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1 RESEARCH PAPER

2 **Comparison of intratesticular lidocaine, sacrococcygeal epidural lidocaine and**
3 **intravenous methadone in cats undergoing castration: a prospective, randomized,**
4 **investigator-blind clinical trial**

5

6 Rocio Fernandez-Parra*, Luca Zilberstein*, Cyril Fontaine* & Chiara Adami§

7

8 *Department of Veterinary Anesthesiology and Critical Care, Ecole Nationale Vétérinaire
9 d'Alfort, Paris, France.

10 §Department of Clinical Sciences and Services, Royal Veterinary College, Hatfield, UK.

11

12 **Correspondence:** Rocio Fernandez-Parra, Department of Veterinary Anesthesiology and
13 Critical Care, Ecole Nationale Vétérinaire d'Alfort, Paris, France. 7 Avenue du General de
14 Gaulle, 94704 Maisons-Alfort, France. E-mail: rocio.fernandez@vet-alfort.fr

15

16 Running head: Locoregional anaesthesia or methadone in cats

17 **Abstract**

18 **Objective** The objective of this study was to compare three analgesic protocols for
19 feline castration.

20 **Study design** Prospective, randomized clinical study.

21 **Animals** Forty-nine client-owned cats.

22 **Methods** Cats were injected intramuscularly with dexmedetomidine ($15 \mu\text{g kg}^{-1}$) and
23 alfaxalone (3 mg kg^{-1}) and assigned randomly to one of three treatment groups. Group
24 ITL ($n = 15$) received intra-testicular 2% lidocaine (0.05 ml each testicle), group SCL (n
25 $= 15$) a sacro-coccygeal epidural injection of 2% lidocaine (0.1 mL kg^{-1}), and group
26 IVM ($n = 19$) intravenous methadone (0.3 mg kg^{-1}), before surgery. Cardiorespiratory
27 variables were recorded. In case of autonomic nociceptive response, intravenous
28 fentanyl ($2 \mu\text{g kg}^{-1}$) was administered. During recovery, time from intramuscular
29 atipamezole ($75 \mu\text{g kg}^{-1}$, administered at the end of surgery) to sternal recumbency and
30 to active interaction was recorded. Quality of recovery was assessed with a simple
31 descriptive scale (SDS). Postoperative analgesia was evaluated with a visual analogue
32 scale (VAS) and the UNESP-Botucatu multidimensional composite pain scale (MCPS)
33 at return of active interaction and then 1, 2 and 3 hours later.

34 **Results** The three analgesic protocols were comparable in terms of intraoperative
35 fentanyl and propofol requirement. Cardiorespiratory variables stayed within normal
36 ranges in the majority of the cases, although group IVM had the lowest intraoperative
37 respiratory rate ($p = 0.0009$).
38 No significant differences were detected between groups in UNESP-Botucatu MCPS
39 scores ($p = 0.21$). However, group ITL showed higher VAS score than group IVM ($p =$
40 0.001). Four cats enrolled in group ITL, as well as three of group SCL and one of group
41 IVM, required rescue analgesics before the completion of pain assessment.

42 **Conclusion and clinical relevance** Intratesticular and sacrococcygeal epidural
43 lidocaine injections could be regarded as good alternatives to systemic opioids in cats
44 undergoing castration, although the benefits of these techniques seem to be of shorter
45 duration than intravenous methadone.

46

47 **Introduction**

48 Neutering of client-owned cats is a common procedure in veterinary practice. Traditionally,
49 when performing castration of male cats the majority of French veterinarians prefer
50 injectable anaesthetic techniques to inhalation anaesthesia. The reasons behind this choice
51 may be a lack of familiarity with feline tracheal intubation, as well as the potential for
52 complications associated with this procedure (Brodbelt et al. 2007).

53 Ideally, an intramuscular (IM) anaesthetic protocol for castration should be safe for the
54 animal, inexpensive, and provide reliable unconsciousness, muscle relaxation and analgesia.
55 Combinations of alpha 2-adrenoreceptor agonists, induction agents suitable for IM
56 administration and opioids are used for this purpose (Adami et al. 2015).

