

Oxygen Monitoring: Brain vs Fingertip (or Tissue)?

(Narcotic-Induced Respiratory Depression)

John M Murkin MD, FRCPC, FAI (hon)
Dept of Anesthesiology and Perioperative Medicine
Schulich School of Medicine
University of Western Ontario
London, Ontario
Canada

Acknowledgement/Disclosures:



Excellence in Safety Research Symposium
Honolulu, Hawaii
Sunday, March 21, 2010
12:45 - 4:45 PM

Welcome & Overview: Moderator Sorin Brull, MD

Innovation & Introductions of Featured Speakers
Dr. David B. Goodale

1. Excellence in Education for Patient Safety

Hospital Transparency after Medical Errors
Timothy McDonald, MD JD

2. Excellence in Anesthesia Management

Lipid Bolus for Local Anesthetic Toxicity
Guy Weinberg, MD

Hawaiian Break with Experts

3. Excellence in Device Innovations: Cerebral Oximetry

Optimizing Perfusion Improves Outcomes in Cardiac Surgery
John Murkin, MD

Critical Anesthetic Variables in Beach Chair Orthopedic Position
Meg A. Rosenblatt, MD

4. Excellent Preclinical Studies: Implications for Future Safety Initiatives
Anesthetics & Alzheimer's: Something to remember or forget?
Gregory Crosby, MD

Summary & Conclusions: Moderator Sorin Brull, MD

No relevant commercial affiliations

PSI funded grant 84-41:

(Tissue NIRS in assessment and management of critically ill patients)

WAIKIKI BEACH

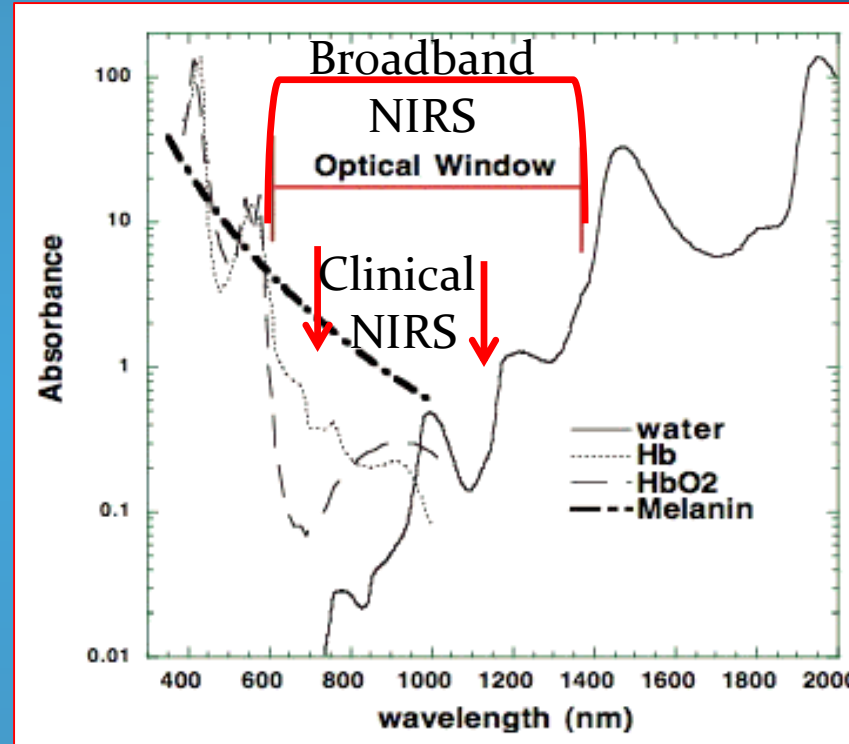
NIR Spectroscopy

Water highly absorptive except within range 600-1350 nm “optical window”

