Volume 09, Issue 04, 2022

A comparative study of spinal anesthesia with hyperbaric ropivacaine plus dexmedetomidine and hyperbaric bupivacaine plus dexmedetomidine in lower abdominal and lower limb surgeries

¹Dr. T Anusha, ²Dr. Yalala Shamili, ³Dr. Kiran Kumar Suggala

¹Assistant Professor, Mamata Medical College, Khammam, Telangana, India
²PG Final Year, Mamata Medical College, Khammam, Telangana, India
³Professor and HOD, Mamata Medical College, Khammam, Telangana, India

Corresponding Author: Dr. Yalala Shamili

Abstract

Aims: The aim of the study is to compare the onset, duration of sensory and motor block, and hemodynamic changes between equipotent doses of 0.75% Ropivacaine H with dexmedetomidine & 0.5% Bupivacaine H with dexmedetomidine.

Materials and Methods: This was a comparative study conducted on ASA grade I and II patients, aged between 18 and 60 years, scheduled for elective lower abdominal and lower limb surgeries. Minimum of 60 patients, divided into two groups, Group R, and Group B (30 patients in each Group). Group R patients received hyperbaric ropivacaine 0.75% 3cc + 10 mcg dexmedetomidine and Group B patients received hyperbaric bupivacaine 0.5% 3cc + 10 mcg dexmedetomidine. Monitoring of vitals and observation for the block parameters were carried out and compared. Categorical data are compared using the Chi-Square test. Continuous variables are compared using a student t-test. P-value < 0.05 is considered significant.

Results: Time taken for sensory onset, maximum sensory block, and complete motor block was faster in group B. Time taken for sensory regression to S1 and duration of motor block was longer in group B. However, subjects in group B experienced lesser mean systolic, diastolic, and mean arterial pressures when compared to group R. Duration of sensory and motor block was increased with the addition of adjuvant Dexmedetomidine.

Conclusion: Spinal anesthesia with intrathecal 0.75% Ropivacaine H with dexmedetomidine has characteristically delayed onset, with a shorter duration of action on the sensory as well as motor nerve roots when compared to 0.5% Bupivacaine H with dexmedetomidine. Complications like hypotension and bradycardia were less in the Ropivacaine group. Compared to the bupivacaine group, better alternative for spinal anesthesia in the geriatric population. With a shorter recovery profile, Ropivacaine is a useful agent for Spinal Anesthesia for the intermediate duration of surgeries and for ambulatory surgeries.

Keywords: 0.75% Hyperbaric Ropivacaine, 0.5% hyperbaric bupivacaine, Dexmedetomidine, Spinal anesthesia

Introduction

Spinal anesthesia is widely used because of its fast onset and effective sensory and motor blockade for surgeries. Bupivacaine is available as a racemic mixture of its enantiomers, dextro-Bupivacaine, and levobupivacaine ^[1]. In the last few years, its pure S-enantiomer Ropivacaine has been introduced into clinical practice because of its lower toxic effects on the heart and central nervous system ^[2, 3-5]. Thus, ropivacaine, with its efficacy, lower propensity for motor block, and reduced potential for CNS toxicity and cardiotoxicity, appears to be an important option for regional anesthesia and management of postoperative pain ^[6].

Alpha-2 adrenergic receptor agonists such as clonidine and dexmedetomidine have been the focus of interest for their sedative, analgesic, and perioperative sympatholytic properties. Dexmedetomidine $\alpha 2:\alpha 1$ selectivity is eight times higher than that of Clonidine ^[7]. It prolongs the duration of both sensory and motor blockade induced by local anesthetics thereby prolonging the duration of analgesia.

Objective

The aim of the study is to compare the onset, duration of motor and sensory block, and hemodynamic changes between equipotent doses of 0.75% Ropivacaine H with Dexmedetomidine and 0.5% Bupivacaine H with Dexmedetomidine.

Inclusion criteria

Patient willing for study and who has given informed and written consent, belonging to ASA classes 1 & 2 between the age of 18 and 60 years with no local infection and neurological deficit.