57 Systemic full μ -opioid agonists are commonly employed to provide perioperative analgesia.
58 Unfortunately, they are controlled drugs and their use requires detailed record keeping; a
59 drawback which can prevent practitioners from using them on a regular basis (Hugonnard et
60 al. 2004). As an alternative to systemic analgesia, locoregional anaesthesia is becoming
61 increasingly popular in veterinary medicine, and its use is widespread not only by board-
62 certified anaesthetists, but also between general practitioners.

63 Intratesticular injection of local anaesthetics has been successfully used to provide
64 perioperative analgesia for castration in dogs (Huuskonen et al. 2012), piglets (Haga et al.
65 2006a), horses (Haga et al. 2006b), alpacas (Nickell et al. 2015) and people undergoing
66 testicular biopsies (Kamal et al. 2002).

67 Sacrococcygeal epidural injection of lidocaine is widely used in horses and ruminants to
68 desensitize the perineum and the pelvic organs without a loss of motor function of the pelvic
69 limbs. This technique has also been reported to relieve the pain associated with urethral
70 catheterization in cats with an onset of action of about five minutes (O'Hearn et al. 2011).
71 Both intratesticular and sacrococcygeal epidural injections of local anaesthetics may be used
72 to desensitize the testicles and the spermatic cord in cats.
73 The aim of this study was to compare three analgesic protocols: systemic administration of
74 methadone, sacrococcygeal epidural lidocaine, and intratesticular lidocaine injection, in
75 terms of quality and duration of analgesia in male cats undergoing castration.
76 Our hypothesis was that the three protocols would result in comparable propofol
77 requirements and quality of intraoperative analgesia in cats undergoing elective castration.

78

79 **Materials and methods**

80 **Animals**

81 Forty-nine client-owned male cats undergoing elective castration were included in the
82 study. The number of participants was established on the basis of a sample size
83 calculation using a commercial software program (SigmStat and SigmaPlot 12). It was
84 performed by setting the power at 80%, the level of significance at 5% and the end point
85 as a postoperative Visual Analogue Scale (VAS) pain score difference between groups
86 of 10 mm with a standard deviation of 5 mm.

87 Cats underwent a routine preanaesthetic physical examination in order to assess the health
88 status. Exclusion criteria were: presence of systemic disease, impaired cardiovascular
89 function, and age above 8 years. Food, but not water, was withdrawn 12 hours prior to
90 surgery. The study was performed under approval of the ethical committee of the Faculty of
91 Veterinary Medicine of Alfort, France, and informed owner consent.

92 Procedures

93 All cats were injected IM into the dorsolumbar muscles with dexmedetomidine ($15 \mu\text{g kg}^{-1}$)
94 (Dexdomitor; Orion Pharma, Finland) and alfaxalone (3 mg kg^{-1}) (Alfaxan; Jurox, Australia)
95 by the anaesthetist in charge for evaluating intraoperative nociception, depth of anaesthesia,
96 postoperative pain and quality of recovery. The drugs were combined in the same syringe
97 and if the total injection volume exceeded one mL, it was split into two injection sites. The
98 doses were established on the basis of previous pilot work.

99 The times from injection to sternal recumbency (defined as a position with the pelvic limbs
100 tucked under the body) and to lateral recumbency (defined as the cats lying on the side) were
101 recorded, as well as the time of induction of general anaesthesia. The latter was defined as
102 absence of righting reflex when the cats were positioned in dorsal recumbency, and
103 unresponsiveness to vocal and tactile stimulation. If general anaesthesia was not induced 30
104 minutes after the injection, the cats were injected IM with half of the initial doses of both
105 dexmedetomidine and alfaxalone, and excluded from the study.

