



SAOA 2023, Spring Meeting
April 1st 2023, Lucerne

Use of neuraxial dexmedetomidine in OB anesthesia

Rita Al khoury M.D.

OB-GYN anesthesia division

rial@hcuge.ch

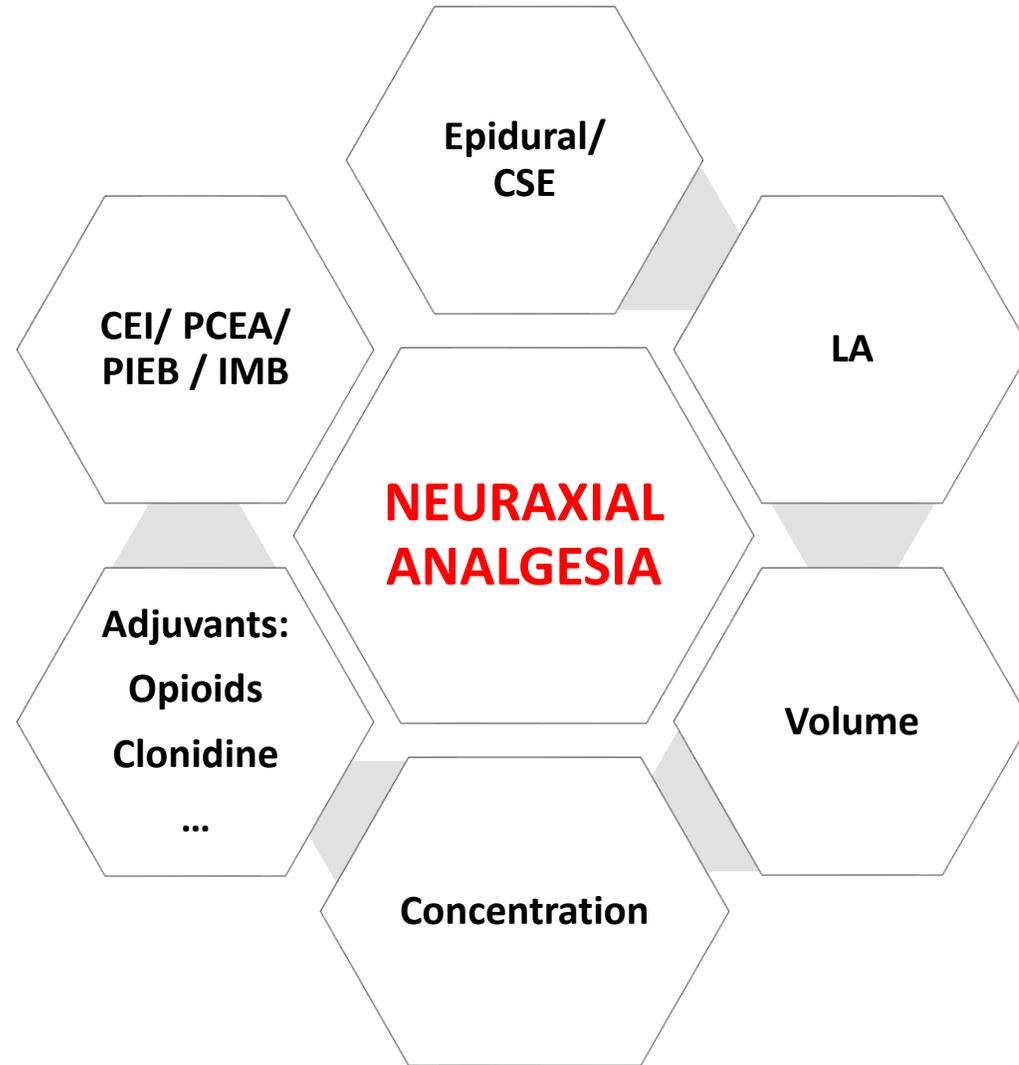




Optimal analgesia
Quality of block
Minimal top ups
↓ LA concentration



Pruritus
Nausea
Vomiting
Respiratory depression



Motor block
Maternal hypotension
Urinary retention
↑ second stage of labor
↑ instrumental delivery

OPIOID FREE ANALGESIA

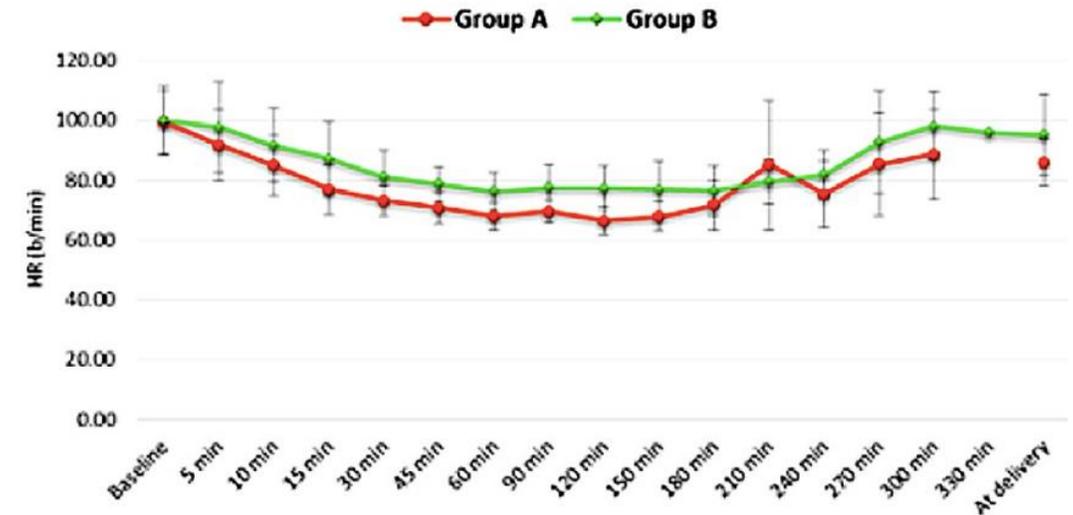
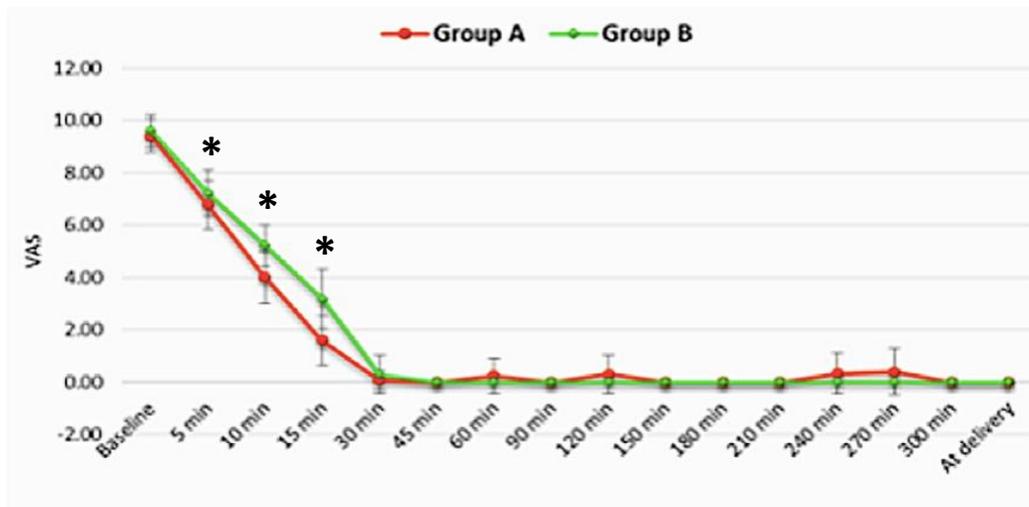
Adjuvants: α 2 agonists

