

JAN BLÁHA

**Klinika anesteziologie, resuscitace
a intenzivní medicíny**
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Všeobecná fakultní nemocnice v Praze



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JAK NA TO...

AKUTNÍ SEKCE

Střet zájmů:

S P A A



AKUTNÍ nebo **AKUTNÍ!** ?

- bradykardie plodu
- prolaps pupečníku
- masivní krvácení
- eklampsie
- ...

Volba anestezie dle času, který má anesteziolog k dispozici !

Čas do incize (min.)	
elektivně	EPID, SAB, CSE ... (CA)
15-20	EPID s rychle nasedající epidurální směsí (2% lidocain), SAB, CSE, (CA)
10-15	SAB, (CSE), (CA)
5-10	SAB (zkušený anesteziolog + dobré anatomické podmínky, jinak spíše CA)
0-5	CELKOVÁ ANESTEZIE
	Poznámka: je nutno vzít v úvahu i čas nutný k zpolohování rodičky, natažení směsi a punkci epidurálního/subarachnoidálního prostoru, nikoli pouze čas nasednutí účinku lokálního anestetika. Současně je nutno odhadnout čas svolání a umytí operačního týmu, desinfekci a zarouškování rodičky, ...



Epidurální směs: lidocain 2% 18 ml + sufentanyl 10 µg/2 ml + adrenalin 0,1 ml



VS.



CELKOVÁ ANESTEZIE je u císařského řezu indikována pouze tehdy, je-li **REGIONÁLNÍ ANESTEZIE KONTRAINDIKOVÁNA.**

PROČ **NECHCEME** CELKOVOU ANESTEZII ?



Retrospektivní 3letá studie srovnávající celkovou, epidurální a spinální anestézii z hlediska respiračního distresu plodu.

Není signifikantní rozdíl mezi jednotlivými typy anestézie.

Sigalas et al: Clin Exp Obstet Gynecol. 2006; 33(1):10-12

Authors' conclusions



Trusted evidence.
Informed decisions.
Better health.

There is no evidence from this review to show that RA is superior to GA in terms of major neonatal outcomes. Further research to evaluate neonatal morbidity and maternal outcomes, such as satisfaction with technique, will be useful.

Afolabi BB. Cochrane Database Syst Rev. 2012 Oct 17;10:CD004350

Anaesthesia for Caesarean section and neonatal acid-base status: a meta-analysis★

F. Reynolds¹ and P. T. Seed²

¹ Emeritus Professor of Obstetric Anaesthesia, Department of Anaesthesia, St Thomas' Hospital, London SE1 7EH, UK

² Lecturer in Medical Statistics, Division of Reproductive Health, Endocrinology and Development, King's College, London SE1 7EH, UK

Table 8 Difference between umbilical artery acid-base values with type of anaesthesia for Caesarean section: results of meta-analysis.

	Comparison	All studies				Randomised trials only			
		#	Difference	95% CI	p	#	Difference	95% CI	p
pH	spinal – general	13	-0.015	-0.029 to -0.001	0.038	5	-0.027	-0.051 to -0.002	0.034
	spinal – epidural	11	-0.013	-0.024 to -0.002	0.025	7	-0.010	-0.022 to 0.01	0.074
	epidural – general	13	-0.006	-0.016 to 0.005	0.317	4	0.001	-0.023 to 0.025	0.938
Base deficit (mEq.l ⁻¹)	spinal – general	7	1.109	0.434 to 1.784	0.001	2	1.235	-0.821 to 3.290	0.239
	spinal – epidural	7	0.910	0.222 to 1.598	0.010	4	0.834	-0.192 to 0.859	0.111
	epidural – general	8	0.137	-0.198 to 0.471	0.424	2	-0.018	-1.026 to 0.990	0.972

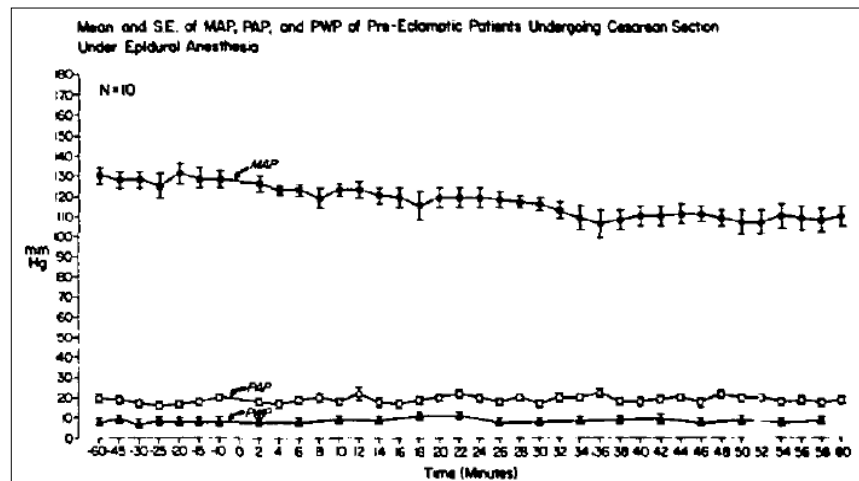
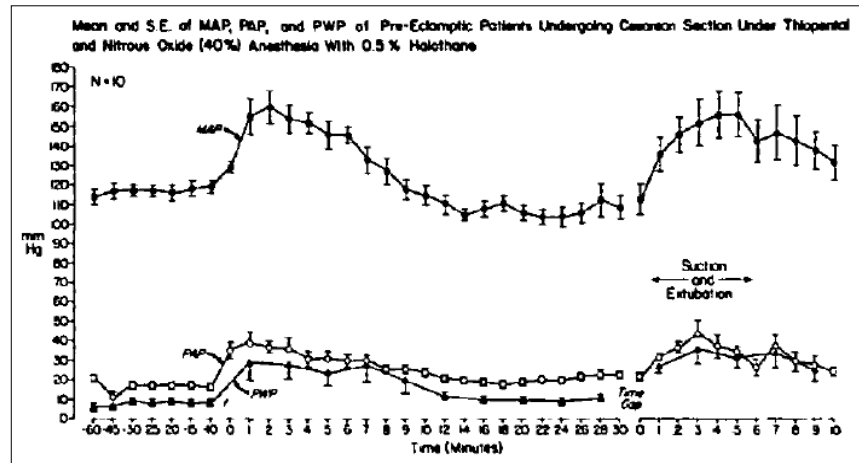
= number of studies.

Summary

Spinal anaesthesia is generally preferred for Caesarean section, but its superiority for the baby is often assumed. Umbilical artery acid-base status provides a valid index of fetal welfare. Twenty-seven studies reporting neonatal acid-base data with different types of anaesthesia were used to compare umbilical artery or vein pH and base deficit, using random-effect meta-analysis. Cord pH was significantly lower with spinal than with both general and epidural anaesthesia. Larger doses of ephedrine contributed to the latter effect ($p = 0.023$). Sixteen studies reported a base deficit, which was significantly higher for spinal than for general and epidural anaesthesia.

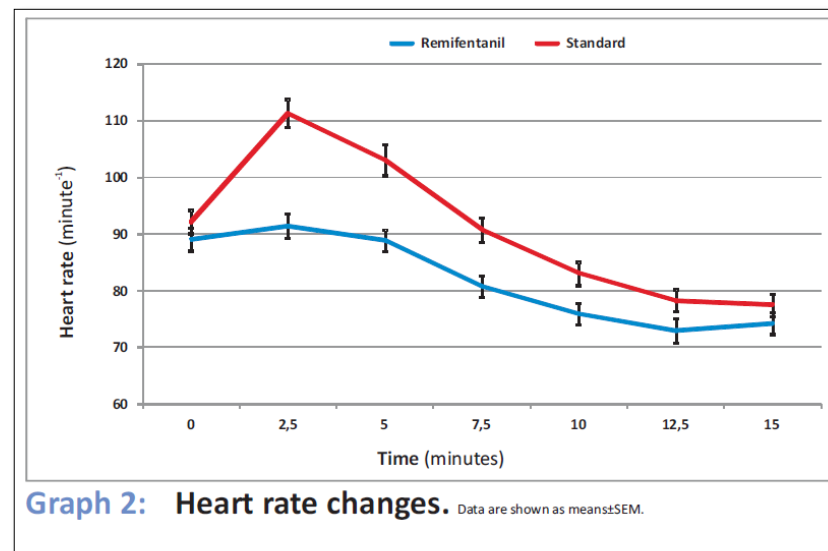
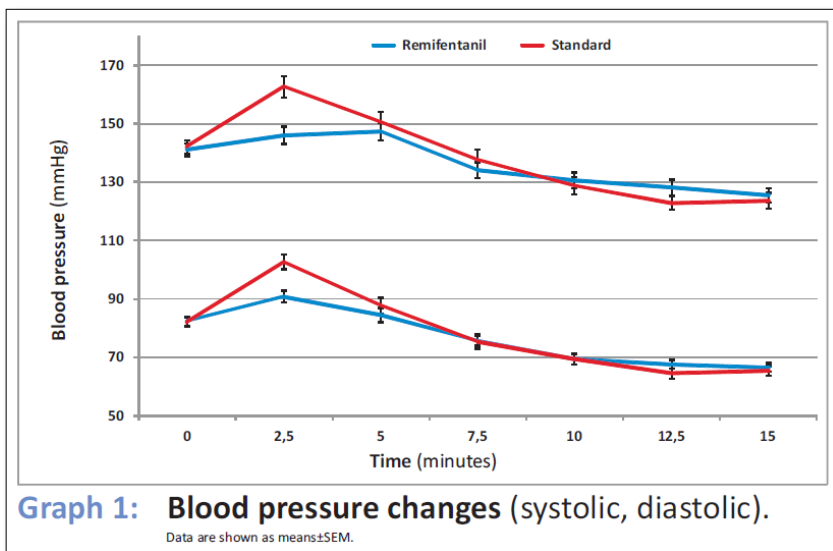
Spinal anaesthesia cannot be considered safer than epidural or general anaesthesia for the fetus.

PROČ NECHCEME CELKOVOU ANESTEZII ?

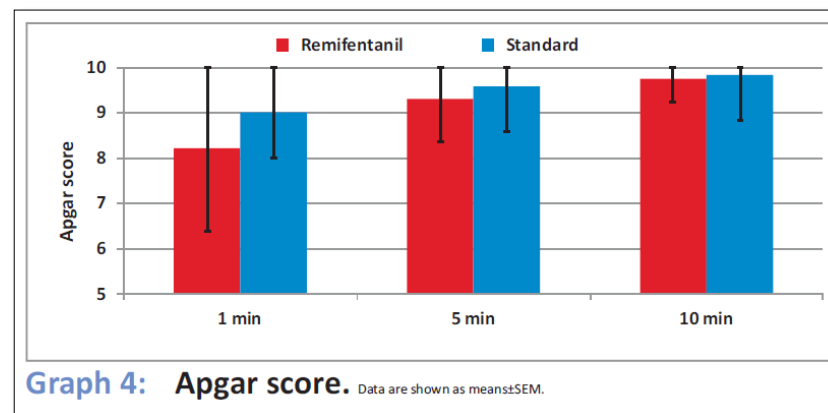


Hodgkinson et al. Can J Anesth 1980 27: 389-394.

Remifentanil 1 µg/kg před úvodem do celkové anestezie



Nosková, Bláha et al. BMC Anesthesiology 2015 (in press).

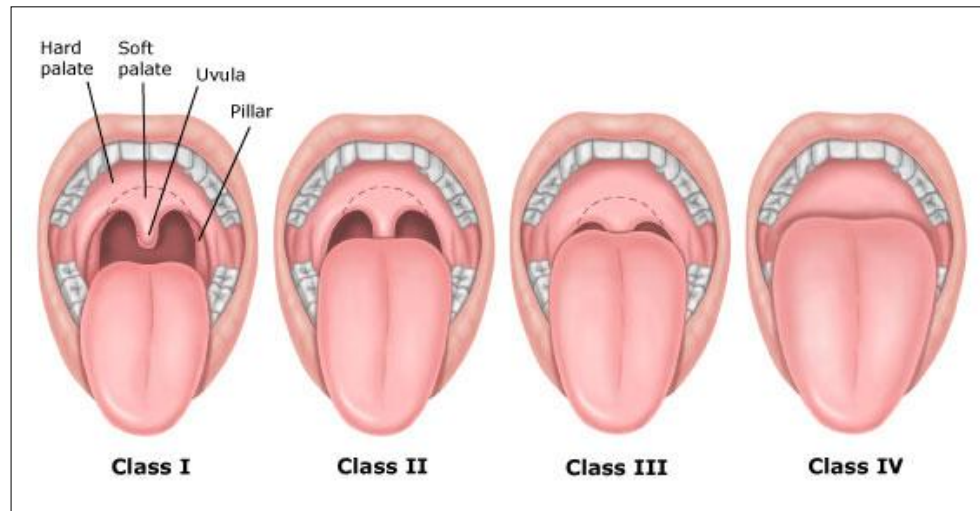


PROČ **NECHCEME** CELKOVOU ANESTEZII ?



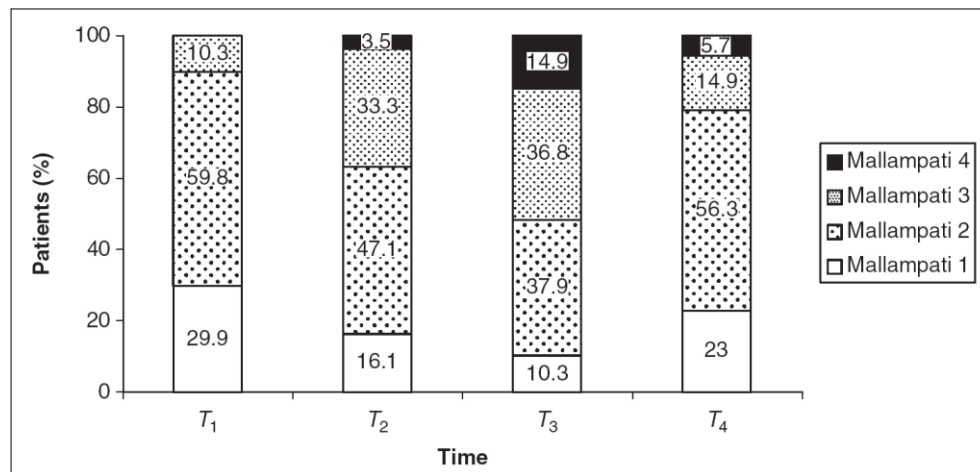
= vždy riziko obtížné intubace !

OBTÍŽNÁ INTUBACE V TĚHOTENSTVÍ



10x

vyšší riziko obtížné intubace



se v průběhu porodu ještě dále zvyšuje!

Fig 1 The Mallampati classes at different time points. T₁, 8 months of pregnancy; T₂, during labour; T₃, 20 min after delivery; T₄, 48 h after delivery. The percentages of patients with Mallampati class 3 or 4 changed significantly: T₁ vs T₂, P=0.0000; T₂ vs T₃, P=0.0005; T₃ vs T₄, P=0.0000; T₄ vs T₁, P=0.0062.