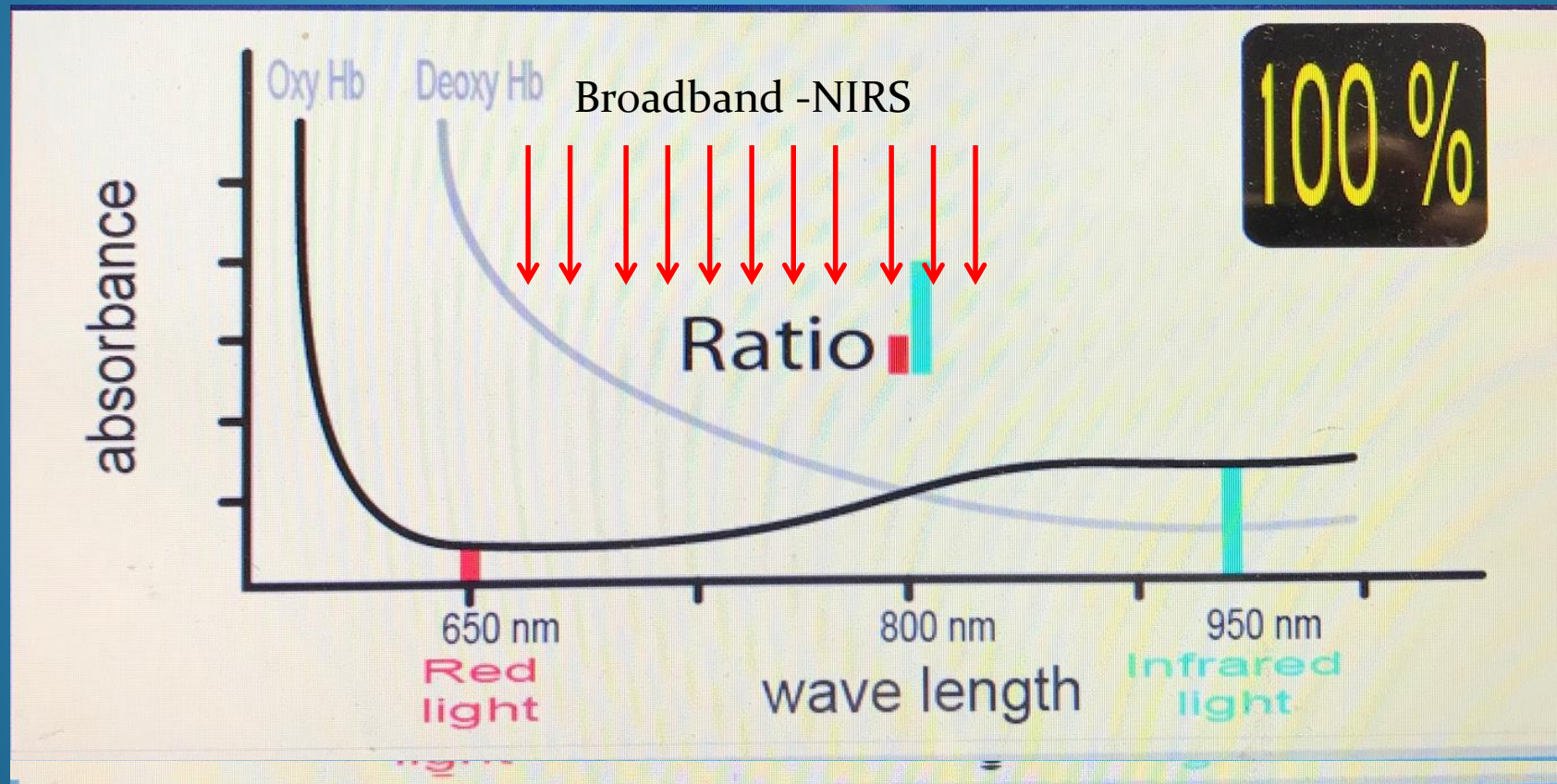
Facilitates measurement of key species of HbO₂ and oxidative metabolism

Conventional NIRS: 2-4 wavelengths

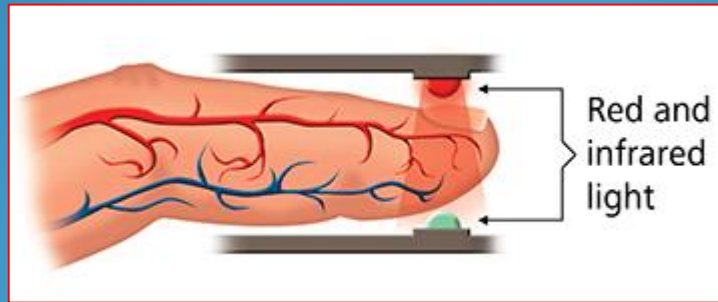
Broadband NIRS: multi wavelengths



NIRS hemoglobin O₂ saturation: differential absorption of various wavelengths between oxygenated and deoxygenated Hemoglobin