	CLONIDINE	DEXMEDETOMIDINE	
SELECTIVITY α 2/1	220/1	1600/1	8 X
T $\frac{1}{2}$ β	8h	2h	
Route administration	Oral, IV, patch, epidural	IV	
Primary indication	Antihypertensive	Sedative	
Other indication	Analgesic	Analgesic	
Maternal/<u>fetal index</u>	0.85	0.68 - 0.77	<u>Liposolubility ++</u>

❖ Thus Dex is an interesting drug to study in labor pain

A Comparison Between Dexmedetomidine and Clonidine as Adjuvants to Levobupivacaine in Labour Analgesia

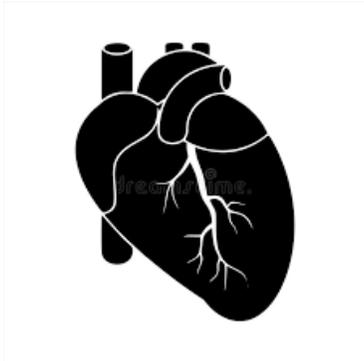
10 mL bolus: 0.125% levobupi + 0.5 ug/kg dex (A) vs 1 µg/kg clo (B) , PCA 10 ml/h, rescue 5 ml / 10 min



OUTCOMES	DEX	CLO
ONSET min	11 ± 1.3	15 ± 2
MAXIMUM min	14.6 ± 1.2	18 ± 2

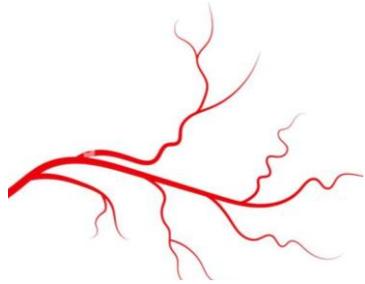
- ❖ **Faster onset with DEX**
- ❖ **Fall in HR significantly greater**
- ❖ **NO difference in obstetric, neonatal outcomes**
- ❖ **NO difference in adverse reactions**

DEXMEDETOMIDINE



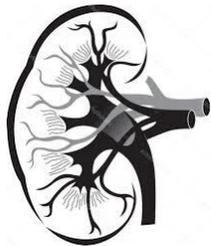
α 2a
vagomimetic action

α 2b
Decrease tachycardia
Blocks cardioaccelerator nerve

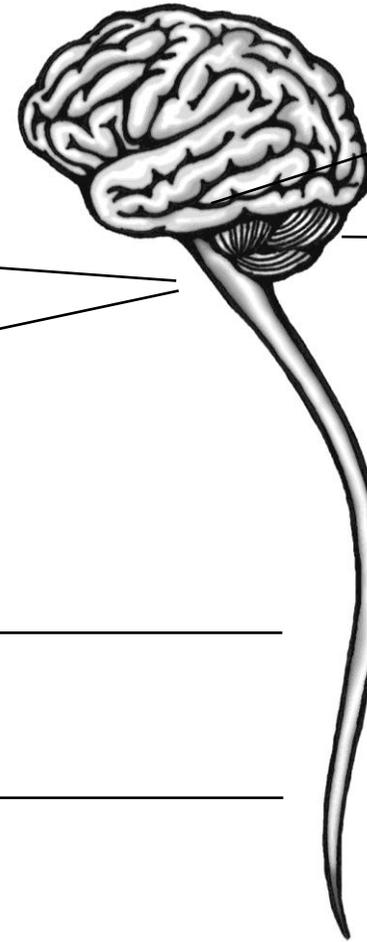


α 2b vasoconstriction
antishivering

α 2a vasodilation



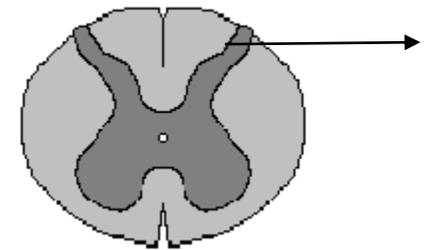
α 2b diuresis



α 2a sedation

α 2c anxiolysis

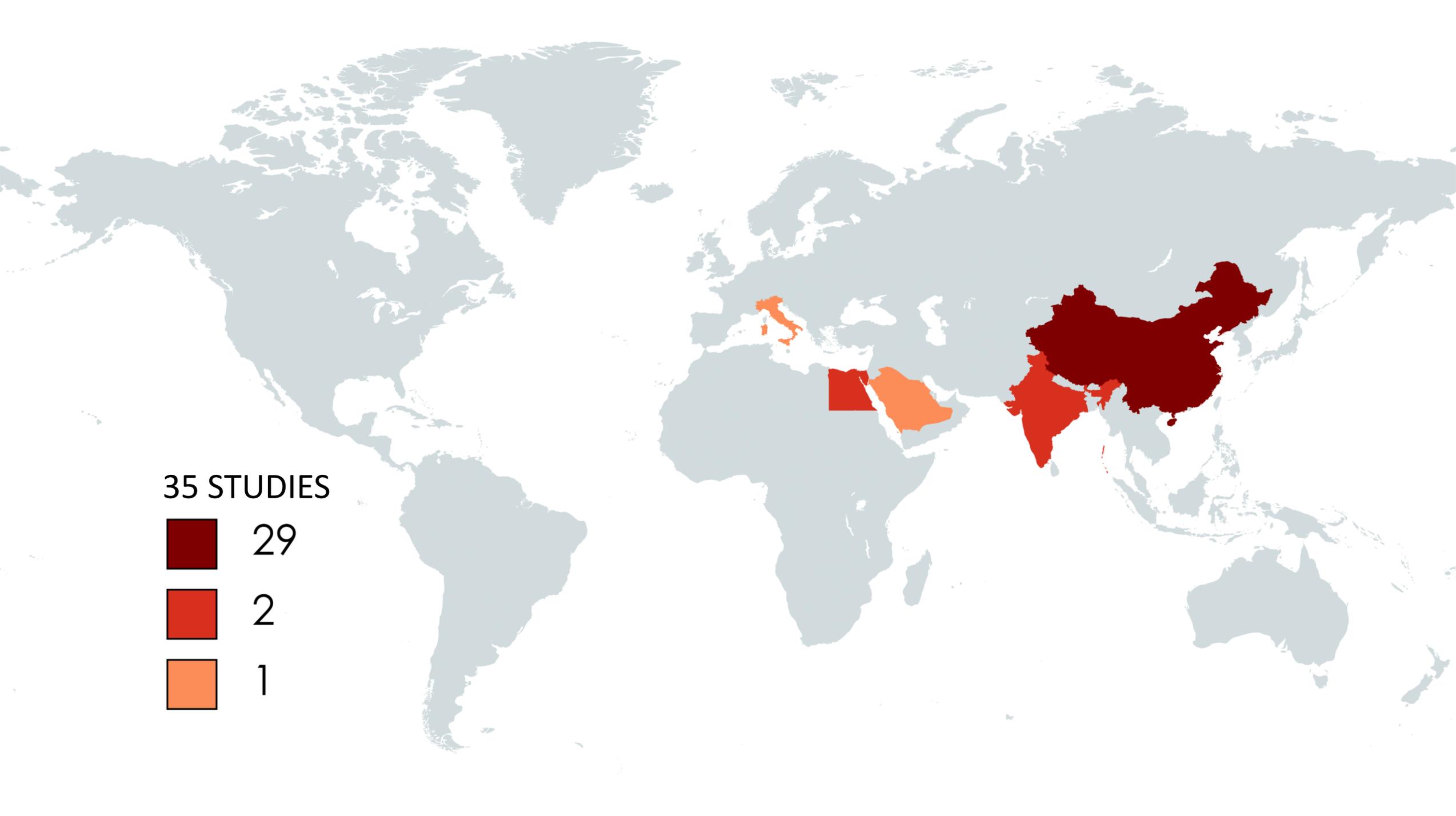
- \downarrow NE release
- Sympatolysis



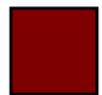
α 2a analgesia

- α 2-independent mechanism
- limit firing and rate of substance P release
- Direct inhibition of the signaling pathway

