ions down the electrical and chemical gradient which leads to channel inactivation. Impulse generation along a nerve causes rapid movement of sodium ions inwards and potassium ions outwards. An action potential is generated when the membrane potential increases to a threshold of -55 mV. Thereafter, the sodium/potassium pump restores the excitable state of the resting membrane potential (Malamed, 2019).

Chemistry and Pharmacology of Local Anaesthetics

The local anaesthetic molecule consists of 3 components: a lipophilic aromatic ring, an intermediate ester or amide chain, and a terminal hydrophilic amine (Columb and MacLennan, 2007; Taylor and McLeod, 2019). Each of these contribute distinct properties to the molecule (see fig 1). The intermediate linkage provides a basis for classification and also determines the pattern of biotransformation: amino-esters are hydrolysed by plasma esterases, whereas amino-amides are bio transformed hepatically. The terminal amine may exist in a lipid-soluble tertiary form (3 bonds) or as a positively charged, water-soluble quaternary form (4 bonds) (Becker, 2006).

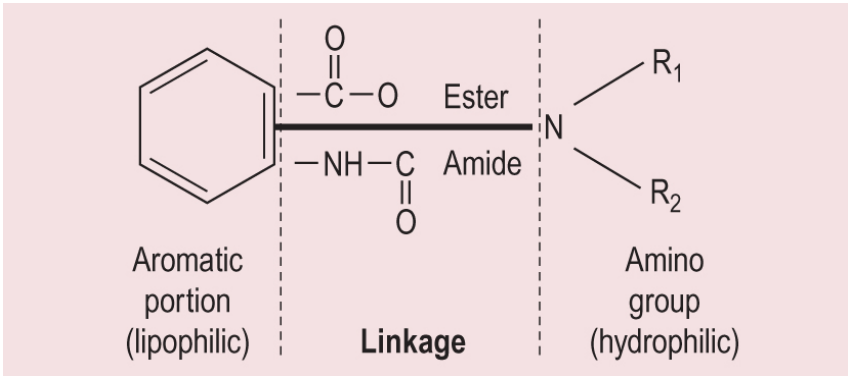
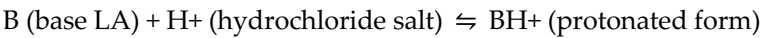



Figure 1: basic structure a local anaesthetic (Metcalfe and Reilly, 2010)

For the local anaesthetic base to be stable in solution, it is formulated as a (water-soluble) hydrochloride (acid) salt.   This is represented by:

B (base LA) + H+ (hydrochloride salt) ⇋ BH+ (protonated form)

Following the diffusion of the non-ionised base through the neural (cell) membrane into the axoplasm, it re-protonates and enters the voltage-gated sodium channels to suppress action potentials by inhibiting the influx of sodium ions, preventing the ion channel from assuming an active or ''open'' state (Becker, 2006; Taylor and McLeod, 2019).   The quaternary form of a LA will not penetrate the neuron and therefore the time for onset of anaesthesia is predicated on the proportion of molecules that convert to the tertiary, lipid-soluble structure when exposed to physiologic pH (7.4) (Becker 2006).

The pKa of a molecule represents the pH at which 50% of the molecules exist in the tertiary, lipid-soluble form and 50% in the quaternary, water-soluble form.   The Henderson-Hasselbalch equation (Log [non-ionized drug]/[ionized drug] = pH – pKa) allows us to calculate the ratio of those two states (Tsuchiya, 2007).   The acidic environment of inflamed tissues favours the quaternary, water-soluble configuration and accounts for difficulty in anesthetising inflamed or infected tissues as fewer molecules exist as tertiary lipid-soluble form that penetrates nerves.   In these situations, M.HCl with pKa of 7.6 (this pH value varies in different sources) - compared to bupivacaine with a pKa of 8.1 - is more likely to provide effective anaesthesia (Becker, 2006).