Exclusion criteria

Patient's refusal and belonging to ASA grade III and IV with baseline heart rate < 60 bpm and baseline BP < 100/50 mmHg.Patients who are allergic to protocol drugs and on anticoagulants. Patients with cardiorespiratory, hepatic, and renal problems and those who are undergoing emergency surgeries

Materials and Methods

This was a comparative study conducted on ASA grade I and II patients, aged between 18 and 60 years, scheduled for elective lower abdominal and lower limb surgeries in Mamata General Hospital for a period of one year after obtaining approval from the institutional ethics committee. Minimum of 60 patients, divided into two groups, Group R, and Group B (30 patients in each Group). Group R patients received hyperbaric ropivacaine 0.75% 3cc + 10 mcg dexmedetomidine and Group B patients received hyperbaric bupivacaine 0.5% 3cc + 10 mcg dexmedetomidine. Patients in the study underwent thorough preoperative evaluation which included history, Hb, PCV, BT, CT, RFT, Blood sugar, ECG, CXR, Platelet count, Blood grouping, and cross-matching done. Hypotension, tachycardia, and bradycardia were noted. Assessment of sensory blockade by pinprick and motor blockade by Modified Bromage scale was done.

Assessment of sensory block

- 1. Time of onset of sensory block (Time taken to attain T10 dermatome).
- 2. Time taken for maximum Sensory block. (Time taken to attain T6 dermatome).
- 3. Sensory block duration. (Time taken to regress up to S1 dermatome)

Assessment of motor block

- 1. Time taken for a complete motor block. (Time taken to achieve Bromage score 3).
- 2. Motor block duration. (Time of regression to Bromage score 0).
- 3. Grade 3-unable to move feet or knees (complete).
- 4. Grade 2-able to move feet only (almost complete).
- 5. Grade 1-just able to move knees (partial).
- 6. Grade 0-full flexion of feet and knees (none).

Results

Demographics

The demographic characteristics of the two groups did not differ significantly. Table 1 displays the results.

Parameters	Group B (n=30)	Group R (n=30)	P-value
Age	32.67±8.12	35.12±7.45	0.76
Gender (M: F)	20:10	22:8	0.56
Weight (Kgs)	60.87±8.12	59.76±12.21	0.12
Height	$158.43{\pm}10.45$	160.24±7.12	0.25
ASA (I/II)	13/17	14/16	0.54

Table 1: Demographic characteristics of the study participants

Time taken for onset of sensory block (minutes)

Time taken for onset of sensory block was observed to be longer in Group R compared to group B (4.32 ± 0.48 vs 2.18 ± 0.19 mins; p=0.001). Table 2 displays the results.

Table 2: Time tak	en for onset of s	ensory block betweer	the two groups
-------------------	-------------------	----------------------	----------------

Parameters	Group B (n=30)	Group R (n=30)	P value
Time taken for onset of sensory block (mins)	2.18±0.19	4.32±0.48	0.001

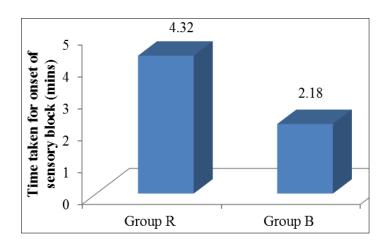


Fig 1: Time taken for onset of sensory block between the two groups 3858

Time taken to attain maximum sensory blockade

The time taken to attain maximum sensory blockade was observed to be significantly longer in group R as compared to group B (15.25 ± 4.12 vs 7.56 ± 2.48 mins; p=0.006). Table 3 displays the results.

Table 3: Time taken to attain maximum sensory blockade between the two groups

Parameters Group	р в (n=эv)(Group R (n=30)	P value
The time taken to attain maximum sensory blockade (mins) 7.5	6±2.48	15.25±4.12	0.006

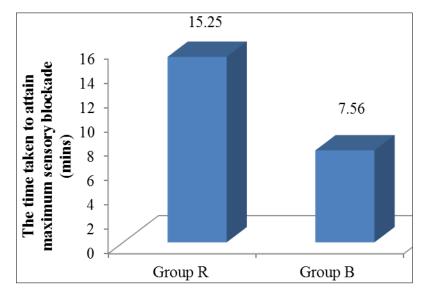


Fig 2: Time taken to attain maximum sensory blockade between the two groups

Time taken for regression of sensory block to S1

The time taken for regression of sensory block to S1 was observed to be significantly longer in group B as compared to group R (320.56 ± 38.34 vs 218.33 ± 30.36 min; p=0.006). Table 4 displays the results.