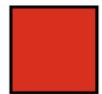
ANALGESIC OUTCOMES



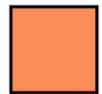
35 STUDIES



29



2



1

INTENSITY

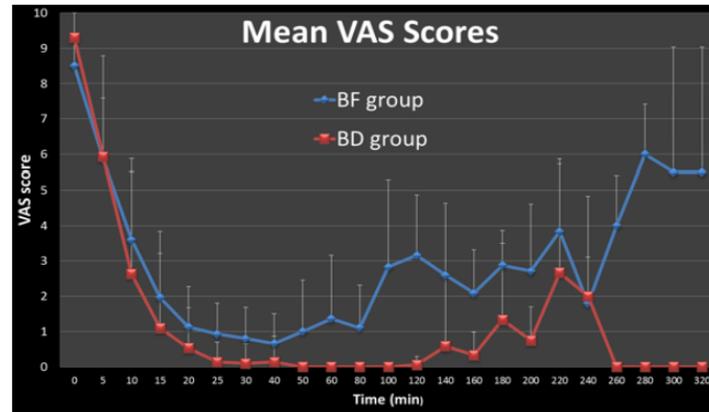
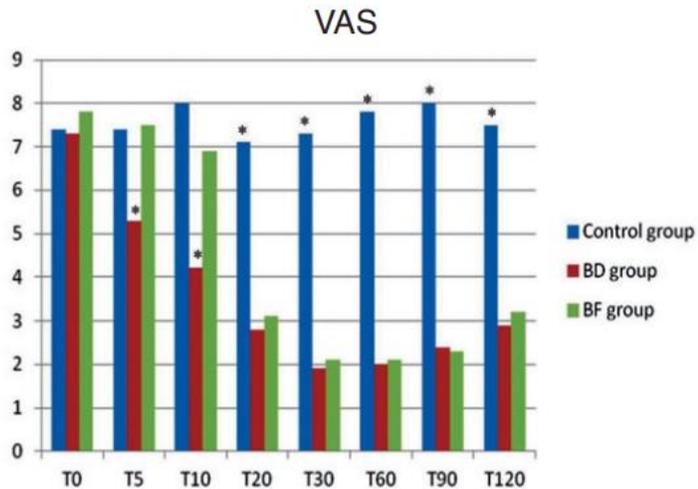
BUPI + FENTA vs DEX

❖ BETTER QUALITY OF PAIN RELIEF WITH DEX

17 ml: 0.25% B ± 1µg/kg D or F

15 ml: 0.0625 % B + 1.5 µg/ml D or 2 µg/ml F
VAS > 4 bolus 5ml

14 ml: 0.25% B + 1µg/kg D or F
Bolus if pain



Variables	Group D (n=85)	Group F (n=85)	P-value
Pain verbal scale			
0	58	45	0.041
1	13	4	0.021
2	2	8	0.050
3	9	12	0.484
4	3	16	0.002

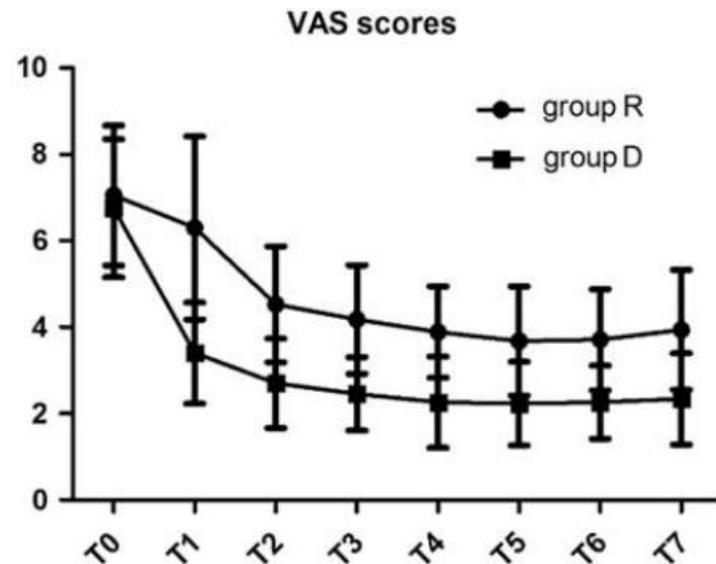
INTENSITY

ROPI ± DEX

ROPI + SUF vs DEX

❖ LOWER VAS WITH DEX WHEN CERVICAL DILATION > 3

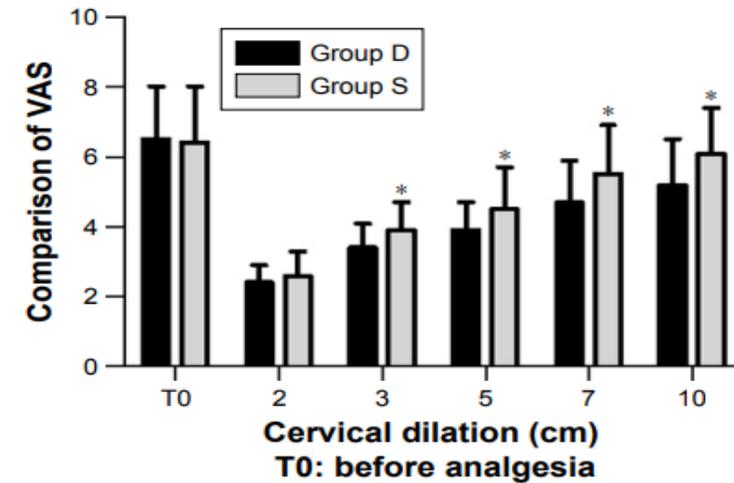
0.125% ropivacaine ± dex (0.5 mg/kg as bolus only)
n = 40



Yang Zhao et al. Clin J Pain Volume 33, Number 4, April 2017

10 ml bolus: 0.1% ropi + suf vs dex (0.5µg/ml)

PCEA: 6ml/h, 6ml /20min ; n= 70



Zhang et al. Drug Design, Development and Therapy 2019:13 1171–1175

OTHER VARIABLES

BUPI + FENTA vs DEX

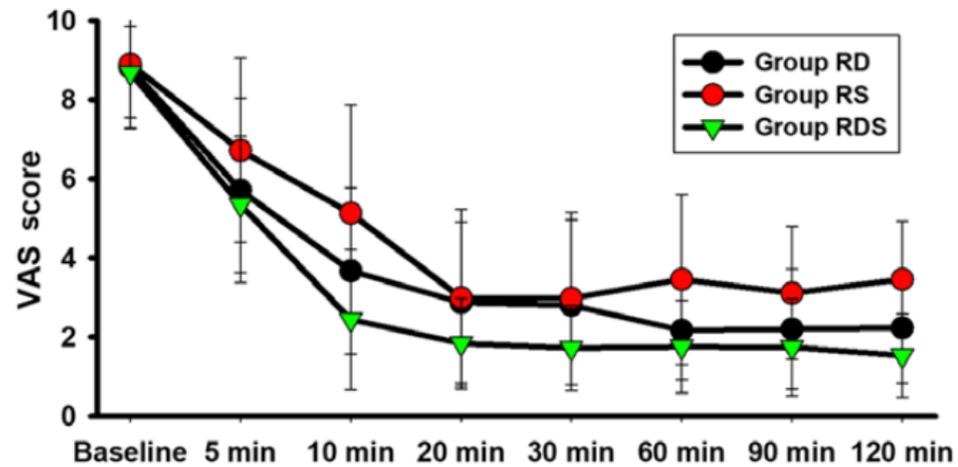
❖ BETTER ONSET, DURATION, AND ANESTHETIC REQUIREMENTS

