

Effect of bupivacaine and adjuvant drugs on skeletal muscle tissue oximetry and blood flow: an experimental study

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Ann-Kristin Schubert¹
Stefan Müller¹
Hinnerk Wulf¹
Thorsten Steinfeldt^{1,2}
Thomas Wiesmann¹

¹Department of Anesthesiology and Intensive Care Medicine, University Hospital Marburg, Philipps University of Marburg, Marburg, Germany;

²Department of Anesthesiology and Intensive Care Medicine, Diakoniekrankenhaus Schwäbisch Hall, Schwäbisch Hall, Germany

Background: Skeletal muscle microvascular blood flow plays a critical role in many myopathologies. The influence of bupivacaine and adjuvants on skeletal muscle microvascular perfusion and tissue oximetry is poorly understood but might be a relevant risk factor for myopathies after local anesthetic administration. The aim of this experimental study was to determine the effects of bupivacaine alone or in combination with epinephrine or clonidine on skeletal muscle perfusion and tissue oximetry.

Methods: Combined tissue spectrophotometry and Laser-Doppler flowmetry and tissue oximetry were used to assess local muscle blood flow in anesthetized pigs after topical administration of test solutions (bupivacaine, bupivacaine with epinephrine or clonidine, saline). Measurements were performed for up to 60 mins.

Results: The application of bupivacaine alone did not alter relative muscle blood flow significantly, whereas the addition of epinephrine or clonidine to bupivacaine resulted in a significant reduction of relative muscle blood flow at T30 and T60. However, bupivacaine resulted in a significant decrease of tissue oximetry values when compared to saline control group at T30 and T60. The application of bupivacaine combined with clonidine or epinephrine resulted in no significant reduction of tissue oximetry when compared to bupivacaine alone.

Conclusion: Bupivacaine alone results in a significant decrease of tissue oximetry in skeletal muscle which is not increased by the addition of epinephrine or clonidine despite further reductions of microcirculatory perfusion. Overall, bupivacaine alone or with adjuvants does produce local muscle ischemia for which pathological consequences need to be addressed in further studies.

Keywords: bupivacaine, adjuvants, tissue perfusion, tissue oxygenation, ischemia

Introduction

Local anesthetics (LAs) reversibly inhibit specific Na⁺ channels in neurons and prevent propagation of action potential along peripheral nerves. In regional anesthesia, LAs are injected in proximity to the target nerve in the surrounding tissue. In most peripheral nerve blockades, the surrounding tissue consists of skeletal muscle. After injection, the LA molecules spread and diffuse through the tissue and finally reach their intracellular target structure within neural tissue. A larger proportion of the LA spreads through the surrounding tissue and exhibits myotoxic and vasoactive effects.¹⁻⁵ The consequences of this inadvertent spread are known as myotoxic effects of LAs on skeletal muscle which are well described especially for bupivacaine^{1,3,6} by increased intracellular calcium levels and a breakdown of intracellular homeostasis⁷ as well as inhibition of

Correspondence: Thomas Wiesmann
Department of Anesthesiology and Intensive Care Medicine, University Hospital Marburg, Philipps University of Marburg, Baldingerstraße, Marburg 35033, Germany
Tel +49 6 421 586 9362
Email wiesmann@med.uni-marburg.de

mitochondrial function.⁸ Interestingly, the long-known vasoactive effects of LAs and adjuvants are still poorly understood. Regarding the vasoactive potential, LAs as well as the adjuvant epinephrine both have a negative impact on nerve blood flow.⁹ The question as to whether LAs also have a negative impact on skeletal muscle blood flow has not been adequately addressed yet. Skeletal muscle vasculature is rich in $\alpha 1$ and $\alpha 2$ adrenoreceptors that both mediate a profound vasoconstrictive response and resulting tissue ischemia after activation. Thus, epinephrine as well as the widely used adjuvant clonidine might result in profound muscle vasoconstriction and resulting tissue hypoxemia. These potential alterations in blood might be a potential additional risk factor for muscle damage after peripheral regional anesthesia.

The aim of this study is to evaluate the effect of bupivacaine alone or in combination with the adjuvants epinephrine and clonidine on skeletal muscle blood flow and tissue oximetry using a large-animal model.