106 Vomiting, hypersalivation, tremors, myoclonus and/or increased muscular tone were
107 considered adverse events and were recorded. After induction of anaesthesia, a 22 gauge
108 catheter (Delta Med, Italy) was placed in one cephalic vein. All cats received 7 mL kg^{-1}
109 hour¹ intravenous crystalloids (NaCl 0.9%; B. Braun, Germany) during the anaesthetic.
110 Amoxicillin (Clamoxyl; GlaxoSmithKline, UK), 20 mg kg^{-1} , was administered IV 30
111 minutes before the start of surgery. A multiparametric module (Monitor BSM-2301K; Nihon
112 Kohden, Japan) was used to monitor cardiorespiratory variables. Electrocardiography was
113 used to detect heart rate (HR), visual observation of the chest movements to detect
114 respiratory rate (f_R), pulse oximetry for pulse rate and arterial oxygen saturation (SpO_2), and
115 oscillometry to measure systolic, mean and diastolic arterial pressures (SAP, MAP and

116 DAP). An appropriate size cuff (width equal to 40% of limb circumference) was placed over
117 the radial artery.

118 The cats breathed room air. When intraoperative HR was lower than 100 beats minute⁻¹
119 atipamezole (75 µg kg⁻¹, Alzane, Zoetis, NJ, USA) was administered IM, and the cats were
120 excluded from the study. Cats with $f_R < 6$ breaths minute⁻¹ required endotracheal intubation
121 to allow manually assisted ventilation, and were excluded from the study. Hypotension,
122 defined as MAP values below 60 mmHg, was treated with a 3 ml kg⁻¹ bolus of crystalloids.
123 If the fluid bolus failed to increase the MAP above the cut-off value of 60 mmHg, the
124 crystalloid's rate of infusion was increased to 10 ml kg⁻¹ h⁻¹. Unresponsive hypotension was
125 treated with a bolus of hydroxyethyl starch (5 ml kg⁻¹; Voluven 6%, Fresenius Kabi, France).
126 If hypotension still persisted and the anaesthetist considered the administration of
127 vasopressors or anticholinergic appropriate, then the cats were excluded from the study.
128 Animals with SpO₂ values below 94% received supplemental oxygen at a rate of 2 L minute⁻¹
129 ¹, delivered via face mask. If SpO₂ failed to normalize, endotracheal intubation was
130 performed to allow manually assisted ventilation and administration of 100% oxygen, and
131 the cats were excluded from the study. If during the anaesthetic the rectal body temperature
132 decreased below 36.5 °C, a forced air warmer (Warm Touch, Mallinckrodt Medical, Ireland)
133 was used.

134 The end of surgery was defined as completion of the last suture knot (deferens and/or blood
135 vessels), at which time atipamezole was administered IM (75 µg kg⁻¹, Alzane, Zoetis, NJ,
136 USA). Times to sternal recumbency and to active interaction (defined as responsiveness to
137 vocal calls, alertness and interest in the surrounding), were recorded.

138 At the end of the assessments (T8; Fig. 1), 0.2 mg kg⁻¹ subcutaneous meloxicam (Metacam,
139 Boehringer-Ingelheim, Germany) was administered to all cats. Subcutaneous buprenorphine

140 (20 $\mu\text{g kg}^{-1}$) was administered to all cats who had no prior buprenorphine administered as a
141 rescue analgesic.

142 **Treatment groups**

143 The cats were randomly assigned to receive one of three treatments. A manual
144 randomization technique, based on drawing pieces of paper from an envelope, was used.

145 Group ITL received 2% lidocaine at a volume of 0.05 mL kg^{-1} per testicle. Gentle
146 aspiration before injection was used to exclude intravenous needle placement. Group
147 SCL received a sacrococcygeal epidural injection of 2% lidocaine, at the dose of 2 mg
148 kg^{-1} , corresponding to a volume of 0.1 mL kg^{-1} . Lack of resistance to injection and
149 subsequent relaxation of the anal sphincter were used to confirm the correct location of
150 the epidural injection. Group IVM received an intravenous (IV) injection of methadone
151 at the dose of 0.3 mg kg^{-1} .