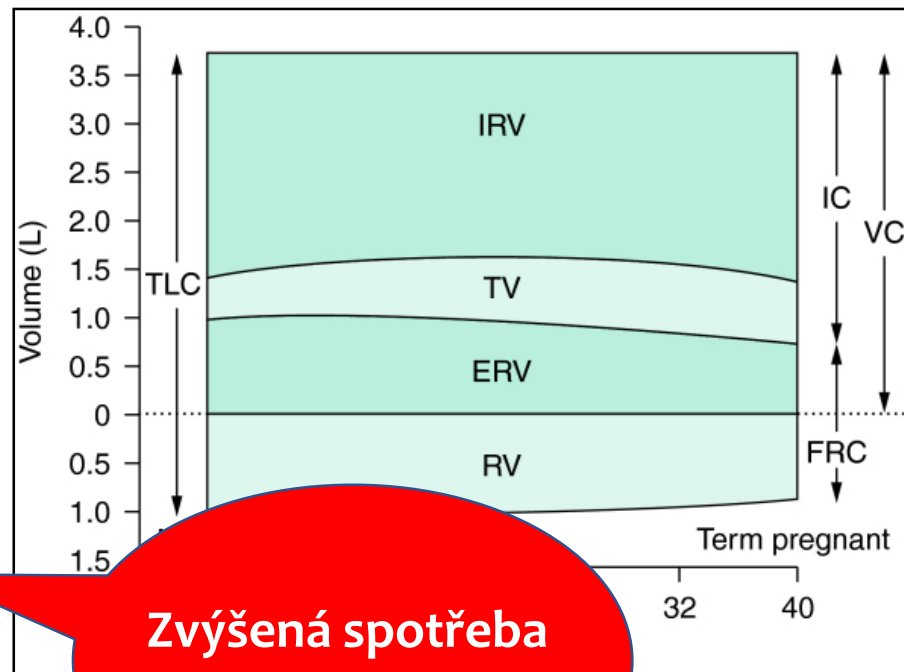
FYZIOLOGICKÉ ZMĚNY V TĚHOTENSTVÍ



- růst dělohy = zvýšená poloha bránice
- zvýšená senzitivita k CO_2 v respiračním centru (vliv progesteronu)
- **vyšší spotřeba kyslíku** (zvýšený metabolismus matky + plod)

Relativní změna

Spotřeba O_2	+40%
Dechová frekvence	mírně zvýšena
Minutová ventilace	+45%
Alveolární ventilace	+45%
Difúze přes alv.-kap. membránu	-15%
Dechový objem	+45%
Vitální kapacita	beze změny
Funkční reziduální kapacita	-20%
Poloha bránice	o 4 cm výše



**Zvýšená spotřeba
a snížená zásoba O_2 !!!**

Conklin KA. Semin Anesth 1991; 10:221-34.

and Practice, 4th Edition, 2009

PREOXYGENACE !!!

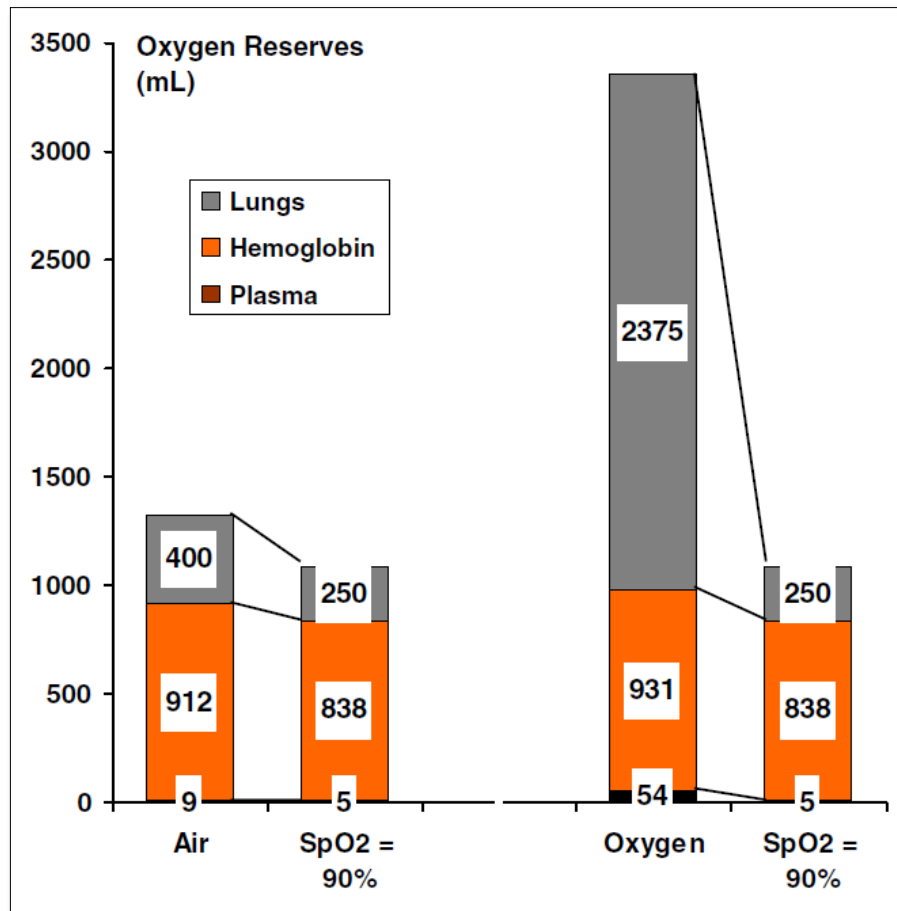


Fig. 1 Oxygen reserves in a normal healthy adult when breathing room air (left), after breathing 100% oxygen (right), at onset of apnea, and when reaching an oxygen saturation (SpO₂) of 90%. In this example, the oxygen available for consumption during the apneic period amounts to 228 mL when breathing air and 2267 mL when breathing oxygen. Calculations are based on a functional residual capacity of 2500 mL, hemoglobin concentration 140 g · L⁻¹, SpO₂ = 98% on air, SpO₂ = 100% on oxygen, and blood volume 5 L. In this example, a subject with an oxygen consumption of 250 mL · min⁻¹ could sustain a period of apnea of 228/250 = 0.9 min after breathing air and 2267/250 = 9 min after breathing oxygen

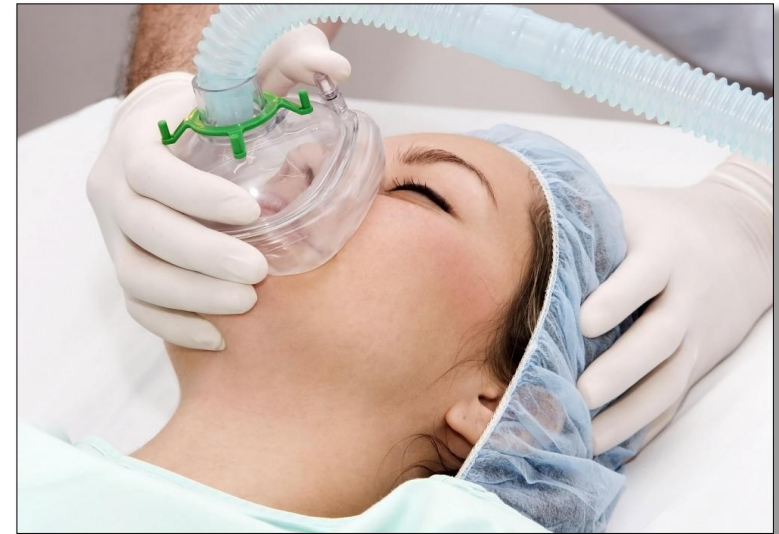
Tanoubi I. Can J Anesth/J Can Anesth (2009) 56:449–466

PREOXYGENACE !!!



A. lehká obličejová kyslíková maska

5-8 minut dýchání 100% O₂
normálním objemem



B. plně těsnící obličejová kyslíková maska

3-8 vdechů 100% O₂
v objemu vitální kapacity

THE ASPIRATION OF STOMACH CONTENTS INTO THE LUNGS DURING OBSTETRIC ANESTHESIA*

CURTIS L. MENDELSON, M.D., NEW YORK, N. Y.

(From the Department of Obstetrics and Gynecology, Cornell University Medical College and
New York Hospital)

Am J Obstet Gynecol 1945;49:554-66.

Summary

Sixty-six cases of aspiration of stomach contents into the lungs during obstetric anesthesia are analyzed. The incidence of this complication is 0.15 per cent in 44,016 pregnancies at the New York Lying-In Hospital from 1932 to 1945.

Table 7 Reported incidence of aspiration in obstetric and general surgical populations

Study	No. of cases	Patient group characteristics	Incidence of aspiration [no. of cases]
This study	1870	Obstetric; peripartum; nonintubated	0.053% [1]
Kranz & Edwards [3]	37 282	Obstetric; vaginal delivery; nonintubated	0.013% [5]
Kranz & Edwards [3]	3076	Obstetric; Caesarean section; intubated	0.228% [7]
Olsson <i>et al.</i> [2]	2643	Obstetric; Caesarean section; intubated	0.15% [4]
Olsson <i>et al.</i> [2]	111 215	General surgery; nonintubated	0.018% [20]
Olsson <i>et al.</i> [2]	74 143	General surgery; intubated	0.085% [63]
Cohen <i>et al.</i> [5]	112 000	General surgery; intubated and nonintubated	0.064% [72]
Kallar [6]	529 150	Outpatients; intubated and nonintubated	0.017% [90]
Warner <i>et al.</i> [4]	13 427	General surgery; emergency	0.112% [15]
Warner <i>et al.</i> [4]	202 061	General surgery; elective	0.0257% [52]

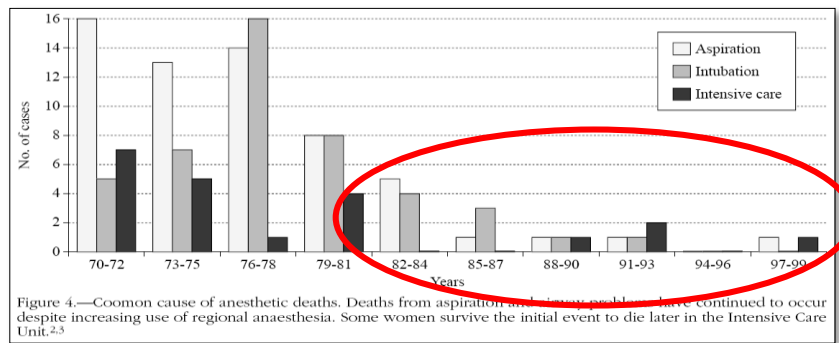
Ezri *et al.* Anaesthesia 2000; 55:421-426

ANESTEZIE A MATEŘSKÁ MORTALITA

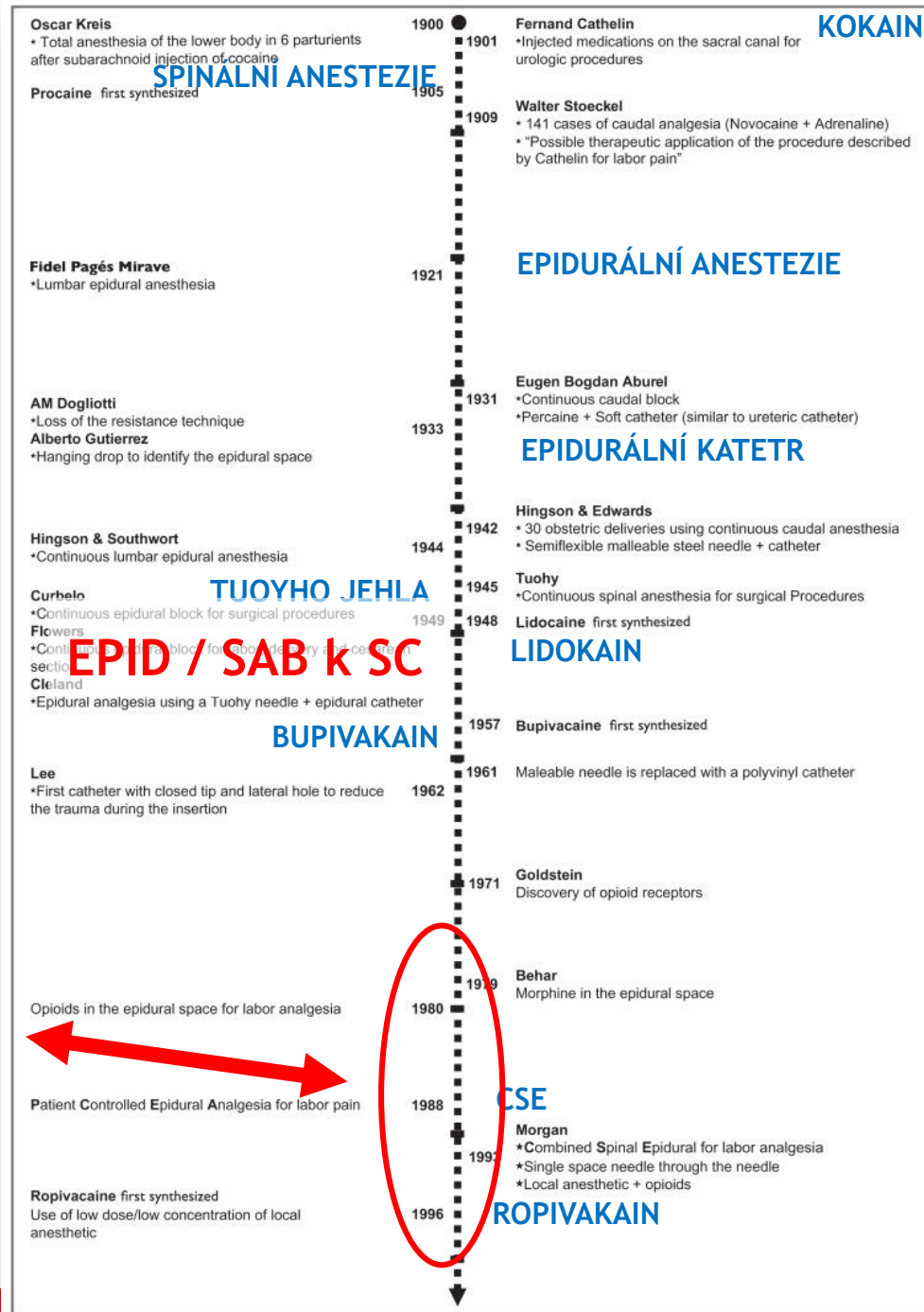
70./80. léta celková anestézie má **17x** vyšší mortalitu než regionální

90. léta již "pouze" **6x** vyšší mortalita CA proti RA

21. století **není rozdíl** mezi celkovou a regionální anestezií



Silva Mt al. Local Reg Anesth. 2010; 3: 143–153
 Hawkins JL, Anesthesiology 1997;86:277-84
 Hawkins JL, Clin Obstet Gyn 2003; 46: 679-87
 Cochrane Database of Systematic Reviews 2012



Oral sodium citrate increases nausea amongst elective Cesarean delivery patients

[Le citrate de sodium oral augmente les nausées pendant la césarienne réglée]

Klaus Kjaer MD, Michele Comerford MD, Linda Kondilis BA, Lauren DiMaria BA, Sharon Abramovitz MD, Michael Kiselev MD, Jon Samuels MD, Farida Gadalla MD, Barbara L. Leighton MD

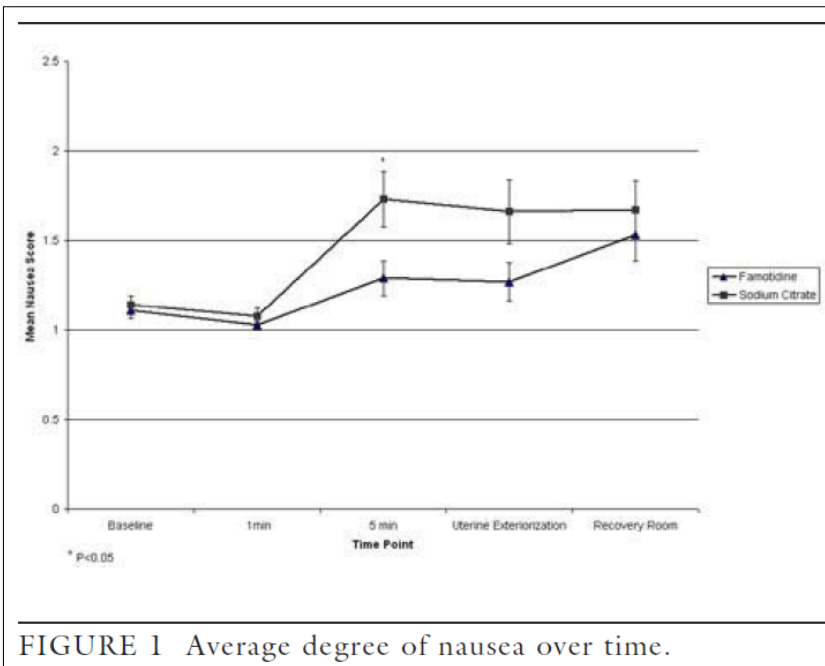


FIGURE 1 Average degree of nausea over time.