Pulse oximetry: O₂ substrate delivery



Concerns:

malposition
motion artifact
peripheral v/c (cold, ischemia)
ambient light

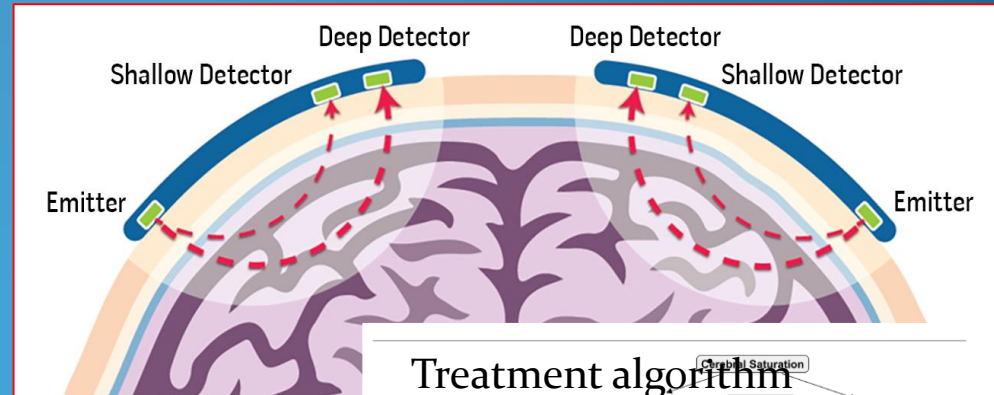


'alarm fatigue'

'Change' in absorptance = arterial saturation

Tissue oximetry: delivery/metabolism

Cerebral NIRS



Sensitive to global ischemic events
Amenable to physiologic interventions

BUT

Variable extracerebral contamination (5-15%)

Analysis algorithm assumes fixed arterial/venous partitioning (30%/70% - change with ischemia/ PaCO_2)

Measures very small sample frontal cortex (1cc)

High-Risk Cardiac Surgery

...ture, MD,* Antoine Rochon, MD,*

Assessing Regional ation Frequency During and Responsiveness to an

...n, BS, †† Maria Fritock, MD, § Rebecca Y. Klinger, MD, ||
...ie Huffmyer, MD, ** Michelle Parish, BSN, ††

Gayane Yenokyan, PhD, †† and Charles W. Hogue, MD ††

Cerebral NIRS: new developments-

Photo-acoustic coupling

Intraoperative Cerebral Autoregulation Assessment Using Ultrasound-Tagged Near-Infrared-Based Cerebral Blood Flow in Comparison to Transcranial Doppler Cerebral Flow Velocity: A Pilot Study

John M. Murkin, MD, FRCPC,* Moshe Kamar, MD,† Zmira Silman, MSc,† Michal Balberg, PhD,† and Sandra J. Adams, RN*

Objective: This was a pilot study comparing the ability of a new ultrasound-tagged near-infrared (UT-NIR) device to detect cerebral autoregulation (CA) in comparison to transcranial Doppler (TCD).

Design: An unblinded, prospective, clinical feasibility study.

Setting: Tertiary-care university hospital cardiac surgical operating rooms.

Participants: Twenty adult patients undergoing cardiac surgery with cardiopulmonary bypass (CPB).

Interventions: There were no clinical interventions based on study monitoring devices, but a continuous correlation analysis of digital data from transcranial Doppler (TCD) velocity was compared with a novel UT-NIR device and correlation analysis of change signals versus mean arterial pressure was performed in order to detect presence or absence of intact CA and for determination of the lower limit of cerebral autoregulation during CPB.

Measurements and Main Results: Similar and highly significant concordance ($\kappa = 1.00$; $p < 0.001$) was demonstrated between the 2 methodologies for determination of CA, indicating good correlation between the 2 methodologies. Intact CA was absent in 2 patients during CPB, and both devices were able to detect this.