BUPI 0.25%	FENTA 1µg/kg	DEX 1µg/kg	Statistic significance
ONSET min	7 – 13	4 - 9	+
DURATION min	110 - 118	125 - 185	+
SECOND DOSE	1/3	1/7	+

BUPI 0.0625% + 2 F or 1.5 D			
Group	BF	BD	p value
Onset (mins.)	6.00±2.034	5.33±1.269	0.133
Duration of analgesia (mins.)	85.33±22.512	131.83±45.760	<0.0001
Number of top ups	1.80±1.518	0.17±0.461	<0.0001
Episiotomy/suturing without local anaesthetic (number)	1	28	<0.0001

Combination of sufentanil, dexmedetomidine and ropivacaine to improve epidural labor analgesia effect: A randomized controlled trial

❖ RD, RDS > RS



Bolus 10 ml + PCEA
RD: 0.1% + 0.5 µg/ml
RS: 0.1% + 0.5 µg/ml
RDS: 0.1% + 0.25 µg/ml + 0.25 µg/ml

Variable	Group RS (n=35)	Group RD (n=36)	Group RDS (n=36)	P-value
Onset time (min)	15.50±2.67	12.97±3.13	9.68±1.26 ^{a,c}	0.037
Total volume of anesthetic solution (ml)	65.44±5.64	42.65±6.44	50.34±6.56 ^a	0.043
Bolus frequency	2.80±0.92	0.10±0.31 ^a	0.80±0.78 ^a	0.026

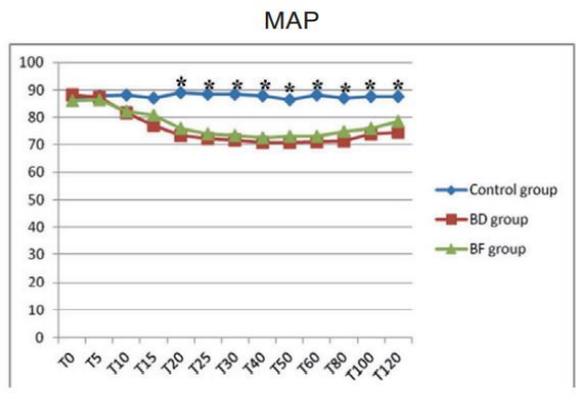
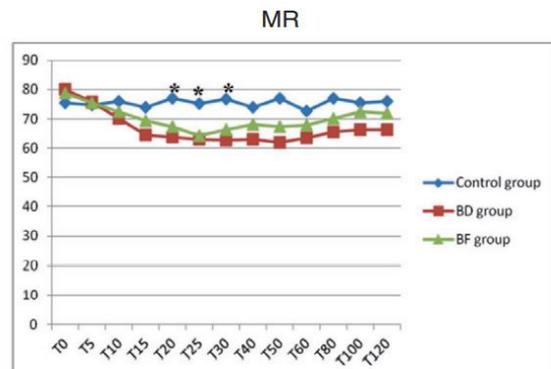
HEMODYNAMIC MEASUREMENTS

BLOOD PRESSURE ; HR

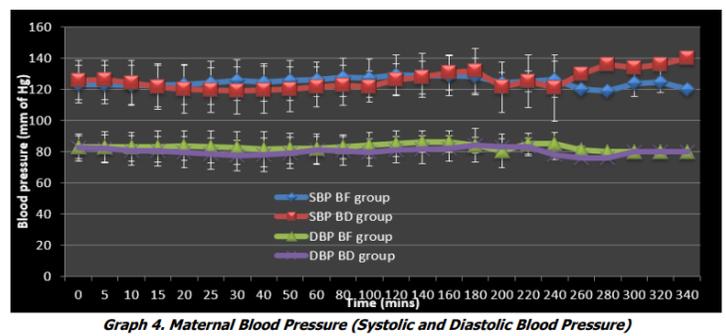
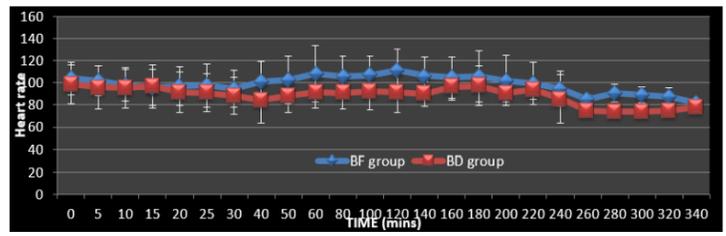
BUPI + FENTA vs DEX

- ❖ RISK OF MATERNAL HYPOTENSION OR BRADYCARDIA ?
- ❖ NO TREATMENT NEEDED

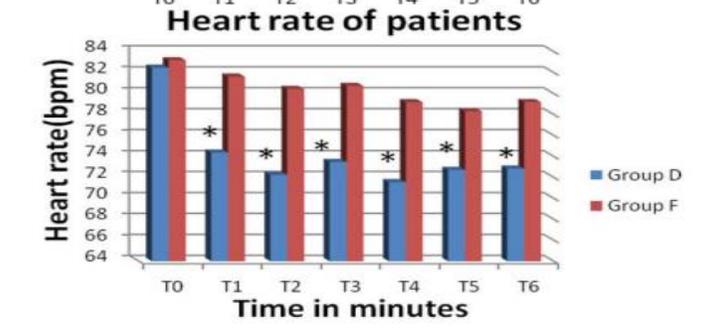
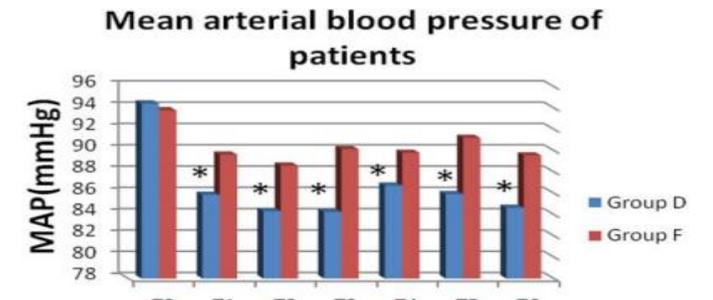
1 bolus 17 ml: 0.25% B + 1µg/kg D or F



15 ml: 0.0625 % B + 1.5 µg/ml D or 2 µg/ml F
VAS > 4 bolus 5ml



14 ml: 0.25% B + 1µg/kg D or F
Bolus if pain



BLOOD PRESSURE ; HR

ROPI + SUF vs DEX

❖ **NO SIGNIFICANT DIFFERENCE**

Event	Group RS (n=35) (%)	Group RD (n=36) (%)	Group RDS (n=36) (%)	P-value
Hypotension	0 (0.0)	1 (2.8)	0 (0.0)	1.000
Bradycardia	0 (0.0)	1 (2.8)	0 (0.0)	1.000

Li et al Experimental and therapeutic medicine 20: 454-460, 2020

	Group D (n=36)	Group S (n=34)	P-value
Hypotension	0	0	
Maternal bradycardia	0	0	

Zhang et al. Drug Design, Development and Therapy 2019:13 1171-1175

OBSTETRIC AND NEONATAL OUTCOMES

BUPI + FENTA vs DEX

ROPI + SUF vs DEX

- ↓ Duration of 1st stage of labor (30 - 100min, statistically significant with ropi)*

❖ No clinical nor statistical significant difference

- Duration of 2nd stage of labor
- Instrumental delivery
- C- section

- Fetal heart rate
- APGAR 1 / 5 min
- Umbilical cord pH
- Uretro placental blood flow