152 All the analgesic treatments (either one of the two locoregional techniques or the systemic
153 administration of methadone) were performed five minutes before the surgical incision, by a
154 co-investigator not involved in the assessments. In order to prevent the primary investigator
155 from recognizing the treatment group, the sacrococcygeal area was clipped and surgically
156 prepared in all the cats enrolled in the study.

157

158 **Intraoperative evaluation of nociception**

159 All the assessments were carried out by the primary investigator who was unaware of the
160 treatment allocation. The surgeries were performed by junior clinicians under the supervision
161 of a senior surgeon. Depth of anaesthesia was evaluated based on the following descriptors:
162 spontaneous blinking (yes or no); movements during surgical stimulation (yes or no); and
163 adequate muscle relaxation (yes or no). If depth of anaesthesia was too light the cat received
164 propofol (0.5 mg kg^{-1} IV) (Propovet; Abbot, UK).

165 For each cat, baseline values for HR, f_R and MAP were established after induction of
166 anaesthesia and before surgical stimulation (T0, baseline values). The above listed variables
167 were then measured and recorded at the following time points: first surgical incision (T1),
168 traction of the first testicle (T2), second surgical incision (T3) and traction of the second
169 testicle (T4). Intraoperatively, any increase in two of three parameters (HR, f_R or MAP) of
170 30% above baseline was considered indicative of nociception. When such an increase was
171 observed for at least two of the three physiological variables, $2 \mu\text{g kg}^{-1}$ fentanyl (Fentanyl
172 Mylan $50 \mu\text{g ml}^{-1}$, PA, USA) was administered IV. The requirement for fentanyl during
173 surgery was used to evaluate intraoperative antinociception.

174 **Assessment of postoperative pain and quality of recovery**

175 After atipamezole injection, a simple descriptive scale for the assessment of recovery quality
176 was used with (0) defined as a very smooth recovery, (1) a smooth recovery, (2) a poor
177 recovery and (3) a very poor recovery, as soon as the cats regained sternal recumbency.
178 Postoperative pain was evaluated with a VAS, where 0 mm was labelled as “no pain” and
179 100 mm as “worst possible pain” (Jensen et al. 2003). Additionally, a modified version of the
180 UNESP-Botucatu MCPS (Brondani et al. 2013) was used. The subscale named
181 “physiological change” was excluded from the evaluation so that the maximum total score
182 was 24 (severe pain) instead of 30. Pain assessments were performed during recovery, when
183 the cats were observed to interact actively (T5) with the investigator, and then 1 (T6), 2 (T7)
184 and 3 (T8) hours later as shown in Fig. 1. The intervention levels for administration of
185 additional analgesia (buprenorphine $20 \mu\text{g kg}^{-1}$ IV, Vetergesic, Sogeval, France) were the
186 following: a score greater than 2 for the descriptor “expression of pain”, or a score greater
187 than 3 for the descriptor “psychomotor changes” on the UNESP-Botucatu MCPS, or a score
188 exceeding 40 mm on the VAS.

189 The post-operative pain assessments were carried out by the same investigator who evaluated
190 intraoperative nociception and who was unaware of the treatment allocation.

191

192 **Statistical analysis**

193 Data are presented as means \pm standard deviation or as medians (range) where applicable.
194 Normality of data distribution was assessed with the Shapiro-Wilk test and with the
195 Kolmogorov-Smirnov test. Age, body weight, number of propofol and fentanyl boluses
196 administered intraoperatively in each group, and time from atipamezole injection to recovery
197 were analyzed with a non-parametric test (Kruskal Wallis test, followed by Kruskal-Wallis
198 multiple comparison Z value test). Repeated measures ANOVA, followed by Tukey
199 Kramer's multiple comparison test, was used to compare the intraoperative physiological
200 variables (HR, f_R and MAP), as well as the postoperative pain scores, between treatments
201 and between time points. Duration of anaesthesia and time to active interaction were
202 analysed with a one-way ANOVA, followed by Bonferroni multiple comparison test. The
203 Fisher exact test was used to compare the number of animals within each group requiring
204 rescue buprenorphine before the completion of the last pain assessment. Statistical analyses
205 were performed using commercially available software (NCSS, 2007). Values of $p < 0.05$
206 were considered statistically significant.