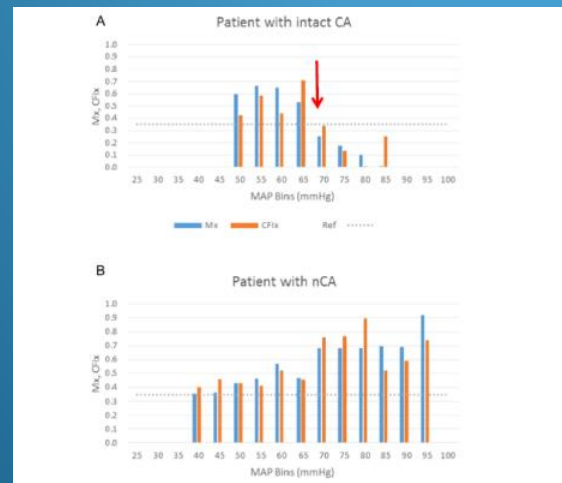
Conclusions: To the authors' knowledge this is the first clinical report of a UT-NIR device that shows promise as a clinically useful modality for detection of CA and the lower limit of cerebral autoregulation. The utility of UT-NIR was demonstrated further during times at which extensive usage of electrocautery or functional absence of the transcranial window rendered TCD uninterpretable.

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KEY WORDS: cerebral blood flow, CBF, transcranial Doppler, TCD, ultrasound-tagged near-infrared device, UT-NIR, cardiopulmonary bypass, CPB, cerebral autoregulation



U/S focus beam
'tags' photons at depth
Discriminates deep cerebral tissue

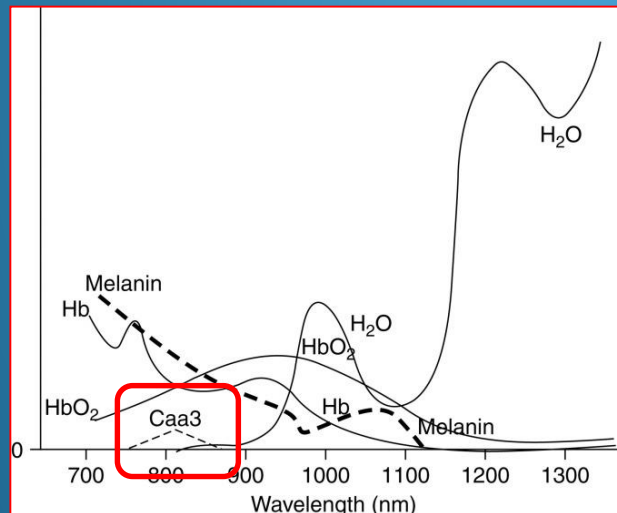


“UT-NIRS detects presence/impairment of cerebral autoregulation”

Broadband-NIRS:

Cytochromeaa3 - measurement of energy substrates

Continuous monitoring
CBF and CMRO₂
Detect onset ischemia



Development of a combined broadband near-infrared and diffusion correlation system for monitoring cerebral blood flow and oxidative metabolism in preterm infants

Mamadou Diop,^{1,2} Jessica Kishimoto,^{1,2} Vladislav Toronov,³ David S. C. Lee,⁴ and Keith St. Lawrence^{1,2,*}

¹Department of Medical Biophysics, University of Western Ontario, London, ON, Canada

²Imaging Division, Lawson Health Research Institute, London, ON, Canada

³Department of Physics, Ryerson University, Toronto, ON, Canada

⁴Department of Neonatology, London Health Sciences Centre, London, ON, Canada

*kstlaw@lawsonimaging.ca

Abstract: Neonatal neuromonitoring is a major clinical focus of near-infrared spectroscopy (NIRS) and there is an increasing interest in measuring cerebral blood flow (CBF) and oxidative metabolism (CMRO₂) in addition to the classic tissue oxygenation saturation (StO₂). The purpose of this study was to assess the ability of broadband NIRS combined with diffusion correlation spectroscopy (DCS) to measure changes in StO₂, CBF and CMRO₂ in preterm infants undergoing pharmaceutical treatment of patent ductus arteriosus. CBF was measured by both DCS and contrast-enhanced NIRS for comparison. No significant difference in the treatment-induced CBF decrease was found between DCS (27.9 ± 2.2%) and NIRS (26.5 ± 4.3%). A reduction in StO₂ (70.5 ± 2.4% to 63.7 ± 2.9%) was measured by broadband NIRS, reflecting the increase in oxygen extraction required to maintain CMRO₂. This study demonstrates the applicability of broadband NIRS combined with DCS for neuromonitoring in this patient population.