ADVERSE REACTIONS

GENERAL

BUPI + FENTA vs DEX

ROPI + SUF vs DEX

- ❖ **Significantly less reactions with BD compared BF**
- ❖ **Dex is effective in reducing pruritus when associated with suf**

%	BF	BD	RS	RD	RDS
Nausea	12-20	4-5			
Vomiting	7	2			
Pruritus	10-14	0-2	5	0	0
Respiratory depression	0-5	0			
shivering	12	4			
Dry mouth	4	12			

M. F. Selim et al. Journal of Prenatal Medicine 2012; 6 (3): 47-54
Soliman et al. Journal of Anesthesiology & Clinical Science 2016
Harsoor et al. J. Evid. Based Med. Healthc Vol. 3/Issue 91/Nov. 14, 2016
Zhang et al. Drug Design, Development and Therapy 2019:13 1171–1175
Li et al. EXPERIMENTAL AND THERAPEUTIC MEDICINE 20: 454-460, 2020

NEUROLOGICAL

BUPI + FENTA vs DEX

ROPI + SUF vs DEX

❖ **Dex ↑ motor block and sedation**

❖ **No excessive sedation**

Variable	Group BD (n=44)	Group BF (n=43)
Motor block (maximum bromage score) 0-1-2-3(n)	40-4-0-0	38-6-0-0
Sedation score of 1-2-3-4 (n)	20-24-0-0*	33-10-0-0

M. F. Selim et al. Journal of Prenatal Medicine 2012; 6 (3): 47-54

Group	BF (Number)	BD (Number)	p value
Bromage Scores (0-1-2-3)	28-2-0-0	17-12-1-0	0.004
Ambulation	26	3	<0.0001

Harsoor et al. J. Evid. Based Med. Healthc., Vol. 3/Issue 91/Nov. 14, 2016

Variable	Group RS (n=35)	Group RD (n=36)	Group RDS (n=36)	P-value
Bromage score (1/2/3/4)	27/2/1/0	9/19/2/0 ^b	22/8/0/0 ^d	0.000

Li et al. EXPERIMENTAL AND THERAPEUTIC MEDICINE 20: 454-460, 2020

Variables	Group D (n=85)	Group F (n=85)	P-value
Motor block	14	3	0.004
Sedation	9	0	0.002

Soliman et al. Journal of Anesthesiology & Clinical Science 2016

NEUROTOXICITY

Local anesthetics neurotoxicity?

The efficacy and neurotoxicity of dexmedetomidine administered via the epidural route

Konakci, S.*; Adanir, T.*; Yilmaz, G.*; Rezanko, T.†

[Author Information](#) ©

European Journal of Anaesthesiology 25(5):p 403-409, May 2008. | DOI: 10.1017/S0265021507003079

21 New Zealand white rabbits
Epidural catheter
Lido (2%) / Lido (2%) + dex (5 µg) / Dex (10 µg)

- No differences between the groups for ischaemia of the medulla spinalis
- Evidence of demyelination of the oligodendrocytes in the white matter in Group D > L ($P = 0.035$)

Evaluation of the neurotoxicity of intrathecal dexmedetomidine on rat spinal cord (electromicroscopic observations)

0.1 ml Intrathecal, rats
10 µg dex / 0.9% NaCl

Saudi Journal of Anesthesia / Volume 12 / Issue 1 / January-March 2018

- A single intrathecal dose of dexmedetomidine did not lead to histologic neurotoxicity

Efficacy and safety of dexmedetomidine-ropivacaine versus sufentanil-ropivacaine for epidural labor analgesia: a randomized controlled trial

Mei Fan et al.

❖ No difference in clinical practice ...

- 10 mL loading dose
- RD: 0.1% ropivacaine + 0.5 µg/mL dex (80)
- VS
- RS: 0.1% ropivacaine + 0.5 µg/mL suf (80)
- CEI rate of 6 mL/h
- PCEA: bolus 6 mL / 20 min

Respiratory rate (breaths/min)	RD (n=80)	RS (n=80)	P value
Heart rate (beats/min)	82.6±8.0	86.4±11.5	0.016 ^a
Body temperature (°C)	36.6±0.2	36.6±0.2	0.747 ^a

Comparison of dexmedetomidine and lipophilic opioids as adjuvants to local anesthetics for epidural labor analgesia: a meta-analysis of randomized controlled trials

Shi-ke Yang^{1,*}, Min Liu¹, Jie Chen¹, Yuan-yuan Yang¹, Fang-zheng Zhuan¹, Wen-qun Sun¹, De-zhi Mao¹

12 Studies

Study	Participants	Intervention (sample size)	Control (sample size)	Anesthetic administration	Outcomes
Selim MF 2012 [9] Egypt	Primiparas and multiparas	0.25% Bupivacaine 12 mL + DEX 1 µg/kg = 17 mL (44)	0.25% Bupivacaine 12 mL + Fentanyl 1 µg/kg = 17 mL (43)	Bolus injection + rescue	1, 3, 4, 5, 7.
Karuna H 2016 [17] Bangalore	Primiparas	0.0625% Bupivacaine + DEX 1.5 µg/mL = 15 mL (30)	0.0625% Bupivacaine + 2 µg/mL Fentanyl = 15 mL (30)	Bolus injection + rescue	1, 3, 4, 5.
Zhang T 2019 [23] China	Primiparas	0.1% Ropivacaine + DEX 0.5 µg/mL (36)	0.1% Ropivacaine + Sufentanil 0.5 µg/mL (34)	CEI + PCEA	1, 2, 3, 4, 5, 6.
Cheng Q 2019 [24] China	Primiparas	0.125% Ropivacaine + DEX 0.5 µg/mL (40)	0.125% Ropivacaine + Sufentanil 0.5 µg/mL (40)	CEI + PCEA	1, 2, 3, 4, 5, 6.
Cheng Q 2019 [24] China	Primiparas	0.08% Ropivacaine + DEX 0.5 µg/mL (40)	0.08% Ropivacaine + Sufentanil 0.5 µg/mL (40)	CEI + PCEA	1, 2, 3, 4, 5, 6.
Soliman R 2016 [25] Saudi Arabia	Primiparas and multiparas	0.25% Bupivacaine 13 mL + DEX 1 µg/kg = 15 mL (85)	0.25% Bupivacaine 13 mL + Fentanyl 1 µg/kg = 15 mL (85)	Bolus injection + rescue	3, 4, 5, 6.
Huang Y 2016 [26] China	Primiparas	0.1% Levobupivacaine + DEX 0.5 µg/mL (60)	0.1% Levobupivacaine + Sufentanil 0.5 µg/mL (60)	CEI + PCEA	1, 2, 3, 4, 5.
Zhu X 2018 [27] China	Primiparas	0.1% Ropivacaine + DEX 2 µg/mL (56)	0.1% Ropivacaine + Sufentanil 0.5 µg/mL (56)	CEI + PCEA	1, 2, 5, 6.
Mao S 2017 [28] China	Primiparas	0.1% Ropivacaine + DEX 0.5 µg/mL (40)	0.1% Ropivacaine + Sufentanil 0.5 µg/mL (40)	CEI + PCEA	1, 2, 3, 4, 5, 6.
Shen S 2020 [29] China	Primiparas	0.1% Ropivacaine + DEX 1 µg/mL (60)	0.1% Ropivacaine + Fentanyl 2 µg/mL (60)	PIEB + PCEA	1, 2, 3, 4, 5.
Tang Y 2019 [30] China	Primiparas	0.09% Ropivacaine + DEX 0.5 µg/mL (33)	0.09% Ropivacaine + Sufentanil 0.5 µg/mL (30)	PIEB + PCEA	1, 2, 3, 4, 5, 7.
Yu C 2020 [31] China	Primiparas	0.1% Ropivacaine + DEX 0.5 µg/mL (30)	0.1% Ropivacaine + Fentanyl 2 µg/mL (30)	PIEB + PCEA	1, 2, 4, 5, 7.