Methods

General anesthesia and tissue preparation

Experiments were approved by the local authorities (V 54-19c 2015 (1) MR 20/13 Nr 92/2011, Regional Board of Animal Welfare, Gießen, Germany) and performed in accordance with the current national laws on animal protection. All study results are presented according to the ARRIVE-Guidelines for reporting Animal Studies.¹⁰

Twenty-two young female domestic pigs (*Sus scrofa domestica*, German Landrace) were used in this experimental study. The animals had a mean age of four months and a mean bodyweight of 30 kg (range: 27–41 kg). In the anesthetized pigs, 40 quadriceps femoris muscle areas were exposed surgically and randomly assigned to one of the four groups. Thus, 10 muscles per group were included in the final data analysis.

Standardized general anesthesia and euthanasia were performed as described in previous studies.^{5,11,12} In brief, after premedication consisting of diazepam and ketamine, general anesthesia was induced using propofol (2–3 mg kg⁻¹) and sufentanil 0.2–0.3 mg kg⁻¹. Following endotracheal intubation, general anesthesia was maintained using propofol 0.2 mg kg⁻¹ min⁻¹ and sufentanil 0.5 μ g kg⁻¹ min⁻¹. Mechanical ventilation settings were modified to achieve an end tidal CO₂ of 35–45 mmHg. Inspiratory oxygen fraction (FiO₂) was set to 0.3 to simulate clinical conditions comparable to low-flow oxygen supplementation in spontaneous breathing.

An arterial catheter for invasive blood pressure monitoring was inserted via left femoral artery using ultrasound guidance.

To achieve standardized conditions and minimize motion artifacts, pancuronium bromide 0.1 mg kg⁻¹ was administered repeatedly during the study period.

The pharyngeal temperature was monitored to estimate core body temperature and kept at a constant level using warm blankets. All anesthetized animals were placed supine. Quadriceps femoris muscles were exposed surgically without damaging the muscle fascia. For proper positioning of the probe for combined measurement of spectrophotometry and Laser-Doppler flowmetry, a custom-made application device was fixed with sutures on the surrounding tissue to achieve a constant application pressure and measuring volume during the study period.

At the end of the experiments, euthanasia was performed under general anesthesia using potassium chloride (4 mmol kg⁻¹).

Combined tissue spectrophotometry and Laser-Doppler flowmetry

Measurements of muscle blood flow (flow in arbitrary units (AU)), tissue oxygen saturation (SO₂ in %) and tissue hemoglobin level (rHb in AU) were performed using Laser spectrophotometry (O2C, Lea Medizintechnik, Gießen, Germany) as described before.^{5,13} The system and probe were calibrated according to the manufacturer's protocol.

Tissue blood flow is measured using a Laser Doppler Flowmetry (Figure 1). The tissue spectrophotometry unit calculates hemoglobin content and tissue oximetry. The measuring volume within the tissue is illuminated with coherent Laser light of 500–630 nm wavelength and 30 mW power through a fiber-optic cable. Probe geometry allows for the detection of blood flow at approximately 1–2 mm depth by analyzing backscattered light. By fitting measured spectra with spectra of known tissue saturation values, the percental oxygen saturation of the capillary and post-capillary microvasculature is calculated by the O2C device as SO₂. The tissue hemoglobin value (rHb) represents a hemoglobin concentration per tissue volume (displayed as arbitrary unit, AU). It is dependent from microvessel density, venous filling and microvessel shunting. Thus, the rHb is suitable to analyze venous congestion in the measured tissue volume.