TABLE II Occurrence of nausea and hypotension

	Sodium citrate	Control	P-value
1 min after spinal			
Nausea	4/60 (7%)	2/63 (3%)	0.43
Hypotension	4/59 (7%)	4/61 (7%)	1
5 min after spinal			
Nausea	22/60 (37%)	9/63 (14%)	0.006
Hypotension	13/60 (22%)	11/63 (17%)	0.65
Uterine exteriorization			
Nausea	14/53 (26%)	7/52 (13%)	0.14
Hypotension	1/53 (2%)	2/53 (4%)	1
Recovery room			
Nausea	17/58 (29%)	15/62 (24%)	0.54
Hypotension	5/60 (8%)	9/63 (14%)	0.39

Nausea = 2–5 on a scale of 1–5, with 1 being no nausea and 5 being vomiting. Hypotension = systolic blood pressure \leq 100. Compared with Pearson Chi-square test.

Kjaer K et al. Can J Anaesth. 2006 Aug;53(8):776-80.

Interventions at caesarean section for reducing the risk of aspiration pneumonitis

S. Paranjothy,^a J.D. Griffiths,^b H.K. Broughton,^a G.M.L. Gyte,^c H.C. Brown,^d J. Thomas^c

^a *Department of Primary Care and Public Health, Clinical Epidemiology Interdisciplinary Research Group, School of Medicine, Cardiff University, Cardiff, UK*

^b *Department of Anaesthesia, Royal Women's Hospital, Parkville, Australia*

^c *Cochrane Pregnancy and Childbirth Group, School of Reproductive and Developmental Medicine, Division of Perinatal and Reproductive Medicine, University of Liverpool, Liverpool Women's NHS Foundation Trust, Liverpool, UK*

^d *Maternity Services Department, Royal Sussex County Hospital, Brighton, UK*

Int J Obstet Anesth. 2011 Apr;20(2):142-8

ABSTRACT

Background: Various interventions are used as prophylaxis for aspiration pneumonitis in obstetric anaesthesia. This review, based on a Cochrane systematic review currently being updated, examines whether interventions given before caesarean section reduce the risk of aspiration pneumonitis.

Methods: Twenty-two studies, involving 2658 women providing data in a usable format for meta-analysis were identified.

Results: Compared to no treatment or placebo, there was a significant reduction in the risk of intra-gastric pH <2.5 with antacids (risk ratio (RR) 0.17, 95% confidence interval (CI) 0.09–0.32), H2 antagonists (RR 0.09, 95% CI 0.05–0.18) and proton-pump antagonists (RR 0.26, 95% CI 0.14–0.46). H2 antagonists were associated with a reduced risk of intra-gastric pH <2.5 when compared with proton-pump antagonists (RR 0.39, 95% CI 0.16–0.97), but compared with antacids the findings were unclear. Combined use of antacids plus H2 antagonists was associated with a significant reduction in the risk of intra-gastric pH <2.5 when compared with placebo (RR 0.02, 95% CI 0.00–0.15) or compared with antacids alone (RR 0.12, 95% CI 0.02–0.92).

Conclusion: The quality of evidence was weak and may not reflect a reduction in the risk of aspiration pneumonitis since none of the studies assessed substantive clinical outcomes or potential adverse effects. Further work is required to validate the suitability of surrogate markers of pH and gastric volume for clinical outcomes in the context of aspiration pneumonitis.

RIZIKO ASPIRACE

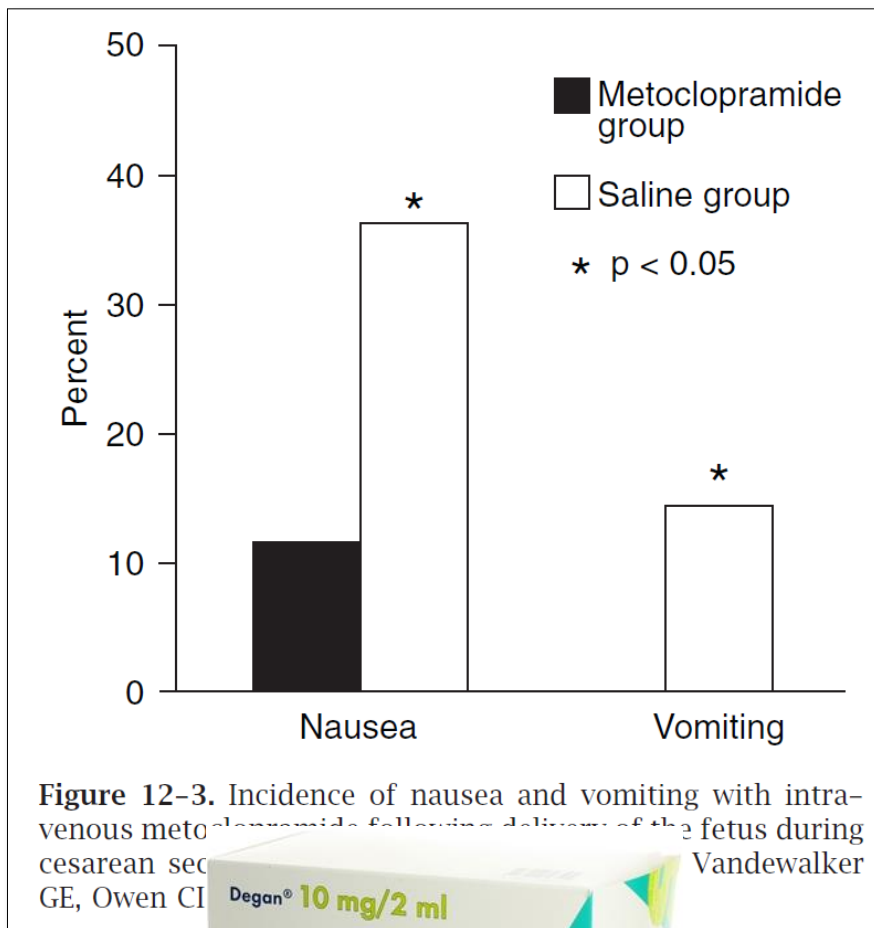


Figure 12-3. Incidence of nausea and vomiting with intravenous metoclopramide following delivery of the fetus during cesarean section. GE, Owen CI; Vandewalker

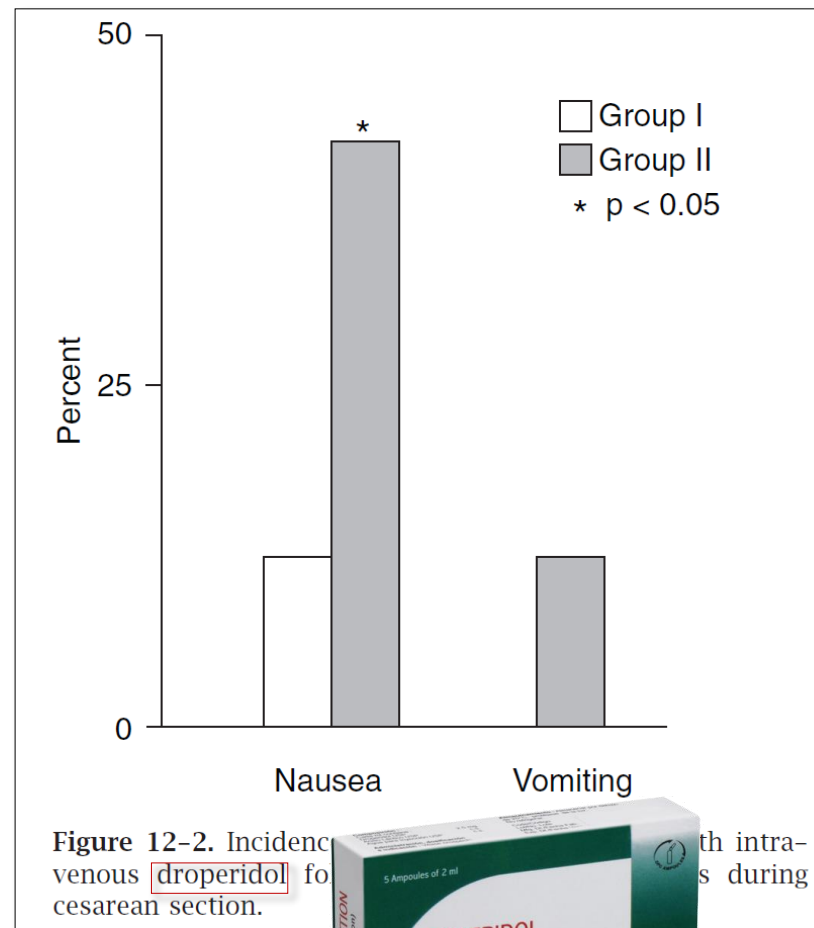
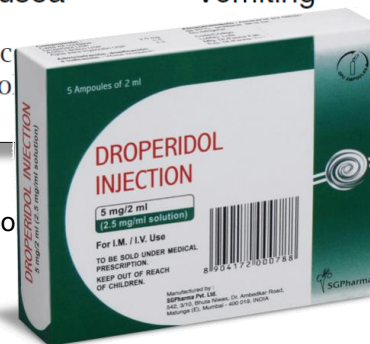


Figure 12-2. Incidence of nausea and vomiting with intravenous droperidol for cesarean section.



Sanjay Datta, ed. Obstetric Anesthesia Handbook

ger, 2006



Table 1. Lower Esophageal Sphincter, Intra-gastric, and Barrier Pressures Obtained before and after Administration of 0.15 mg/kg Intravenous Metoclopramide

	Before Metoclopramide		After Metoclopramide	
	Baseline		Baseline	Cricoid Pressure Applied
Lower esophageal pressure	14.1 ± 2.9		19.6 ± 4.7†	5.0 ± 4.3*
Intra-gastric pressure	4.6 ± 1.4		5.7 ± 1.9	5.8 ± 2.3
Barrier pressure	9.6 ± 3.4		14.1 ± 5.5†	-0.2 ± 5.1*

Data are in mmHg ± SD.

* $P < 0.05$ vs. respective baseline value. † $P < 0.05$ vs. respective pre-metoclopramide value.

Salem et al. Anesthesiology 2008; 109:806–10

TABLE 9. *Side effects of succinylcholine.*

- Massive hyperkalemia in susceptible patients
- Cardiac arrhythmias
- Muscle fasciculations
- Myalgias
- Rhabdomyolysis
- Increased intracranial pressure
- Increased intragastric pressure
- Increased intraocular pressure
- Malignant hyperthermia
- Masseter muscle spasm or jaw rigidity
- Prolonged apnea (1–4 hours), if atypical plasma cholinesterase

From Bevan DR. Complications of muscle relaxants. *Semin Anesth.* 1995;14:63.



THE LANCET

Preliminary Communications

CRICOID PRESSURE TO CONTROL REGURGITATION OF STOMACH CONTENTS DURING INDUCTION OF ANÆSTHESIA

WHEN the contents of stomach or œsophagus gain access to the air-passages during anæsthesia the consequences are disastrous. In spite of modern anæsthetic techniques—or sometimes, regrettably, because of them—regurgitation is still a considerable hazard during the induction of anæsthesia, particularly for operative obstetrics and emergency general surgery.¹⁻⁸

By a simple manœuvre during induction of anæsthesia, regurgitation of gastric or œsophageal contents can be controlled until intubation with a cuffed endotracheal tube is completed. The same manœuvre may also be used to prevent inflation of the stomach (a potent cause of regurgitation) resulting from positive-pressure ventilation

1. De Lee, J. B., Greenhill, J. P. *Principles and Practice of Obstetrics*; p. 255. Philadelphia, 1951.
2. Mendelson, C. L. *Amer. J. Obstet. Gynec.* 1946, **52**, 191.
3. Morton, H. J. V., Wylie, W. D. *Anæsthesia*, 1951, **6**, 190.
4. Coleman, D. J., Day, B. L. *Lancet*, 1956, **i**, 708.
5. Edwards, G., Morton, H. J. V., et al. *Anæsthesia*, 1956, **ii**, 194.
6. *Lancet*, 1956, **i**, 734.
7. *Rep. Publ. Hlth med. Subj., Lond.* no. 97, 1957.
8. *Reports on Confidential Enquiries into Maternal Deaths in England and Wales, 1952-54 and 1955-57.* H.M. Stationery Office