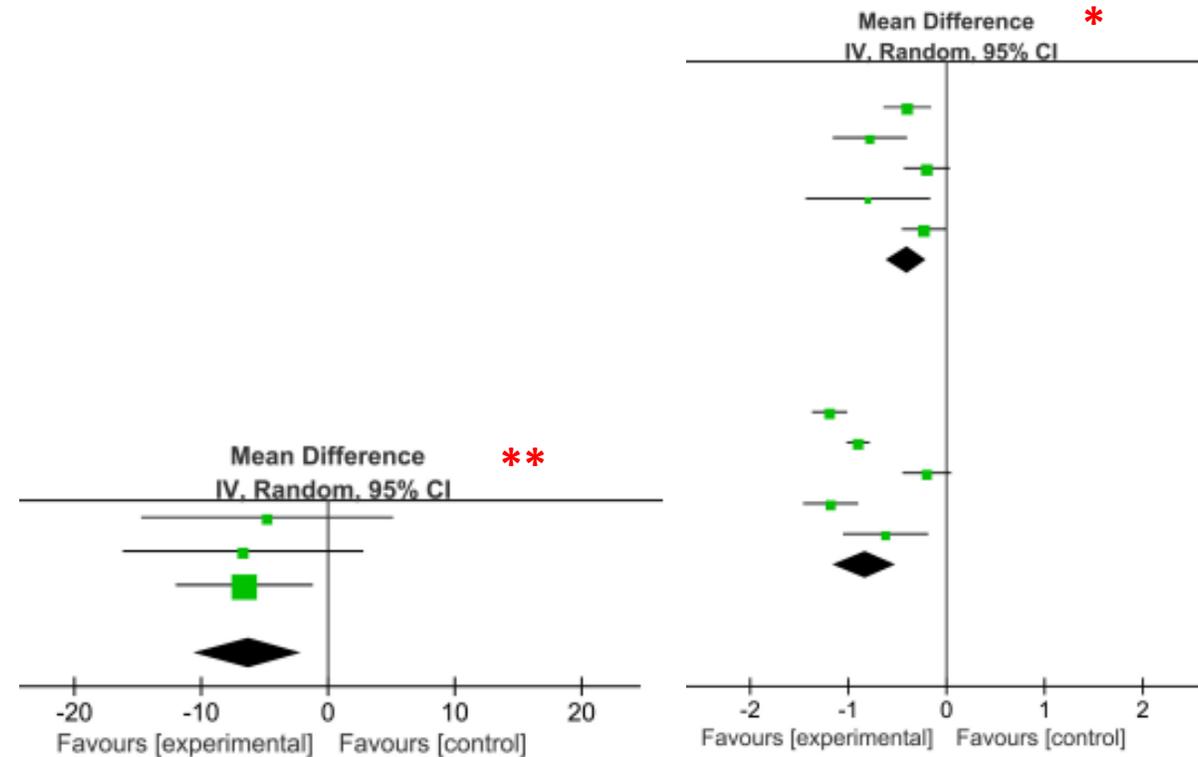
DEX, dexmedetomidine; CEI, continuous epidural infusion; PCEA, patient-controlled epidural analgesia; PIEB, programmed intermittent epidural bolus.

1: visual analogue scale (VAS) scores, 2: duration of labor, 3: mode of delivery, 4: maternal complications, 5: neonatal Apgar scores, 6: umbilical cord blood gas, 7: total analgesic consumption.

Comparison of dexmedetomidine and lipophilic opioids as adjuvants to local anesthetics for epidural labor analgesia: a meta-analysis of randomized controlled trials

Shi-ke Yang^{1,*}, Min Liu¹, Jie Chen¹, Yuan-yuan Yang¹, Fang-zheng Zhuan¹, Wen-gun Sun¹, De-zhi Mao¹

- ❖ ↓ VAS 30 minutes after induction *
- ❖ ↓ VAS on delivery
- ❖ ↓ Analgesic consumption **
- ❖ ↓ Duration of the 1st and 2nd stage of labor
- ❖ ↓ Nausea, vomiting, pruritus, shivering
- ❖ ↑ Bradycardia and motor blockade



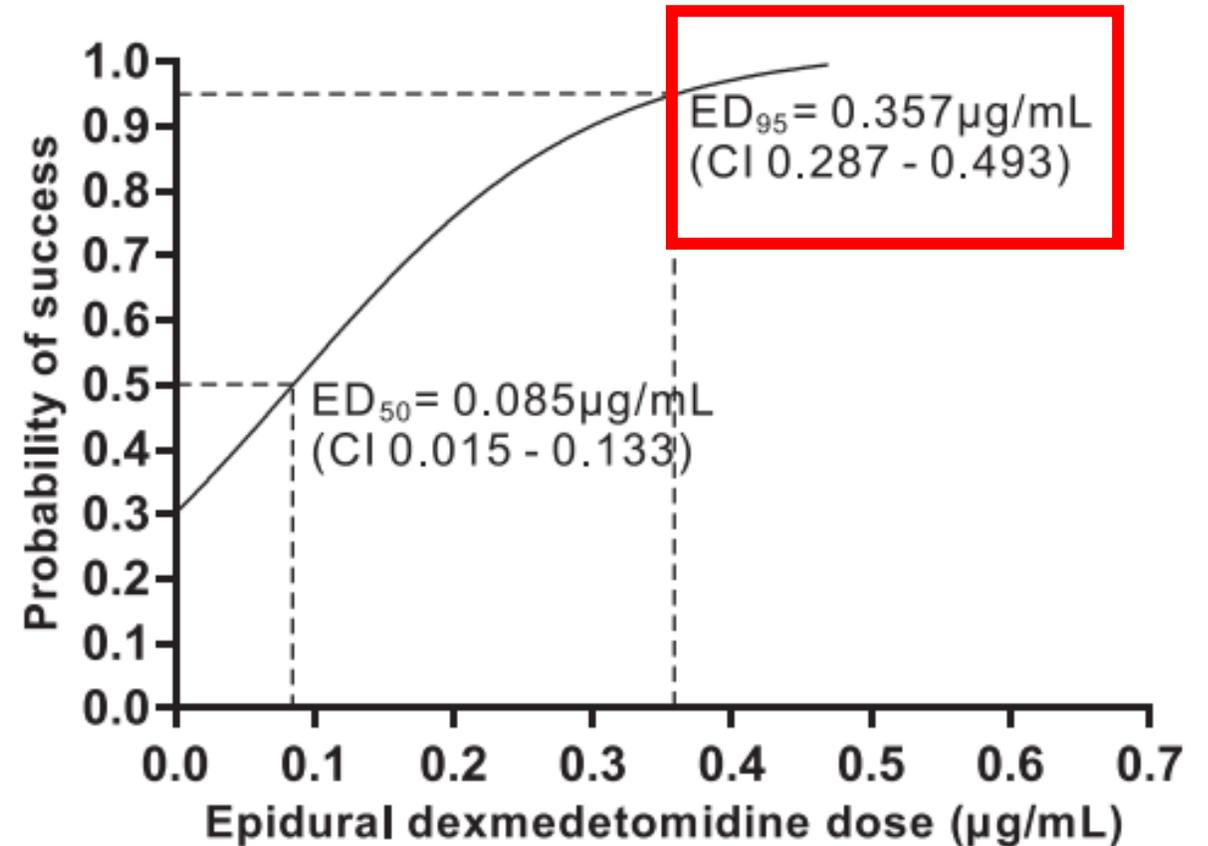
DOSE DETERMINATION

Determination of the Dose-Response Relationship of Epidural Dexmedetomidine Combined with Ropivacaine for Labor Analgesia

Jian-Xin et al.