207 **Results**

208 Fifty-four cats were considered possible candidates for the study, but seven were excluded
209 due to their fractious nature. A total of 49 cats, which were aged 8 (5 – 18) months and
210 weighed 3.8 (2.2 - 6.5) kg, were included. Treatment groups did not statistically differ with
211 respect to age and body weight ($p = 0.07$ and $p = 0.33$, respectively). All the cats enrolled in
212 the study were assigned an American Society of Anaesthesiologists risk classification of I.
213 Anaesthesia was induced in all cats within 30 minutes from IM injection (Table 1). The IM

214 injection exceeded 1 ml volume and was therefore split into two injection sites in 10, 13 and
215 10 cats of groups ITL, IVM and SCL, respectively. No adverse reactions were observed.
216 General anaesthesia (from anaesthetic induction to active interaction) lasted 40 ± 10 , 42 ± 9
217 and 45 ± 8 minutes in groups ITL ($n=15$), IVM ($n=19$) and SCL ($n=15$), respectively. These
218 differences were not significant ($p = 0.3$). The mean duration of surgery (from first incision
219 to the last suture knot) was 8 ± 2 minutes. Physiological variables stayed within acceptable
220 ranges for the species (Table 2), however group IVM had the lowest intraoperative
221 respiratory rates ($p = 0.0009$, Table 2). No statistically significant differences were found
222 between groups and time points for HR ($p = 0.10$ and $p = 0.06$, respectively) and MAP ($p =$
223 0.42 and $p = 0.82$, respectively). The SpO₂ fell below 94% in 4 out of 49 cases, 2 of which
224 were enrolled in group ITL and 2 in group IVM. These cats received oxygen
225 supplementation by mask. None of the animals required endotracheal intubation (Table 2).
226 Intraoperatively, groups ITL, IVM and SCL received 0 (0–1), 0(0–3) and 0 (0–3) doses of
227 propofol and 0 (0–0), 0 (0–0) and 0 (0–1) doses of fentanyl, respectively. These differences
228 were not statistically significant ($p = 0.38$ for propofol and $p = 0.86$ for fentanyl,
229 respectively). Two cats of group ITL and 3 of group SCL received intraoperative propofol,
230 while one animal only, enrolled in group SCL, required rescue fentanyl.
231 Recovery was smooth and uneventful for all cats and time from atipamezole injection to
232 recovery was shorter in group SCL [4 (3–9) minutes] than in group IVM [8 (2–17) minutes;
233 $z = 2.4$], and was 6 (3–95) minutes in group ITL. Time to active interaction was 16 ± 9 , $19 \pm$
234 4 , and 18 ± 6 minutes in groups ITL, IVM and SCL, respectively ($p = 0.86$) (Table 1).
235 Postoperative SDS score was 1 (0–2) in all groups and no statistically significant difference
236 was detected between treatments ($p = 0.7$) (Table 3).
237 With respect to the postoperative pain scores performed repeatedly at four time points, no
238 differences in UNESP-Botucatu MCPS were detected between groups ($p = 0.21$). However,

239 groups SCL and ITL showed higher VAS scores than group IVM, although this difference
240 was statistically significant only for group ITL ($p = 0.001$; Table 3).
241 Regarding the differences between time points, the values recorded during the first post-
242 operative pain assessment (T5: active interaction) were the highest for both the VAS and the
243 UNESP-Botucatu MCPS ($p = 0.008$ and $p = 0.004$, respectively). All the pain scores
244 decreased over time (Table 3). Eight cats, four of which enrolled in group ITL, three enrolled
245 in group SCL, and one of group IVM, received rescue buprenorphine before the completion
246 of pain assessments. This difference was not statistically significant ($p = 0.25$).