Sellick B. *The Lancet* 1961;2:404

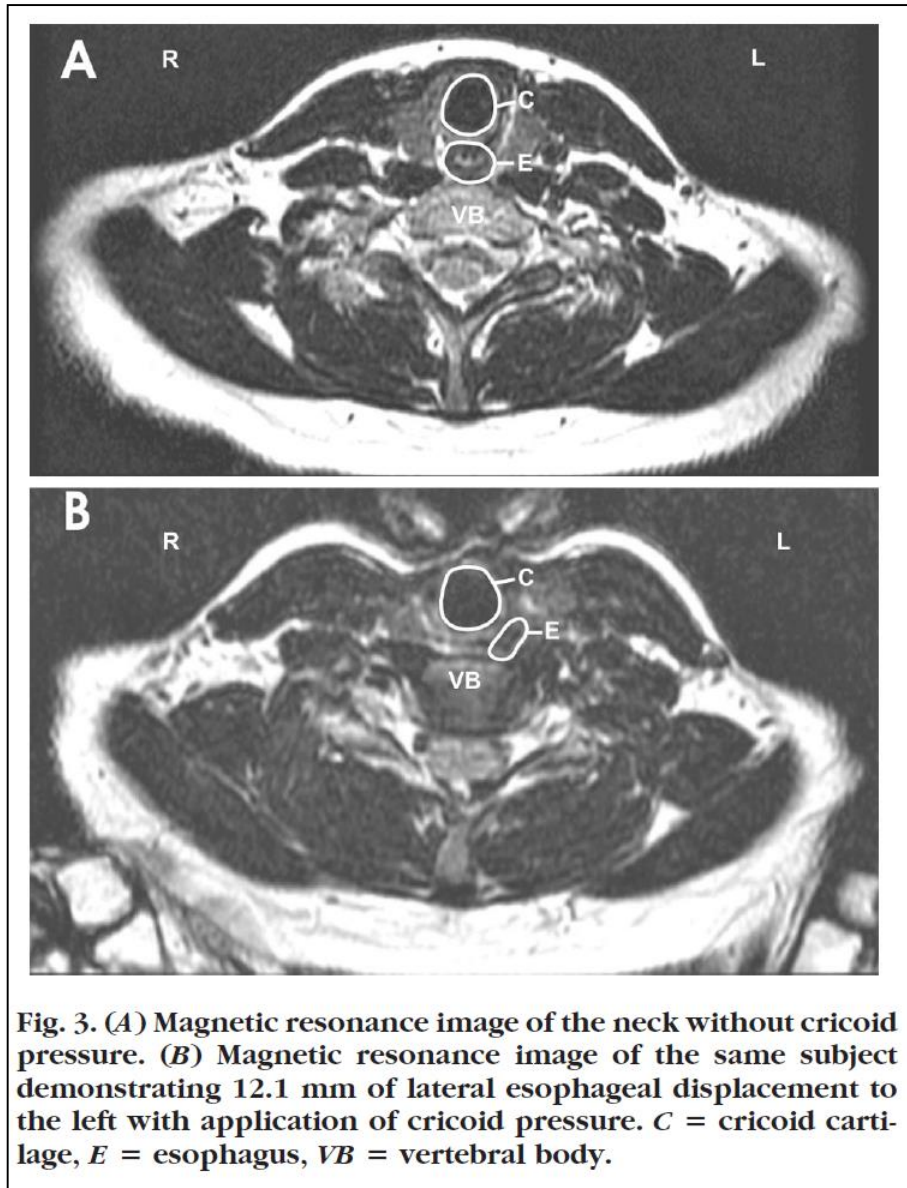
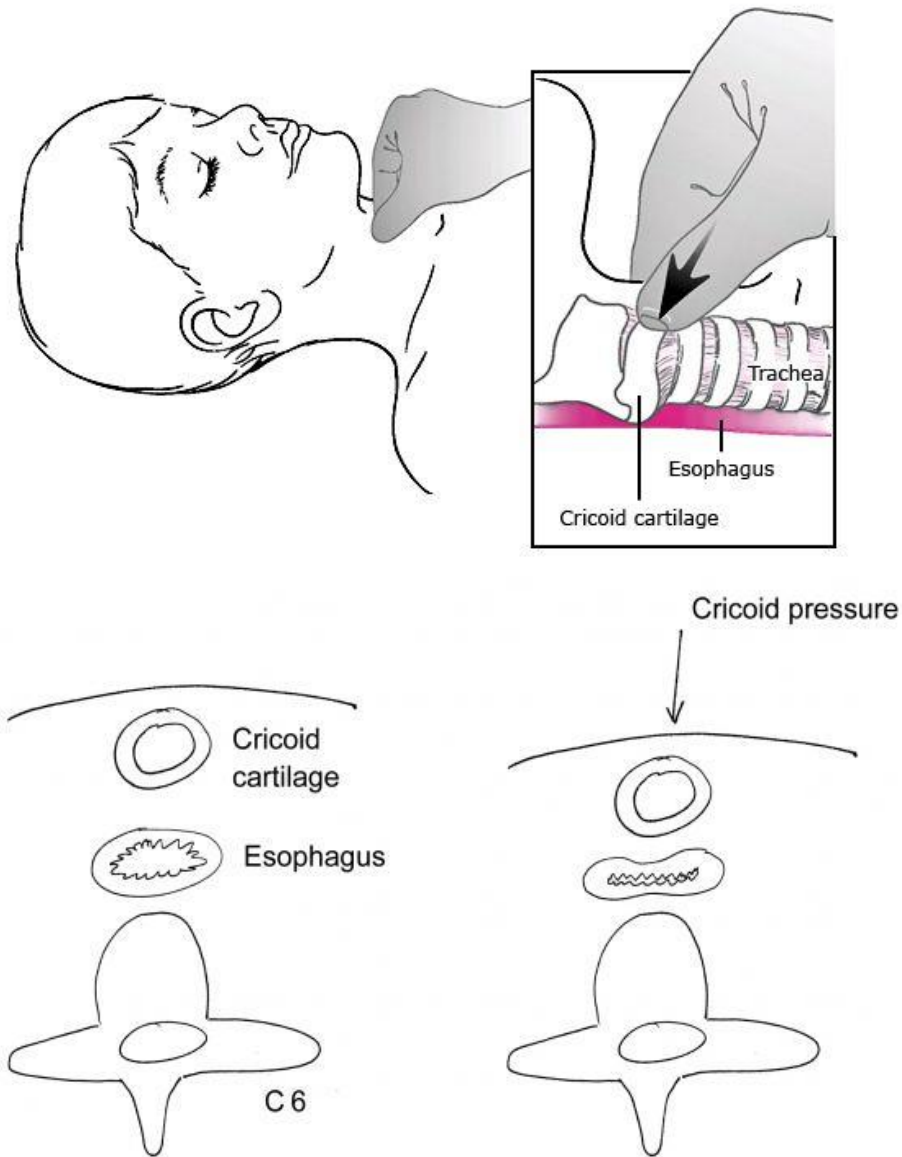
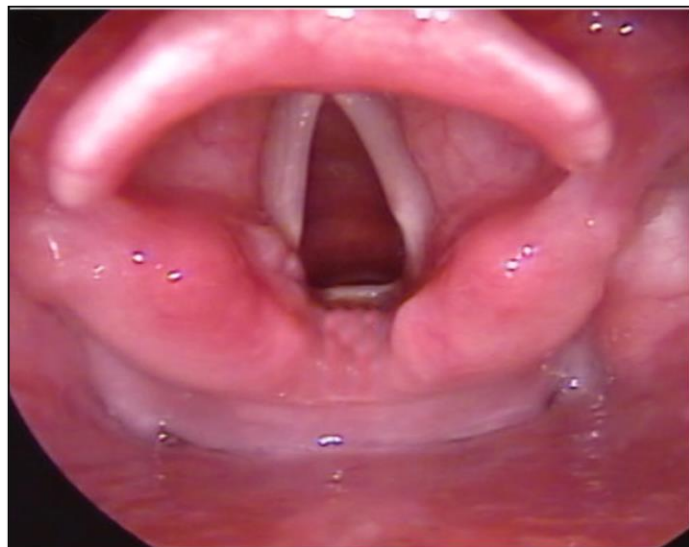


Fig. 3. (A) Magnetic resonance image of the neck without cricoid pressure. **(B)** Magnetic resonance image of the same subject demonstrating 12.1 mm of lateral esophageal displacement to the left with application of cricoid pressure. *C* = cricoid cartilage, *E* = esophagus, *VB* = vertebral body.

Smith KJ et al. *Anesthesiology* 2003; 99:60-4

Rice et al. *Anesth Analg* 2009;109:1546-52



...tlak 30 N na krikoidní chrupavku může zcela 'zrušit' vizualizaci glotis

Haslam et al. *Anaesthesia* 2005; 60: 41-47

Effect of Cricoid Pressure on the Success of Endotracheal Intubation with a Lightwand

R. Eric Hodgson, M.B., Ch.B.(Hons.), F.C.A.(S.A.)(Crit. Care),* P. Dean Gopalan, M.B., Ch.B., F.C.A.(S.A.),* Richard C. Burrows, M.B., Ch.B., F.C.A.(S.A.)(Crit. Care),† Khangelani Zuma, M.Sc.‡

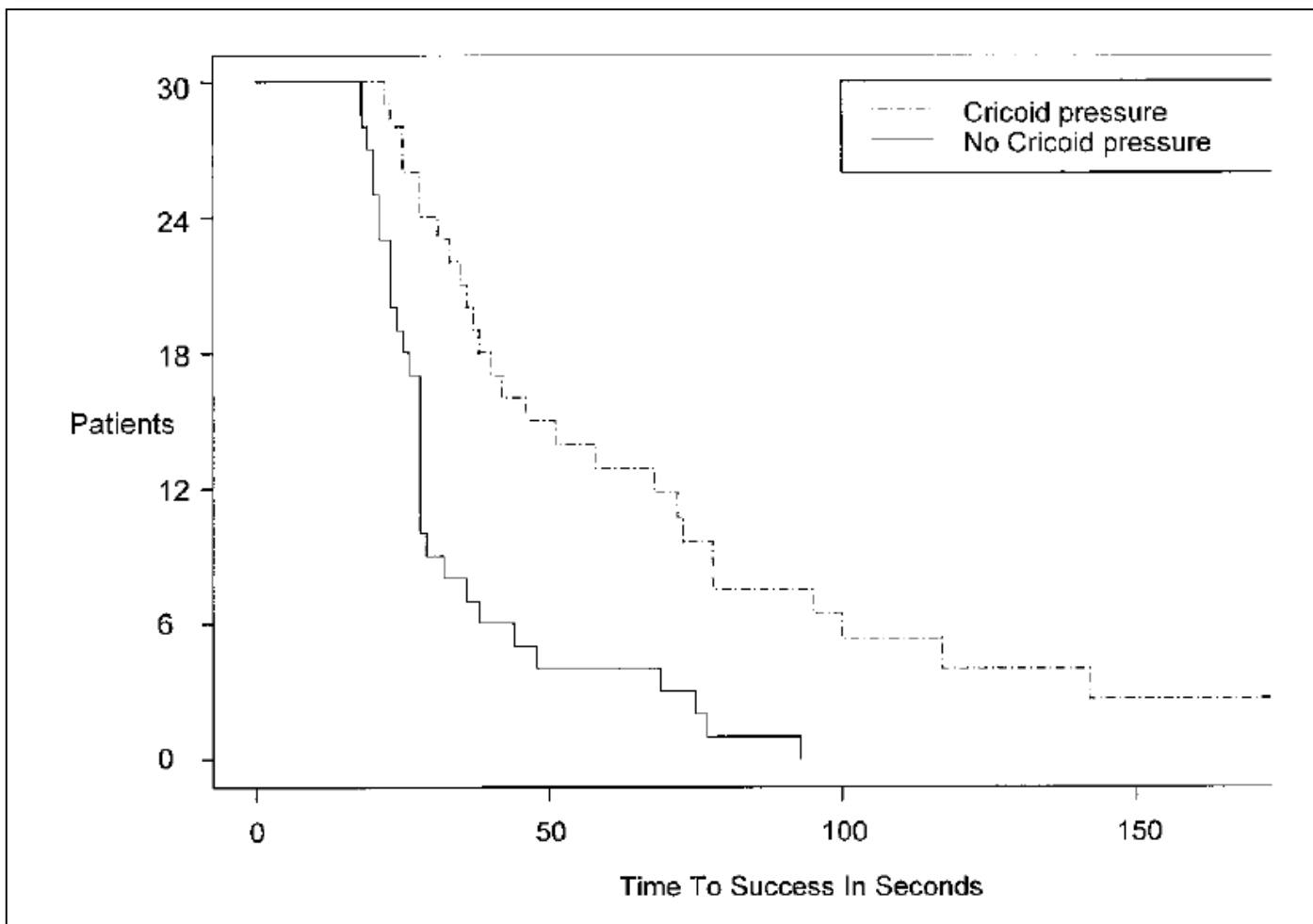
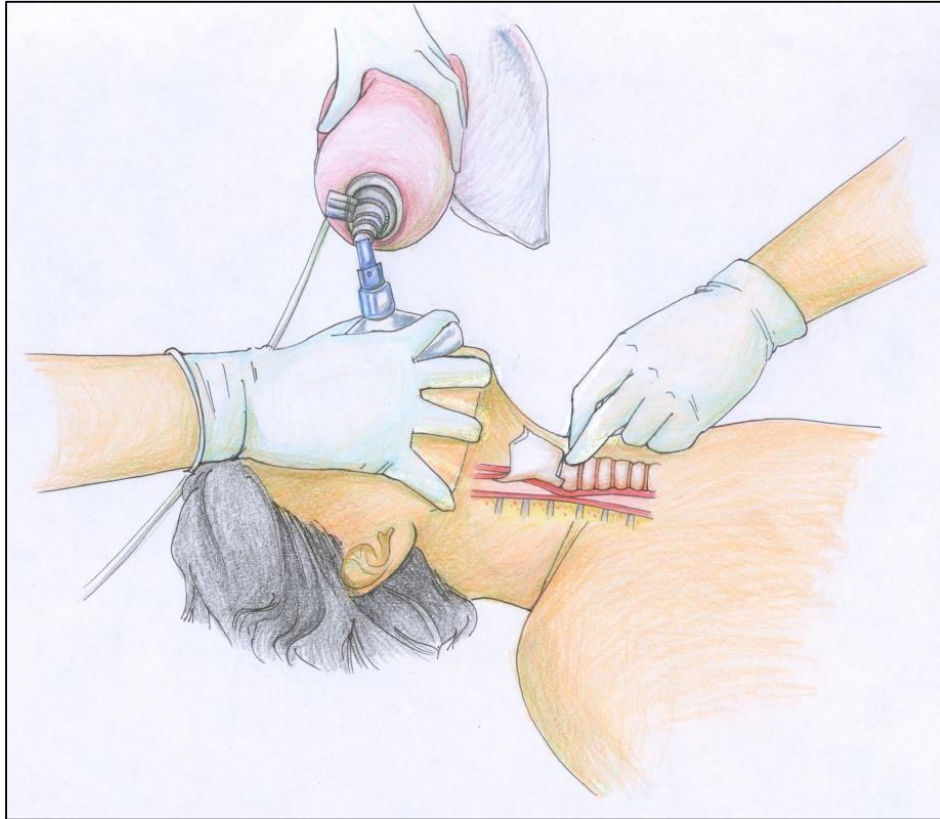


Fig. 1. Kaplan-Meier plot of time to successful intubation for the sixty patients. Thirty patients were intubated with cricoid pressure, with values for four patients requiring multiple attempts censored. Thirty patients were intubated without cricoid pressure. Time to successful intubation was significantly prolonged in the cricoid pressure group ($P = 0.0001$, log-rank test).

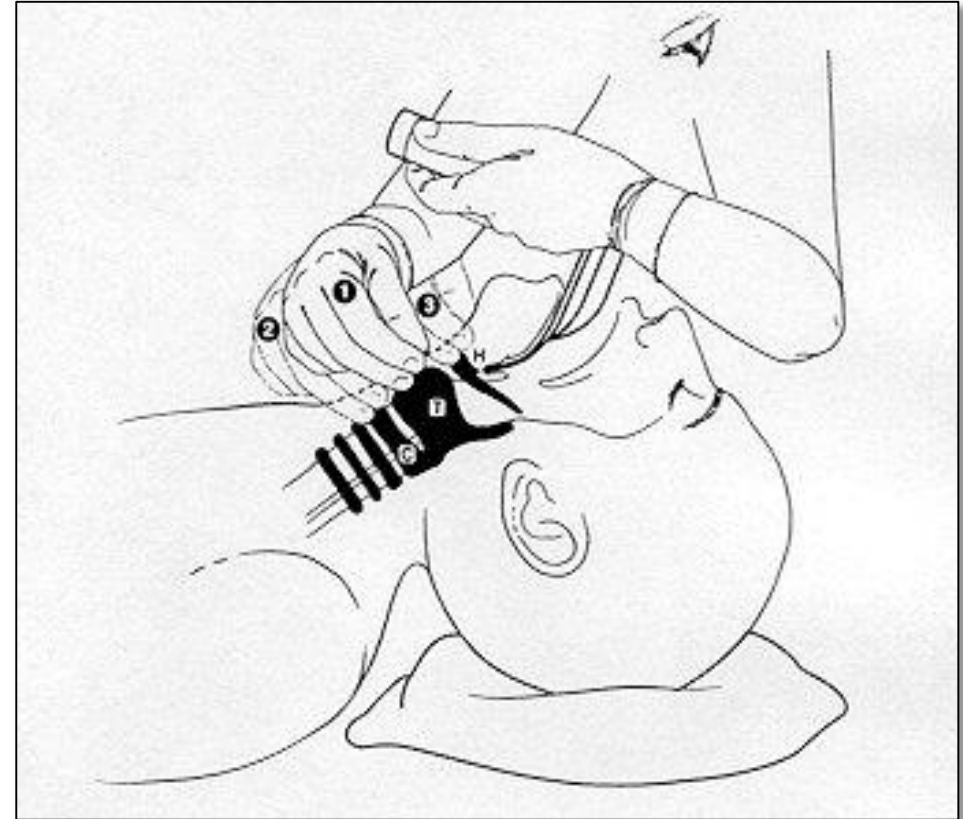
Hodgson. Anesthesiology 2001; 94:259-62

Sellick's Maneuver



“BURP”

Backward, Upward, Rightward Pressure



V 90% případů získáme nejlepší “pohled” tlakem na **štítnou chrupavku**, nikoli krikoidální!

**ZKONTROLOVAT
ODSÁVAČKU !**

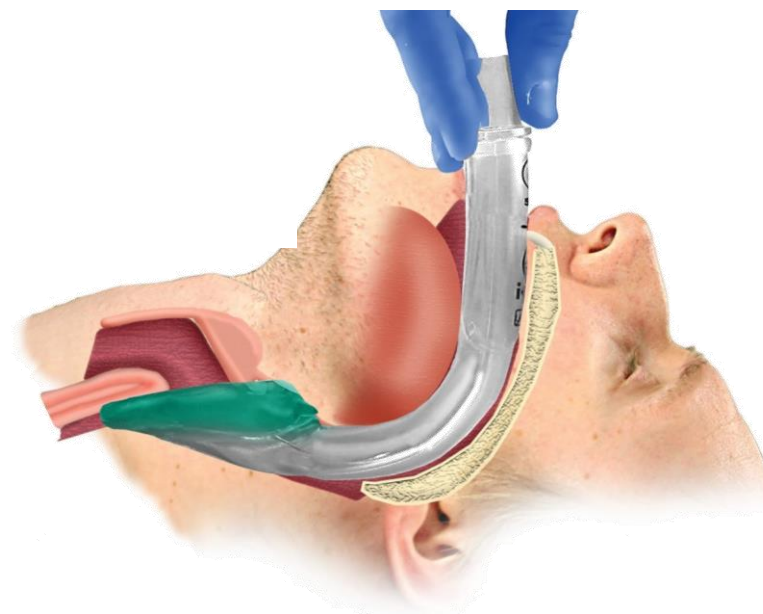


**ZKONTROLOVAT
ODSÁVAČKU !**





PAMATUJ, ŽE OXYGENACE JE DŮLEŽITĚJŠÍ NEŽ INTUBACE.