- 0.075% ropi
+ DEX groups of different concentration (20)
- Primary outcome:
Effective rates of labor analgesia for different doses of dexmedetomidine at 30 min of administration

❖ **Clinically recommended 0.4 µg /ml**



LA CONCENTRATION

10 ml ropivacaine ± 0.5µg/ml dex

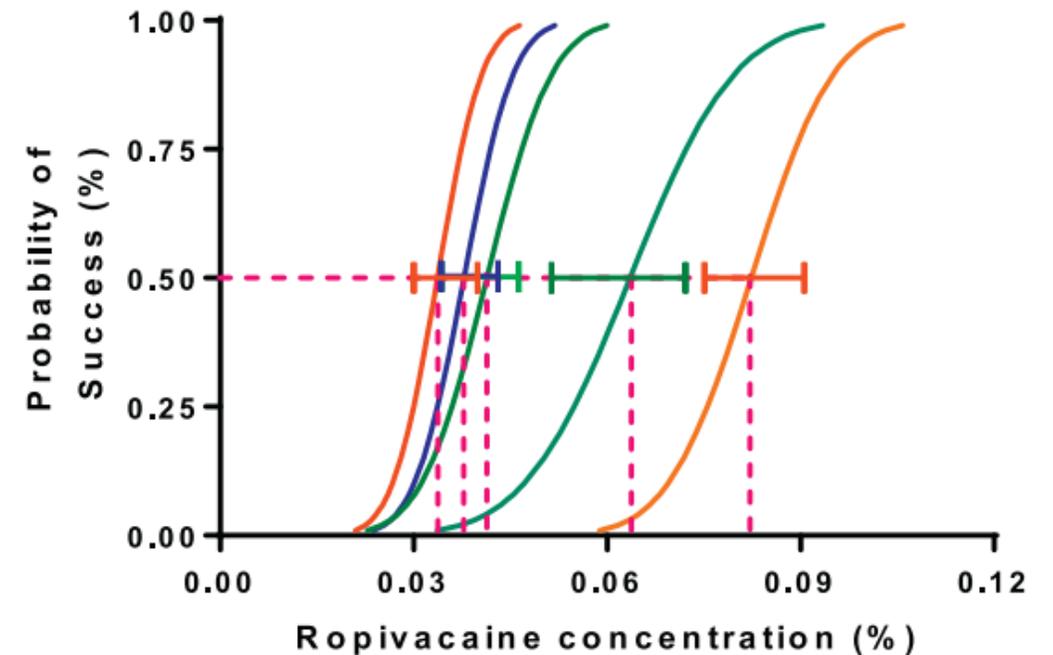
In each group the initial concentration of ropi was 0.1% for the first patient

If analgesic effect of the previous patient was adequate, the concentration of ropivacaine for the next patient was decreased by 0.01%

Outcome: median effective concentration (EC50) of epidural ropivacaine for labor analgesia

Adjuvant	None	Dex 0.5µg/ml	Clo 60 µg	suf 5µg/ml
Ropi EC 50 %	0.083	0.062	0.035	0.023

	Concentration (µg/ml)	Ropi EC 50 (%)	Ropi EC 95 (%)
— group 0	0	0.082	0.095
— group 0.3	0.3	0.064	0.080
— group 0.4	0.4	0.041	0.052
— group 0.5	0.5	0.034	0.041
— group 0.6	0.6	0.038	0.046



Intrathecal dexmedetomidine improves epidural labor analgesia effects: a randomized controlled trial

Gehui Li1 , et al.

- Group C: 1 mL Nacl 0.9%
- Group D: 5 µg Dex
- Group S: 5 µg suf
- PCEA 0.1% ropi + 0.2 µg /ml suf

Time (minutes)	Group C (n = 36)	Group D (n = 36)	Group S (n = 35)	P-value
Baseline	8.91 ± 3.05	9.01 ± 3.11	9.13 ± 2.35	0.964
5	8.53 ± 2.11	4.32 ± 2.98 ^b	4.51 ± 2.15 ^b	<0.001
10	5.22 ± 1.91	2.98 ± 1.99 ^b	2.86 ± 1.63 ^b	<0.001
15	3.56 ± 2.89	2.95 ± 2.12 ^b	2.67 ± 1.87 ^b	<0.001
20	2.91 ± 2.15	2.66 ± 2.33	2.59 ± 2.19	0.770
30	2.64 ± 2.55	2.53 ± 2.16	2.37 ± 1.19	0.847
60	2.74 ± 2.55	4.57 ± 3.55	4.13 ± 2.81	0.054

Onset time (minutes)	19.69 ± 3.11	8.39 ± 3.41 ^D	7.78 ± 2.21 ^D	<0.001*
Duration of intrathecal injection (minutes)	0.00	48.41 ± 2.55 ^b	50.85 ± 2.27 ^b	<0.001*
Total volume of anesthetic solution (mL)	56.65 ± 4.17	43.44 ± 2.14 ^b	44.34 ± 2.33 ^b	<0.001*
Bolus frequency	6.51 ± 2.21	4.50 ± 1.21 ^a	4.62 ± 1.01 ^a	0.001*
Bromage score (1/2/3/4)	36/0/0/0	36/0/0/0	35/0/0/0	1.000 [#]
Hypotension	1 (2.8)	1 (2.8)	0 (0.0)	1.000
Bradycardia	1 (2.8)	2 (5.6)	1 (2.9)	1.000
Nausea	1 (2.8)	0 (0.0)	1 (2.9)	1.000
Vomiting	0 (0.0)	0 (0.0)	0 (0.0)	1.000
Shivering	4 (11.1)	0 (0.0) ^{a,c}	6 (17.1)	0.046
Pruritus	1 (2.8) ^a	0 (0.0) ^{a,c}	5 (14.3)	0.025
Excessive sedation	1 (2.8)	2 (2.8)	2 (5.7)	0.803

C-SECTION

Sufentanil vs. Dexmedetomidine as Neuraxial Adjuvants in Cesarean Section: A Mono-Centric Retrospective Comparative Study



Intrathecal Bupivacaine 0.5% 10 mg + dex 10 µg vs Suf 5 µg

OUTCOMES	DEX (n=24)	SUF (n= 28)	Statistical significance
Time to onset sensory block min	7.3	7.4	
Hypotension	7	8	
Bromage score 1	24	28	
VAS h24	4 ± 2	2 ± 1	*
Flatus time ≤ 12h	24	7	*
Shivering	0	2	
Nausea	1	4	
Vomiting	1	4	
Itching	0	10	*
APGAR score 1 and 5 min > 7	24	28	

❖ Suf ↑ postoperative analgesia

❖ Dex ↓ pruritus

❖ ↔ Hypotension

Addition of dexmedetomidine to epidural morphine to improve anesthesia and analgesia for cesarean section

YANG et al., 2020

12 ml bolus: 0.75% ropi + Mo 2mg ± Dex 0.5 µg/kg
PCEA 48h

OUTCOMES	Mo (40)	Mo + Dex (40)	Statistical significance
Level of sensory block	T6	T6	
NRS at incision	0	0	
Visceral pain	28	19	*
NRS at peritoneal traction	5	0	
Rescue sufenta IV	26	15	*
Ramsay sedation score 0 , 4 and 12h	2	3	*
NRS post operative	4.5	0	

- ❖ ↓ intra and post-operative visceral pain
- ❖ Better sedation during and following delivery
- ❖ No influence on morphine-associated side effects

Comparison of Intrathecal Dexmedetomidine with Morphine as Adjuvants in Cesarean Sections

Xiaofei Qi et al.