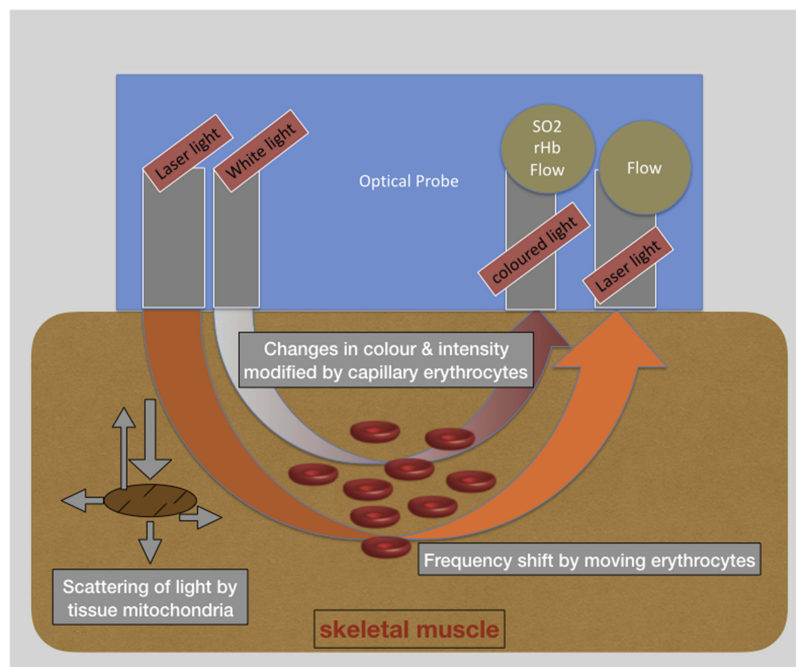


Figure 1 Schematic illustration of O2C device (SO₂, tissue oxygen saturation; rHb, tissue hemoglobin).

The O2C probe was applied to the exposed muscle tissue with a custom-made application device as shown in **Figures 2** and **3** and fixed with sutures for constant application pressure. Probe fixation and measurements were established 10 mins before the intervention to create optimal measurement stability of the system.

After this time span, 1 mL of the randomly assigned test solution was applied to the underlying muscle next to the probe immediately after data were acquired at T0 (timepoint 0 mins).

Study groups

One mL of the test solutions was randomly applied topically to the muscle:

Bupivacaine 0.5% isobaric (group Bupi) (Carbostesin, B. Braun, Melsungen, Germany), Bupivacaine 0.5% isobaric with Epinephrine 1:100.000 (group BupiEpi) (Suprarenin, Sanofi-Aventis, Frankfurt, Germany).

Bupivacaine 0.5% isobaric with 3.75 µg/mL Clonidine (group BupiCloni) (Clonidin ratiopharm, Ratiopharm, Ulm, Germany).

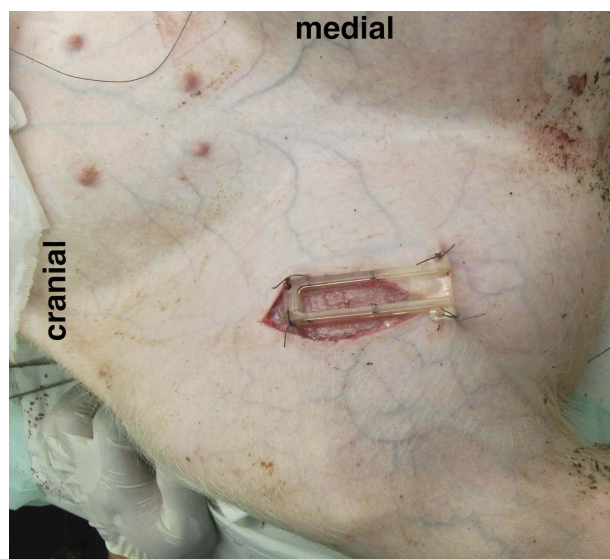


Figure 2 Mounting device (O2C probe removed).

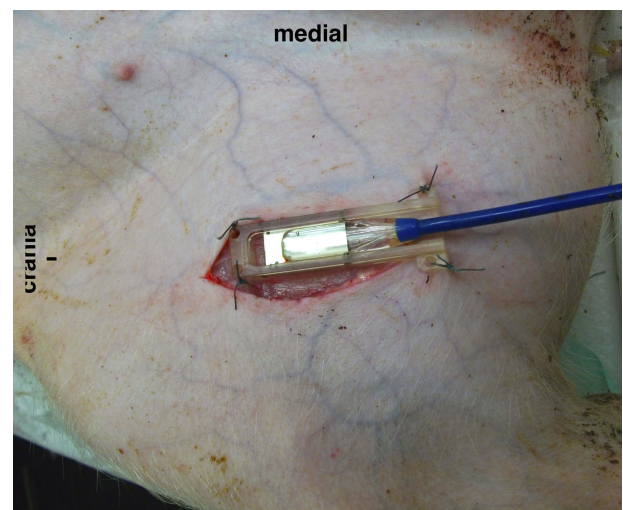


Figure 3 Mounted probe in situ.