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OCIS codes: (170.3660) Light propagation in tissues; (170.3880) Medical and biological imaging; (170.6510) Spectroscopy, tissue diagnostics.

Tissue oximetry

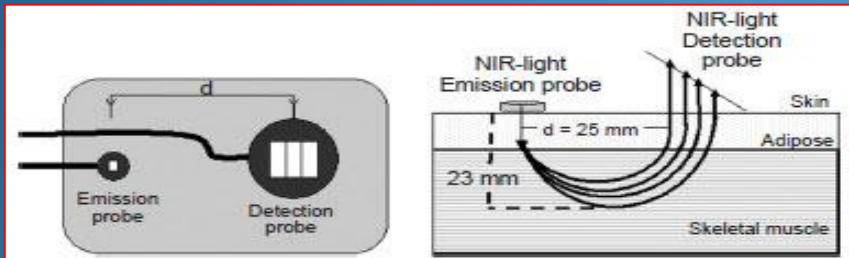


Figure 1- Diagram of a distal tip of the NIRS optical cable (A). With 25 mm spacing (d) between emission and detection probes, approximately 95% of the detected optical signal is from 23 mm of tissue penetration (B). Note the curved shape of the light path (banana shape).



Journal of Critical Care 30 (2015) 315–320

Contents lists available at ScienceDirect

Journal of Critical Care

journal homepage: www.jccjournal.org

Soft tissue oxygenation and risk of mortality (STORM): An early marker of critical illness? 

Biniam Kidane, MD, MSc^{a,b,c}, Sami A. Chadi, MD, MSc^{a,b,c}, Anthony Di Labio, MD^{a,b}, Fran Priestap, MSc^d, Wael Haddara, MD^{d,e}, Tina Mele, MD, PhD^{a,d,e,*}, John M. Murkin, MD^{f,g}

^a Western University, Schulich School of Medicine & Dentistry, General Surgery, London, Canada
^b London Health Sciences Centre, General Surgery, London, Canada
^c McMaster University, Clinical Epidemiology & Biostatistics, Hamilton, Canada
^d London Health Sciences Centre, Critical Care Medicine, London, Canada
^e Western University, Schulich School of Medicine & Dentistry, Critical Care Medicine, London, Canada
^f Western University, Schulich School of Medicine & Dentistry, Department of Anesthesiology, London, Canada
^g Western University, Schulich School of Medicine & Dentistry, Department of Critical Care Medicine, London, Canada

Tissue oxygen saturation did not predict ICU admission but was the only independent predictor of mortality (adjusted odds ratio, 1.06; 95% confidence interval, 1.01-1.12; $P = .04$).

Conclusions: Tissue oxygen saturation may identify critical illness in patients who would not traditionally meet ICU admission criteria and thus may identify patients who benefit from closer monitoring. *Journal of Critical Care* 30 (2015) 315–320

Initial Soft Tissue Oxygenation is Associated with Prolonged ICU Admission

Of 126 consecutive consenting adult patients admitted to ICU:

Primary Diagnosis

| | |
|-----------------------------|-----------------|
| cardiogenic shock (n= 31) | mortality 9/31 |
| hemorrhagic shock (n= 5) | mortality 1/5 |
| neurogenic shock (n=1) | mortality 0 |
| respiratory failure (n= 44) | mortality 10/44 |
| septic shock (n= 45) | mortality 19/45 |



- Initial ICU StO₂ strongly correlated with prolonged ICU admission (>3 d)
- Serial StO₂ trend with prolonged hospitalization (>10d)