The aromatic ring improves the lipid solubility of the compound.   Greater lipid solubility enhances diffusion through nerve sheaths and neural membranes: this correlates with drug potency because a greater portion of an administered dose can enter neurons (Becker, 2006).   Circulating LA is bound by plasma proteins (α1-acid glycoproteins). The duration of action of each LA agent depends on plasma protein binding and correlates with a compound's affinity for the protein of the sodium channels.   Bupivacaine exhibits 95% protein binding compared to 55% or 77% (references vary) for M.HCl, hence the difference in their duration of neural blockade (Becker, 2012).   Both potency and duration of action also correlate with increasing molecular weight but also with an increased onset time and potential for severe systemic toxicity (Butterworth, 2017).   The duration of action can also be affected by:
- Individual response to the drug
- Accuracy in deposition of the drug
- Status of tissues at the site of drug deposition (vascularity, pH)
- Anatomic variation
- Type of injection administered (field or nerve block)
- Addition of a vasoconstrictor (Malamad, 2019)

The addition of a vasoconstrictor, such as epinephrine (adrenaline) prolongs the anaesthetic effect by reducing the systemic distribution of the agent.   Multiple sources state that vasoconstrictors must not be used to produce ring-block of an extremity (e.g.   a finger or toe) because they may cause prolonged ischaemia and gangrene (Broomhead, 2020). The senior author regularly adds 2% lidocaine with 1/200,000 epinephrine to long acting LA agents such as levo-bupivacaine and ropivacaine for peripheral nerve blocks.   This is interesting topic and worthy of a further commentary in another paper (spoiler alert: the risks are likely exaggerated).   The chapter by Broomhead (2002), in the most up to date general podiatric text, has outline on all the common amino-amides available to podiatrists apart from M.HCl.

Amino amides are metabolised by phase I and II hepatic cytochrome P450.   Through phase I, hydroxylation, N-dealkylation and methylation occurs.   Phase II includes conjugation with amino acids such as glycine (Columb and MacLennan, 2007). | 176
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Mepivacaine | 180

In 1956 and 1957, Af Ekenstam et al developed, and later discussed, mepivacaine and bupivacaine (and others) (Af Ekenstam et al., 1956; Af Ekenstam et al., 1957), the former being the subject of this review.   Mepivacaine is a white, crystalline, poorly soluble, odourless powder, with a melting point 1510C and a molecular weight and 246.4 Da; 2610C/285.5 Da as the readily soluble mepivacaine hydrochloride (Gordon, 1960).   Its properties are summarised in table 2.   This is a summary from product monographs, Patient Information Leaflets (PILs, e.g., www.drugs.com) and Summary of Product Characteristics (SPCs) found through the literature search.   As noted below, the primary evidence source is often missing or unclear in such documentation.   PILs and product data are supported by the pharmaceutical industry.   In communication with the Chief Executive Officer of one of these companies, it was stated that these guidelines are for educational purposes only and are not intended for medical advice, diagnosis and treatment (unreferenced on request). | 181

Chemistry	Mepivacaine was the second amide local anaesthetic to be clinically introduced.
	$C_{15}H_{22}N_2O.HCl$: 1-methyl-2', 6'-pipecoloxylidide monohydrochloride is a tertiary amide salt. The hydrochloride is easily soluble in water, stable in solution, very resistant to acid and alkaline hydrolysis, and can be repeatedly boiled without decomposition.
Brand names	Scandonest (UK), Carbocaine (US)
Available concentrations	1%, 1.5%, 2% with epinephrine 1:100,000, 3% solutions
Dose mg/kg and total	Adults: 4.4, 5, 6, or 6.6 (varies on the reference)
	The MSD for healthy adults should not exceed 400mg at one treatment
	Children 3 or 5/6 (varies on the reference)
	Concentrations of less than 2% for those under 3 years of age
Onset time	Medium (10) minutes
pH/pKa	4.5 to 6.8 / 7.6 or 7.7 (varies on the reference)
Onset time	Medium (10) minutes
Duration of anaesthesia	Medium (120-180) minutes
Peak plasma levels	10-20 minutes
Relative toxicity	Low (0.77)
Potency	Medium
Mean half-life	1-2 hours
Protein binding	55 or 77% (varies on the reference)
Contra-indications and precautions	Mepivacaine has been shown to be less toxic (systemic and neural tissue) than lidocaine in adults and provides a somewhat longer duration of action than lidocaine because of a less pronounced vasodilatory effect. Injection of repeated doses of mepivacaine may cause significant increase in blood levels with each repeated dose due to slow accumulation of the drug or its metabolites, or due to slower metabolic degradation than normal. Tolerance varies with the status of the patient.