2 mL: Bupi 0.5% alone vs ± Dex 5 µg vs ± Mo 100 µg

OUTCOMES	BD (40)	BM (40)	B (39)	Statistical significance
Sensory onset (min)	6.5 ± 1.3	7.8 ± 2.3	7.4 ± 2.2	+ ++
Motor onset (min)	4.8 ± 1.3	5.9 ± 2	5.8 ± 1.9	+ ++
Sensory regression (min)	253 ± 42	192 ± 40	188 ± 37	+ ++
Motor regression (min)	226 ± 40	161 ± 40	162 ± 25	+ ++
VAS intraoperative	0.5 ± 0.5	0.5 ± 0.5	0.5 ± 0.5	
VAS postoperative ≤ 24h	2.7 ± 1.2	2.7 ± 1.2	3.5 ± 1.5	+
Time to first analgesic (h)	17 ± 6	16 ± 6	3.5 ± 1.6	+
Total volume analgesics 24h	16 ± 5	19 ± 5		+
Total volume analgesics 48h	103 ± 20	115 ± 32	150 ± 31	+
Shivering	3	12	14	+ ++
Pruritus	0	13	1	+ ++

- ❖ ↑ Sensory and motor blockade
- ❖ ↔ Analgesic effects

- ❖ ↓ Side effects
- ❖ No obvious side effects on neonates

The Anesthetic Effect and Safety of Dexmedetomidine in Cesarean Section: A Meta-Analysis

Gang Pang et al.

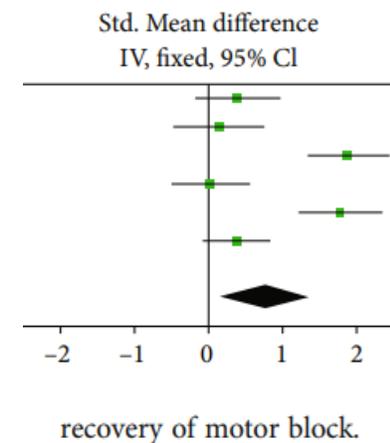
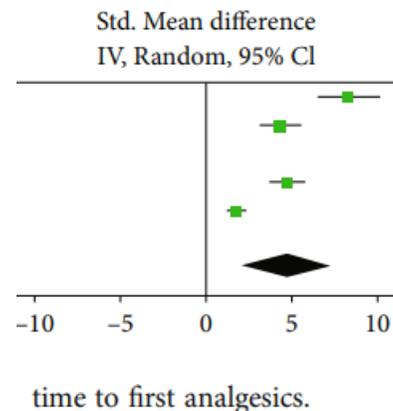
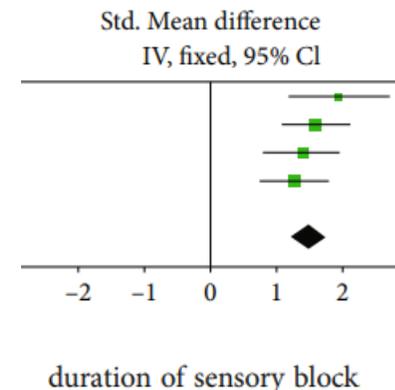
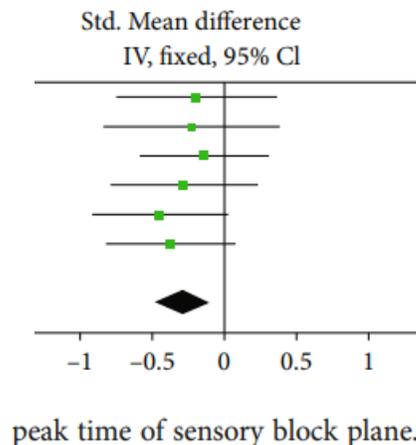
8 studies _(epidural and spinal)

Study (year)	Experimental group (n)	Control group (n)
Hanoura (2013) [18]	2 ml of spinal anesthesia 0.5% bupivacaine+10 ml of epidural injection 0.25% bupivacaine+dexmedetomidine 1 μ /kg and fentanyl 100 μ g (n = 25)	2 ml of lumbar anesthesia 0.5% bupivacaine+10 ml of epidural injection 0.25% bupivacaine+fentanyl 100 μ g (25)
Li (2015) [19]	Lumbar anesthesia 10 mg bupivacaine+10 μ g dexmedetomidine (21)	Lumbar anesthesia 10 mg bupivacaine (21)
Liu (2015) [20]	1.5 ml of lumbar anesthesia 0.5% bupivacaine+0.5 μ g/kg dexmedetomidine (40); 1.5 ml of lumbar anesthesia 0.5% bupivacaine+1 μ /kg dexmedetomidine (40) (n = 80)	1.5 ml of lumbar anesthesia 0.5% bupivacaine+20 ml of 0.9% sodium chloride injection (n = 40)
Nasseri (2017) [21]	Lumbar anesthesia 12.5 mg 0.5% bupivacaine+5 μ g dexmedetomidine (25)	Lumbar anesthesia 12.5 mg 0.5% bupivacaine+0.5 ml of 0.9% sodium chloride injection (25)
Sun (2015) [22]	2 ml of lumbar anesthesia 0.5% bupivacaine+10 μ g dexmedetomidine (30)	2 ml of lumbar anesthesia 0.5% bupivacaine+1.0 ml of 0.9% sodium chloride injection (30)
Yousef (2015) [23]	1.5 ml of spinal anesthesia 0.5% bupivacaine+10 ml of epidural infusion 0.25% bupivacaine+1 ml of dexmedetomidine 0.5 μ g/kg+1 ml of fentanyl 50 μ g (40)	1.5 ml of spinal anesthesia 0.5% bupivacaine+10 ml of epidural injection 0.25% bupivacaine+1 ml of 0.9% sodium chloride injection+1 ml of fentanyl 50 μ g (40)
Qi (2016) [24]	2 ml of 0.5% bupivacaine containing 5 μ g of dexmedetomidine (n = 40)	2 ml of 0.5% bupivacaine alone (n = 40)
Xia (2018) [25]	Bupivacaine+5 mcg dexmedetomidine (45)	Bupivacaine+the same volume of saline (45)

The Anesthetic Effect and Safety of Dexmedetomidine in Cesarean Section: A Meta-Analysis

Gang Pang,¹ Yuanmao Zhu,² Yan Zhou,³ and Shanshan Tong

- ❖ ↓ peak time
- ❖ ↑ Duration of the sensory block
- ❖ ↑ Onset of the first postoperative pain
- ❖ ↓ Postoperative pain
- ❖ ↓ Nausea, vomiting, chills, and fever
- ❖ Stable hemodynamics



LIMITATIONS

LIMITATIONS

- Not licensed for epidural/ intrathecal use (swissmedic/FDA)
- Lack of european or American studies
- Limited number of studies
- Single-center clinical trial
- Small number of patients
- One part of the world
- Limited studies comparing Dex to Clo a frequently used adjuvant

SUMMARY OF OUTCOMES

Intensity of analgesia	++
Onset time	↓↓
Duration of analgesia	↑↑
LA sparing effect	↓↓ vol and concentration
Hypotension	-
Bradycardia	↑ not clinical significant, no treatment required
Nausea	-
Vomiting	-
Shivering	↓↓
Motor block	↑ dependent on LA concentration
Sedation	↑ No excessive sedation
Neonatal outcomes	No difference
Labor outcomes	Shorter 1st stage

TAKE HOME MESSAGES

- Dex seems to be an interesting molecule for labor analgesia
- Effects on the mother and newborn needs further research before clinical promotion for neuraxial use
- Large-scale multicenter phase IV clinical trials with larger sample size

THANK YOU