Narcotic-Induced Respiratory Depression

Oxygen supplementation

Conversion tables

1 Estimating PaO₂ from a given SO₂

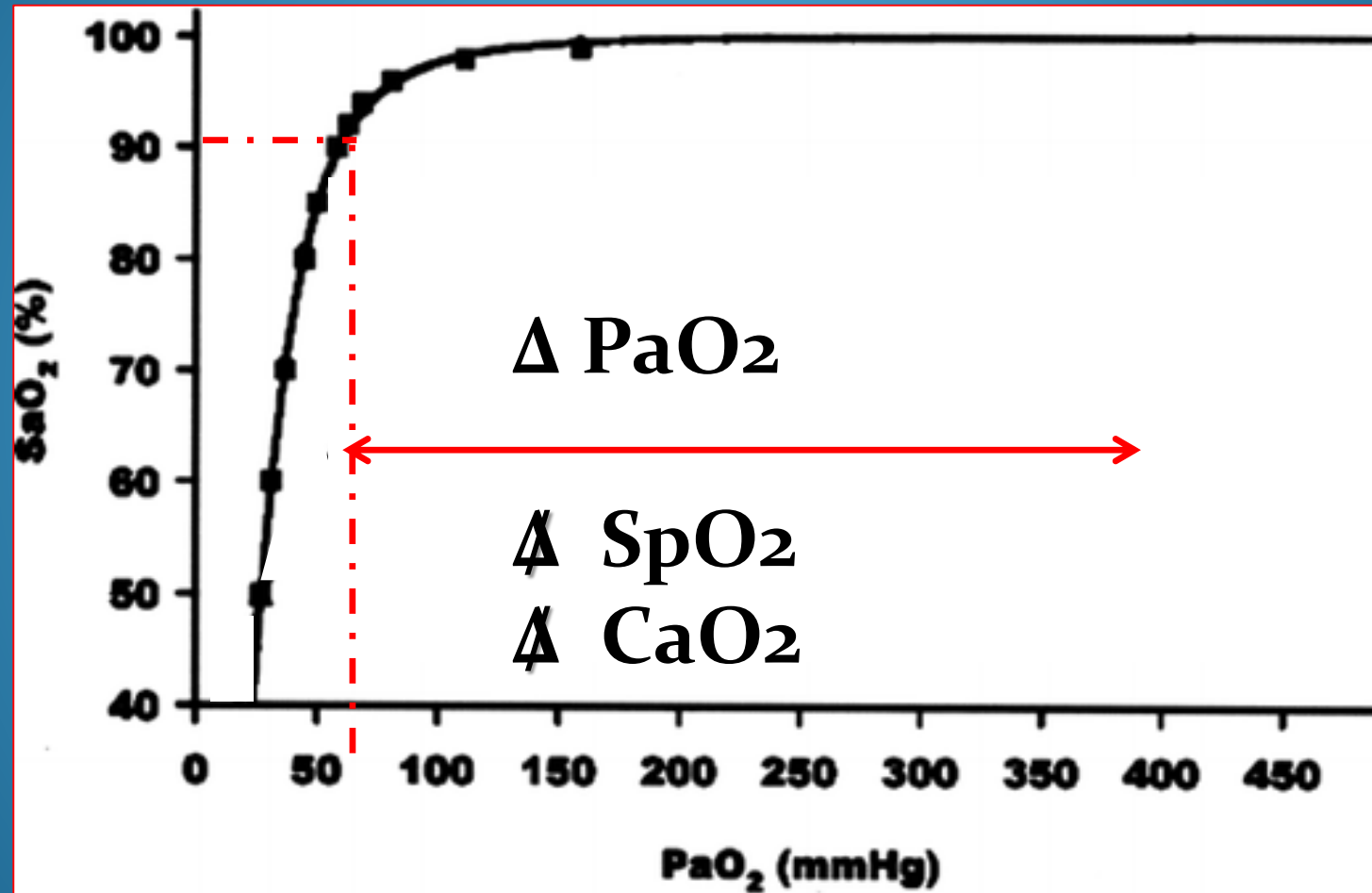
| SO ₂ (%) | PaO ₂ (mmHg) |
|---------------------|-------------------------|
| 80 | 44 |
| 81 | 45 |
| 82 | 46 |
| 83 | 47 |
| 84 | 49 |
| 85 | 50 |
| 86 | 52 |
| 87 | 53 |
| 88 | 55 |
| 89 | 57 |
| 90 | 60 |
| 91 | 62 |
| 92 | 65 |
| 93 | 69 |
| 94 | 73 |
| 95 | 79 |
| 96 | 86 |

