



 **Съезд**
Congress



5-7 сентября 2018 / Санкт-Петербург
September 5-7, 2018 / St. Petersburg



Pre-eclampsia & HELLP syndrome

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Pre-eclampsia & HELLP syndrome

- Antenatal Issues
- Labour Analgesia
- Anaesthesia for Delivery
- High Dependency Care



- **Hypertension**

systolic >140 mmHg or diastolic >90 mmHg

severe: systolic >160 mmHg or diastolic >110 mmHg

- **Chronic hypertension**

pre-pregnancy or diagnosed before 20 weeks

- **Gestational hypertension**

new-onset after 20 weeks

no end-organ damage

- **Pre-eclampsia**

new onset after 20 weeks (early <34 vs. late >34)

end-organ damage

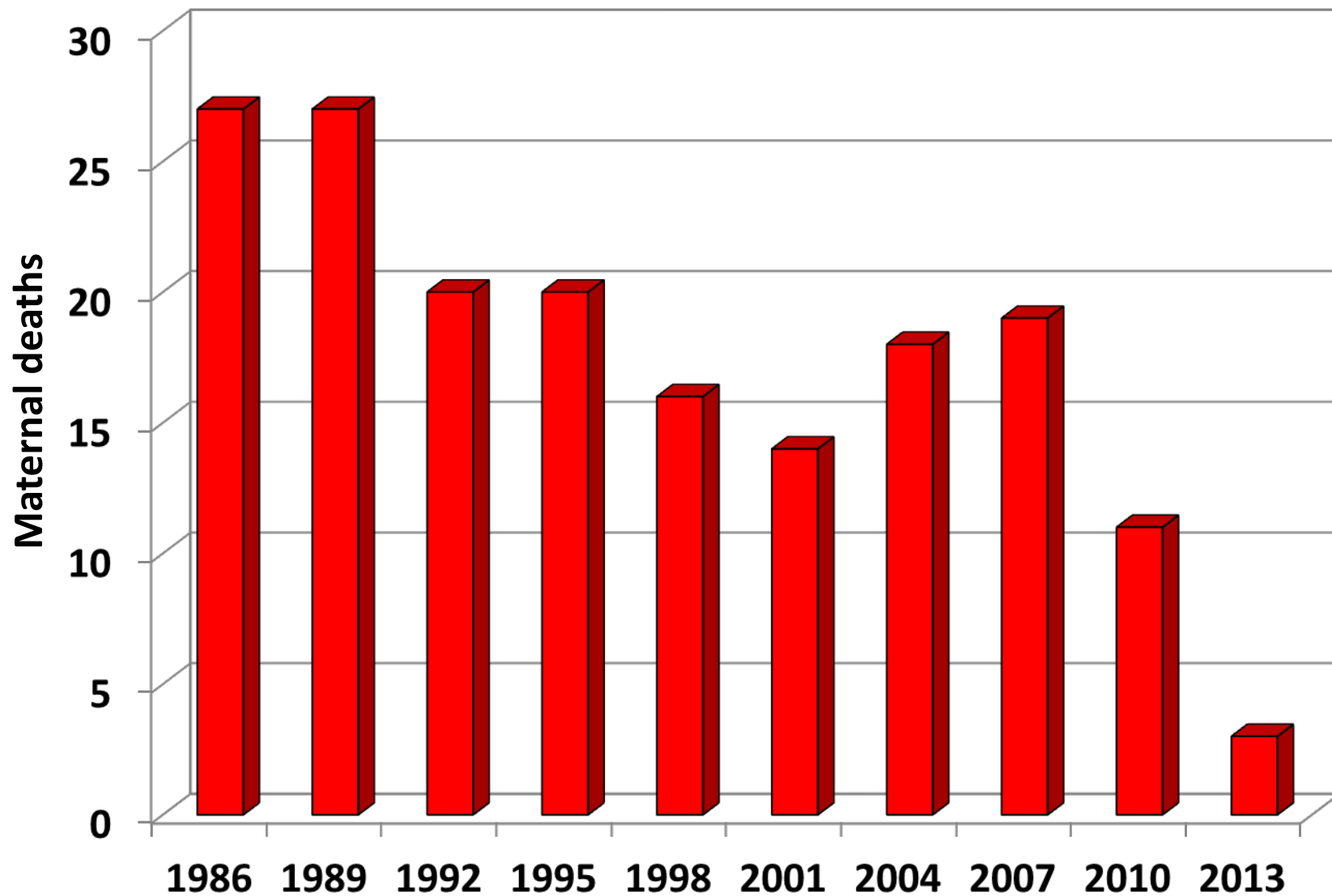
(proteinuria >300 mg / 24 h)

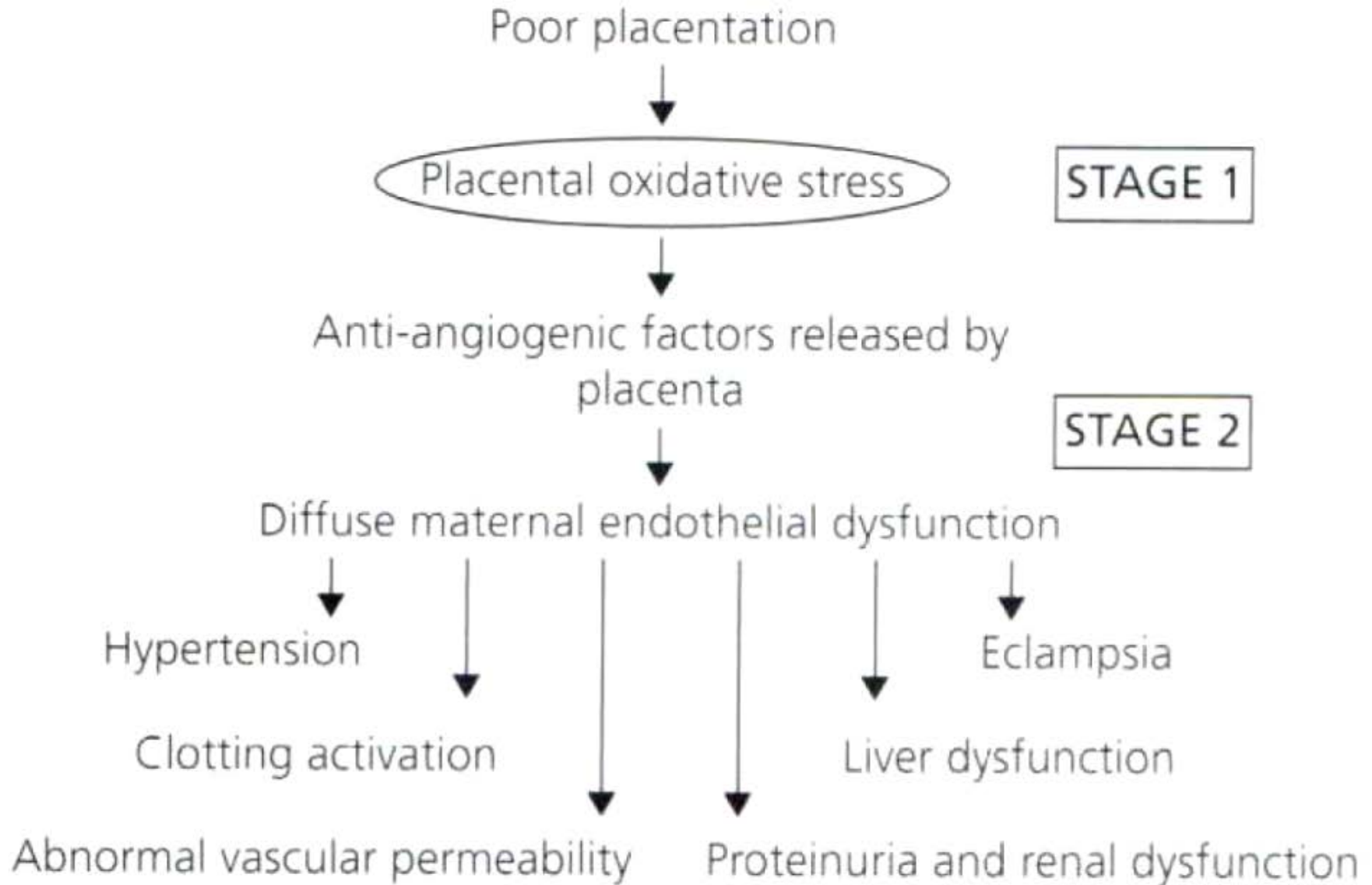
regression 3/12 post delivery

- **HELLP syndrome**

Haemolysis, Elevated Liver enzymes, Low Platelets

UK Maternal Deaths from Pre-eclampsia & Eclampsia





Presentation

- cardiovascular
- central nervous
- respiratory
- haematology
- hepatic
- renal
- fetus



Presentation

- cardiovascular
- central nervous
- respiratory
- haematology
- hepatic
- renal
- fetus
- hypertension
- ↑ cardiac output
- ↑ SVR
- loss of autoregulation
- cerebral haemorrhage
- future risk



Presentation

- cardiovascular
- central nervous
- respiratory
- haematology
- hepatic
- renal
- fetus
- headache
- visual disturbance
- nausea & vomiting
- photophobia
- clonus / hyperreflexia
- eclampsia

Presentation

- cardiovascular
- central nervous
- **respiratory**
- haematology
- hepatic
- renal
- fetus
- **breathlessness**
- **pulmonary oedema**
- **fluid management**
- **laryngeal oedema**
- **difficult intubation**



Presentation

- cardiovascular
- central nervous
- respiratory
- **haematology**
- hepatic
- renal
- fetus
- **thrombocytopenia**
- **haemolysis**
- **fibrinolysis**
- **coagulopathy**
- **thrombosis**



Presentation

- cardiovascular
- central nervous
- respiratory
- haematology
- hepatic
- renal
- fetus
- periportal oedema
- ↑ liver enzymes
- epigastric pain
- hepatic necrosis
- subcapsular haemorrhage
- rupture

Presentation

- cardiovascular
- central nervous
- respiratory
- haematology
- hepatic
- **renal**
- fetus
- **↓ glomerular filtration**
- **proteinuria**
- **↑ creatinine / urea**
- **↑ uric acid**
- **acute renal failure**

Presentation

- cardiovascular
- central nervous
- respiratory
- haematology
- hepatic
- renal
- fetus
- ↓ placental perfusion
- growth restriction
- preterm delivery
- abruption
- intrauterine death

Risk factors

- nulliparity
- age <20 or >40 years
- previous pre-eclampsia
- family history
- multiple pregnancy
- anaemia
- obesity
- hypertension, diabetes, renal disease
- autoimmune disease
- assisted reproduction
- high altitude



Diagnosis & Prognosis

- **blood pressure**
- **proteinuria**
- **urinary protein : creatinine ratio**
- **creatinine / uric acid / liver enzymes**
- **uterine artery Doppler studies**
- **SFlt-1 : Placental Growth Factor ratio**

NICE National Institute for Health and Care Excellence

NICE Pathways

NICE Guidance

Standards and indicators

Evidence services

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Home > NICE Guidance > Conditions and diseases > Cardiovascular conditions > Hypertension

Hypertension in pregnancy: diagnosis and management

Clinical guideline [CG107] Published date: August 2010 Last updated: January 2011 [Register as a stakeholder](#) [Uptake of this guidance](#)

Guidance

Tools and resources

Information for you

For your library

Overview

Introduction

Woman-centred care

Key priorities for implementation

1 Guidance

2 Notes on the scope of the guidance

3 Implementation

4 Research recommendations

5 Other versions of this guideline

6 Related NICE guidance

7 Updating the guideline

Appendix A: The Guideline Development Group and NICE project team

Guidance

1 Guidance

[Definitions](#)

- [1.1 Reducing the risk of hypertensive disorders in pregnancy](#)
- [1.2 Antenatal diagnosis and management of hypertension](#)
- [1.3 Assessment of proteinuria in hypertensive disorders of pregnancy](#)
- [1.4 Management of pregnancy with gestational hypertension](#)
- [1.5 Management of pregnancy with pre-eclampsia](#)
- [1.6 Fetal monitoring](#)
- [1.7 Intrapartum care](#)
- [1.8 Management of severe pre-eclampsia in a critical care setting](#)
- [1.9 Delivery](#)
- [1.10 Advice and follow-up care at transfer to community care](#)

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The following guidance is based on the best available evidence. The [full guideline](#) gives details of the methods and the evidence used to develop the guidance.



“Women with severe pre-eclampsia need effective team care based on clear communication and common understanding. There should be early involvement of intensive care specialists where appropriate.”

Published by
Association of Anaesthetists of Great Britain & Ireland
Obstetric Anaesthetists' Association

June 2013

Volume 118, Supplement 1, March 2011

BJOG

An International Journal of
Obstetrics and Gynaecology

Reviewing maternal deaths to make
not England, 2006-2008

March 2011

Report on the Confidential Enquiries into Maternal
Deaths in the United Kingdom



CMACE

Centre for Maternal and Child Enquiries
Improving the health of mothers, babies and children

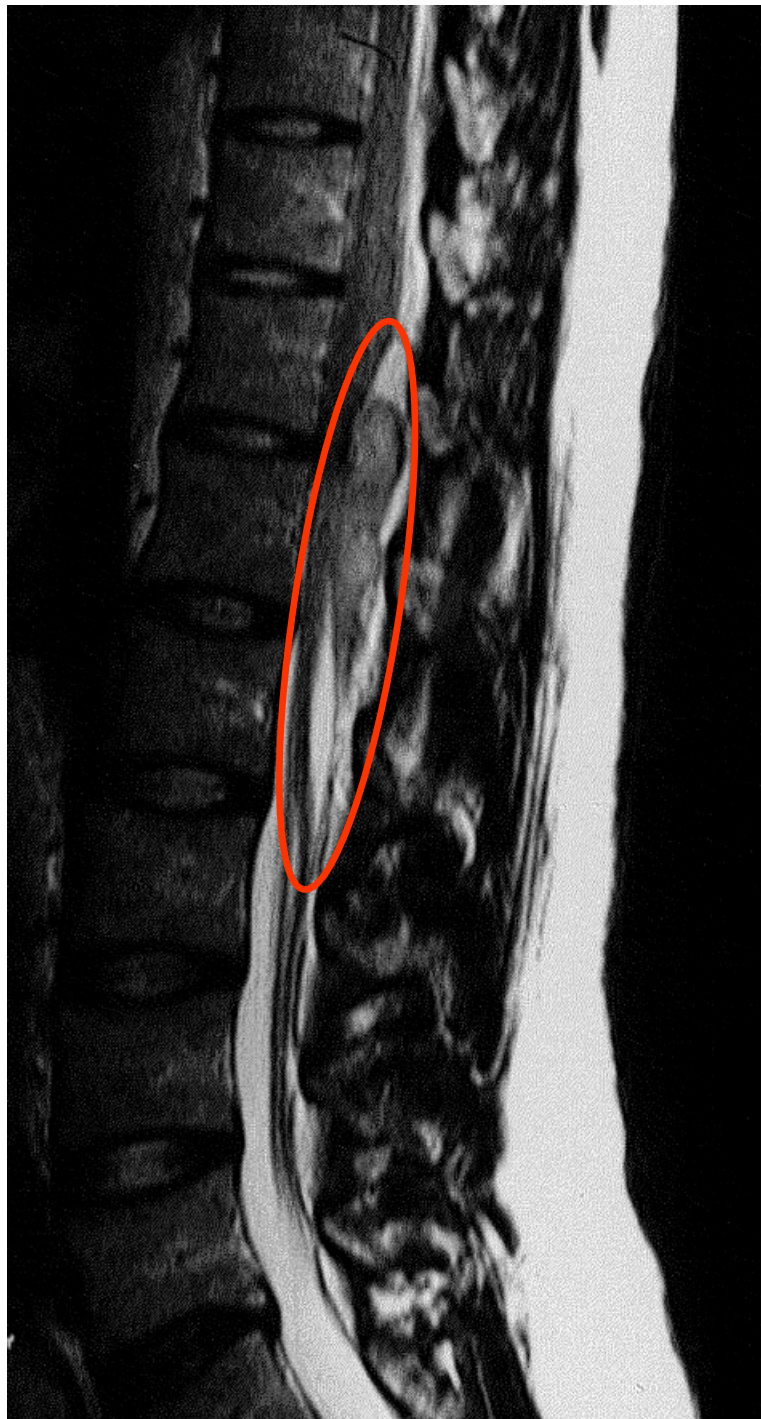
 **WILEY-
BLACKWELL**



Pre-eclampsia & HELLP syndrome

- Antenatal Issues
- **Labour Analgesia**
- Anaesthesia for Delivery
- High Dependency Care



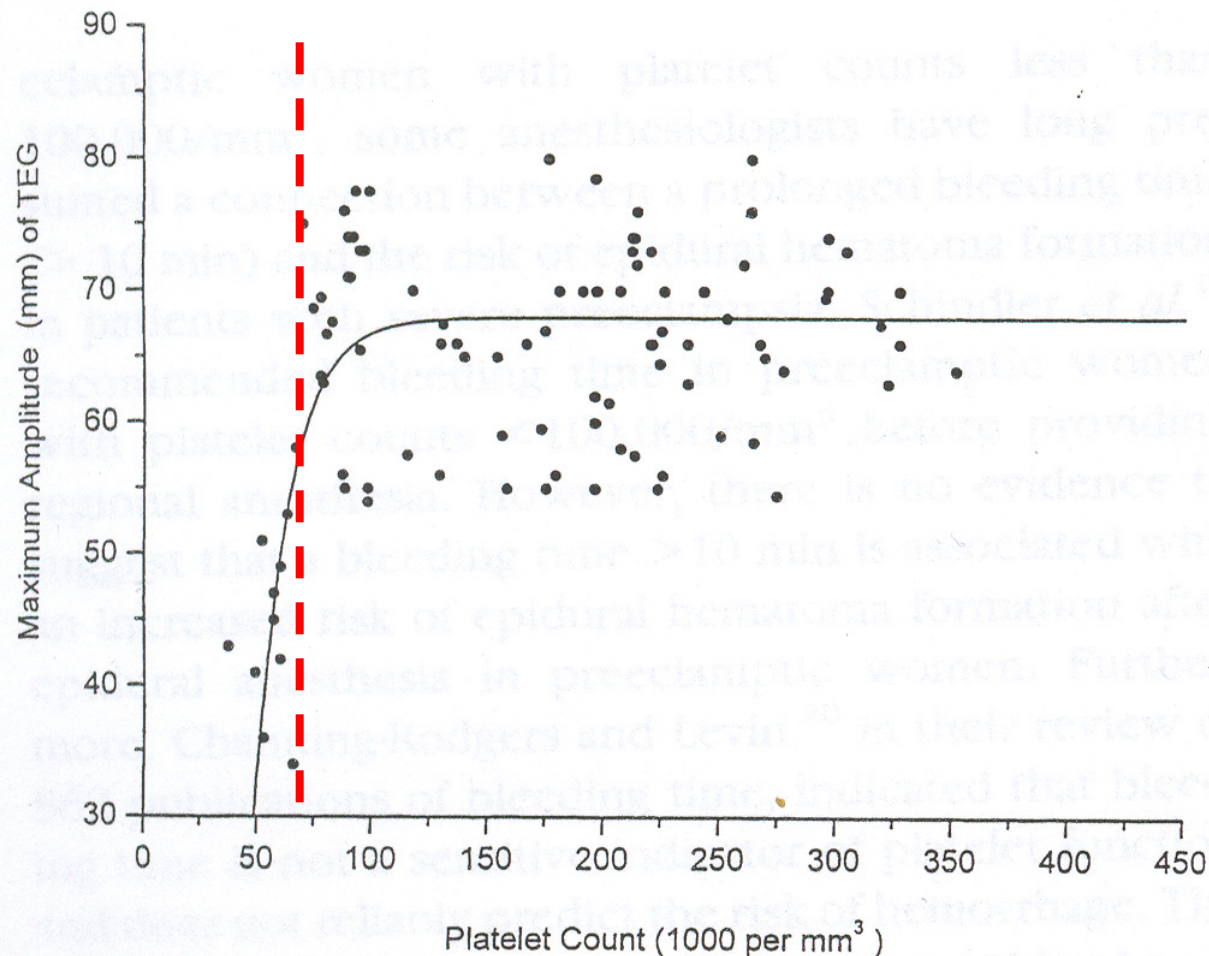




Tests of Coagulation

- **bleeding time**
- **platelet count**
- **clotting studies**
- **factor assays**
- **TEG / ROTEM**

Thromboelastography & pre-eclampsia



Sharma et al. Anesthesiology 1999



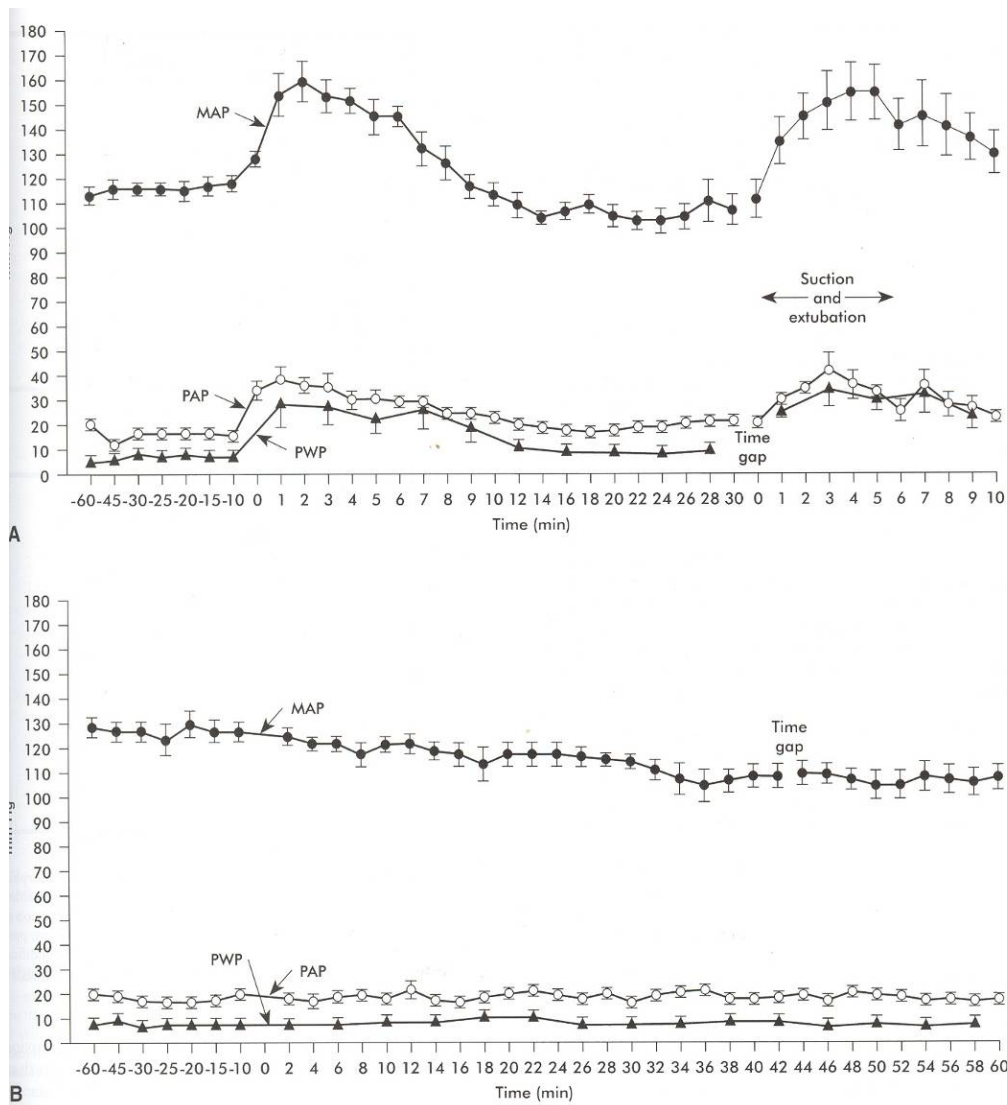
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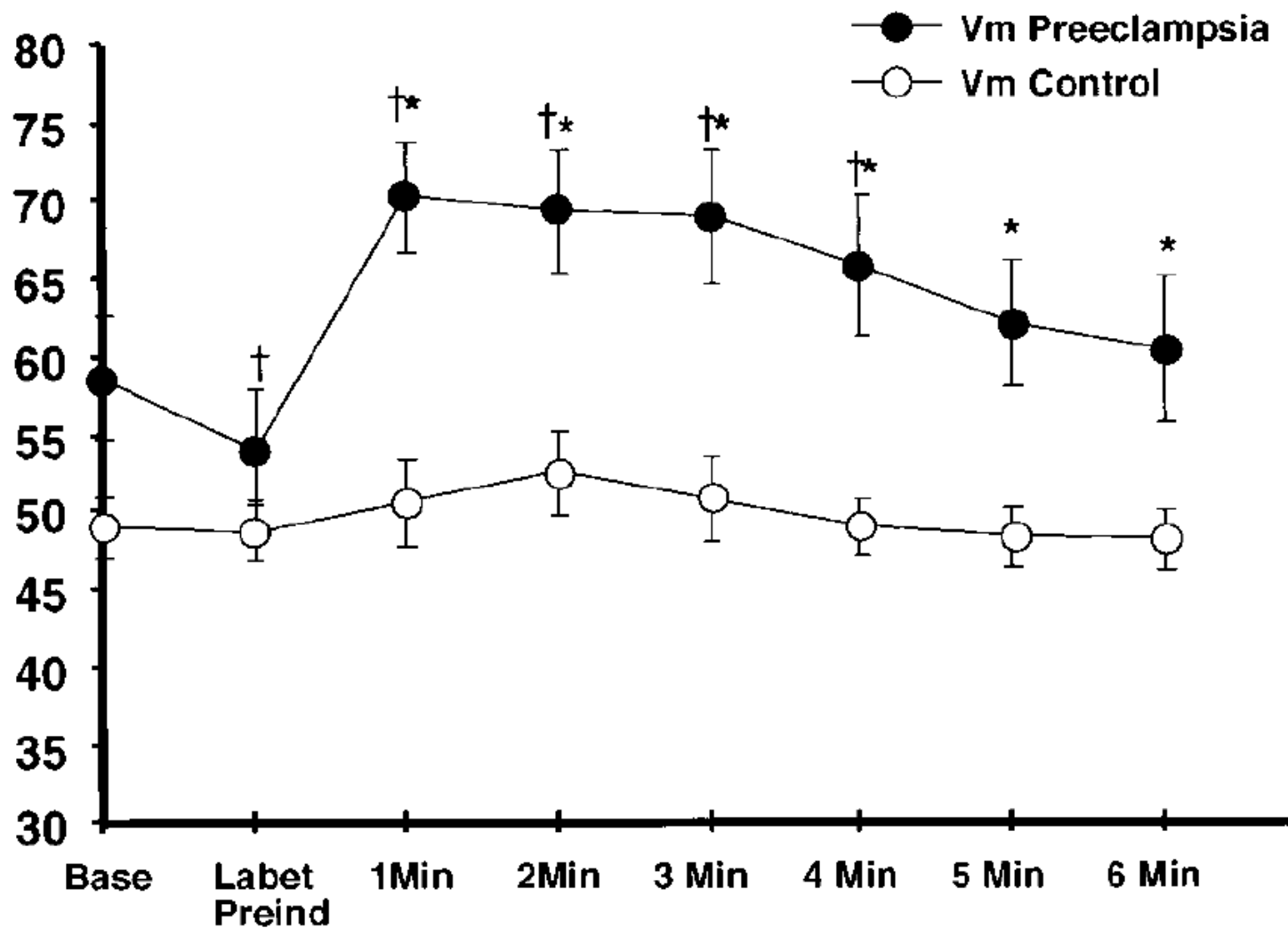


Hypertensive response to laryngoscopy



Hodgkinson et al. Can Anaesth Soc J 1980

Middle cerebral artery blood flow & pre-eclampsia



Ramanathan et al. Anesth Analg 1999



“Anaesthetists should anticipate an additional rise in BP at intubation in women with severe pre-eclampsia who are undergoing caesarean section under GA and take measures to avoid a speed that compromises maternal wellbeing, even when there are concerns about fetal wellbeing.”

December 2007

The Seventh Report of the Confidential Enquiries
into Maternal Deaths in the United Kingdom

Saving Lives, Improving Mothers' Care

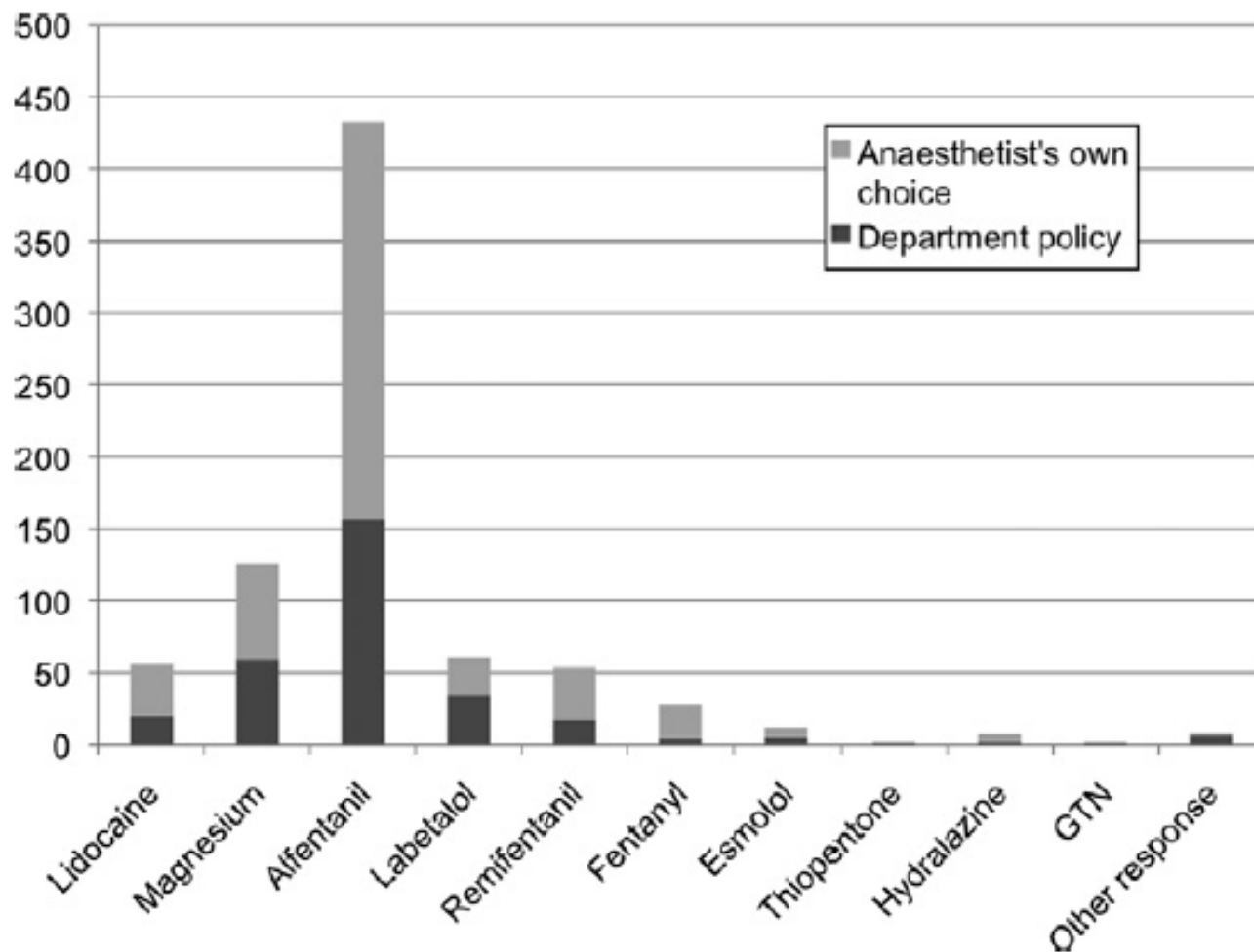
Surveillance of maternal deaths in the UK 2012–14 and
lessons learned to inform maternity care from the UK
and Ireland Confidential Enquiries into Maternal Deaths
and Morbidity 2009–14

“Whilst intubation may be required for airway control, maternal stabilisation and blood pressure control is vital prior to intubation in order to minimise maternal risk.”

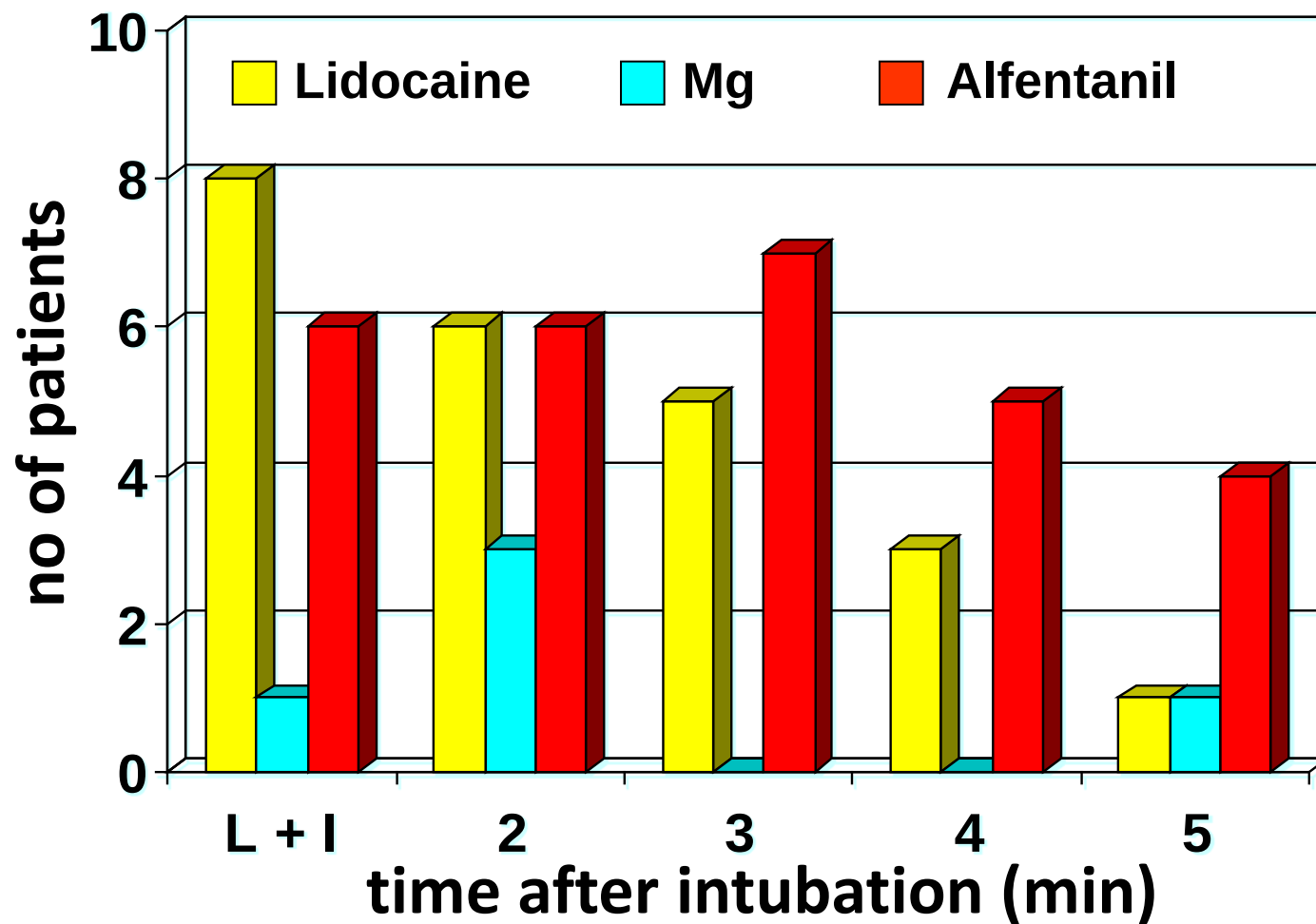


December 2016

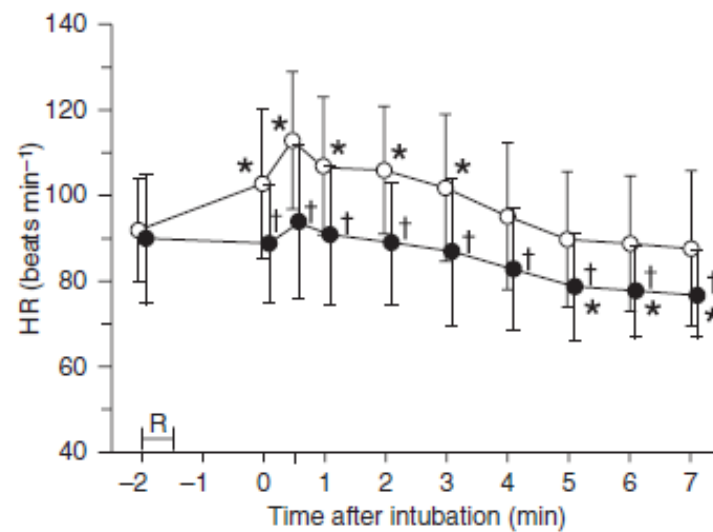
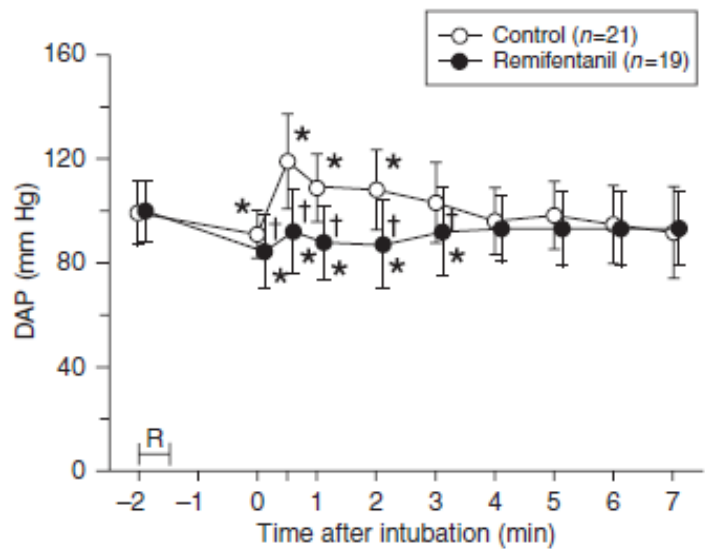
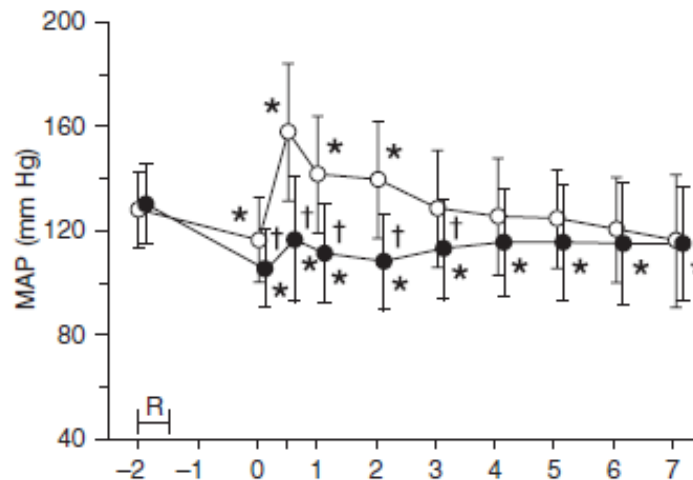
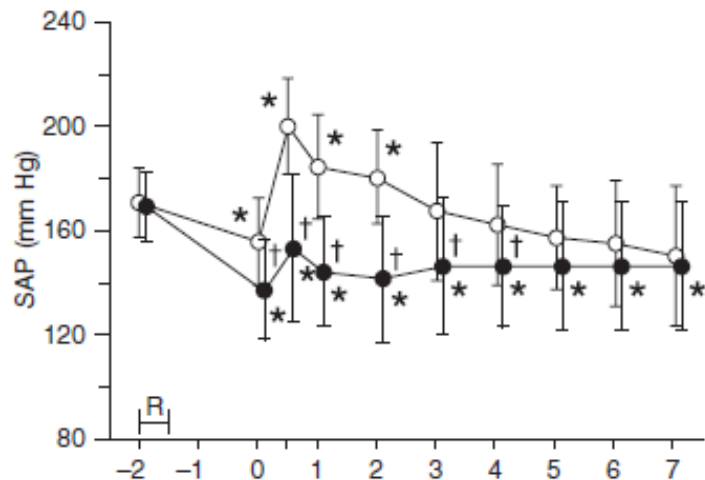
Drugs to attenuate pressor response



Attenuation of pressor response Systolic BP > 180 mmHg



Remifentanyl & pressor response



Remifentanyl & pressor response

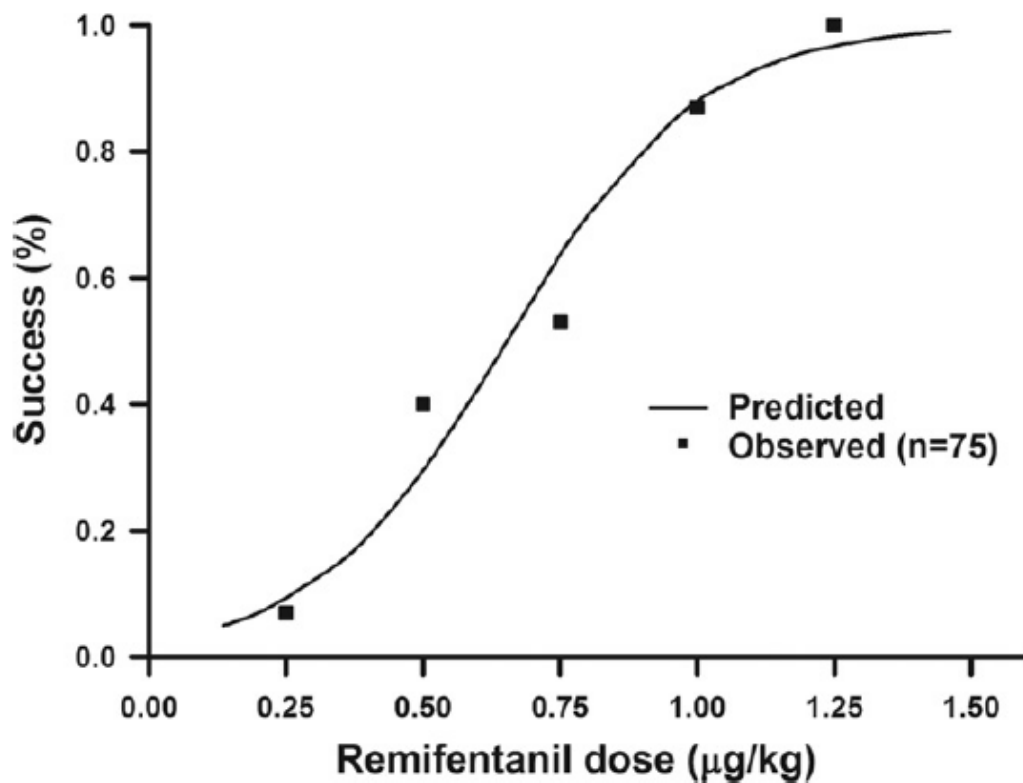
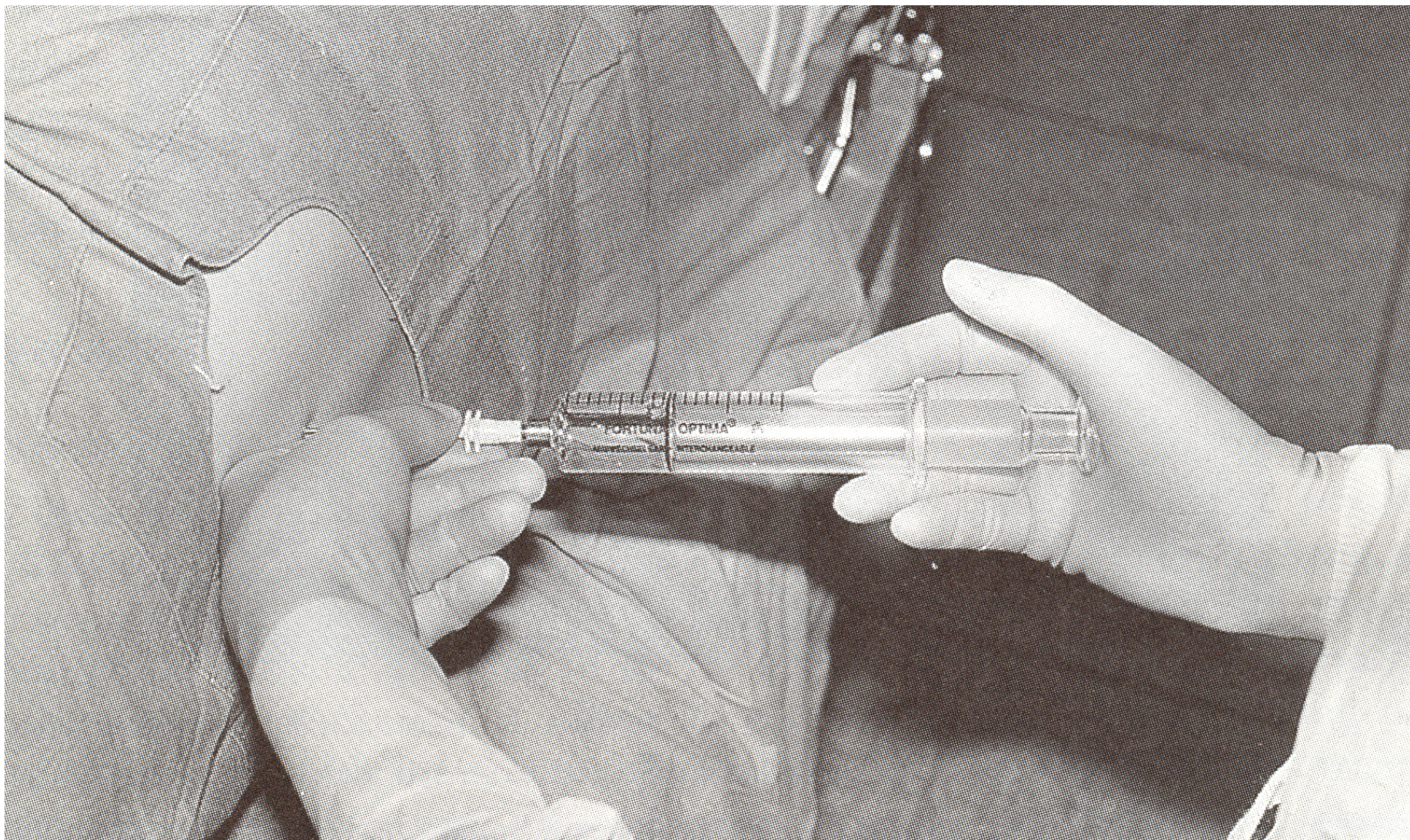
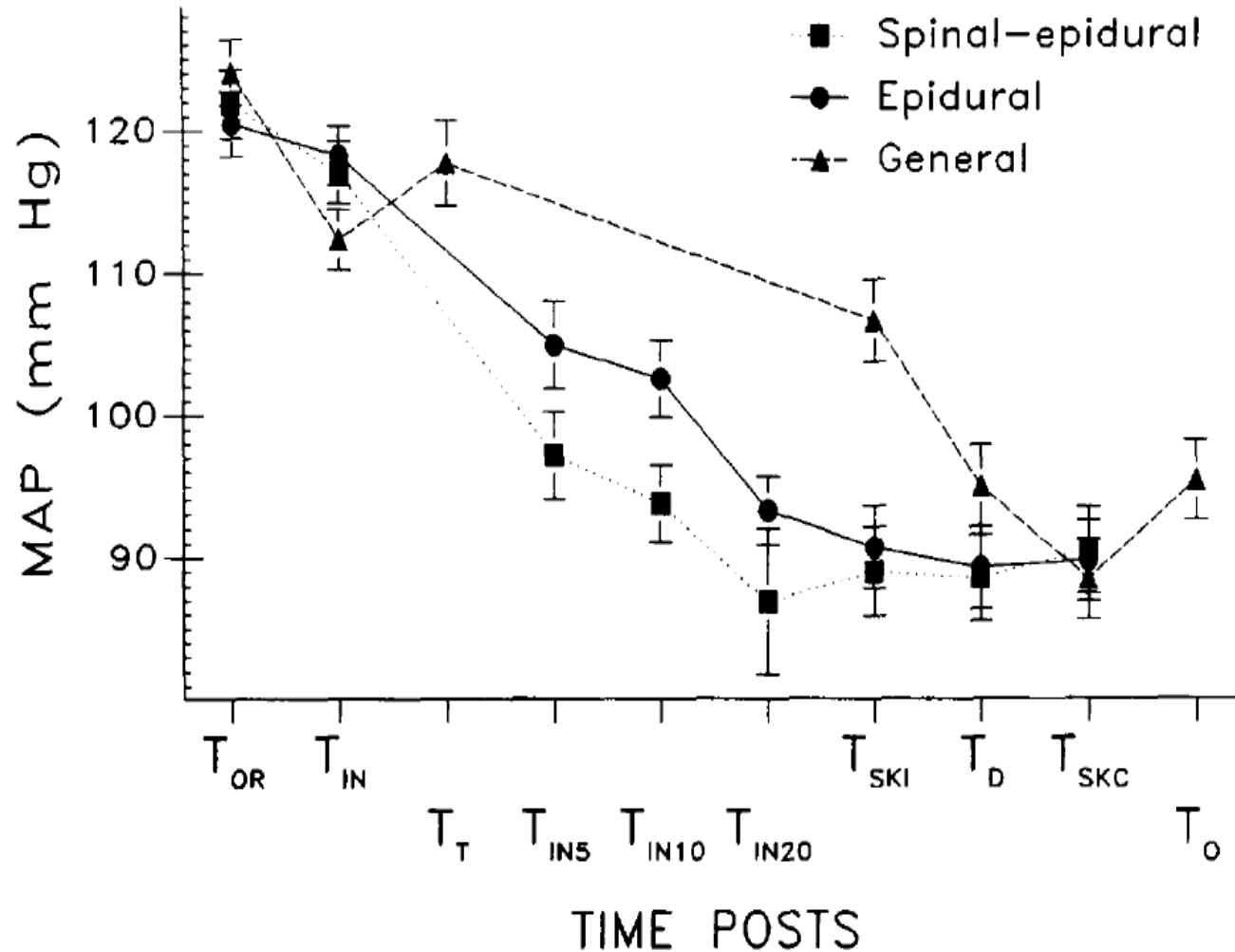


Fig. 3 Dose-response curve for success of intravenous remifentanyl in attenuating the pressor response to tracheal intubation.



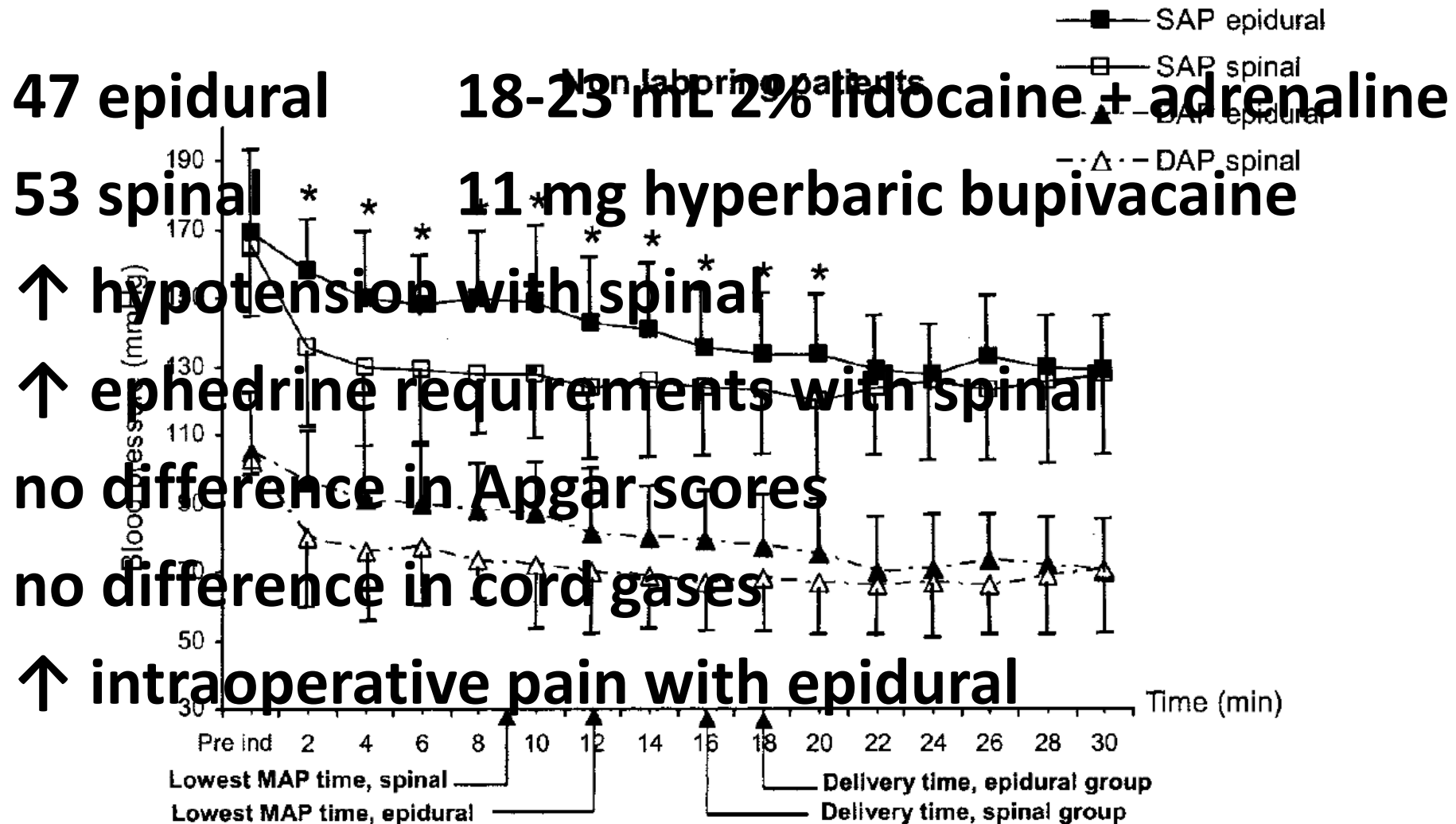


General vs. regional anaesthesia in pre-eclampsia



Wallace et al. Obstet Gynecol 1995

Spinal vs. epidural anaesthesia in pre-eclampsia



CLINICAL INVESTIGATIONS

Anesthesiology 2008; 108:802-11

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Hemodynamic Changes Associated with Spinal Anesthesia for Cesarean Delivery in Severe Preeclampsia

Robert A. Dyer, F.C.A. (S.A.),* Jenna L. Piercy, F.C.A. (S.A.),† Anthony R. Reed, F.R.C.A.,† Carl J. Lombard, Ph.D.,‡ Leann K. Schoeman, F.C.O.G. (S.A.),§ Michael F. James, Ph.D.¶

Background: Hemodynamic responses to spinal anesthesia (SA) for cesarean delivery in patients with severe preeclampsia are poorly understood. This study used a beat-by-beat monitor of cardiac output (CO) to characterize the response to SA. The hypothesis was that CO would decrease from baseline values by less than 20%.

Methods: Fifteen patients with severe preeclampsia consented to an observational study. The monitor employed used pulse wave form analysis to estimate nominal stroke volume. Calibration was by lithium dilution. CO and systemic vascular resistance were derived from the measured stroke volume, heart rate, and mean arterial pressure. In addition, the hemodynamic effects of phenylephrine, the response to delivery and oxytocin, and hemodynamics during recovery from SA were recorded. Hemodynamic values were averaged for defined time intervals before, during, and after SA.

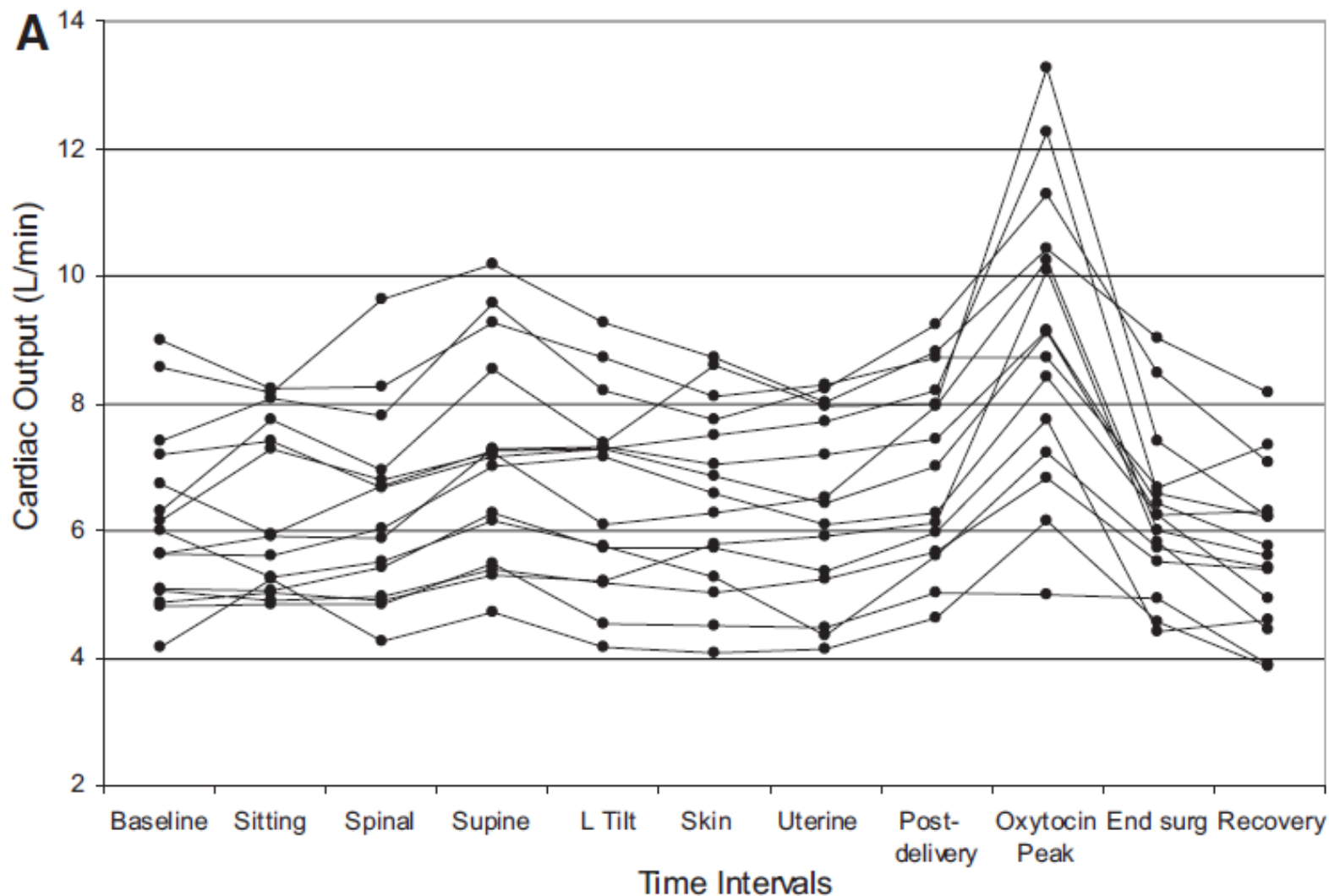
Results: Cardiac output remained stable from induction of SA until the time of request for analgesia. Mean arterial pressure and systemic vascular resistance decreased significantly from the time of adoption of the supine position until the end of surgery. After oxytocin administration, systemic vascular resistance decreased and heart rate and CO increased. Phenylephrine, 50 µg, increased mean arterial pressure to above target values and did not significantly change CO. At the time of recovery from SA, there were no clinically relevant changes from baseline hemodynamic values.

Conclusions: Spinal anesthesia in severe preeclampsia was associated with clinically insignificant changes in CO. Phenylephrine restored mean arterial pressure but did not increase maternal CO. Oxytocin caused transient marked hypotension, tachycardia, and increases in CO.

delivery in severe preeclampsia was published,¹ has spinal anesthesia (SA) been considered an option in this high-risk group of patients. As recently as 1998, an editorial recommended that epidural anesthesia is preferable to SA for cesarean delivery,² even if the patient has not received epidural anesthesia in labor. Many recent studies suggest that SA is safe in the absence of contraindications to regional anesthesia.³⁻⁶ Some studies have shown less hypotension and lower vasopressor requirements than during SA in healthy parturients. One investigation found less hypotension during SA in severe preeclamptics than in preterm women in whom fetal weights were similar.⁴ This eliminated the possibility that the more minor degree of hypotension was due to a lesser degree of aortocaval compression in preeclamptic patients. Nevertheless, hypotension and placental underperfusion remain a risk,⁷ and SA may be associated with more neonatal acidosis than general anesthesia.⁸

Most studies have used heart rate (HR) and blood pressure measurements as surrogate markers of maternal cardiac output (CO). Although pulse and blood pressure measurements are of value in assessing the safety of an anesthetic technique, the true goal of SA for cesarean delivery is to maintain maternal CO and uteroplacental blood flow. In healthy patients, the maximum change in

Spinal anaesthesia & cardiac output



Spinal anaesthesia & vasopressors

OBSTETRIC ANESTHESIA
SECTION EDITOR
DAVID J. BIRNBACH

Pre-eclamptic group

Patients with Severe Preeclampsia Experience Less

- **↓ IV fluid**
Hypotension During Spinal Anesthesia for Elective Cesarean Delivery than Healthy Parturients: A Prospective

- **↑ bupivacaine**
Cohort Comparison

Antoine G. M. Aya, MD, PhD, Roseline Mangin, MD, MSc, Nathalie Vialles, MD,

Jean-Michel Ferrer, MD, Colette Robert, MD, Jacques Ripart, MD, PhD, and

Jean-Emmanuel de La Coussaye, MD, PhD

Division of Anesthesiology, Pain Management, Emergency and Critical Care Medicine, University Hospital, Nîmes, France

- **↓ frequency hypotension**
- **↓ severity hypotension**

- **↓ vasoconstrictor requirement**

In this prospective cohort study, we compared the incidence and severity of spinal anesthesia (SA)-associated hypotension in severely preeclamptic (n = 20) versus healthy (n = 30) parturients undergoing cesarean delivery. After the administration of IV fluids, SA was performed with hyperbaric 0.5% bupivacaine, sufentanil, and morphine. Blood pressure (BP) was recorded before and at 2-min intervals for 30 min after SA. Clinically significant hypotension was defined as the need for ephedrine (systolic BP decrease to <100 mm Hg in healthy parturients or 30% decrease in mean BP in both

groups). Despite receiving a smaller fluid volume (1653 ± 331 mL versus 1895 ± 150 mL; $P = 0.005$) and a larger bupivacaine dose (2.5 ± 0.9 mg versus 1.2 ± 0.7 mg; $P = 0.019$), the severely preeclamptic patients had a less frequent incidence of clinically significant hypotension (16.6% versus 53.3%; $P = 0.006$), which was less severe and required less ephedrine. The risk of hypotension was almost six times less in severely preeclamptic patients (odds ratio, 0.17; 95% confidence interval, 0.05–0.58; $P = 0.006$) than that in healthy patients.

(Anesth Analg 2003;97:867–72)



Pre-eclampsia & HELLP syndrome

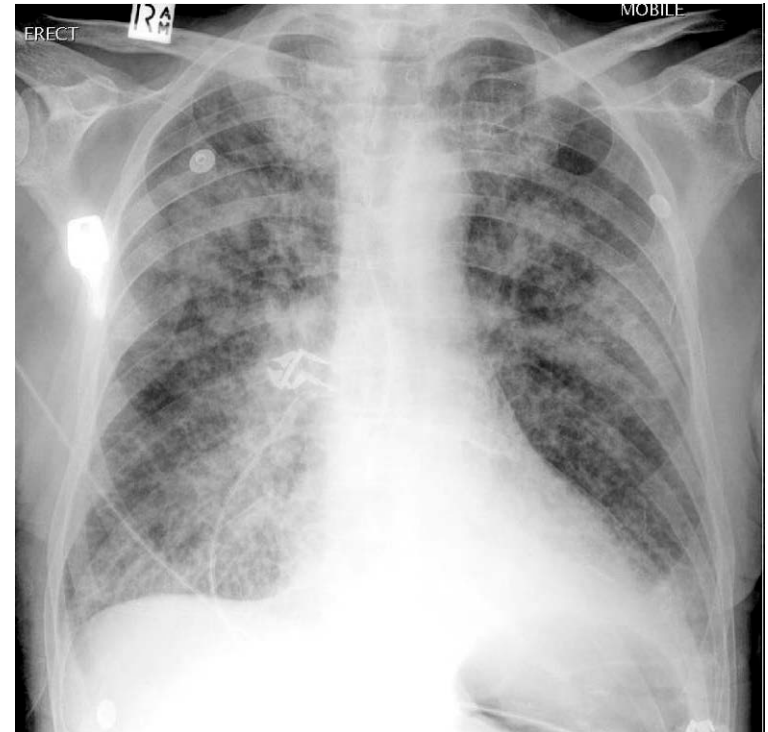
- Antenatal Issues
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Pulmonary oedema

- pulmonary capillary leak
- ↓ colloid osmotic pressure
- left ventricular dysfunction
- excess iv fluid administration
- inadequate monitoring





Indications for increased monitoring

- **severe refractory hypertension**
- **pulmonary oedema**
- **oliguria unresponsive to fluid therapy**
- **anaesthesia**
- **haemorrhage**



Cardiac Output Monitoring

- **Pulse waveform analysis**

LiDCO[®]

PiCCO[®]

FloTrac-Vigileo[®]

- **Ultrasound**

Oesophageal Doppler

Suprasternal Doppler

Transthoracic echocardiography

Oesophageal echocardiography

- **Electrical resistance**

Bioimpedence

Bioreactance

Electrical velocimetry

- **Others**

Modelflow

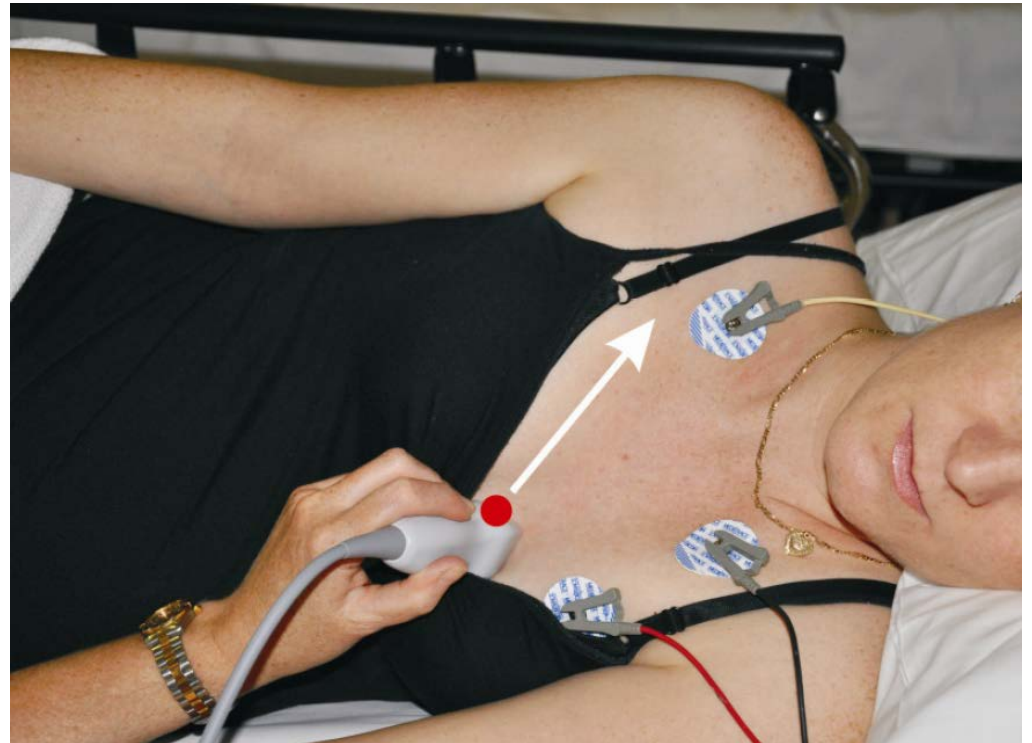
Pleth variability index

Dye densitometry

Pulmonary artery catheter

Cardiac Output Monitoring

- Accuracy & precision
- Applicability
- Dynamic vs. static
- Cost
- Training
- Complications





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