Parameters	Group B (n=30)	Group R (n=30)	P value
The time taken for regression of sensory block to S1 (mins)	320.56±38.34	218.33±30.36	0.002
The time taken for regression of sensory block to 31 (finits) up to the taken for regression of sensory block to 31 (finits) 213.33 100 - 213.33 213.33 Group R	320.30±38.34	218.33±30.30	0.002

Table 4: Time taken for regression of Sensory block to S1 between the two groups

Fig 3: Time taken for Sensory regression to S1 between the groups

Time taken to attain complete motor block

The time taken to attain complete motor block was observed to be significantly longer in group R compared to group B (6.21 ± 1.65 vs 3.65 ± 0.76 mins; p=0.000). Table 5 displays the results.

Table 5: Time taken to attain complete motor block between the two groups.

Parameters	Group B (n=30)	Group R (n=30)	P value
The time taken to attain complete motor block (min)	3.65±0.76	6.21±1.65	0.000

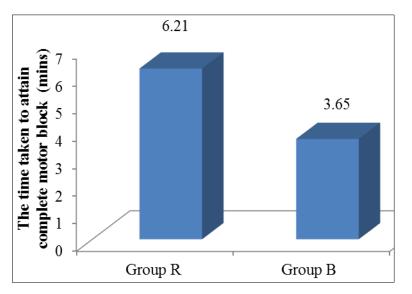


Fig 4: Time taken to attain complete motor block

Motor block duration

Motor block duration was observed to be significantly shorter in group R as compared to group B and it was found to be significant (189.45 \pm 24.65 vs 312 \pm 35.25 mins; p= 0.006). Table 6 displays the results.

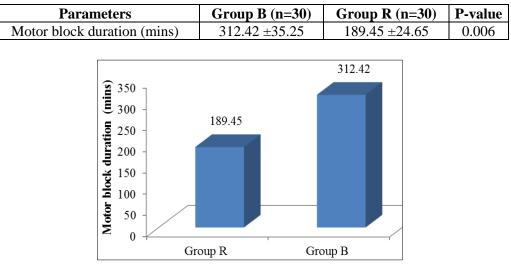


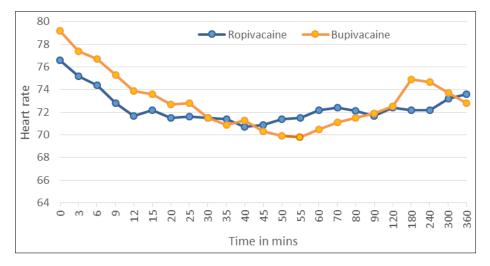
Table 6: Motor block duration between the two groups

Fig 5: Motor block duration between the two groups

T:	Group Bupivacaine (N=30)		Group Ropivac	Group Ropivacaine (N=30)		
Time	Mean HR (/min)	SD	Mean HR (/min)	SD	P value	
0 min	79.2	7.8	76.6	8.8	0.329	
3 min	77.4	7.8	75.2	8.9	0.421	
6 min	76.7	7.8	74.4	8.6	0.381	
9 min	75.3	7.4	72.8	9.0	0.343	
12 min	73.9	6.7	71.7	9.1	0.390	
15 min	73.6	5.4	72.2	8.6	0.527	
20 min	72.7	5.8	71.5	9.0	0.617	
25 min	72.8	4.3	71.6	9.0	0.593	
30 min	71.5	5.5	71.5	8.1	0.982	
35 min	70.9	5.6	71.4	7.5	0.831	
40 min	71.3	5.6	70.7	7.2	0.770	
45 min	70.3	4.9	70.9	6.9	0.753	
50 min	69.9	5.2	71.4	7.1	0.436	
55 min	69.8	6.0	71.5	6.9	0.399	
60 min	70.5	4.9	72.2	6.6	0.358	
70 min	71.1	5.6	72.4	6.4	0.482	
80 min	71.5	5.9	72.1	6.3	0.758	
90 min	71.9	6.0	71.7	6.2	0.918	
120 min	72.5	6.1	72.4	5.9	0.958	
180 min	74.9	7.5	72.2	6.0	0.225	
240 min	74.7	7.7	72.2	6.5	0.274	
300 min	73.7	7.5	73.2	5.8	0.833	
360 min	72.8	5.8	73.6	7.5	0.709	

Table 7: Comparison of mean heart rate between the two study groups at various points of time(n=60)

Two groups showed no significant difference in the mean heart rate throughout the intraoperative period.