2 Estimating FiO₂

| Method | O ₂ flow (l/min) | Estimated FIO ₂ (%) |
|---------------|-----------------------------|--------------------------------|
| Nasal cannula | 1 | 24 |
| | 2 | 28 |
| | 3 | 32 |
| | 4 | 36 |

| | | |
|--------------------------|-----|----|
| | 6-7 | 50 |
| | 7-8 | 60 |
| Face mask with reservoir | 6 | 60 |
| | 7 | 70 |
| | 8 | 80 |
| | 9 | 90 |
| | 10 | 95 |

Decrease in SpO₂ is a late indicator of hypoxemia



Since ScO₂ preserved:

SpO₂, ScO₂, StO₂ all decrease late

Oxygen supplementation

British Journal of Anaesthesia 110 (5): 837-41 (2013)
Advance Access publication 4 January 2013 - doi:10.1093/bja/aes494

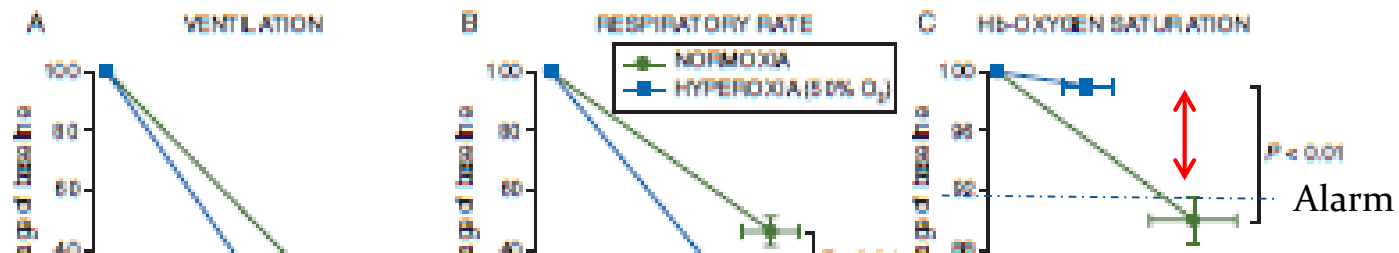
20 healthy volunteers BJA

RESPIRATION AND THE AIRWAY

50 ug remifentanyl

FiO₂ 0.21 vs 0.5

High-inspired oxygen concentration further impairs



O₂ supplementation can delay SpO₂ desaturation by several minutes

Hyperoxia decreases peripheral (~80%) and central (~20%) chemoceptor activity (exacerbates apnea since decrease CO₂ responsivity)

PaCO₂ increases 3-4 mmHg/min ($\Delta\text{PaCO}_2 = 10 \rightarrow \Delta\text{pH} = 0.08$)

4 min \rightarrow pH \approx \downarrow 7.28

Acidosis, non-invasive ventilation and mortality in hospitalised COPD exacerbations

C M Roberts,^{1,2} R A Stone,^{1,3} R J Buckingham,¹ N A Pursey,¹ D Lowe,¹ On behalf of the National Chronic Obstructive Pulmonary Disease Resources and Outcomes Project (NCROP) implementation group *Thorax* 2011;**66**:43–48. doi:10.1136/thx.2010.153114

Table 4 Mortality

| | Inpatient mortality: | | | | Fishers Exact test (NIV versus not NIV) |
|-----------------------|------------------------------------|----------|--------------------------------|----------|-----------------------------------------|
| | For ALL Patients not receiving NIV | | For ALL Patients receiving NIV | | |
| | % | N | % | N | |
| ALL PATIENTS | 5 | 475/8639 | 25 | 270/1077 | <0.001 |
| Acidotic patients | 14 | 165/1174 | 26 | 249/969 | <0.001 |
| Non-acidotic patients | 4 | 246/5994 | 10 | 8/78 | 0.02 |

Narcotic-Induced Respiratory Depression

?

OPEN

REVIEW ARTICLE**Improving detection of patient deterioration in the general hospital ward environment**

Jean-Louis Vincent, Sharon Einav, Rupert Pearse, Samir Jaber, