#### first announcement

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#### L'ANESTHÉSIE **R**egional Anesthesia RÉGIONALE Its Technic and Clinical Application Perth Regional Anaesthesia Group Gaston Labat, M. D. Gaston Labat, M. D. (PRAG Furtion Regional Anesthesia at The New York University; Laureate of the Facility of Sciences, University of Montpellier; Laureate of the Faculty of Medicine, University of Paris; Formerly Special Lecturer on Regional GASTON LABAT. De in Faculte de Medecine de Paris, Laureat de la Faculté des Sciences de Montpelli avo Foundation, University of Minnesota Journal Club Meeting TROISIÈME Mafeitzeral Mamat Avec 308 figures dans le texte. Anaesthesia & Pain Medicine riginal Illustrations Royal Perth Hospital 3 July 2012 PARIS LIBRAIRIE OCTAVE DOIN GASTON DOIN, ÉDITEUR 8, PLACE DE L'ODÉON, S PHILADELPHIA AND LONDON W. B. SAUNDERS COMPANY 1922 23676



#### Randomized Study of the Effect of Local Anesthetic Volume and Concentration on the Duration of Peripheral Nerve Blockade

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PERTH REGIONAL ANAESTHESIA GROUP (PRAG) JOURNAL CLUB MEETING 3 JULY 2012

### Background

- Ultrasound guided technique had reduce the volume of local anaesthetic agent used to achieve a successful block.
- ? Unclear of block duration

#### Hypotheses

# Block duration depends on volume and concentration.



# Study Design

- Dual Centre
- Prospective
- Randomized
- Observer-Blinded trial

# Ethics approval

- Northern Y Regional Ethics Committee, Hamilton, New Zealand.
- Australian and New Zealand Clinical Trials Registry (ACTRN12611000155998, February 2011).
- Written & oral informed consent as per Helsinki Declaration

### Recruitment

Introduction Objectives Method Results Discussion Summary

#### • ASA I , II , III

- 16-80 year old
- Shoulder Surgeries
- February December 2011
  Southern Cross Brightside Hospital
  North Harbour Hospital

#### **Study Interventions**

Interscalene Block

Ropivucaine

- 0.75% 5,10,20 ml
- 0.375% 20, 40 ml



### **End Points**

#### • Primary:

Time to first shoulder pain

• Secondary:

Numerical Rated Pain Tramadol Consumption Numerical Rated numbness Adverse effect in 24 hours

#### Placement of catheter

- Initial modified Superficial Cervical Plexus block given
- Ultrasound & Nerve Stimulator Needle used
- End point ultrasound: 10ml Dextrose
  5% spread
- End point neurostimulator: Elicitation of motor response < 0.5mA</li>
- Catheter advanced 2-3cm from needle tip blindly

### Anaesthesia & Analgesia

- GA with spontaneous breathing
- Desflurane Laryngeal Mask
- Ropivucaine infusion given before surgical incision
- Acetaminophen & Parecoxib
- Rescue Alfentanil for RR> 25/min

## Post Op

Introduction Objectives Method Results Discussion Summary

- PACU strict exclusion criteria
- Elastomeric Ropivucaine infusion (PCRA) at the point of first shoulder pain.
- Multimodal Acetominophen, Diclofenac, Tramadol.

### Results

Introduction Objectives Method Results Discussion Summary

- 185 patients enrolled
- 61 excluded from analysis
- n=40 for 5ml group excluded (30% failure rate)
- Multivariate Regression Analysis Cox proportional Hazards model

#### TABLE 1. Patient and Surgical Characteristics (n = 185)

	5 mL 0.75% (n = 40)	10 mL 0.75% (n = 41)	20 mL 0.375% (n = 35)	20 mL 0.75% (n = 33)	40 mL 0.375% (n = 36)
Male sex	24 (60)	30 (73)	22 (63)	24 (73)	29 (81)
Age, y	49 (12)	48 (15)	49 (16)	49 (14)	46 (18)
Weight, kg	83 (53-121)	85 (47-115)	83 (52-134)	88 (64-125)	87 (61-125)
Body mass index, kg/m <sup>2</sup>	28 (19-45)	29 (19-41)	27 (18-47)	29 (21-38)	29 (20-39)
Surgery					
Open rotator cuff repair	6	12	6	11	9
Arthroscopic rotator cuff repair	12	9	2	8	6
Arthroscopic stabilization	5	8	9	8	11
Arthroscopic lateral clavicle resection	4	3	2	1	2
Arthroscopic acromioplasty	8	6	9	3	3
Arthroscopic capsular release	2	1	1	0	1
Total shoulder joint replacement	1	1	4	0	2
Other	2	1	2	2	2

Values are mean (SD), mean (range), or n.

	5 mL 0.75% (n = 40)	10 mL 0.75% (n = 41)	20 mL 0.375% (n = 35)	20 mL 0.75% (n = 33)	40 mL 0.375% (n = 36)	<b>P</b> *
Ultrasound needle end point	38 (97)	40 (98)	32 (91)	32 (97)	34 (94)	0.77
Stimulated motor response: deltoid/biceps/triceps/none	11/9/4/16	11/10/2/18	3/8/6/18	8/6/3/16	11/5/2/18	0.30
Minimum stimulation threshold, mA	0.65 (0.5-0.80)	0.70 (0.50-0.80)	0.60 (0.3-0.7)	0.70 (0.50-0.80)	0.70 (0.39-0.80)	0.66
Intraoperative alfentanil bolus ≥1	3 (8)	4 (10)	1 (3)	0 (0)	4 (11)	0.26
Surgery duration	75 (60-90)	80 (60-90)	80 (60-105)	80 (75-90)	75 (60-91)	0.73
PACU						
Exclusions						
PACU local anesthetic bolus	12 (30)	5 (12)	1(3)	2 (6)	3 (8)	0.006†
PACU catheter failure/reinsertion	0	1	0	0	2	
Protocol violation	0	0	1	0	1	
Lost to follow-up	0	2	1	1	1	

TABLE 2. Catheter Placement and Intraoperative and PACU Interventions (n = 185)

Values are n (%), n, or median (interquartile range).

\*P values refer to a 5-group comparison.

†With group 5 mL excluded, P = 0.50.

 Probability of pain as a function of time was associated with not only dose, but also volume corrected for concentration and concentration corrected for volume:

hazard ratio (95% confidence interval)

- for dose = 0.992 (0.987-0.997) (P = 0.002),
- volume = 0.959 (0.937-0.982) (P = 0.001),
- concentration = 0.852 (0.743-0.976)
  (P = 0.021).

### Volume

Introduction Objectives Method Results Discussion Summary Increasing the volume of ropivacaine 0.375% from 10 to 40 mL :

 increased median (quartiles) block duration from

**10.0** (9.5-11.5) to **15.0** (10.75-21) hours

#### Concentration

Increasing the concentration of 20 mL ropivacaine from 0.375% to 0.75% :

 Increased the median (quartiles) block duration from

**10.75** (9.75-14.0) to **13.75** (10.5-21.0) hours.

#### TABLE 3. Postoperative Outcomes (n = 152)

	5 mL 0.75% (n = 28)	10 mL 0.75% (n = 33)	20 mL 0.375% (n = 32)	20 mL 0.75% (n = 30)	40 mL 0.375% (n = 29)	<b>P</b> *
Tramadol consumption	0 (0-1)	0 (0-1)	0 (0-1)	0 (0-1)	0 (0-1)	0.50
Ropivacaine boluses	3 (2-4)	3 (1-5)	3 (2-4)	3 (2-5)	2 (1-4)	0.62
Worst shoulder pain NRS	3 (2-6)	5 (3-6)	4 (3-6)	4 (3-6)	3 (3-5)	0.42
"Average" shoulder pain NRS	1 (0-3)	2 (0-3)	2 (1-3)	2 (1-3)	2 (1-3)	0.98
Hand numbness NRS	9 (6-10)	8 (5-10)	7 (5-8)	8 (6-10)	8 (7-10)	0.17
Hand weakness NRS	8 (5-10)	7 (3–9)	7 (5-9)	7 (5-10)	8 (5-10)	0.85
Adverse effects**	9 (32)	12 (36)	9 (28)	17 (56)	12 (41)	0.14
Satisfaction NRS	9 (8-10)	10 (8-10)	10 (8-10)	9 (7–10)	10 (9–10)	0.16

Values are n (%) or median (interquartile range).

\*P values refer to comparisons of the 4 groups excluding 5 mL 0.75%. Respective P values were similar with inclusion of the 5 mL group.

NRS indicates numerical rating score (0-10, 0=0= no pain, hand numbness/weakness, very unsatisfied; 10 = worst imaginable pain, hand numbness/weakness, very satisfied).

\*\*Adverse effects included "breathlessness" or "difficulty taking a deep breath."

#### Discussion

- The first study to assess duration of block associated with volume & concentration.
- Interest due to reported cases and studies of ultra-low dose volume used. However, non analysed duration as outcome.

#### Discussion

 Despite clear association with volume & concentration, clinical relevance can be questioned.

#### Discussion

Introduction Objectives Method Results Discussion Summary theoretical reduction in the local anaesthetic systemic toxicity risk from lower volumes and concentrations outweighs the downside of shorter block duration, even though published clinical evidence does not support this principle.

#### Discussion

The relatively modest effect of both volume and concentration could be interpreted to mean that the only way to significantly prolong block duration is through perineural catheter placement

## Critique

Introduction Objectives Method Results Discussion Summary

- Exclusion of the 5ml group
- Technique of local anaesthetic agent deposition
- Secondary outcome analysis not powered to justify

#### Summary

In summary, this study found a clear association between local anesthetic volume, concentration (and dose), and the duration of interscalene block, findings that have particular relevance for the current trend in ultrasound-guided regional anesthesia of administering low local anesthetic volumes.

#### **THANK YOU**