247 **Discussion**

248 The main finding of this study is that the administration of systemic methadone,
249 sacrococcygeal epidural lidocaine and intratesticular lidocaine resulted in comparable
250 propofol requirements and intraoperative analgesia in male cats undergoing castration.
251 Our results are in agreement with those previously obtained by other authors, who found that
252 intratesticular lidocaine injection prior to castration decreased intraoperative response to
253 noxious stimuli in dogs (Huuskonen et al. 2012) and in cats (Moldal et al. 2013). Portier and
254 colleagues (2009) reported similar results in horses. In the current study, intratesticular
255 injection did not result in adverse effects and could be easily and quickly performed without
256 requiring high level of expertise in locoregional anaesthesia. Conversely, a sacrococcygeal
257 epidural caused relaxation of the tail and of the anal sphincter, which could be regarded as an
258 undesirable side effect, and was technically more challenging than intratesticular injection.
259 Moreover, the failure rate of epidural anaesthesia was found to be 9% in cats (Troncy et al.
260 2002), and complications and undesired effects, namely development of abscesses at the site
261 of injection or systemic absorption of drugs, have been reported in this species (O'Hearn et
262 al. 2011). Although cats with epidural lidocaine had intraoperative analgesia, it is unknown
263 whether the epidural at the volume and dosages used would also result in desensitization of

264 the nerves in the spermatic cord. These drawbacks, together with the concern that the time
265 required for sacrococcygeal epidural injection may even exceed the duration of such a short
266 surgical procedure, may prevent practitioners from performing it for routine feline castration.
267 Whilst all the three analgesic treatments seemed to provide antinociception of sufficient
268 duration to cover the intraoperative period, the cats enrolled in the groups treated with
269 locoregional anaesthesia had higher postoperative pain scores and also required
270 postoperative rescue buprenorphine earlier than the cats which received methadone.
271 Cats in the methadone group took a longer time to recover after atipamezole administration
272 although this did not affect the quality of the recovery. This may be attributed to enhanced
273 and prolonged sedative effects of dexmedetomidine when the latter is combined with
274 methadone (Menegheti et al. 2014).

275 The dexmedetomidine-alfaxalone combination was suitable for IM administration and
276 resulted in reliable induction and maintenance of anaesthesia in the majority of cats.
277 However, although the doses used in the trial had been established based on a preliminary
278 investigation, five cats needed additional propofol to maintain unconsciousness during
279 surgery. This may be explained by inter-individual pharmacokinetic variability, and possibly
280 also by small variations, between cats, in the site of injection, within the fascia or in the
281 lumbodorsal muscles.

282 Alfaxalone is registered for IM use in cats in Australia but not in Europe. Potential concerns
283 for IM alfaxalone administration in feline patients are the less predictable anaesthetic effects
284 compared to the IV route and pain upon injection, when large volumes are administered. In
285 the cats enrolled in this study induction of anaesthesia was achieved after IM injection.
286 However, in most of the cases the volumes exceeded one mL and had to be split into two
287 injection sites. Large IM injections volumes are impractical and can increase the stress of the
288 patient related to handling and restraint.

289 In this study the treatment groups were not composed of the same number of animals. The
290 reason for this was that a simple randomization technique was used instead of block
291 randomization, which would have allowed a more even distribution of the cats within
292 groups.

293 In order to emulate protocols used in first-opinion veterinary practices in France, which
294 perform more elective castrations than teaching hospitals, it was decided not to supplement
295 inspired oxygen unless specifically needed.

296 The combination of dexmedetomidine and alfaxalone, with or without the addition of
297 methadone, did not result in an appreciable decrease in respiratory rate and less than 10% of
298 the cats enrolled in the study required oxygen supplementation. Additionally, although
299 bradycardia did occur in some cases, the heart rate always stayed above 100 beats minute⁻¹;
300 hence, according to the study protocol, none of the cats needed atipamezole administration.
301 These results seem to indicate that the anaesthetic protocol used in this study does not causes
302 dramatic changes in commonly monitored cardiorespiratory variables.