	Debilitated, elderly patients, acutely ill patients, and children should be given reduced doses commensurate with their weight and physical status.
	Mepivacaine should be used with caution in patients with a history of severe disturbances of cardiac rhythm or heart block. Young children should be given minimal doses of each agent. Changes in sensorium such as excitation, disorientation, drowsiness, may be early indications of a high blood level of the drug and may occur following inadvertent intravascular administration or rapid absorption of mepivacaine.
	Mepivacaine is contraindicated in patients with a known hypersensitivity to amide-type local anaesthetics. The binding affinity of mepivacaine in serum is reduced in the presence of bupivacaine. Displacement of mepivacaine by bupivacaine was observed when an $\alpha1$ acid glycoprotein solution was studied via classic competitive inhibition. Allergic-type reactions are rare and may occur as a result of sensitivity to the local anaesthetic or to other formulation ingredients. These reactions are characterized by signs such as urticaria, pruritus, erythema, angioneurotic oedema (including laryngeal oedema), tachycardia, sneezing, nausea, vomiting, dizziness, syncope, excessive sweating, elevated temperature, and possibly, anaphylactic-like symptoms (including severe hypotension). Patients allergic to methylparaben or para aminobenzoic acid derivatives (procaine, tetracaine, benzocaine, etc. have not shown cross-sensitivity to amino-amides agents.
	Mepivacaine crosses the placenta: there is considerable transfer of mepivacaine across the placenta after maternal doses and the ratio of foetal to maternal concentrations is about 0.7. Although neonates have a very limited capacity to metabolise mepivacaine it appears they are able to eliminate the drug. On the basis of long usage, anaesthetics of the mepivacaine type are considered to be reasonably safe for use on pregnant women.
	Retrospective studies of pregnant women receiving local anaesthetics for emergency surgery early in pregnancy have not shown that local anaesthetics cause birth defects. However, no controlled studies have been carried out in pregnant women. Moreover, no animal reproduction studies have been performed with mepivacaine. Therefore, caution should be taken before administering this anaesthetic during early pregnancy. The mutagenic potential of mepivacaine has not been evaluated. It is not known whether local anaesthetics are excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when mepivacaine is administered to a nursing woman.
Drug interactions	https://www.drugs.com/drug-interactions/mepivacaine.html https://www.medicinescomplete.com/#/content/bnf/_390463595_interactions.

Table 2: M.HCl Summary - Metcalfe and Reilly, 2010; Multrum, 2020, and PILs/SPCs   195

M.HCl is available in various percentages for dental, epidural and peripheral nerve 197
blockade (Fenten et al, 2015; Lopez et al, 2015; Maladad, 2019). After deposition of the 198
drug as close to the nerve as possible, the solution diffuses three-dimensionally according 199
to prevailing concentration gradients. A portion of the injected local anaesthetic diffuses 200
into the nerve. M.HCl is rapidly metabolized, with only a small percentage of the anaes- 201
thetic (5-10%) being excreted unchanged in the urine. Because of its amide structure it is 202
not detoxified by the circulating plasma esterases; the liver is the principal site of metab- 203
olism with over 50% of the administered dose being excreted into the bile as metabolites. 204
Most of the metabolized mepivacaine is resorbed in the intestine and then excreted into 205

the urine since only a small percentage is found in the faeces. The principal route of excretion is via the kidney (Brockmann, 2013; Meffin, 1973).

Systemic adverse reactions involving the central nervous system (CNS) and the cardiovascular system (CVS) usually result from high plasma levels due to excessive dosage, rapid absorption, or inadvertent intravascular injection. Reactions involving the CNS are characterized by excitation and/or depression. Nervousness, dizziness, blurred vision, or tremors may occur followed by drowsiness, convulsions, unconsciousness, and possibly respiratory arrest. Since excitement may be transient or absent, the first manifestations may be drowsiness merging into unconsciousness and respiratory arrest. (Septodont, 2009).