ProSeal



Classic



Supreme



LT



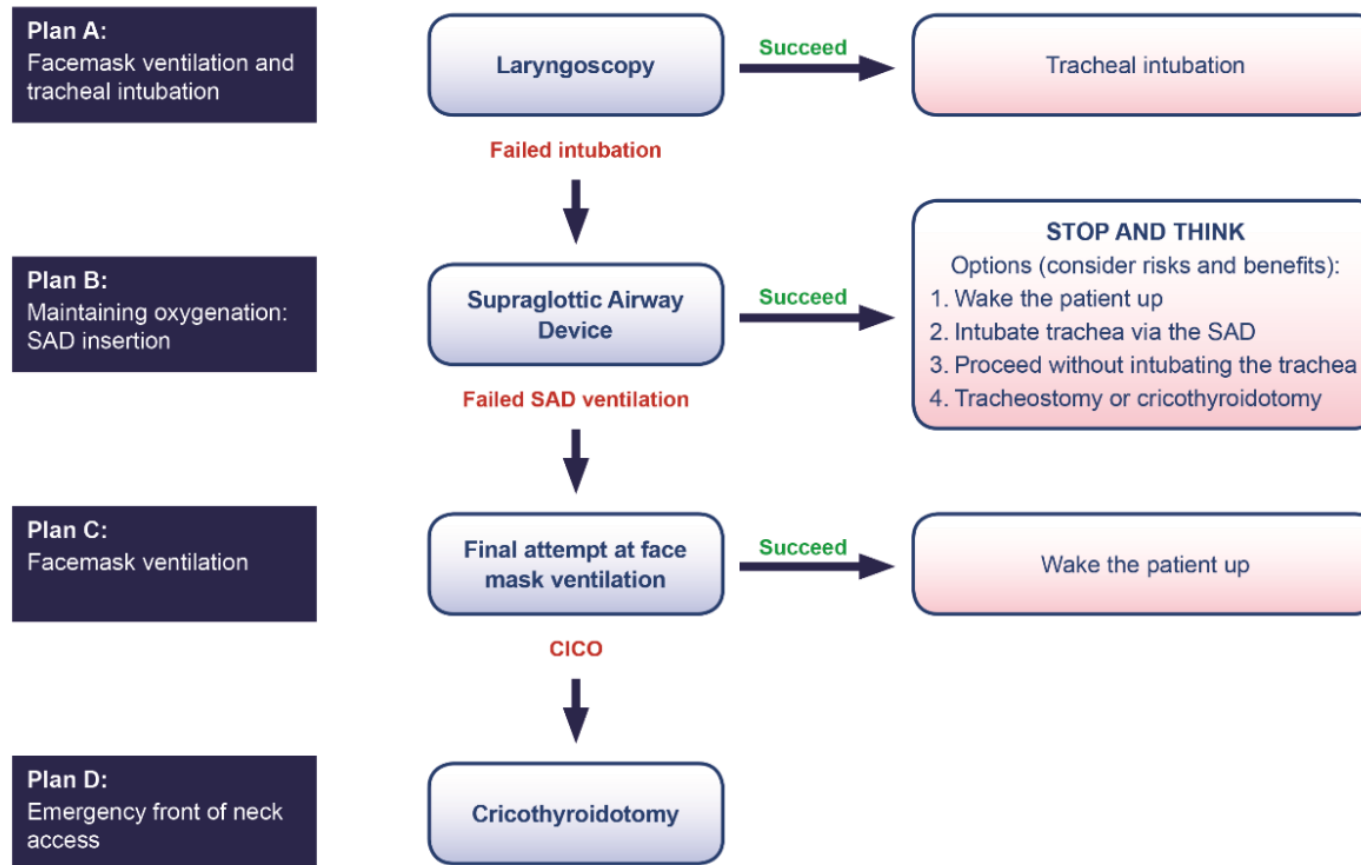
i-gel



air-Q



DAS Difficult intubation guidelines – overview



This flowchart forms part of the DAS Guidelines for unanticipated difficult intubation in adults 2015 and should be used in conjunction with the text.

CELKOVÁ ANESTEZIE

The salient characteristics of RSI were delineated by Stept and Safar in 1970 [3].

- Preoxygenation
- Predetermined doses of thiopental and SCh
- Cricoid force
- Avoidance of ventilation by bag and mask
- Tracheal intubation

Thiopental

mg/mL

PROPOFOL

mg/mL

Sharp LM, Levy DM. Current Opinion in Anaesthesiology 2009, 22:357-361

CELKOVÁ ANESTEZIE

The salient characteristics of RSI were delineated by Stept and Safar in 1970 [3].

- Preoxygenation
- Predetermined doses of thiopental and SCh
- Cricoid force
- Avoidance of ventilation by bag and mask
- Tracheal intubation



Sharp LM, Levy DM. Current Opinion in Anaesthesiology 2009, 22:357-361

The Response of Newborns to Succinylcholine and d-Tubocurarine

Leonard F. Walts, M.D., and John B. Dillon, M.D.†*

Anesthesiology. 1969 Jul;31(1):35-8.

Results

Mean age of the 60 adult patients was 41 years. The group given succinylcholine received an average of 68 mg (range 54–83) of drug. All patients had 100 per cent depression in twitch force. Recovery times to 10, 50 and 90 per cent of control values averaged 7.0, 8.5, and 10 minutes, respectively.

SVALOVÁ RELAXACE

SUKCINYLCHOLIN

- ❖ Nejrychlejší nástup účinku
- ❖ Výborné intubační podmínky
- ❖ Neprochází placentou
- ❖ **Doporučená dávka 1-1,5 mg/kg**



Table 3. Onset Times and Durations of Neuromuscular Block

Succinylcholine dose (mg/kg)	Onset time(s)	Duration of block (min)	<i>n</i>
0.3	72 ± 30	4.4 ± 1.4	13
0.5	68 ± 44	5.2 ± 1.8	27
1.0	53 ± 23	5.9 ± 1.9†	30
1.5	56 ± 31	7.2 ± 2*	30
2.0	52 ± 21	7.5 ± 1.7*	30

Values are means ± SD.

**P* < 0.01 versus succinylcholine 0.3, 0.5, and 1.0 mg/kg groups; †*P* < 0.05 versus succinylcholine 0.3 mg/kg group.

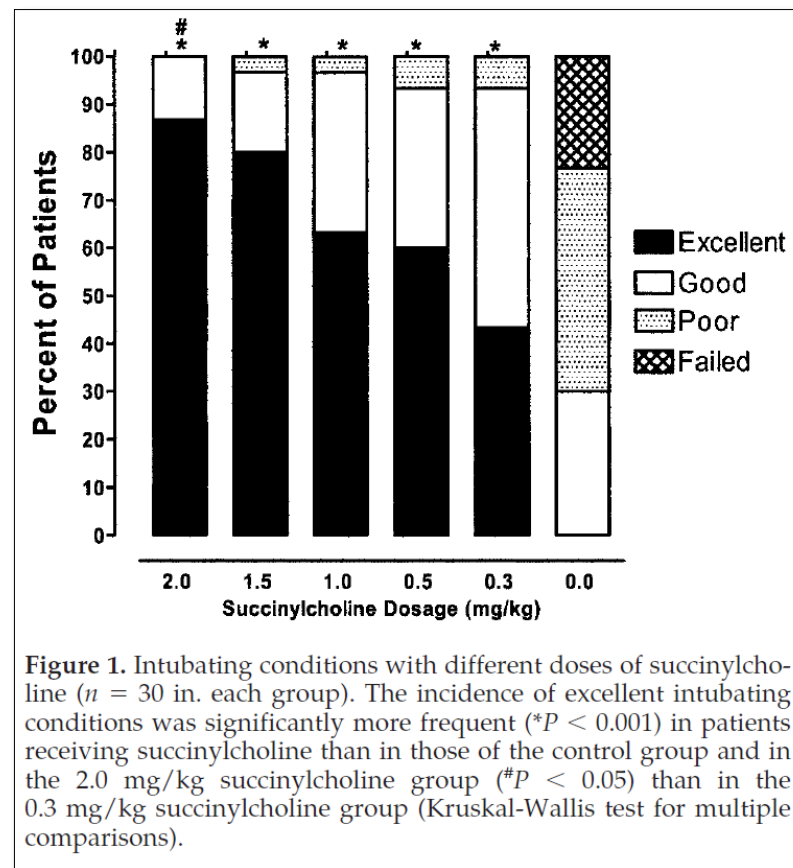
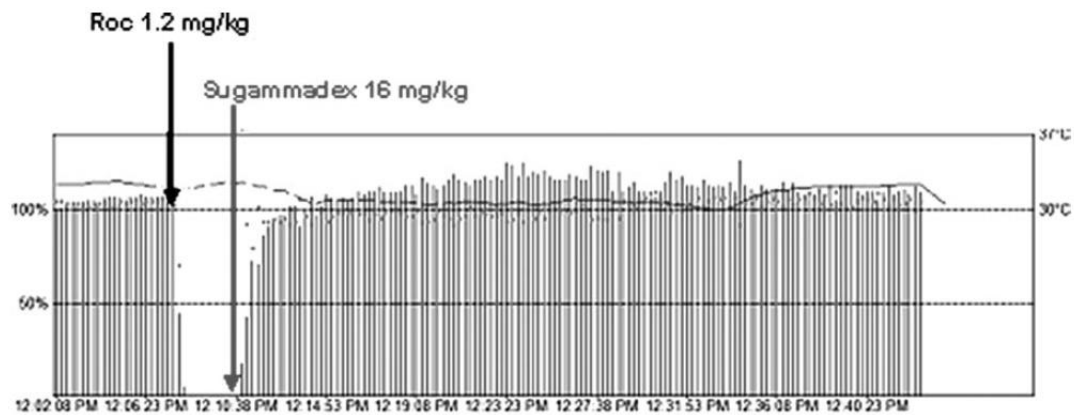
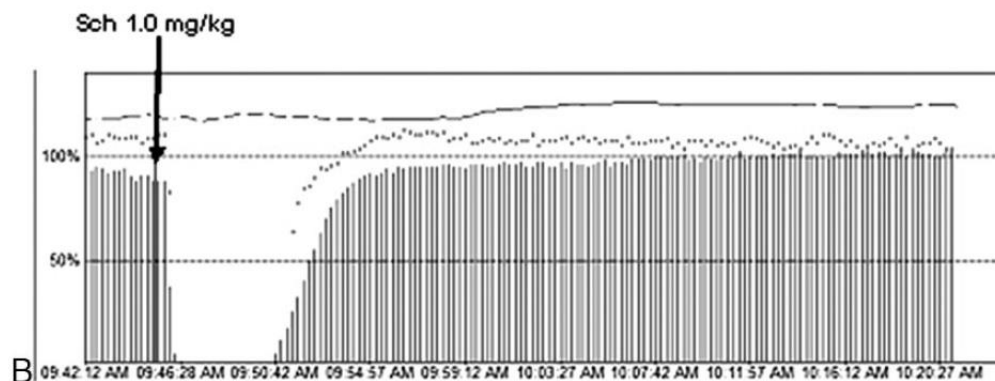


Figure 1. Intubating conditions with different doses of succinylcholine (*n* = 30 in each group). The incidence of excellent intubating conditions was significantly more frequent (**P* < 0.001) in patients receiving succinylcholine than in those of the control group and in the 2.0 mg/kg succinylcholine group (*#P* < 0.05) than in the 0.3 mg/kg succinylcholine group (Kruskal-Wallis test for multiple comparisons).

Naguib M et al. Anesth Analg 2006;102:151-5



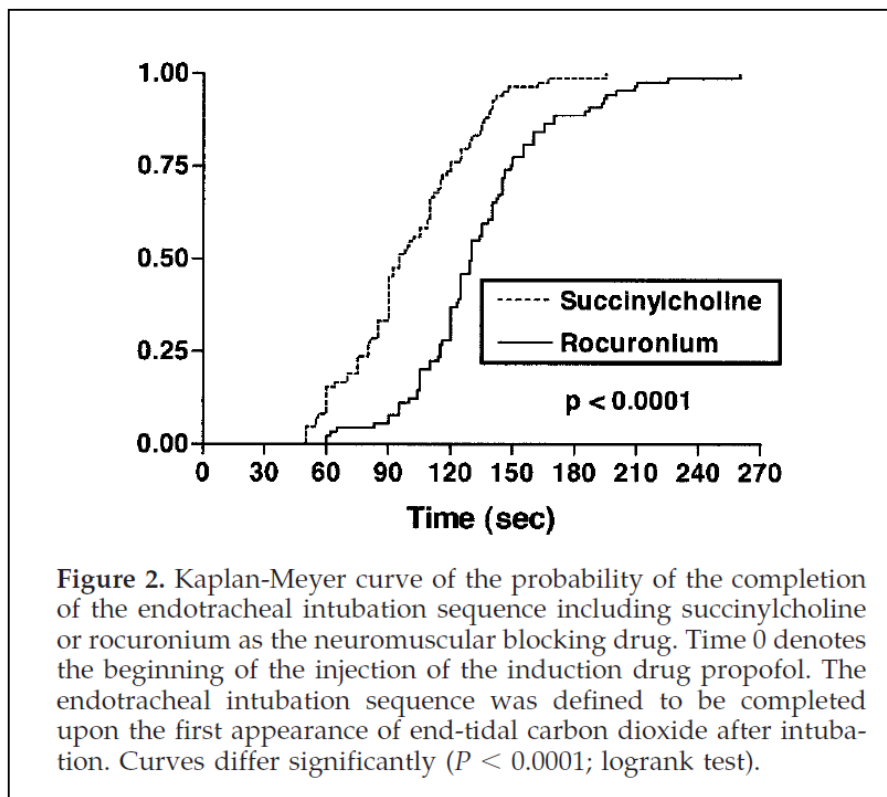
A
3 min



B
09:42 12 AM 09:46 28 AM 09:50 42 AM 09:54 57 AM 09:59 12 AM 10:03 27 AM 10:07 42 AM 10:11 57 AM 10:16 12 AM 10:20 27 AM

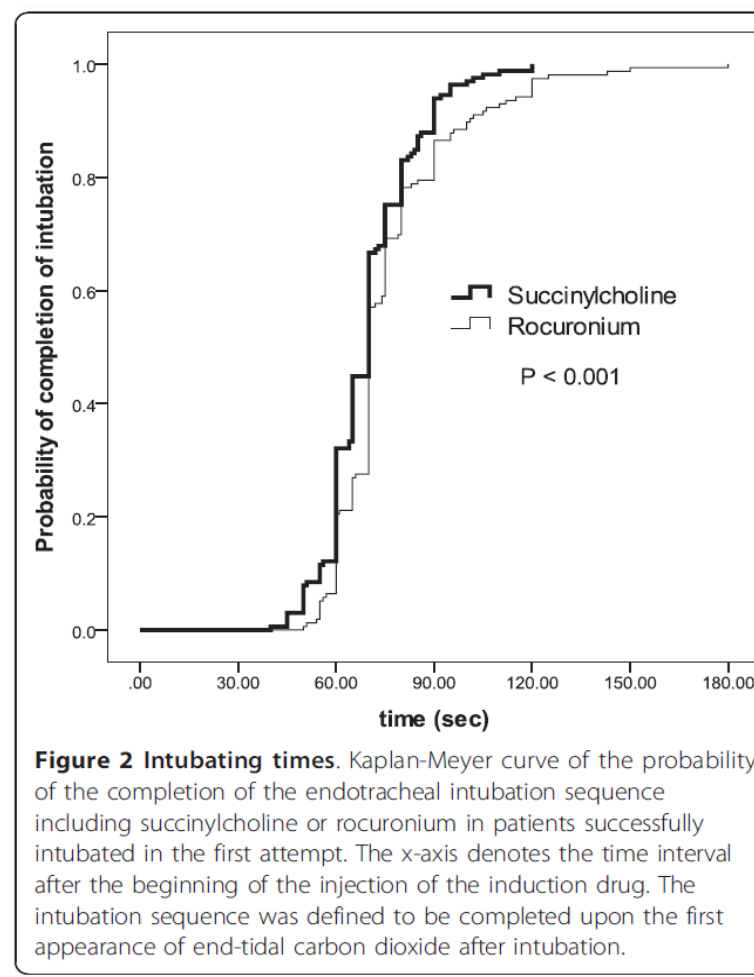
Figure 4. Panel A shows the recovery of the twitch height and train-of-four (TOF) ratio after administration of 1.2 mg/kg rocuronium followed 3 min later by 16 mg/kg sugammadex, both given IV. Recovery to a first twitch height (T1) of 90% and a TOF ratio of 0.94 occurred 110 s later. The onset-offset time with this sequence (i.e., the time from the end of the injection of rocuronium to a T1 recovery to 90%) was 4 min 47 s. Panel B shows the effects of administering 1.0 mg/kg succinylcholine (Sch) with spontaneous recovery to a T1 of 90% occurring after 9 min and 23 s.