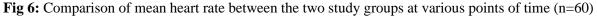


Table 8: Comparison of mean systolic blood pressure (SBP) between the two study groups at various
points of time (n=60)

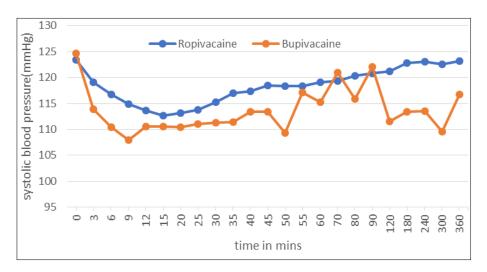
Time	Group Bupivacaine (N=30)		Group Ropivacaine (N=30)		P value
Time	Mean SBP (mmHg)	SD	Mean SBP (mmHg)	SD	r value
0 min	124.7	10.1	123.4	11.0	0.710
3 min	113.9	8.2	119.1	9.5	0.072

ISSN 2515-8260

Volume 09, Issue 04, 2022

6 min	110.4	6.9	116.8	10.4	0.027
9 min	107.9	6.1	114.9	9.3	0.008
12 min	110.5	6.0	113.6	7.7	0.164
15 min	110.5	4.5	112.6	7.7	0.299
20 min	110.4	4.1	113.2	7.5	0.151
25 min	111.0	3.9	113.8	6.0	0.088
30 min	111.3	5.7	115.2	7.6	0.074
35 min	111.4	5.6	117.0	6.9	0.007
40 min	113.4	5.3	117.3	6.6	0.047
45 min	113.4	4.5	118.5	6.9	0.009
50 min	109.3	21.9	118.4	7.2	0.085
55 min	117.1	6.3	118.4	6.7	0.530
60 min	115.3	25.4	119.1	6.8	0.523
70 min	120.9	8.0	119.4	7.1	0.536
80 min	115.9	24.9	120.3	6.8	0.445
90 min	122.1	6.2	120.8	7.3	0.546
120 min	111.5	5.1	121.2	7.5	< 0.001
180 min	113.4	4.6	122.8	6.9	< 0.001
240 min	113.5	4.2	123.0	7.4	< 0.001
300 min	109.6	22.1	122.5	7.8	0.018
360 min	116.7	6.2	123.2	7.9	0.006

Two groups showed significant differences in mean Systolic blood pressure at several points of time. Lesser systolic blood pressures were observed in subjects of the Bupivacaine group than in subjects of the Ropivacaine group.



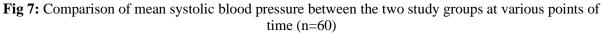


Table 9: Comparison of mean diastolic blood pressure (DBP) between the two study groups at
various points of time (n=60)

Time	Group Ropivacaine (N=30)	Group Bupivacaine (N=30)		P value
Ime	Mean DBP (mm Hg)	SD	Mean DBP (mm Hg)	SD	r value
0 min	82.6	7.6	73.6	6.7	0.000
3 min	72.8	17.1	69.7	5.8	0.446
6 min	75.2	6.4	67.8	5.9	0.001
9 min	74.9	7.0	67.9	5.8	0.001
12 min	75.9	7.1	66.6	4.9	0.000
15 min	74.3	6.2	66.1	4.1	0.000

European Journal of Molecular & Clinical Medicine

ISSN 2515-8260

Volume 09, Issue 04, 2022

20 min	73.9	7.5	67.8	4.9	0.004
25 min	73.2	7.5	67.9	4.1	0.009
30 min	70.7	7.8	68.4	4.2	0.253
35 min	72.4	7.0	69.5	4.4	0.126
40 min	72.8	6.5	69.7	4.7	0.092
45 min	72.4	7.0	70.8	4.2	0.390
50 min	72.4	8.0	70.7	4.1	0.403
55 min	71.6	8.3	71.8	4.0	0.923
60 min	74.3	7.4	72.2	4.3	0.280
70 min	73.9	6.8	72.6	4.5	0.478
80 min	74.1	7.5	71.7	4.3	0.221
90 min	75.4	7.9	71.4	4.1	0.054
120 min	75.9	7.2	72.6	4.6	0.091
180 min	76.2	6.8	73.2	5.4	0.130
240 min	76.6	8.0	73.4	5.5	0.149
300 min	76.6	7.9	73.9	5.2	0.211
360 min	74.6	8.5	73.4	4.9	0.587

Two groups showed significant differences in mean Diastolic blood pressure at the initial period of the intraoperative period. Lesser diastolic blood pressures were observed in subjects of the Bupivacaine group than in subjects of the Ropivacaine group.

