Isoflurane anaesthesia in an African wild dog, Lycaon pictus

G F Stegmann^a

ABSTRACT

Anaesthesia was required in a captive female African wild dog (*Lycaon pictus*) for surgical wound treatment. After it was immobilised with a medetomidine-ketamine combination, bradycardia, hypothermia, systolic hypertension and metabolic acidosis were observed. Surgical anaesthesia was maintained with a 1 % end-tidal isoflurane concentration. A decrease in the arterial blood pressure, rectal temperature and pH occurred during maintenance of anaesthesia.

Key words: acidosis, anaesthesia, hypertension, isoflurane, ketamine, *Lycaon pictus*, medetomidine, wild dog.

Stegmann G F Isoflurane anaesthesia in an African wild dog, Lycaon pictus. Journal of the South African Veterinary Association (2000) 71(4): 246 (En.). Department of Companion Animal Surgery, Faculty of Veterinary Science, University of Pretoria, Private Bag X04, Onderstepoort, 0110 South Africa.

Anaesthesia for surgical treatment was required in a captive, 1-year-old, female African wild dog (*Lycaon pictus*) of approximately 17 kg body weight. The animal was immobilised with a mixture of 0.7 mg medetomidine (41 µg/kg) (Domitor, Novartis) and 40 mg ketamine hydrochloride (2.3 mg/kg) (Anaket, Kyron Laboratories), and transported in a crate to the Onderstepoort Veterinary Academic Hospital (OVAH), Faculty of Veterinary Science, University of Pretoria.

The perioperative cardiopulmonary variables are presented in Table 1, and the arterial blood-gas analysis in Table 2. Cardiopulmonary variables were monitored with the Cardiocap II and Capnomac Ultima (Datex, Helsinki). Arterial blood pressure was measured from the dorsal pedal artery. After endotracheal intubation, anaesthesia was maintained with a 1 % end-tidal isoflurane (Forane, Abbott) concentration on a circle anaesthetic machine with carbon dioxide absorption.

Medetomidine is an alpha₂ adrenergic agonist with potent sedative-hypnotic, muscle relaxant and analgesic effects, and was administered as an anaesthetic adjunct to ketamine to facilitate muscle relaxation and anaesthesia. Before inhalation anaesthesia, bradycardia (35 beats/min) associated with unexpectedly prolonged high systolic blood pressure (28.2 kPa, 60 min after immobilisation) was present in the wild dog. Medetomidine is reported to exert a biphasic blood pres-

Received: June 2000. Accepted: August 2000.

Table 1: Mean (SD) cardiopulmonary variables in an African wild dog after medetomidine/ketamine immobilisation, and during isoflurane anaesthesia.

Variable ^a	M/K	ISO
HR (beats/min)	35	59.0 (10)
SYS (kPa)	28.2	20.0 (1.9)
DIA (kPa)	10.4	8.5 (0.8)
MAP (kPa)	13.6	10.5 (1.0)
F (breaths/min)	14	14 (1.0)
VT (mℓ)	543	303 (29)
VΕ (<i>l</i>)	7.6	4.2 (0.03)
ETCO ₂ (%)	_	5.6 (0.1)
Etiso (%)	-	1.0 (0.2)

^aHR = heart rate; SYS = systolic blood pressure; DIA = diastolic blood pressure; MAP = mean arterial blood pressure; F = ventilation rate; VT = tidal volume; VE = minute volume; ETCO₂ = end-tidal carbon dioxide concentration; Etiso = end-tidal isoflurane concentration; M/K = midazolam-ketamine; ISO = isoflurane.

Table 2: Arterial blood-gas variables in an African wild dog after immobilisation with medetomidine/ketamine and during iso-flurane anaesthesia.

Variable ^a	M/K	ISO
pH units	7.41	7.34
PaCO₂ kPa	4.1	4.8
$TCO_2 \text{ mmol}/\ell$	20.4	20.8
PaO ₂ kPa	12.8	60.4
HCO₃ mmol/ℓ	19.4	19.6
SBE mmol/ ℓ	-4.4	-5.7
SBC mmol/t	21.1	19.7
SAT (%)	97.4	99.9
$O_2CT \text{ mmol}/\ell$	15.4	18.2

^aPaCO₂ = arterial partial pressure of carbon dioxide; TCO₂ = total carbon dioxide; PaO₂ = arterial partial pressure of oxygen; HCO₃ bicarbonate concentration; SBE = standard base excess; SBC = standard bicarbonate; SAT = oxyhaemoglobin saturation; O₂CT = oxygen content; M/K = midazolam-ketamine; ISO = isoflurane.

sure response, characterised by an initial transient increase, followed by a prolonged decrease. The increase in blood pressure is the result of an increase in systemic vascular resistance⁴. The low heart rate may have been a reflex physiological response to the high blood pressure, or the result of hypothermia (37 °C). In midazolam/ketamine-anaesthetised domestic dogs, a systolic and mean arterial blood pressure of 22.3 and 16.5 kPa has been reported². The decrease in blood pressure during maintenance of anaesthesia was probably the result of a decrease in the peripheral vascular resistance due to the isoflurane³.

Medetomidine and hypothermia decrease the dose of inhalation agents required for anaesthesia^{1,6}, and therefore probably contributed to the low end-tidal isoflurane concentration (1%) required to maintain surgical anaesthesia in the wild dog. Medetomidine decreases body temperature⁵. Low ambient temperatures during the winter, and exposure to air-conditioning, probably contributed to the further decrease in temperature (35.5 °C) during treatment. Metabolic acidosis was present, and was probably the result of anaesthesia and hypothermia.

REFERENCES

- Maier C, Steinberg G K, Sun GH 1993 Neuroprotection by the alpha₂-adrenoreceptor agonist dexmedetomidine in a focal model of cerebral ischaemia. *Anesthe*siology 79: 306–310
- Serteyn D, Coppens P, Jones R, Verstegen J, Philippart C, Lamy M 1993 Circulatory and respiratory effects of the combination medetomidine-ketamine in beagles. *Journal* of Veterinary Pharmacology and Therapeutics 16: 199–206
- Stevens W C, Cromwell T H, Halsey M J 1971 The cardiovascular effects of a new inhalation anaesthetic, Forane, in human volunteers at constant arterial carbon dioxide tension. *Anesthesiology* 35: 8–11
- Tranquilly W J, Thurmon J C, Paul A J, Benson G J 1985 Influence of nifedipine on xylazine-induced acute pressor response in halothane anaesthetized dogs. American Journal of Veterinary Research 46: 1892–1895
- Verstegen J, Fargetton X, Donnay I, Ectors F 1991 An evaluation of medetomidine/ ketamine and other drug combinations for anaesthesia in cats. Veterinary Record 128: 32–35
- Vitez T S, White P F, Eeger E I 1974 Effects of hypothermia on halothane MAC and isoflurane MAC in the rat. Anesthesiology 41: 80–84

^aDepartment of Companion Animal Surgery, Faculty of Veterinary Science, University of Pretoria, Private Bag X04, 0110 Onderstepoort, South Africa.