0,6 mg/kg



Sluga M et al. Anesth Analg 2005;101:1356 –61

1 mg/kg



Stephan C Marsch, et al. Crit Care. 2011;15(4):R199-R199

Desaturation following rapid sequence induction using succinylcholine vs. rocuronium in overweight patients

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Background: Rapid sequence induction may be associated with hypoxemia. The purpose of this study was to investigate the possible difference in desaturation during rapid sequence induction in overweight patients using either succinylcholine or rocuronium.

Methods: Sixty patients with a body mass index (BMI) between 25 and 30 kg/m², American Society of Anesthesiologists class I or II, undergoing general anesthesia were randomly divided into a succinylcholine group and a rocuronium group. After a 3-min preoxygenation, patients received rapid sequence induction of general anesthesia with midazolam–fentanyl–propofol and succinylcholine (1.5 mg/kg) or rocuronium (0.9 mg/kg). Ventilation was not initiated until oxygen saturation reached 98%, 96%, 94% and 92%. Safe Apnea Time was defined as the time from administration of neuromuscular blocking drugs to oxygen saturation fell to 92%. The recovery period was defined as the time from initiation of

ventilation until oxygen saturation was 97%. Arterial blood gases were taken at baseline, after preoxygenation and at 92% oxygen saturation.

Results: The mean Safe Apnea Time (95% CI) was 283 (257–309) s in succinylcholine vs. 329 (303–356) s in rocuronium ($P = 0.01$). The mean recovery period (95% CI) was 43 (39–48) s in succinylcholine vs. 36 (33–38) s in rocuronium ($P = 0.002$). Blood gas analysis showed no difference between the two groups.

Conclusions: Succinylcholine was associated with a significantly more rapid desaturation and longer recovery of oxygen saturation than rocuronium during rapid sequence induction in overweight patients.

Accepted for publication 29 October 2010

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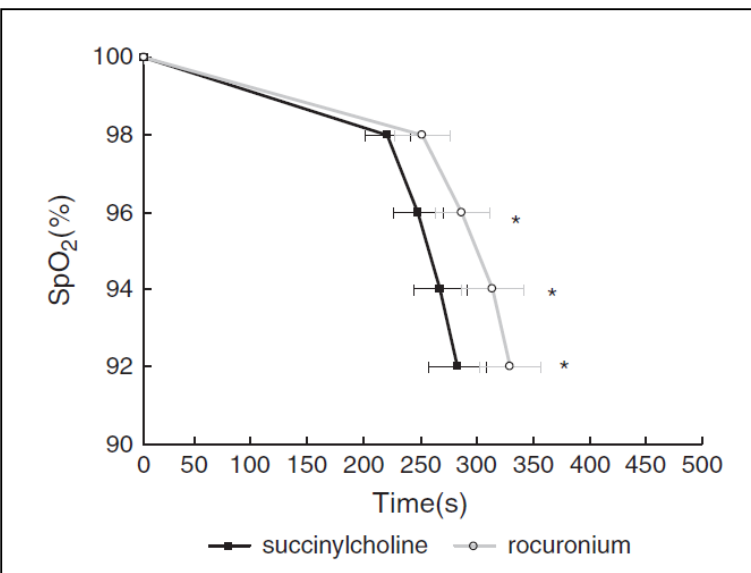


Fig. 2. Changes in oxygen saturation (S_pO_2) with time during non-hypoxic apnea in the succinylcholine or the rocuronium group. Mean values (points) for both groups are shown. The vertical lines indicate 95% CI. The curves show smooth before S_pO_2 reach 98%, but afterward fall straightly to 92% S_pO_2 . * $P < 0.05$ compared with succinylcholin.

ORIGINAL ARTICLE

Effect of suxamethonium vs rocuronium on onset of oxygen desaturation during apnoea following rapid sequence induction

S. K. Taha,¹ M. F. El-Khatib,² A. S. Baraka,³ Y. A. Haidar,⁴ F. W. Abdallah,⁵ R. A. Zbeidy⁴ and S. M. Siddik-Sayyid¹

¹ Associate Professor, ² Professor, ³ Emeritus Professor, ⁴ Chief Resident, ⁵ Fellow, Department of Anesthesiology, American University of Beirut, Beirut, Lebanon

Summary

This study investigates the effect of suxamethonium vs rocuronium on the onset of haemoglobin desaturation during apnoea, following rapid sequence induction of anaesthesia. Sixty patients were randomly allocated to one of three groups. Anaesthesia was induced with lidocaine 1.5 mg.kg⁻¹, fentanyl 2 µg.kg⁻¹ and propofol 2 mg.kg⁻¹, followed by either rocuronium 1 mg.kg⁻¹ (Group R) or suxamethonium 1.5 mg.kg⁻¹ (Group S). The third group received propofol 2 mg.kg⁻¹ and suxamethonium 1.5 mg.kg⁻¹ only (Group SO). The median (IQR [range]) time to reach S_pO_2 of 95% was significantly shorter in Group S (358 [311–373 [215–430]] s) than in Group R (378 [370–393 [366–420]] s; $p = 0.003$), and shorter in Group SO (242 [225–258 [189–370]] s) than in both Group R ($p < 0.001$) and Group S ($p < 0.001$). When suxamethonium is administered for rapid sequence induction of anaesthesia, a faster onset of oxygen desaturation is observed during the subsequent apnoea compared with rocuronium. However, time to desaturation is prolonged whenever lidocaine and fentanyl precede suxamethonium.

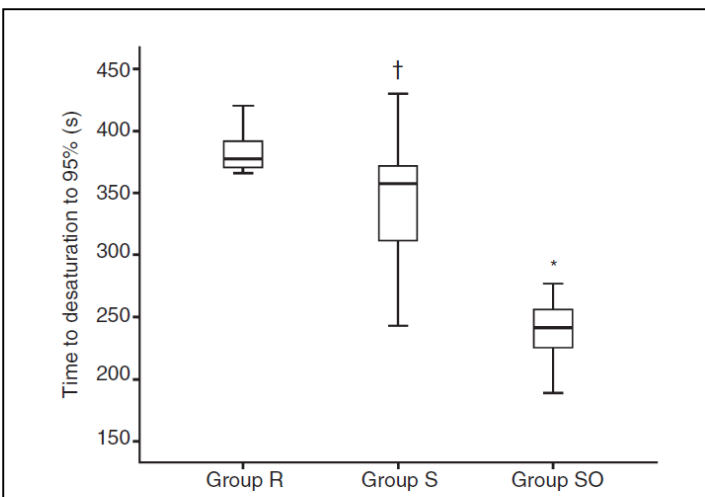


Figure 1 Time to reach S_pO_2 of 95% during apnoea following induction of anaesthesia with lidocaine/fentanyl/propofol/rocuronium (Group R), lidocaine/fentanyl/propofol/suxamethonium (Group S), or propofol/suxamethonium (Group SO).

Rocuronium and sugammadex for rapid sequence induction of obstetric general anaesthesia

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Liverpool Women's Hospital, Liverpool, UK

Background: Many anaesthetists use rocuronium in place of suxamethonium for rapid sequence induction (RSI). This is less common in obstetric anaesthesia as the duration of action of an effective dose of rocuronium exceeds most obstetric procedures. Sugammadex offers the possibility of rapidly reversing profound rocuronium neuromuscular blockade at the end of surgery. We aimed to determine whether rocuronium 1.2 mg/kg used for RSI in the obstetric population would provide good intubating conditions at 60 s and would be effectively reversed by sugammadex at the end of surgery.

Methods: We present a prospective series of 18 patients who received rocuronium 1.2 mg/kg at induction of anaes-

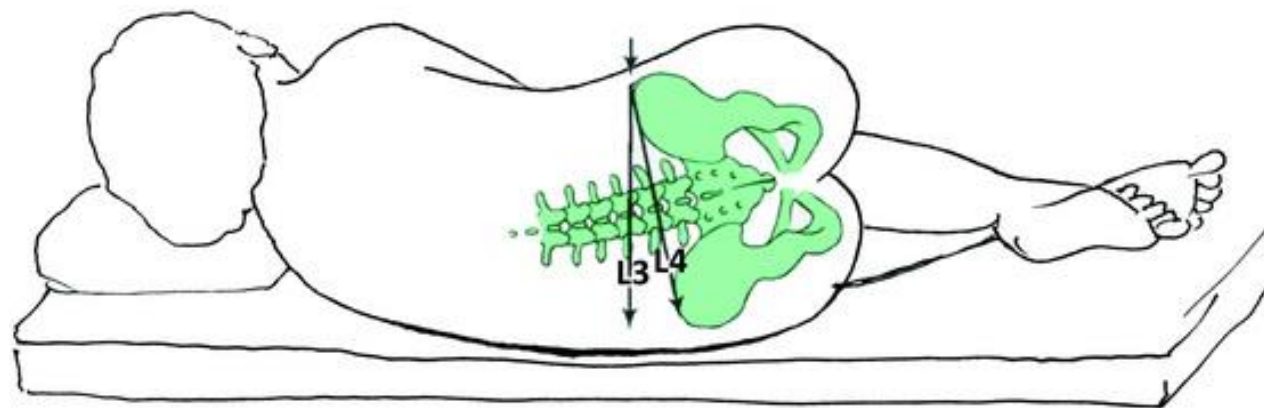
thesia, monitored with a train-of-four ratio (TOF)-Watch SX[®], and reversed using sugammadex 4 mg/kg.

Results: The mean (95% CI) onset time of rocuronium was 71 (56–86) s, and the mean (95% CI) time to recovery of the TOF to $\geq 90\%$, after the administration of sugammadex 4 mg/kg at the end of surgery, was 86 (69–104) s.

Conclusion: Rocuronium 1.2 mg/kg reversed by sugammadex appears to be effective in the obstetric population.

Accepted for publication 26 February 2011

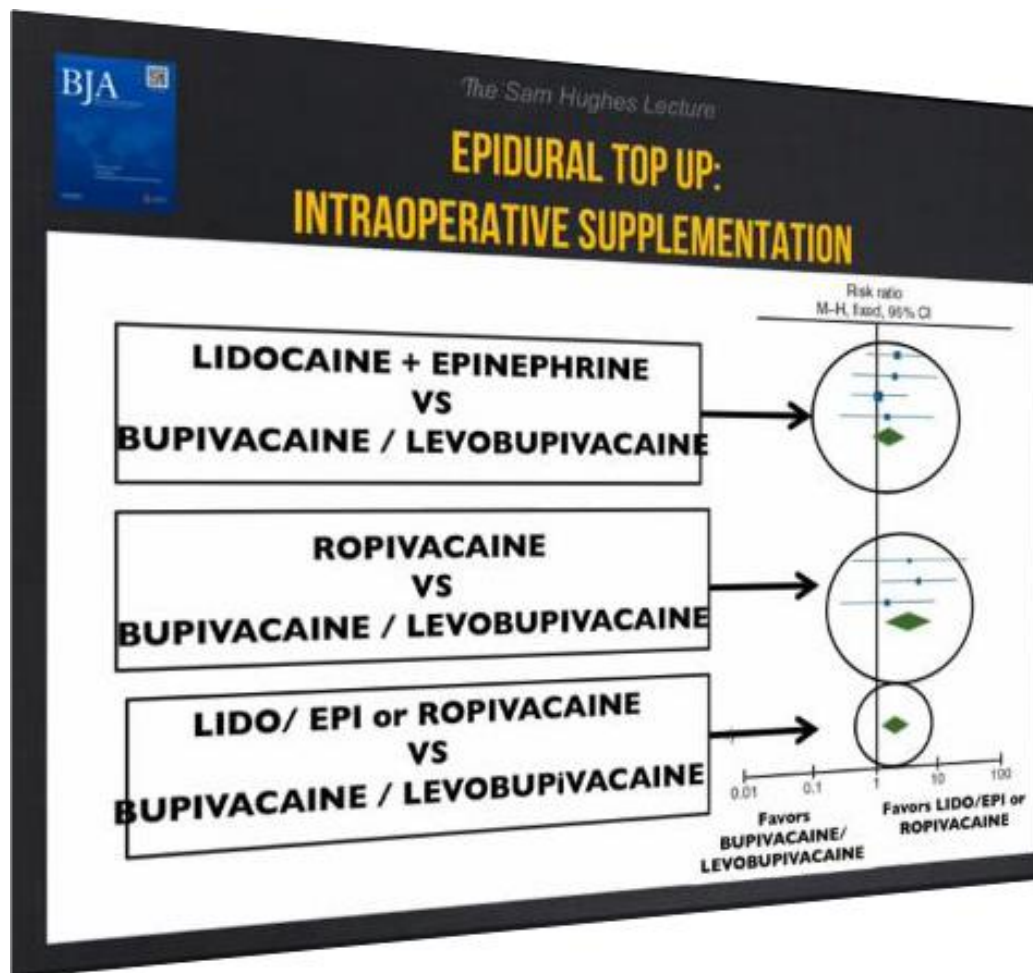
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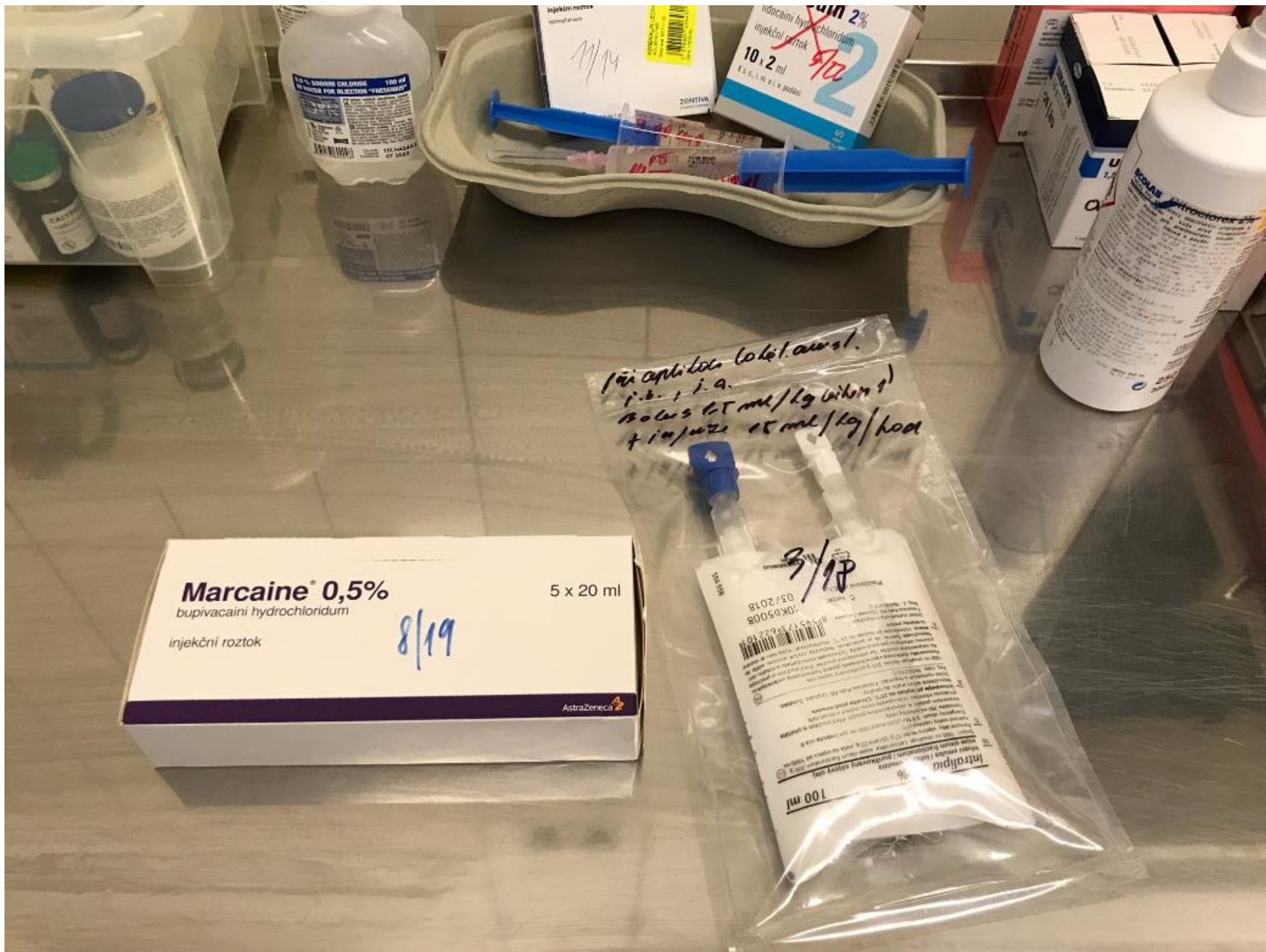
Obr 1. Projekce spojnice obou iliakálních krist u ležící těhotné ženy.