Saline was used as a negative control group: (saline 0.9% isotonic (NaCl, B. Braun, Melsungen, Germany).

All test solutions were prepared fresh immediately before the experiments in a standardized manner using single-dose vials. The concentration of epinephrine and clonidine resemble the highest concentrations of both drugs used as adjuvants to LAs in clinical setting.¹⁴

Outcome parameters

The O2C data concerning tissue blood flow (Flow, AU in arbitrary units), tissue oximetry (SO₂, in percent) and hemoglobin (rHb, in AU) were recorded at the time points T0, T5, T10, T15, T20, T25, T30, T40, T50 and T60 mins and transferred to an Excel sheet (Microsoft Excel for Mac, Redmond, USA). Systemic hemodynamic parameters as mean arterial blood pressure, heart rate and oxygen saturation were collected at timepoints T10, T30 and T60.

Statistical analysis

Obtained data were analyzed by using SPSS (IBM SPSS version 24.0). Values for SO₂ are presented as percentage (%) and values for rHb are given in arbitrary units (AU) according to the manufacturer¹⁵ and presented as absolute values at the given time points. Values for blood flow are given as relative flow compared to baseline flow measurement, according to previous studies.^{5,16,17}

Data of relative tissue blood flow (AU), rHb (AU) and SO₂ (%) at T0, T30 and T60 of bupivacaine (standard group) were compared with either bupivacaine/clonidine or bupivacaine/epinephrine (experimental groups) using the Mann-Whitney U-test using a hierarchical predefined statistical testing strategy. The saline group served as a negative control group and was compared with bupivacaine only. Statistical tests for differences (two-sided) were deemed significant at a level of $p=0.05$. Hodges-Lehmann (HL) estimators with 95% confidence intervals (CI) were calculated to analyze the differences of medians between microcirculation parameters (muscle blood flow, SO₂ and rHb) between bupivacaine and each group, respectively. Statistical testing for differences (two-sided) was deemed significant at a level of $p=0.05$. To exclude bias of microcirculatory variations and identify systemic effects of the study drugs, vital parameters between groups at T0, T30 and T60 were compared. HL estimators with 95% CI for heart rate (bpm), mean arterial blood pressure (MAP, mmHg) and SpO₂ (%) were calculated between the bupivacaine group and each of the other groups.

Differences of less than 15 bpm or heart rate >50, 10 mmHg in MAP or 3% in SpO₂ were assumed to be clinically irrelevant.

Results

Vital parameters

Vital parameter data are presented in Table 1. Comparison of HL estimators for mean arterial pressure, heart rate and systemic pulse oximetry revealed no clinically relevant differences between the four groups at T0 (Table 1). At T30 and T60 mins, HL estimators for mean arterial pressure, heart rate and systemic pulse oximetry showed no clinically relevant differences between the four groups.

Microcirculation

Data of relative muscle flow parameters are presented in Table 2. The bupivacaine group showed no significant difference in relative flow parameters compared with the saline group as control group at T30 ($p=0.853$) and T60 ($p=0.218$). However, a significant decrease in relative blood flow was revealed when bupivacaine with clonidine was compared with bupivacaine alone at T30 ($p=0.011$) and T60 ($p=0.043$). Comparison of bupivacaine with bupivacaine/epinephrine showed a significant decrease in relative flow at T30 ($p=0.03$) and T60 ($p=0.002$). Analysis of HL estimators between bupivacaine and BupiCloni or BupiEpi revealed a relevant difference of relative muscle blood flow reductions at T30 and T60.

Tissue oximetry

Data for SO₂ measurements are presented in Table 3. The Mann-Whitney test was performed for T0, T30 and T60 comparing each group with bupivacaine. At T0, no statistically significant differences were revealed between the four groups. Statistical analyses show significant differences comparing bupivacaine with saline at T30 ($p=0.01$) and T60 ($p=0.01$). Comparison of bupivacaine with bupivacaine/clonidine showed no significant differences at T0 ($p=0.247$), T30 ($p=0.315$) and T60 ($p=0.912$). Comparison of bupivacaine with bupivacaine/epinephrine revealed no statistically significant differences at T0 ($p=0.529$), T30 ($p=0.123$) and T60 ($p=0.280$). HL estimators (and 95% CI) for comparisons between bupivacaine group and the three other groups were calculated for T0, T30 and T60 and are presented in Table 3.