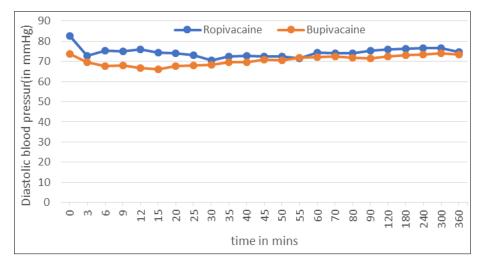


Fig 8: Comparison of mean diastolic blood pressure between the two study groups at various points of time (n=60)

Table 10: Comparison of mean arterial blood pressure (MAP) between the two study groups at
various points of time (n=60)

Time	Group Ropivacaine (N=30)		Group Bupivacaine (N=30)		P value
Time	Mean MAP (mm Hg)	SD	Mean MAP (mm Hg)	SD	r value
0 min	97.9	6.1	90.3	7.9	0.002
3 min	89.0	7.2	86.1	6.6	0.193
6 min	86.9	5.8	82.6	4.6	0.014
9 min	85.9	5.9	82.3	4.6	0.035
12 min	82.4	18.1	82.3	3.5	0.971
15 min	86.1	4.7	81.7	4.8	0.007
20 min	86.3	5.5	82.8	4.8	0.038
25 min	85.7	5.4	83.4	4.5	0.158
30 min	84.4	6.0	83.9	4.6	0.769
35 min	85.4	5.0	85.3	4.2	0.919

40 min	85.7	4.0	85.5	4.7	0.913
45 min	86.3	4.5	86.1	4.4	0.888
50 min	86.5	5.6	86.2	4.3	0.850
55 min	86.7	5.9	87.4	4.4	0.672
60 min	89.7	5.1	87.7	4.4	0.194
70 min	85.8	18.8	88.0	4.2	0.604
80 min	89.6	5.1	88.2	4.5	0.359
90 min	90.0	5.3	87.9	4.2	0.174
120 min	91.0	5.1	89.0	4.5	0.207
180 min	84.8	4.6	89.1	4.7	0.005
240 min	84.4	4.6	89.6	4.8	0.001
300 min	84.9	3.7	89.9	5.4	0.002
360 min	85.9	3.5	90.0	4.7	0.003

Two groups showed significant differences in mean arterial pressure both in the initial period and towards the end of the period. Lesser mean arterial pressures were observed in subjects of the Bupivacaine group than in subjects of the Ropivacaine group.

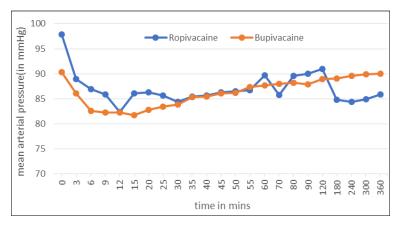


Fig 9: Comparison of mean arterial blood pressure between the two study groups at various points of time (n=60)

Discussion

Ropivacaine is a pure enantiomer and long-acting amide local anesthetic. Its lipid solubility is low making it less likely to penetrate large myelinated motor fibers which relate to the blocking of nerve fibers involved in the transmission of pain (A δ and C fibers) to a greater degree, compared to that of controlling motor functions (A β fibers). This feature is useful when the motor blockade is undesirable ^[7].

Sanchez *et al.* in 2009 compared the effects of intrathecal isobaric Ropivacaine (IR) versus isobaric Bupivacaine (IB) in a dose ratio of 3:2 in non-ambulatory urologic and orthopedic surgery. They concluded that the motor blockade was shorter in the IR Group (226.4 \pm 22.3 min) when compared to the IB Group (266.5+/- 29.5) p < 0.001 ^[8]. Similarly, we noted that motor block duration was shorter in the Ropivacaine group when compared to the Bupivacaine group.