epidurální směs: lidocain 2% 18 ml + sufentanyl 10 µg/2 ml + adrenalin 0,1 ml



epidurální směs: lidocain 2% 18 ml + sufentanyl 10 µg/2 ml + adrenalin 0,1 ml





ELSEVIER

www.obstetanesthesia.com

ORIGINAL ARTICLE

The extension of epidural blockade for emergency caesarean section: a survey of Scandinavian practice

K. Wildgaard,^a F. Hetmann,^b M. Ismaiel^{a,c}

^aDepartment of Anaesthesiology, Næstved Hospital, Næstved, Denmark

^bDepartment of Nursing, Oslo and Akershus University College of Applied Sciences, Oslo, Norway

^cDepartment of Anaesthesiology, Malmö Central Sykehus, Skånes Universitetssjukhus, Malmö, Sweden

48

Epidural top-ups for caesarean section

Table 1 Location of epidural top-up in Denmark, Norway and Sweden

	Denmark (n=43)	Norway (n=43)	Sweden (n=59)	Total (n=145)	Trainee recommendation (n=138)
Full dose in labour ward	6 (14.0%)	5 (11.6%)	3 (5.1%)	14 (9.7%)	10 (6.9%)
Test dose in labour ward	9 (20.9%)	7 (16.3%)	18 (30.5%)	34 (23.4%)	26 (17.9%)
Transfer to theatre before giving local anaesthetic	22 (51.2%)	29 (67.4%)	36 (61.0%)	87 (60.0%)	95 (65.5%)
No top-up					
Alternative not described	0	0	1 (1.7%)	1 (0.7%)	0
Spinal anaesthesia	2 (4.7%)	1 (2.3%)	0	3 (2.1%)	3 (2.1%)
Unclear*	4 (9.3%)	1 (2.3%)	1 (1.7%)	6 (4.1%)	4 (2.8%)

Data are number (%). *Dependent on clinical presentation.

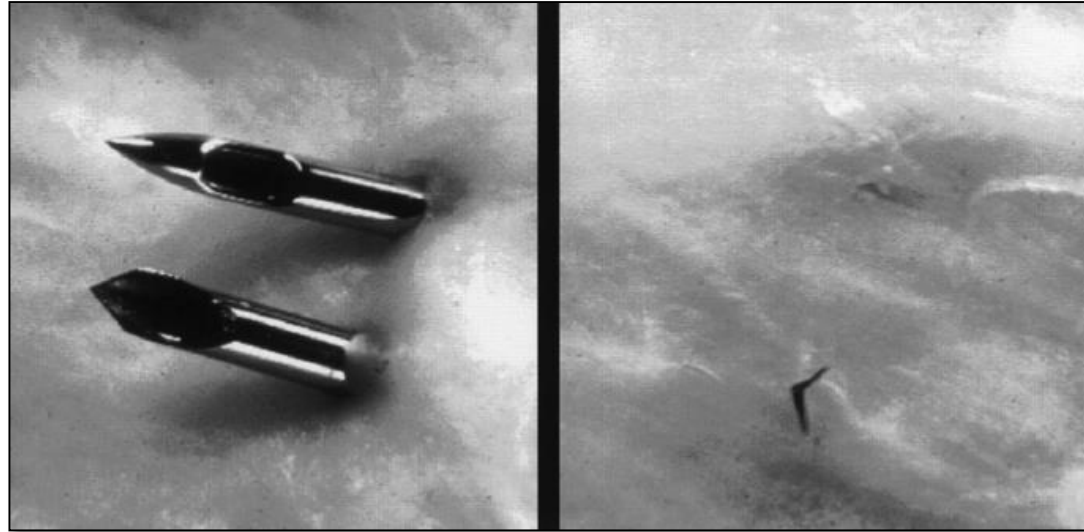


FIGURE 1. Dural puncture holes made by cutting and noncutting needles (Reproduced with permission from Strupp, et al. *Neurology*. 2001; 57:2310–2312).



HUBER POINT



Edward B. Tuohy
(1908–1959, USA)



Robert F. Husted
(1928–2008, USA)

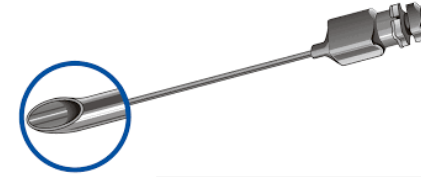
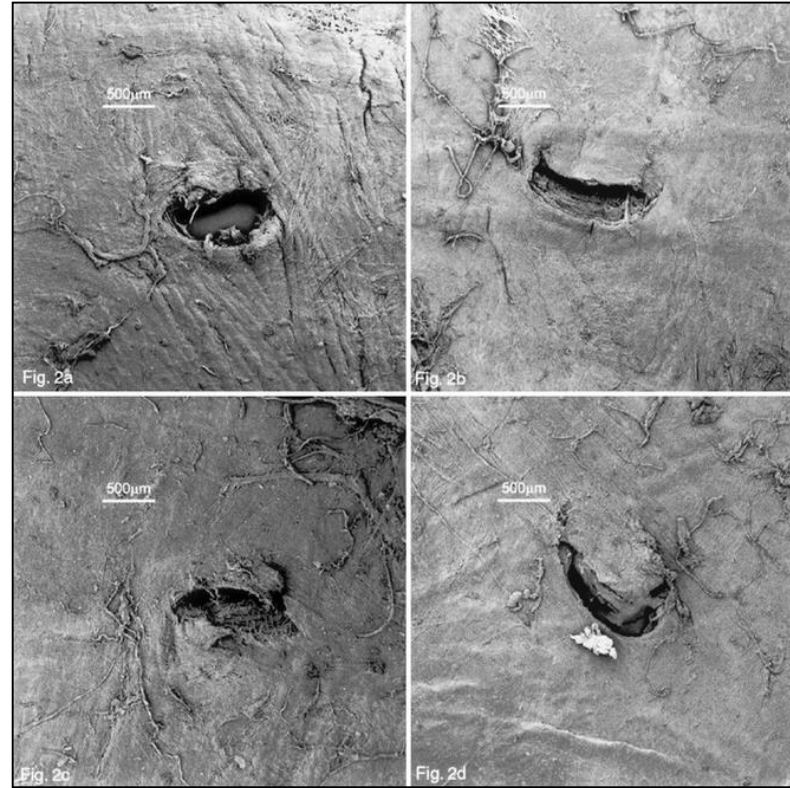
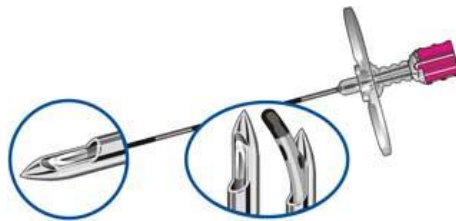


Fig. 2. Scanning electron microscopic images of (a) a 17-gauge Husted epidural needle puncture (bevel parallel, 90° angle), (b) a 17-gauge Tuohy epidural needle puncture (bevel parallel, 90° angle), (c) an 18-gauge Special Sprotte® epidural needle puncture (90° angle), and (d) an 18-gauge Crawford epidural needle puncture (bevel parallel, 90° angle).

Pamela J. Angle et al. Anesthesiology. 2003;99(6):1376-1382



Table 2. Effect of Epidural Needle Design on CSF Leak (90° Punctures, Bevel Parallel), Cadaver n = 10

Epidural Needles	17-Gauge Hustead	17-Gauge Tuohy	18-Gauge Tuohy	20-Gauge Tuohy	18-Gauge Special Sprotte®	18-Gauge Crawford
17-Gauge Hustead	516 ± 319	0.3668	0.2922	0.0018*	0.2078	0.1326
17-Gauge Tuohy		405 ± 209	0.8812	0.0024*	0.6468	0.4312
18-Gauge Tuohy			420 ± 191	0.0003*	0.4324	0.2707
20-Gauge Tuohy				100 ± 112	0.8182	0.0001*
18-Gauge Special Sprotte®					360 ± 208	0.9698
18-Gauge Crawford						356 ± 121

Part 1 results are presented in the form of a *P* value matrix. Mean ± SD cerebrospinal fluid (CSF) leak rates are found on the diagonal for each needle in ml/15-min interval. The table may be read in the following way: Mean ± SD leak for the 17-g Hustead = 516 ± 319 (17-g Hustead [row] vs. 17-g Hustead [column]). Mean ± SD leak rate for the 17-g Tuohy (row) vs. 17-g Tuohy (column) = 405 ± 209. *P* value for differences in leak for the 17-g Hustead (row) vs. 17-g Tuohy (column) = 0.3668. *P* value required to reach statistical significance, corrected for multiple testing = 0.003.

* Statistically significant *P* values.

Pamela J. Angle et al. Anesthesiology. 2003;99(6):1376-1382

Original Article

Influence of needle diameter on spinal anaesthesia puncture failures for caesarean section: A prospective, randomised, experimental study

Fausto Fama^{b,1,*}, Cecile Linard^{b,1}, Damien Bierlaire^a, Maria Gioffre'-Florio^b, Jacques Fusciardi^a, Marc Laffon^a^a University Hospital of Tours, Department of Anaesthesiology and Intensive Care, Hôpital Bretonneau, 2, rue de la Croix Verte, 37000 Tours cedex 9, France^b University Hospital of Messina, Department of Human Pathology, Via Consolare Valeria, 1

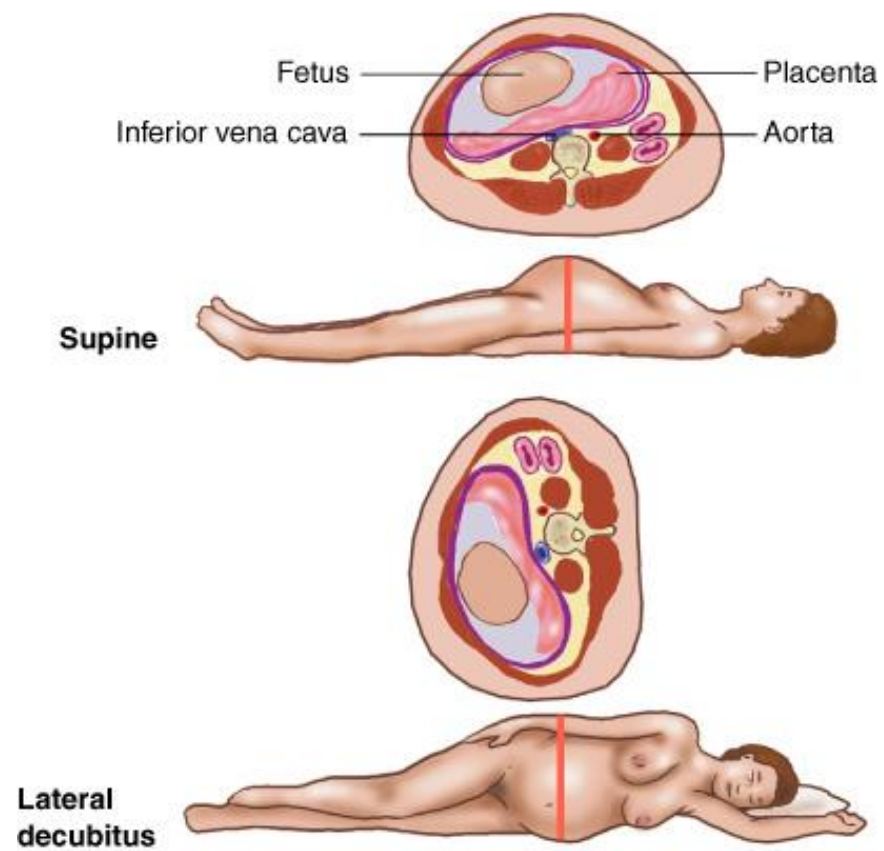

27 G více selhání,
26 G více PDPH

Table 2

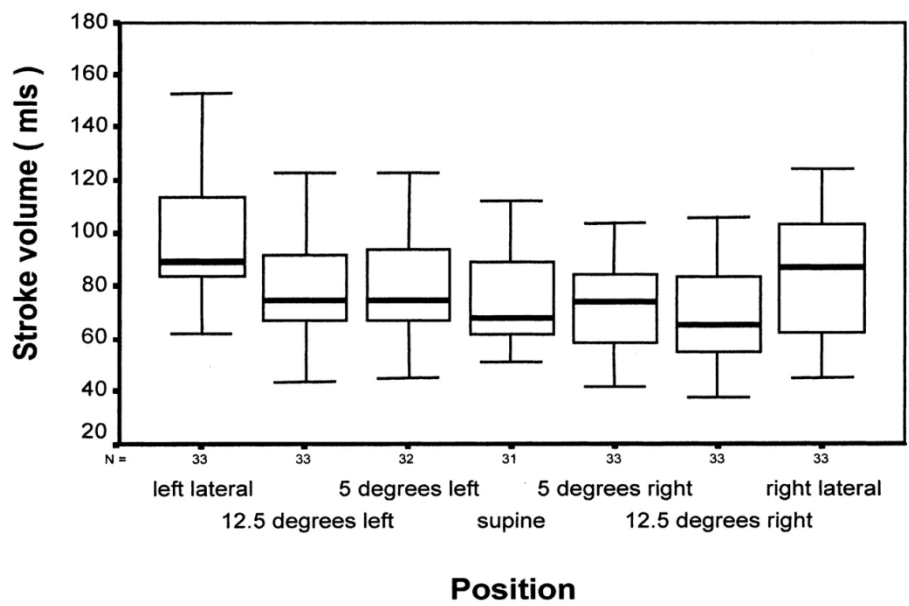
Spinal puncture failures and incidence of postdural puncture headache. The number of puncture failures was statistically significant in the 27 G group ($P=0.006$ versus the 25 G group, $P<0.001$ versus the 26 G group). No statistically significant difference was found between the 25 G and 26 G groups ($P=0.606$). Only 2 general anaesthesia procedures were carried out after 25 G attempt failures.

Group	25 G	26 G	27 G
Number of patients: <i>n</i>	109	121	98
Failure: <i>n</i> (%)	2 (1.8) ^a	1 (0.9) ^a	12 (10.9)
Headache: <i>n</i> (%)	5 (4.6)	3 (2.5)	2 (2.0)
Blood patch: <i>n</i>	1	1	0

^a $P<0.05$, 27 G vs. 25G and 26 G.