Table 1 Vital parameters

Group	Parameter	T0	HL (95% CI)	T30	HL (95% CI)	T60	HL (95% CI)
Bupivacaine N=10	Heart rate	87.5 (78–103)	n/a	95 (78–104)	n/a	90.5 (81–102)	n/a
	MAP	79.5 (71–95)	n/a	78 (66–91)	n/a	76.5 (68–87)	n/a
	SpO ₂	100 (99–100)	n/a	100 (100–100)	n/a	100 (99–100)	n/a
Saline N=10	Heart rate	83 (68–101)	-6 (-20; 10)	81.5 (73–102)	-11 (-24; 4)	81.5 (70–102)	-12 (-26; 4)
	MAP	68 (63–78)	-8.5 (-24; 4)	66 (60–78)	-9 (-28; 10)	72 (65–84)	-4 (-21; 10)
	SpO ₂	100 (99–100)	0 (0;0)	99 (99–100)	0 (-1; 0)	100 (100–100)	0 (0; 1)
Bupivacaine/Clonidine N=10	Heart rate	78.5 (69–89)	10 (-3; 23)	75.5 (72–89)	14 (2; 27)*	76.5 (69–84)	14 (1; 28)*
	MAP	75.5 (66–86)	7 (-10; 24)	71.5 (64–77)	6.5 (-11; 25)	71.5 (65–73)	8.5 (-4; 27)
	SpO ₂	100 (100–100)	0 (-1; 0)	100 (100–100)	0 (0; 0)	100 (100–100)	0 (-1; 0)
Bupivacaine/Epinephrine N=10	Heart rate	82 (75–92)	5.5 (-8; 20)	81.0 (76–90)	12.5 (-2; 25)	78.5 (75–90)	12 (-1; 26)
	MAP	73.5 (67–87)	4 (-12; 21)	74 (73–89)	-2 (-22; 16)	78 (68–89)	-0.5 (-16; 15)
	SpO ₂	100 (98–100)	0 (0; 2)	100 (99–100)	0 (0; 0)	100 (100–100)	0 (-1; 0)

Notes: Heart rate expressed in bpm; MAP in mmHg; and SpO₂ in %. All data are presented as median (25th–75th percentile) and each group was compared with bupivacaine at the same time point with differences shown as HL estimator, with 95% CI. * Significant.

Abbreviations: Bupi, bupivacaine; BupiCloni, bupivacaine with clonidine; BupiEpi, bupivacaine with epinephrine; CI, confidence interval; HL, Hodges-Lehmann; MAP, mean arterial pressure; n/a, not applicable; Saline, saline control group; SpO₂, pulse oximetry; T0, time point 0 mins; T30, time point 30 mins; T60, time point 60 mins.

Table 2 Relative muscle blood flow

Group	T5	T10	T15	T20	T25	T30	HL 95% CI	T40	T50	T60	HL 95% CI
Bupivacaine	0.782 (0.630–0.964)	0.656 (0.524–0.950)	0.751 (0.519–1.034)	0.665 (0.524–1.050)	0.870 (0.512–1.103)	0.736 (0.444–2.077)	n/a	0.668 (0.446–0.862)	0.694 (0.488–1.300)	0.653 (0.444–0.850)	n/a
Saline	0.804 (0.762–0.964)	0.838 (0.655–0.930)	0.784 (0.450–0.907)	0.801 (0.500–1.00)	0.769 (0.670–1.00)	0.726 (0.667–0.808) ^{NS}	-0.93 (-0.374; 0.306)	0.829 (0.738–0.885)	0.803 (0.750–1.055)	0.812 (0.690–0.860) ^{NS}	0.107 (-0.160; 0.380)
BupiCloni	0.503 (0.401–0.791)	0.402 (0.357–0.773)	0.389 (0.286–0.682)	0.349 (0.226–0.682)	0.473 (0.250–0.591)	0.388 (0.190–0.513)*	0.368 (0.58; 0.758)	0.347 (0.252–0.727)	0.265 (0.214–0.636)	0.300 (0.179–0.500)*	0.296 (0.25; 0.624)
BupiEpi	0.298 (0.182–0.480)	0.265 (0.222–0.357)	0.175 (0.139–0.320)	0.217 (0.159–0.280)	0.193 (0.136–0.278)	0.196 (0.143–0.333)*	0.422 (0.126; 0.868)	0.140 (0.103–0.184)	0.206 (0.154–0.286)	0.235 (0.182–0.310)*	0.336 (0.151; 0.660)

Notes: All data are presented as a median (25th–75th percentile) and each group was compared with bupivacaine at the same time point, with differences shown as HL estimator, with 95% CI. Relative values of flow to baseline flow at T0. Statistical significance was tested for T0, T30 and T60 comparing bupivacaine with the respective group. *Significant, NS not significant. Level of significance was set $p < 0.05$.