A small number of reactions may result from hypersensitivity or diminished tolerance to normal dosage on the part of the patient. Allery to mepivacaine, though rare, has been reported (Diaz, 1975; Hiyoshi, 1978; Falace and Hill, 1985; Gonzalez-Delgado et al, 2006; Venemalm et al, 2008; Estrada et al, 2011; Sharma, 2013). Such reactions are characterized by cutaneous lesions of delayed onset or urticaria, oedema and other manifestations of allergy.

Maximum Safe Dose

As with all medications and drug use, a full medical history needs be obtained prior to deliverance to provide safe and effective administration as to avoid systemic toxicity and adverse drug reactions (ADRs) (Kihm, 2011; Columb and MacLennan, 2007; Rosenberg, 2004). The safety and effectiveness of mepivacaine depend upon proper dosage, correct technique, adequate precautions, and readiness for emergencies. The lowest dose that results in effective anaesthesia should be used to avoid high plasma levels and possible adverse effects. The adverse effects of LA may be:

- Local (e.g. transient neurologic syndrome or local trauma)
- Regional (e.g. hypotension after extensive neuraxial block)
- Systemic, after accidental intravenous injection or cumulative overdose (Reynolds, 2005):

For any given LA agent, the dose given is equal to the volume of drug given multiplied by its concentration:

Dose (mg) = volume (ml) x concentration (mg/ml^{-1})

A pharmacological maximum safe dose (MSD) is the largest dose of a medicine or drug consistent with safety: one thought to have an acceptable risk:benefit profile (Merriam-Webster, 2020). Maximum amounts were originally recommended as endpoint values, for example 200mg of lidocaine plain (incidentally this is much lower than in the US where the MSD is 300mg) but manufacturer package inserts now also include LA dose recommendations that take the patient's body mass into consideration (Kihm, 2011) - often without reference to an original source (Columb and MacLennan, 2007; Rosenberg, 2004). Much of evidence for MSD and toxicity is based on extrapolation from animal studies and case reports rather than from higher level evidence from clinical trials (Moore et al., 1977; De Jong, 1994).

UK Podiatry departments often produce local guidelines to direct and support clinical practice. Such protocols may include such statements as:

"When administering local anaesthesia, the weight and maximum safe dosage must be calculated and recorded on the nail surgery treatment record using the following to assist: adult maximum safe dose is 400mg (i.e. 6.5 x 2.2ml cartridges) or 6mg/kg in 24

hours.   3% solution contains 30mgs/ml therefore 13.3ml of 3% solution = 400mg" (Torbay and South Devon NHS Foundation Trust, 2017).

6.5 x 2.2ml cartridges would in fact give a total dose of 429mg, but, in practice, it is almost impossible to expel all the solution via a dental syringe (see fig 2).
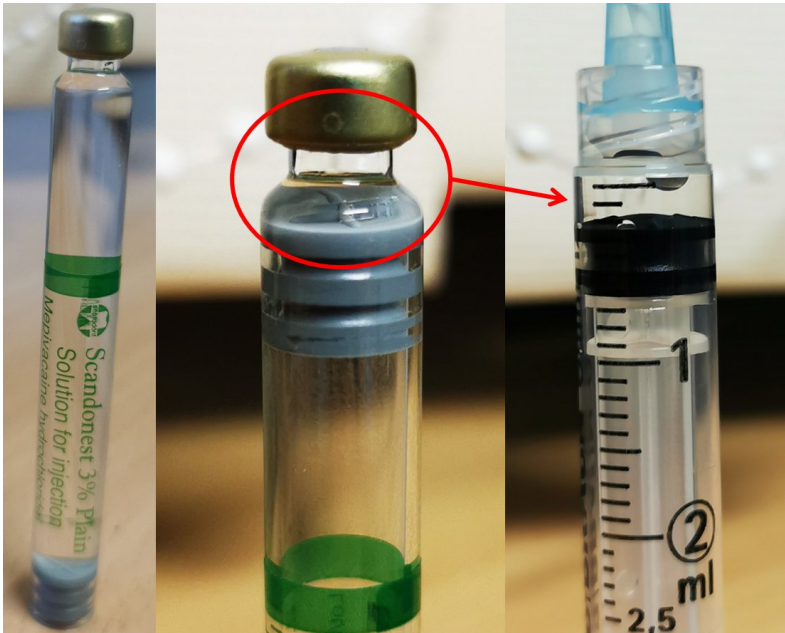
Figure 2: the last 0.2ml of a 2.2 ml Scandonest cartridge extracted via a Luer syringe