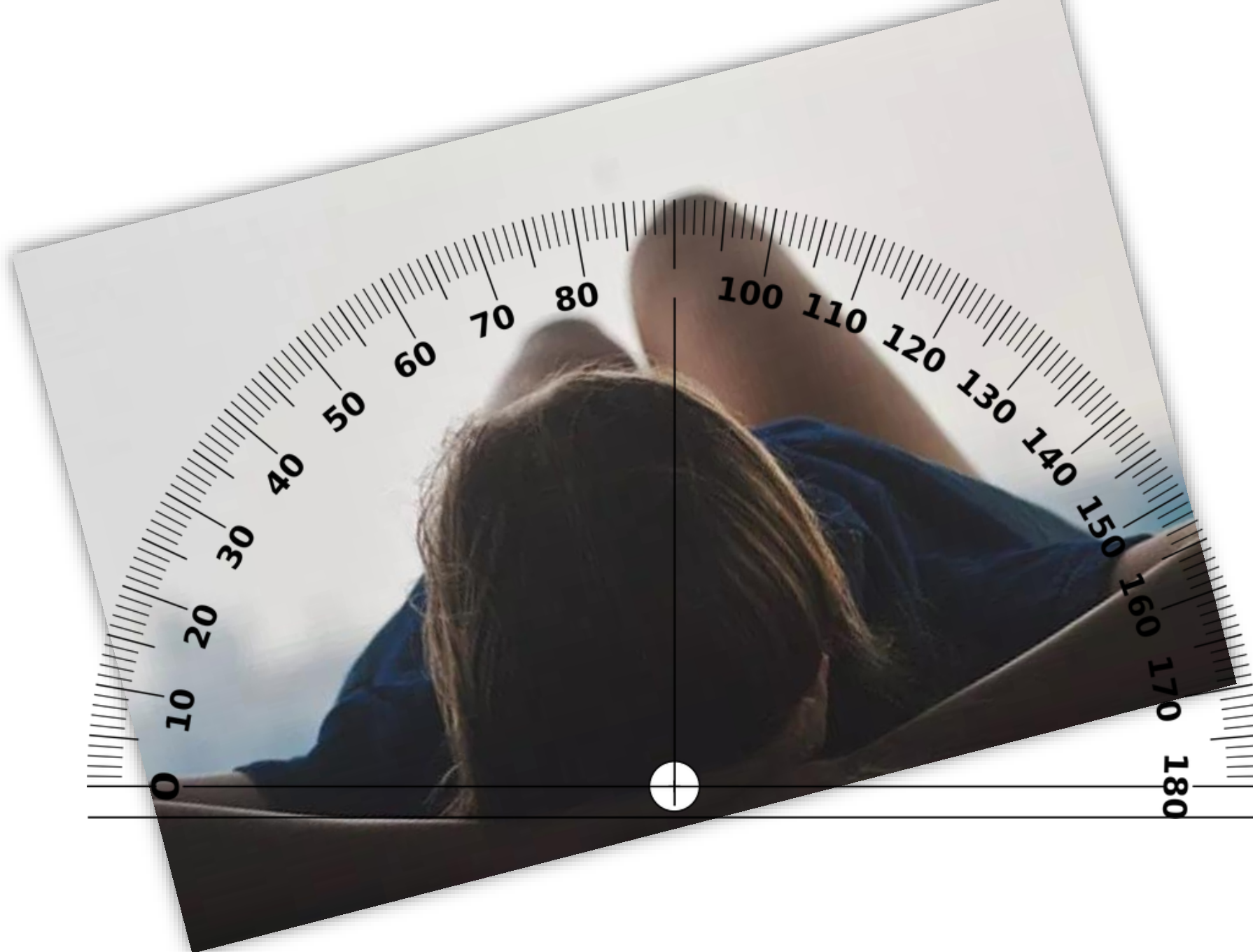


SEMILATERÁLNÍ POLOHA
 naklonění trupu o 5-15 stupňů
 = prevence aortokavální komprese



Bamber, J. H. et al. Anesth Analg 2003;97:256-258





ORIGINAL ARTICLE

A comparison of the haemodynamic effects of lateral and sitting positions during induction of spinal anaesthesia for caesarean section

B.I. Obasuyi, S. Fyनेface-Ogan, C.N. Mato

Department of Anaesthesia, University of Port Harcourt Teaching Hospital, Port Harcourt, Rivers State, Nigeria

ABSTRACT

Background: Hypotension during spinal anaesthesia occurs commonly in parturients. The speed of onset of sensory block and thus the haemodynamic effects of spinal anaesthesia for caesarean section using plain bupivacaine may be influenced by the position of the patient.

Objective: To compare the haemodynamic effects of spinal anaesthesia for caesarean section using plain bupivacaine, 10 mg, in the lateral position (Group L) and the sitting position (Group S). Using a 20% decrease in systolic blood pressure as the primary end point, we hypothesized that the sitting position would result in a faster onset of hypotension.

Methods: A prospective, randomized, controlled study was conducted in the operating theatre. Parturients were recruited from the obstetric ward and were allocated to either the lateral position (Group L) or the sitting position (Group S). Spinal anaesthesia was induced with plain bupivacaine, 10 mg, in either position. The time to onset of hypotension was defined as the time from the start of the spinal anaesthetic to the first 20% decrease in systolic blood pressure. The time to maximum hypotension was defined as the time from the start of the spinal anaesthetic to the lowest recorded mean arterial pressure. The incidence of hypotension was defined as the number of patients who had a 20% decrease in systolic blood pressure at any time during the procedure. The time to onset of hypotension was compared between the two groups using the Mann-Whitney U-test. The time to maximum hypotension was compared between the two groups using the Mann-Whitney U-test. The incidence of hypotension was compared between the two groups using the chi-square test. A p-value of < 0.05 was considered statistically significant.

Results: The time to onset of hypotension was significantly shorter in the sitting position (Group S) compared to the lateral position (Group L) (p < 0.05). The time to maximum hypotension was also significantly shorter in the sitting position (Group S) compared to the lateral position (Group L) (p < 0.05). The incidence of hypotension was significantly higher in the sitting position (Group S) compared to the lateral position (Group L) (p < 0.05).

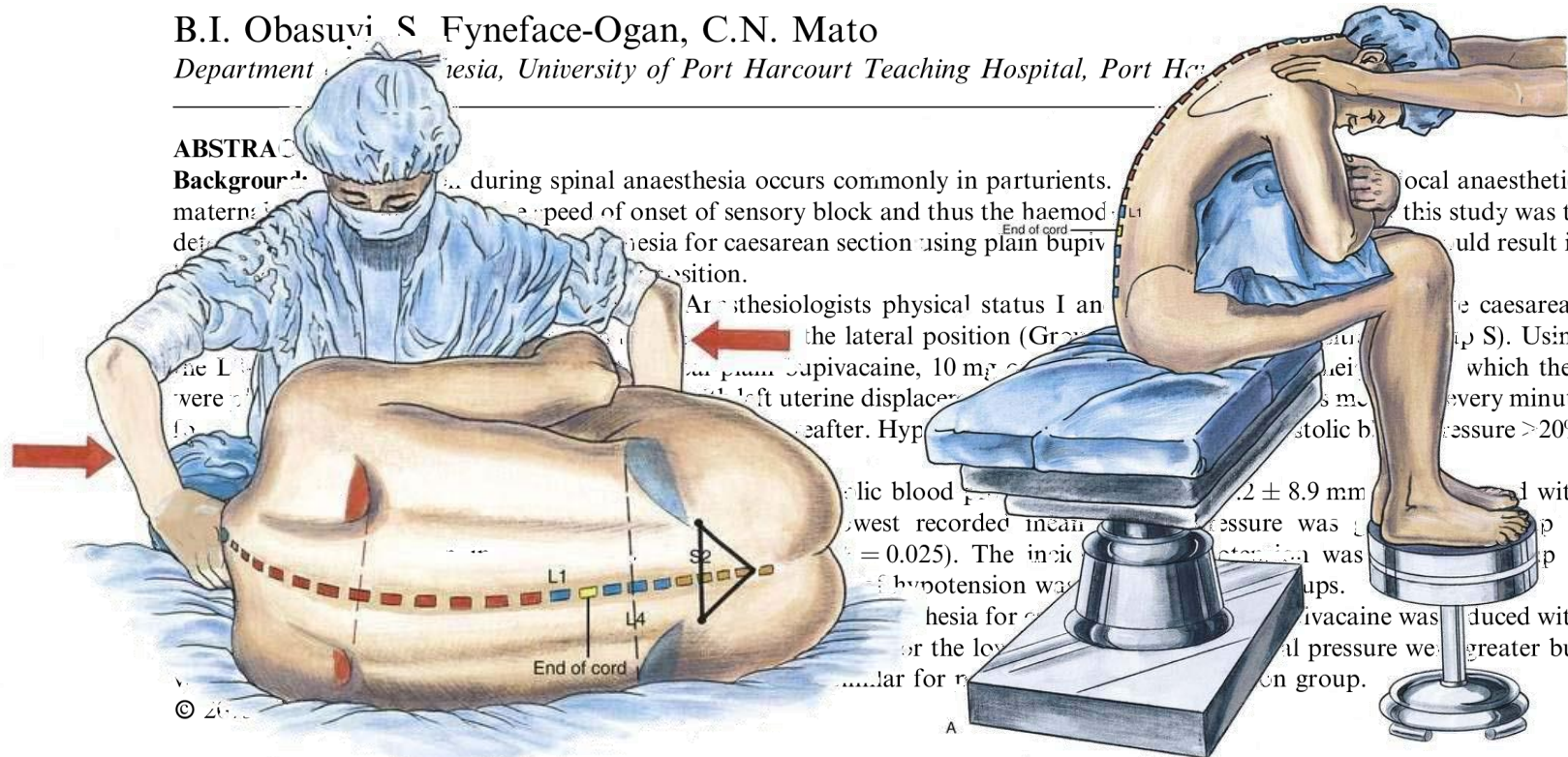


Table 3 Haemodynamic data, ephedrine use and intraoperative blood loss

	Group L (n = 50)	Group S (n = 50)	<i>P</i> value
Baseline SBP (mmHg)	122.4 ± 8.6	124.2 ± 9.9	0.3
Baseline MAP (mmHg)	93.0 ± 7.8	91.8 ± 8.9	0.4
Baseline heart rate (beats/min)	91.4 ± 8.5	92.3 ± 11.4	0.6
Incidence of hypotension	17 (34%)	28 (56%)	0.027
Time from IT injection to first hypotension (min)	11.8 ± 10.7	9.8 ± 8.2	0.5
Lowest SBP within 30 min of IT injection (mmHg)	99.2 ± 8.9	95.4 ± 12.3	0.08
Lowest MAP within 30 min of IT injection (mmHg)	72.9 ± 11.2	68.2 ± 9.6	0.02
Lowest heart rate within 30 min from IT injection (beats/min)	83 ± 11	79 ± 10	0.05
Incidence of ephedrine use	3 (6%)	5 (10%)	0.4
Total dose of ephedrine (mg)	5 ± 0	5 ± 0	1
SBP <90 mmHg	7 (14%)	14 (28%)	0.08
Blood loss (mL)	631 ± 171	697 ± 241	0.1

Data are mean ± SD or as number (%). SBP: systolic blood pressure; MAP: mean arterial pressure; IT: intrathecal.

Table 5 Incidence of complications

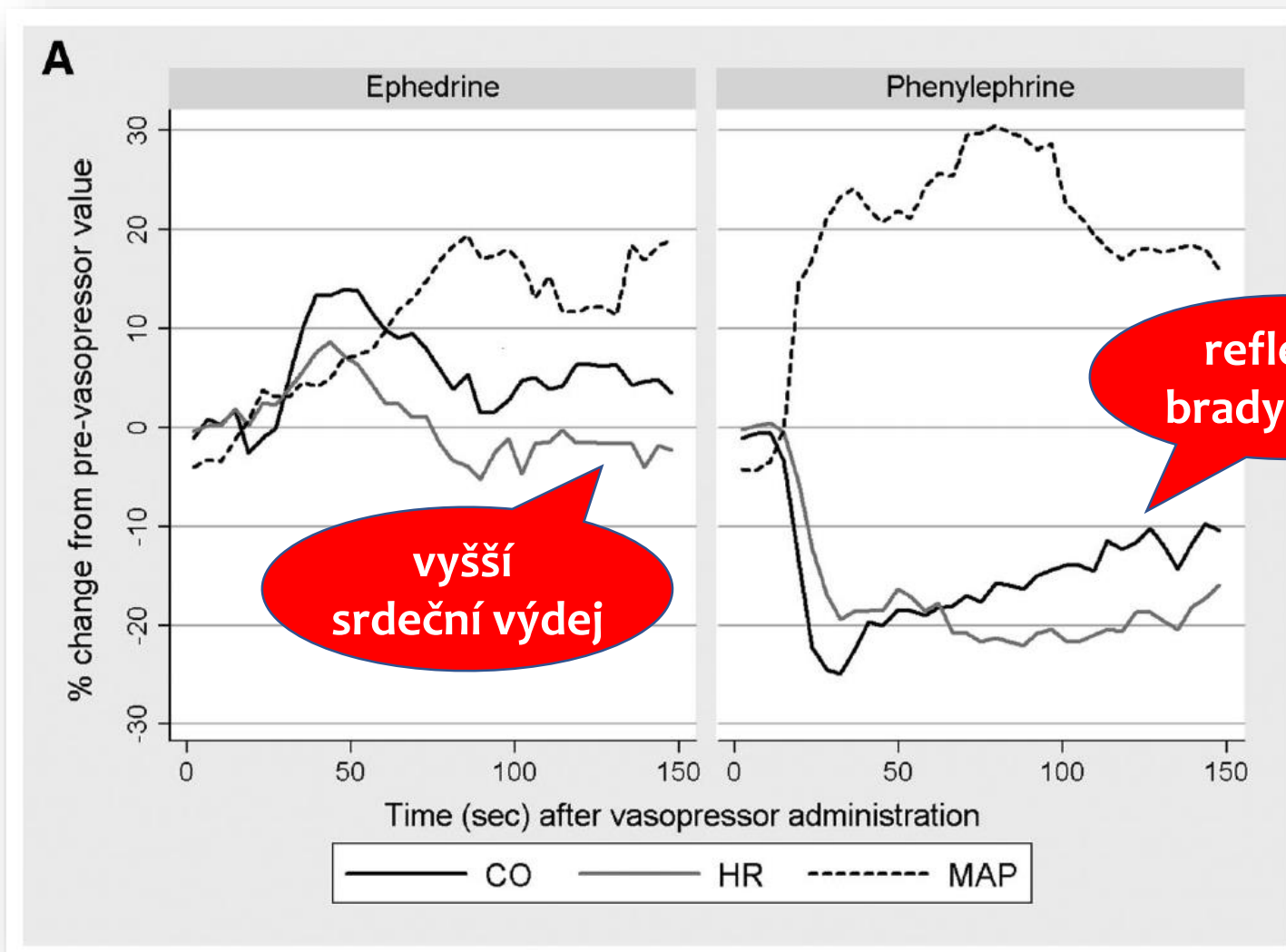
	Group L (n = 50)	Group S (n = 50)	<i>P</i> value
Nausea	2 (4%)	4 (8%)	0.4
Vomiting	0 (0%)	1 (2%)	0.3
Shivering	7 (14%)	11 (22%)	0.2
Dizziness/sleepiness	3 (6%)	5 (10%)	0.4
Respiratory distress	2 (4%)	7 (14%)	0.08

Data are number (%).

**SAB aplikovaný
v sedě má častější
výskyt hypotenze
než při aplikaci
na boku**

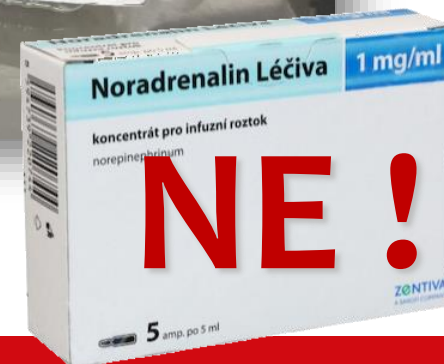
HYPOTENZE NA PORODNÍM SÁLE





Dyer RA et al. Anesthesiology. 2009 Oct;111(4):753-65

HYPOTENZE NA PORODNÍM SÁLE



NE! (zatím)

1. Odhadnout své možnosti !
2. Mít záložní plán (a nebát se ho použít)
3. Prevence hypotenze u neuroaxiální anestezie
4. Preoxygenace u celkové anestezie
5. Oxygenace je cílem...

S P A A