Abbreviations: Bupi, bupivacaine; BupiCloni, bupivacaine with clonidine; BupiEpi, bupivacaine with epinephrine; CI, confidence interval; HL, Hodges-Lehmann; n/a, not applicable; Saline, saline control group; T5–T60, time point 5 mins to 60 mins.

Table 3 Tissue oximetry

Group	T0	HL (95% CI)	T5	T10	T15	T20	T25	T30	HL (95% CI)	T40	T50	T60	HL (95% CI)
Bupivacaine	95.5 (89–98)	n/a	53 (40–62)	32 (20–44)	33 (12–58)	31 (19–50)	28 (9–43)	30.5 (13–46)	n/a	24 (12–47)	16.5 (8–52)	16 (7–52)	n/a
Saline	91 (88–97) ^{NS}	-0.5 (-7; 8)	94 (90–97)	88 (87–94)	89 (82–94)	89.5 (85–94)	87.5 (85–96)	87.5 (86–94)*	57 (35–74)	88.5 (82–90)	86 (80–94)	89 (82–95)*	62 (32; 80)
BupiCloni	89 (87–94) ^{NS}	4 (-4; 9)	72 (51–94)	59.5 (33–80)	54 (37–65)	48.5 (33–62)	48 (27–59)	45 (25–63) ^{NS}	-12 (-41; 12)	48.5 (6–67)	43.5 (6–66)	38.5 (2–64) ^{NS}	-1.5 (-43; 17)
BupiEpi	90 (89–96) ^{NS}	1 (-7; 8)	46 (16–57)	14 (6–29)	11.5 (8–21)	10 (10–19)	11 (9–15)	11 (9–17) ^{NS}	16.5 (-4; 34)	10 (6–17)	9.5 (3–19)	9 (6–18) ^{NS}	6 (-3; 34)

Notes: All data are presented as median (25th–75th percentile) and each group was compared with bupivacaine at the same time point, with differences shown as HL estimator, with 95% CI. SO₂ was measured in %. Statistical significance was tested for T0, T30 and T60 comparing bupivacaine with the respective group. *Significant, NS not significant. Level of significance was set $p < 0.05$.

Abbreviations: Bupi, bupivacaine; BupiCloni, bupivacaine with clonidine; BupiEpi, bupivacaine with epinephrine; CI, confidence interval; HL, Hodges-Lehmann; n/a, not applicable; Saline, saline control group; SO₂, tissue oximetry; T0–T60, time point 0 mins to 60 mins.

Hemoglobin content

Analysis of relative hemoglobin content as a parameter of venous congestion is presented in Table 4. No statistically significant differences of rHb could be detected comparing bupivacaine with saline at T0 ($p=0.684$), T30 ($p=0.739$) and T60 ($p=0.631$). Moreover, comparison of rHb of the bupivacaine group with the bupivacaine/clonidine group revealed no statistically significant differences at T0 ($p=0.063$), T30 ($p=0.280$) and T60 ($p=0.393$). Comparison of the bupivacaine group with the bupivacaine/epinephrine group showed no significant differences at T0 ($p=0.912$), T30 ($p=0.853$) and T60 ($p=0.631$). Results of HL estimator and 95% CI were calculated for T0, T30 and T60 as shown.

Discussion

Our experimental data showed that the application of bupivacaine results in a significant reduction of measured muscle blood flow at T30 and T60 (Table 4) when bupivacaine was compared to the saline control group. Interestingly, the addition of clonidine as well as epinephrine to bupivacaine results in pronounced additional blood flow reductions.