303 This study has some limitations. Junior clinicians performed the surgeries and this
304 considerably increased the duration of the procedures compared to private practice, where
305 experienced operators routinely perform feline castration. This might have increased the
306 intra-operative propofol requirement, which in turn may have affected the assessment of both
307 intra-operative nociception and post-operative pain, by influencing the cardiovascular and
308 respiratory response during surgery and by decreasing the responsiveness to stimulation in
309 the early postoperative period, respectively.

310 In conclusion, both intratesticular and sacrococcygeal epidural injections of lidocaine could
311 be proposed as alternatives to systemic methadone to provide intraoperative analgesia in cats
312 undergoing castration. If the duration of surgery is prolonged, the administration of
313 additional rescue analgesics may be necessary in the early postoperative period.

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317 **Authors' contributions**

318 RF-P: performed data collection and management, interpretation of the data and preparation
319 of the manuscript; LZ: study design, data interpretation and revised the manuscript; CF:
320 performed data collection and management; CA: study design, statistical analysis,
321 interpretation of data and revision of the manuscript.

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369 Figure Legend

370 **Figure 1** Time table for intra- and postoperative assessments. Intraoperatively, cardio-
371 respiratory variables were used to assess nociception and recorded at T0 (before surgical
372 stimulation), T1 (after the first incision), T2 (after traction of the first testicle), T3 (after
373 the second incision) and T4 (after traction of the second testicle). Quality of recovery
374 was evaluated with a simple descriptive scale (SDS) as soon as the cats regained sternal
375 recumbency. Postoperative pain assessments were carried out with a visual analogue
376 scale (VAS) and a multidimensional composite pain scale (MCPS), as soon as the cats
377 showed active interaction (T5) and then one (T6), two (T7) and three (T8) hours after
378 that.

379 **Table 1** Timing data from 49 cats anaesthetized with a combination of intramuscular
 380 dexmedetomidine and alfaxalone and undergoing elective castration.

Timing	Group		
	SLC	ITL	IVM
Injection to sternal recumbency (minutes)	2 ± 1	2 ± 1	2 ± 2
Injection to lateral recumbency (minutes)	3 ± 1	4 ± 2	4 ± 3
Time from injection to anaesthetic induction (minutes)	24 ± 7	21 ± 9	19 ± 7
Surgery time (minutes)	8 ± 2	8 ± 2	8 ± 3
Time from injection to atipamezole (minutes)	40 ± 10	42 ± 8	44 ± 7

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384 SCL, sacrococcygeal lidocaine ($n = 15$); ITL, intratesticular lidocaine ($n = 15$); IVM,

385 intravenous methadone ($n = 19$)

386

387 **Table 2** Mean \pm standard deviation of heart rates (HR), respiratory rates (f_R), mean arterial
 388 pressure (MAP) and haemoglobin oxygen saturation (SpO₂) values of 49 cats undergoing
 389 elective castration and administered three different types of intraoperative analgesia.
 390 Measurements were taken during preanaesthetic physical examination and at five different
 391 time points: T0 (after induction of anaesthesia and before surgical stimulation, baseline), T1
 392 (after first skin incision), T2 (after exteriorization of the first testicle), T3 (after second skin
 393 incision), and T4 (after exteriorization of the second testicle).
 394