Spinal adjuvants decrease the dose of local anesthetics. Dexmedetomidine provides stable hemodynamics, minimal side effects, and good quality intraoperative and postoperative analgesia ^[9]. Whiteside and others observed that the mean time of onset of motor block was 10 min and 15 min and the total duration of motor block was around 180 min and 90 min with a similar dose of hyperbaric Bupivacaine (15mg) and Ropivacaine (15mg) respectively ^[10]. We found the total duration of motor blocks in group B and group R was around 312 ± 35.25 and 189.45 ± 24.65 respectively. Duration of motor block increased with the addition of adjuvant Dexmedetomidine.

We noted that the ropivacaine group had good sensory block than bupivacaine, a shorter time to first micturition, and a favorable recovery profile of motor blockade. These features are beneficial for ambulatory surgery.

Conclusion

Spinal anesthesia for lower abdominal and lower limb surgeries with intrathecal 0.75% Ropivacaine H with dexmedetomidine has delayed onset, with a shorter duration of action on both motor and sensory nerve roots when compared to 0.5% Bupivacaine H with dexmedetomidine. The Ropivacaine group experienced fewer complications like hypotension and bradycardia. Compared to the bupivacaine group, better alternative for spinal anesthesia in the geriatric population. With a shorter recovery profile, Ropivacaine is a useful agent for Spinal Anesthesia for the intermediate duration of surgeries and for ambulatory surgeries.

Source of funding: None.

Conflict of interest: None.

References

- 1. Vanna O, Chumsang L, Thongmee S, Levo Bupivacaine. Bupivacaine in spinal anesthesia for transurethral endoscopic surgery, J. MED. ASSOC. THA. 2006;89(8):1133-9.
- 2. Foster RH, Markham A, Levo Bupivacaine: A review of its pharmacology and local Drugs. 2000;59(3):551-79.
- 3. Markham A, Faulds D, Ropivacaine. A review of its pharmacology and therapeutic use in regional anesthesia DRUGS. 1996;52(3):429-49.
- 4. McClellan KJ, Faulds D, Ropivacaine: An update of its use in regional anesthesia, DRUGS. 2000;60(5):1065-93.
- 5. Milligan KR. Recent advances in local anesthetics for spinal anesthesia, EUR. J. Anaesthesiol. 2004;21:837-847.
- 6. Van Kleef JW, Veering BT, Burm AG. Spinal anesthesia with ropivacaine: A doubleblind study on the efficacy and safety of 0.5% and 0.75% solutions in patients undergoing minor lower limb surgery. Anesth Analg. 1994;78:1125-30.
- Hinojosa-Sánchez O, Alamilla-Beltrán I, Han-Alonso R, *et al.* Subarachnoid blockade with Ropivacaine versus Bupivacaine in urologic and orthopedic surgery. Rev Med Inst Mex Seguro Soc. 2009;47(5):539-44. [PubMed]
- 8. Kuthiala Gaurav, Chaudhary Geeta. Ropivacaine: A review of its pharmacology and clinical use. Indian journal of anaesthesia. 2011;55:104-10. 10.4103/0019-5049.79875.
- Gupta R, Verma R, Bogra J, Kohli M, Raman R, Kushwaha JK. A comparative study of intrathecal dexmedetomidine and fentanyl as adjuvants to Bupivacaine. Journal of Anaesthesiology, Clinical Pharmacology. 2011;27:339-343. DOI: 10.4103/0970-9185.83678.
- 10. Whiteside JB, Burke D, Wildsmith JA. Comparison of ropivacaine 0.5% (in glucose 5%) with bupivacaine 0.5% (in glucose 8%) for spinal anaesthesia for elective surgery. Br J Anaesth. 2003;90:304-8.
- 11. Fettes PD, Hocking G, Peterson MK, Luck JF, Wildsmith JA. Comparison of plain and hyperbaric solutions of ropivacaine for spinal anesthesia. Br J Anaesth. 2005;94:107-11.
- 12. Kallio H, Snäll EV, Tuomas CA, Rosenberg PH. Comparison of hyperbaric and plain ropivacaine 15 mg in spinal anesthesia for lower limb surgery. Br J Anaesth. 2004;93:664-9.
- 13. Hocking G, Wildsmith JA. Intrathecal drug spread. Br J Anaesth. 2004;93:568-78.