Compared to saline, the topical application of bupivacaine results in profound reductions of tissue oximetry values at T30 as well as at T60. However, the addition of clonidine or epinephrine did not show any difference (as an additional risk factor for tissue ischemia) regarding tissue oximetry compared to bupivacaine alone.

To our knowledge, this is the first study investigating the effect of bupivacaine on skeletal muscle microcirculation. We showed a pronounced reduction of muscle perfusion which resulted in consecutive reductions of tissue oximetry within the measured tissue volume. A previous study by Palmer et al. showed no effect of lidocaine 10 mg mL^{-1} on muscle perfusion in an experimental model in rats using the radiolabeled microsphere technique.¹⁸ Interestingly, in the same study by Palmer, the addition of epinephrine ($10 \text{ } \mu\text{g mL}^{-1}$) to lidocaine did result in only modest reductions of muscle blood flow. A previous study by Johns et al. found a dose-dependent vasoconstrictive effect on rat cremaster arterioles.¹⁹

Epinephrine is used as an adjuvant for LAs to reduce systemic absorption and toxic effects. A shortened nerve block onset time and prolonged block duration is assumed when epinephrine is used.¹⁴ It prolongs nerve block duration of LAs due to its reduction of local tissue clearance and enhanced concentration of LA in the perineural tissue.^{9,20,21} Our data is in contrast to the Palmer study, as we were able to

Table 4 Relative hemoglobin content

Group	T0	HL (95% CI)	T5	T10	T15	T20	T25	T30	HL (95% CI)	T40	T50	T60	HL (95% CI)
Bupivacaine	53 (47–64)	n/a	49 (42–65)	53 (45–63)	53 (44–61)	52.5 (46–63)	52 (45–64)	54.5 (47–65)	n/a	55.5 (45–65)	54.5 (42–65)	55.5 (44–65)	n/a
Saline	58.5 (50–69) ^{NS}	2.5 (–14; 16)	52.5 (50–64)	53 (50–63)	52 (49–64)	52.5 (43–62)	53 (44–62)	53 (47–61) ^{NS}	–3 (–16; 12)	56.5 (47–63)	58.5 (47–66)	58.5 (47–64) ^{NS}	3 (–1; 17)
BupiCloni	63 (60–71) ^{NS}	–11.5 (–21; 0)	62.5 (55–70)	60 (55–69)	61 (55–69)	60 (55–69)	61 (54–71)	60.5 (56–70) ^{NS}	–6.5 (–21; 7)	60 (56–69)	59 (55–70)	60.5 (55–70) ^{NS}	–6 (–18; 9)
BupiEpi	54.5 (49–62) ^{NS}	0.5 (–10; 10)	50.5 (43–59)	48.5 (43–58)	50.5 (46–58)	51 (46–58)	51.5 (46–59)	53.5 (47–59) ^{NS}	1 (–10; 14)	56 (49–62)	56 (50–62)	58 (49–62) ^{NS}	–2.5 (–14; 13)

Notes: Data are presented as median (25th–75th percentile), rHb was measured in AU; statistical significance was tested for T0, T30 and T60 comparing bupivacaine with the respective group. Level of significance was set $p<0.05$. **Abbreviations:** AU, arbitrary units; Bupi, bupivacaine; BupiCloni, bupivacaine with clonidine; BupiEpi, bupivacaine with epinephrine; rHb, relative tissue hemoglobin; Saline, saline control group; T0–T60, time points 0 mins to 60 mins; NS not significant.

demonstrate a significant reduction of muscle blood flow compared to saline when bupivacaine alone or in combination with epinephrine or clonidine was used. Clonidine is commonly used as an adjuvant to LA in regional anesthesia.¹⁴ Underlying mechanisms are still unknown. Interestingly, due to α_2 -adrenergic effects of clonidine, one might assume a resulting vasodilatation with the increase of tissue blood flow. Contrary to a previous study by our group⁵ investigating the effect of adjuvants on nerve blood flow showing no effect of clonidine when added to bupivacaine, the addition of clonidine to bupivacaine did result in an additional decrease of muscle tissue blood flow in this study. The extent of additional decrease of tissue perfusion by adding clonidine is comparable with the effect of addition of epinephrine to bupivacaine in the BupiEpi group in our study. These reductions of muscle blood flow resulted in profound reductions of skeletal muscle tissue oximetry in all three groups containing bupivacaine. Between these three groups, there was no difference in the extent of ischemia, even despite a numeric but not statistically significant difference of the BupiCloni group compared with bupivacaine alone or in combination with epinephrine. The microcirculation of skeletal muscle is regulated in a complex system comprising systemic (mediated by hormonal or nervous effects) as well as local factors (myogenic, metabolic or endothelial mechanism) that differ in resting and activated muscles.²²