The maximal adult doses for M.HCl, and the dose per body weight, vary widely from source to source for both total dose and dose by weight (see table 3), with another (unverified) internet source at 500mg as an MSD.   In UK podiatric practice, the figures quoted above of 6mg/kg - and an upper maximal of 400mg - are commonly used but often without reference, and rarely quoting an original reference (Reilly Rule 101 – always find the original source).

Reference	Dose (mg/kg^{-1})	Total Dose (mg)	Notes
Septodont, 2013	4.4	300	0.15ml/kg^{-1}
Crawford, 2012	5	300	7 mg/kg^{-1} with vasoconstrictor
University of Iowa, 2019	4.5-5	400	1g in 24 hours
Taylor and McLeod, 2020	5	Not given	7 mg/kg^{-1} with vasoconstrictor
Metcalfe and Reilly, 2010*	6	400	No original reference
Malamed, 2019	6.6	400	The same with vasoconstrictor
Lee, 2017	Not given	550	Based on FDA figure

Table 3: the variety of adult MSDs (*fails Reilly Rule 101)

Williams and Walker (2014) note that dose limits (mg/kg^{-1}) for local anaesthetics based on body weight are well-established (the points above should be noted), but that the calculation of the maximum safe volume (ml) of a given agent and formulation can be complex and subject to errors.   Working out the MSD can be done using various methods.

Podiatrists commonly use the following 2-step approach (Metcalfe and Reilly, 2010):

1. drug MSD (in mg/kg) x patient weight (in kg) = total dose (MSD in mg)
2. MSD (in mg) = volume (in ml) x drug percentage (as w/v) x 10

A worked example using 3% M.HCl for a 50kg patient would be:

MSD (mg): 50 (kg) x 6 (mg/kg-1) = 300mg

300mg = unknown volume x 3% x 10

MSD (volume) = $\frac{300}{3 \times 10}$ = **10ml**

An alternative method is to use the drug concentration in a one-step method (Williams and Walker, 2014):

$$maximal\ volume\ (ml) = \frac{body\ weight\ (kg)\ x\ maximum\ dose\ (mg/kg)}{concentration\ (mg/ml)}$$

A worked example using 3% M.HCl (a 3% solution is a 30mg/ml^{-1} concentration), for our 50kg patient would be:

$$\frac{50\ x\ 6}{30} = \textbf{10ml}$$

The upper patient weight for drug calculations is usually considered to be 70kg. As Metcalfe and Reilly note: "…metabolic capacity is not increased in a linear manner with weight. Although calculations based upon body surface area would provide a more accurate measure of an appropriate dose, body weight is the normal factor against which MSDs are calculated in podiatry. As a guide, in healthy patients a maximal body weight of 70 kg is taken for the calculation of maximum safe dose of local anaesthetic. A person weighing 140 kg does not have a liver mass twice that of someone weighing 70 kg". The British National Formulary (BNF, 2020) states that doses should be calculated based on ideal body weight, which for most adults is 60–75 kg, depending on height and sex.

M.HCl use in children is worthy of special note. Various formulas exist to calculate dosing in infants and children. Clark's rule equation is defined as the weight of the patient in pounds divided by the average standard weight of 150lb multiplied by the adult dose of a drug to equal the paediatric medication dose (Delgado et al, 2020). Using the two-step, worked example above for our 50kg patient (110lbs), this would be:

$\left(\frac{110}{150}\right) x\ 400$ = 293mg

293mg = ? volume x 3% x 10

MSD (volume) = $\frac{293}{3 \times 10}$ = **9.7ml**

Young's rule, based on the age of the child, is (Wade and Martinez, 2020):

adult dose x (age ÷ (age+12)) = child's dose

For our 50 kg patient who is 14 years old, this would be:

400 x (14 ÷ (14 + 12) = 215mg

213mg = ? volume x 3% x 10

MSD (volume) = $\frac{215}{3x10}$ = **7.2ml**

Septodont (2013) recommended that "*the quantity to be injected should be determined by the age and weight of the child and the magnitude of the operation. The average dosage is 0.75mg/kg = 0.025ml of mepivacaine solution per kg body weight. Maximum recommended dosage: do not exceed the equivalent of 3mg mepivacaine/kg (0.1ml mepivacaine/kg) of body weight.*" They caution against use in children below 4 years of age (i.e. weighing at least 20 kg). Note that this is half of the oft-quoted adult dose. One must also remember the physiological discrepancy between a would-be prop forward 10-year-old and a 15-year-old that needs to run around in the shower to get wet.

The College of Podiatry Nail Surgery guidelines (Gohil, 2019) quote an earlier statement from the then Director of Education, Ashcroft in 2003 and notes that M.HCl is not licenced for podiatric use in children because the manufactures did not apply for such a licence. Ashcroft went on to reassure members that "*The Society would consider the administration of Scandonest, suitably dose adjusted for children, to be acceptable professional practic*e."

4. Discussion

A brief overview of LA theory has been presented together with a more detail update on the drug mepivacaine. It was surprising to see the discrepancies in the literature over the various chemo-physical properties of the drug, as well as variability is MSD values. This cuts across the traditional way LA theory is taught in UK podiatric practice.

Malamed (2019) - in a key dental anaesthetic text - suggests that the properties of an LA include:
- It should not be irritating to the tissue to which it is applied
- It should not cause any permanent alteration of nerve structure
- Its systemic toxicity should be low
- It must be effective regardless of whether it is injected into the tissue or is applied topically to mucous membranes
- The time of onset of anaesthesia should be as short as possible
- The duration of action must be long enough to permit completion of the procedure yet not so long as to require an extended recovery

The choice of LA in peripheral nerve blockade is (or should be) determined by desired speed of onset, block intensity, and duration of anaesthesia and analgesia (Eng, 2014) and therefore Podiatrists should consider the use other agents where indicated (Uddin and Reilly, 2008). It might be argued that this is less important for the humble digital block and therefore in this, 3% M,HCl is a good choice.

One of the factors affecting the duration of peripheral nerve block is the total dose of LA: dose being the product of volume and concentration (Fenten et al, 2015). Their study used 1% and 1.5% mepivacaine preparations. Gordon et al., (1960) used M.HCl on 230 humans as a 2% concentration for peripheral nerve blockade and favoured it over procaine and lidocaine for its quicker onset and duration (3-4 times longer), demonstrating satisfactory and prolonged anaesthesia without the addition of vasoconstrictors. No patients recorded toxic manifestations or tissue irritation in this study.

Maher et al (2008) note that the Department of Health (DoH) expressed concerns re- 384
garding the non-medical administration of local anaesthetics when access for first granted 385
and placed restrictions on the maximum dosage which could be administered in a 24-hour 386
period. They go on to summarise earlier papers that state the Society of Chiropodists 387
successfully fought against this restriction and in 1980 the restriction was lifted by the 388
DoH, allowing podiatrists to use their clinical judgement in deciding dose.   Note that the 389
24-hour restriction still pervades in the Podiatric lexicon (in many, readily searchable Po- 390
diatry Department protocols) some 40 years later without consideration of the true half- 391
life of a given drug: 1-2 hours for M.HCl.   In fact, continuous infusion is common with 392
anaesthetic practice for post-operative pain control.   Buettner et al (1989) report on cath- 393
eter-induced axillary plexus blocks performed in 17 patients using 400 mg of M.HCl every 394
two hours.   One patient had 12 injections of 400mg of mepivacaine in 24 hours - an over- 395
all dosage of 4800mg - without evident signs of central nervous system or cardiovascular 396
toxicity.   Most LAs will not produce CV toxicity until the blood concentration exceeds 397
three times that necessary to produce seizures. 398

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Rosenberg et al (2004) suggest that MSD calculations referring to the total amount of 400
the drug per body mass of the patient do not account for the site of administration and 401
advocated abandoning blanket maximum doses in favour of a therapeutic range.   They 402
emphasized the importance of variables such as age, site and speed of injection, and the 403
presence of systemic disease (especially hepatic, renal, and cardiac disease).   Butter- 404
worth (2017) notes that there is no scientific justification for the exact dose recommenda- 405
tions or the mg/kg MSD guideline, and that any recommendation on the MSD can be valid 406
only in reference to a specific nerve block procedure.   Instead, he advocates more a clin- 407
ically adequate 'safe dose range'.   Moore et al., (1997), for example, conducted a trial on 408
humans using up to 500mg of M.HCl with no clinically significant outcome of systemic 409
toxicity regardless of age, gender and weight.   In contrast, Reynolds (2005), while sup- 410
porting variation based on individual patient factors, still recognized the importance of 411
imposing clear limits. 412

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Rosenberg et al (2004) stated that MSD recommendations are published by the man- 414
ufacturer giving a false assumption that the manufacturers are responsible for the guide- 415
line.   However, as per direct communication with the CEO of IBM Watsons international 416
pharmaceutical industry, this is not the case and the guidelines can be traced to their orig- 417
inal source being transcriptions from medical professionals.   This is not clearly stated 418
upon distribution and raises further debate for MSD recommendations being evidence 419
based.   De Luke et al (2018) highlighted the discrepancy in recommendations for maxi- 420
mum doses and in the teaching of safe maximum doses in dental schools with key texts 421
and manufacturer guidelines giving different figures.   They concluded that students 422
should be made aware that there is more than one standard, and that teaching should 423
emphasize sound medical and pharmacologic principles, a concern echoed in this paper. 424
The order of peak plasma concentration after a single dose is: intrapleural > intercostal > 425
lumbar epidural > brachial plexus > subcutaneous > sciatic > femoral.   (Taylor and 426
McLeod, 2019).   The implication from the literature is that pedal blocks – being even more 427
distal – have slower systemic uptake and reduced potential for acute overdosage. 428

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There are currently no studies suggesting a safe prediction of toxicity of LA in hu- 430
mans (Kihm, 2004) and mepivacaine toxicity is rarely reported, (Luduena, 1960; Manas- 431
sero, et al 2014) however, it is important to understand the properties, mode of action and 432
clinical signs of toxicity.   When administered simultaneously, mepivacaine and bupiva- 433
caine interact synergistically to produce a higher-than-expected free concentration of 434
mepivacaine.   This interaction increases the risk of toxicity (Hartrick, 1984).   Saraghi et 435
al., (2015) provided data specific to LA systemic toxicity (LAST) in children.   They com- 436
pared LA agents and found that using M.HCl 3% plain solution it is easier to exceed the 437

recommended MSD due to its potency. Inclusively, they mention Clark's rule and the Rule of 25 to calculate the MSD in children.

For all the variability in dose calculations, Malamed (2019) states that "local anaesthetics are the safest and the most effective drugs available in all of medicine for the prevention and the management of pain" with the added proviso "when used properly". Koay (2002) also feels that when used properly, LA agents are very effective at producing a transient regional loss of sensation with minimal risks of adverse effects. M.HCl remains the most popular choice amongst UK podiatrists for nail surgery and will likely remain so. M.HCl's 3% presentation makes for a potent and popular LA agent but whose use in UK Podiatry has been largely confined to digital blocks, though some Podiatrists and Podiatric Surgeons have utilised M.HCl for a wider variety of foot and ankle pathology and diagnostic investigative purposes (Maher et al ,2008).

5. Conclusions

In the UK, the podiatric LA drug of choice is mepivacaine hydrochloride. It is a good choice for digital anaesthesia, and its use has transformed the treatment of nail pathology. It is almost equally popular in Dentistry but less so in the world of (medical) anaesthesia where less cardio-toxic agents such as levo-bupivacaine and ropivacaine are the first line agents for peripheral nerve block because of their reduced cardiotoxicity. The effectiveness of a given LA is influenced by the dose, site of administration, additives, temperature, and changes in neural susceptibility: something not emphasised in Podiatric education. A rigid MSD calculation without consideration of patient factors is therefore open to critique and made more difficult by the variability of maximal dosage presented in the literature, and differing calculations available to estimate the same.

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