Variable	Group	Time					
		Pre- anaesthetic physical examination	T0 Baseline	T1	T2	T3	T4
HR (beats minute ⁻¹)	SCL	184 \pm 21	119 \pm 7	116 \pm 7	127 \pm 7	124 \pm 7	136 \pm 7
	ITL	188 \pm 23	109 \pm 7	114 \pm 7	111 \pm 7	103 \pm 7	109 \pm 7
	IVM	176 \pm 23	120 \pm 6	115 \pm 6	122 \pm 6	113 \pm 6	120 \pm 6
f_R (breaths minute ⁻¹)	SCL	64 \pm 17	42 \pm 2	41 \pm 2	43 \pm 2	41 \pm 2	42 \pm 2
	ITL	79 \pm 19	42 \pm 2	44 \pm 2	42 \pm 2	41 \pm 2	42 \pm 2
	IVM	77 \pm 26	40 \pm 2*	39 \pm 2*	38 \pm 2*	34 \pm 2*	37 \pm 2*
MAP (mmHg)	SCL	N/A	93 \pm 4	91 \pm 4	89 \pm 4	91 \pm 4	87 \pm 4
	ITL	N/A	87 \pm 4	83 \pm 4	90 \pm 4	85 \pm 4	88 \pm 4
	IVM	N/A	86 \pm 3	91 \pm 4	92 \pm 3	88 \pm 4	84 \pm 4
SpO ₂ (%)	SCL	N/A	96 \pm 3	95 \pm 3	96 \pm 3	96 \pm 2	96 \pm 4
	ITL	N/A	94 \pm 4	95 \pm 5	94 \pm 5	94 \pm 5	94 \pm 5
	IVM	N/A	91 \pm 3	90 \pm 4	90 \pm 5	91 \pm 5	93 \pm 2

395

396 *Statistically significant difference between groups ($p = 0.009$). N/A: non-applicable.

397 SCL, sacrococcygeal lidocaine ($n = 15$); ITL, intratesticular lidocaine ($n = 15$); IVM,

398 intravenous methadone ($n = 19$)

399 **Table 3** Median (range) of quality of recovery scores [assessed with a simple descriptive scale (SDS)] and postoperative pain [assessed with a Visual
 400 Analogue Scale (VAS) and with the UNESP-Botucatu multidimensional composite pain scale (MCPS)], recorded from 49 cats undergoing elective
 401 castration. Pain assessments were carried out at various time points: as soon as the cats were observed to interact actively with the investigator (T5),
 402 and then 1 (T6), 2 (T7) and 3 (T8) hours after that. SCL, sacrococcygeal lidocaine ($n = 15$); ITL, intratesticular lidocaine ($n = 15$); IVM, intravenous
 403 methadone ($n = 19$).
 404

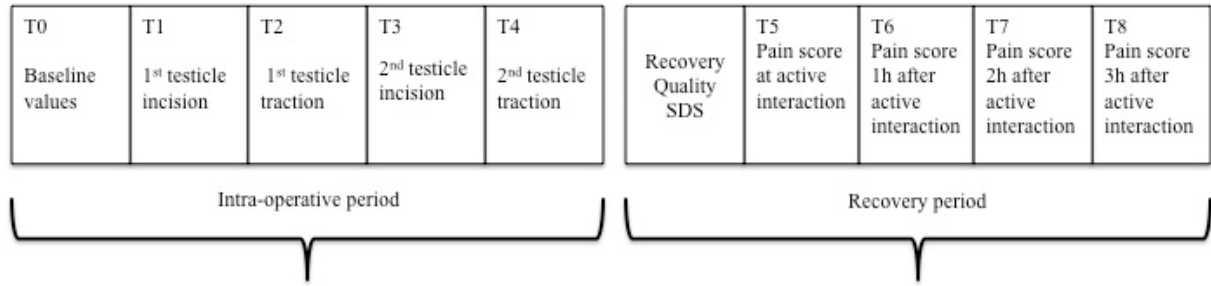
Group	SDS	VAS T5	VAS T6	VAS T7	VAS T8	MCPS T5	MCPS T6	MCPS T7	MCPS T8
SLC	1 (0-2)	20 (0-40)	20 (0-40)	20 (0-20)	20 (0-20)	3 (0-6)	3 (0-4)	1 (0-4)	1 (0-3)
ITL	1 (0-2)	20 (0-80)*	1 (0-60)*	1 (0-60)*	1 (0-30)*	2 (0-10)	3 (0-8)	3 (0-6)	3 (0-5)
IVM	1 (0-2)	20 (0-20)	20 (0-20)	20 (0-20)	20 (0-20)	2 (0-6)	3 (0-5)	3 (1-5)	3 (0-5)

405

406 *Statistically significant difference between ITL and IVM group ($p = 0.001$).

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