In general, arterial (but not venous) branches of microcirculation in skeletal muscles are supplied by adrenergic sympathetic nervous fibers with α_1 and α_2 as well as beta-2 receptors. In previous studies, the postsynaptic α_2 adrenoreceptor activation resulted in profound reductions of skeletal muscle perfusion.^{23,24}

This might be the pharmacological explanation of reductions of tissue perfusion in the BupiCloni group. The clinical impact of our results is unclear as we were only able to measure short-term effects of bupivacaine in our study. However, the ischemic effect of bupivacaine alone or in combination with the adjuvants epinephrine or clonidine might be an additional risk factor to LA-induced myotoxicity after peripheral nerve blocks. LA-induced myotoxicity is induced by bupivacaine-related early and late aberrations in cytoplasmic Ca^{2+} homeostasis.^{7,25} The clinical impact of LA-induced myotoxicity is controversially discussed. Clinical relevant myopathy and myonecrosis following continuous peripheral nerve blocks, infiltration of wound margins and peri- and retrobulbar blocks have been described.² There are several reports of myotoxicity due to LAs. Zink et al. reported that when bupivacaine and ropivacaine were continuously

administered through femoral nerve catheters over six hours, both caused irreversible skeletal damage.³ Neal et al. reported that three individuals who received adductor canal block with lidocaine or mepivacaine bolus followed by ropivacaine for continuous infusion developed progressive weakness of quadriceps muscles.²⁶ Overall, the clinical impact of LA myotoxicity has been controversially debated. However, there is good experimental evidence that LAs caused lesions in skeletal muscle tissue. In a recent systematic review by Hussain et al., the authors concluded that further research needs to be performed to rule out the effect of adjuvants to LAs on myotoxicity.²⁵ We might speculate that combined injury by profound ischemia in addition to the myotoxicity of the LAs used might result in more severe muscle damage.

Limitations

This experimental study has several limitations that should be addressed. Firstly, the differences between the results in this animal experimental study might not be transferred to clinical conditions, due to differences in muscle blood flow and regulative mechanisms. Vascular tone might be influenced by general anesthesia, surgical preparation. Changes in blood pressure, systemic oxygen saturation and end-tidal carbon dioxide might influence skeletal muscle blood flow. We aimed to control this bias by invasive blood pressure monitoring, controlled mechanical ventilation with fixed inspired oxygen fraction and the maintenance of normocapnia with the use of capnometry. Measurements were analyzed as individual cases despite performing up to two measurements (left and right) per animal. This approach is similar to other published studies by our group and others with the aim to reduce animal numbers.

In conclusion, our study demonstrated profound reductions of skeletal muscle blood flow as well as tissue oximetry after topical application of bupivacaine alone in comparison with saline solution as control. The addition of epinephrine or clonidine both resulted in additional reductions of local muscle blood flow but without further reductions of tissue oximetry values.

Abbreviations

AU, arbitrary unit; Ca^{2+} , Calcium (ionized); LA, local anesthetic; MAP, mean arterial blood pressure; n, number; rel, relative; rHb, relative tissue hemoglobin; SO_2 , tissue oxygen saturation; SpO_2 , systemic arterial oxygen saturation; O₂C, "oxygen to see" (Proprietary name of LEA company, Giessen, Germany).

Data sharing statement

Full raw data can be requested from the corresponding author.

Ethical approval

Experiments were approved by the local authorities (V 54-19c 2015 (1) MR 20/13 Nr 92/2011), Regional Board of Animal Welfare, Gießen, Germany.

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Author contributions

All authors contributed to data analysis, drafting or revising the article, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

Disclosure

Thomas Wiesmann reports personal fees from B. Braun and from Pajunk, outside the submitted work. Hinnerk Wulf reports speaker fees from Syntetica, B. Braun, Teleflex and Vygon. Thorsten Steinfeldt reports speaker fees from B. Braun, Teleflex and Vygon. The authors report no other conflicts of interest in this work.

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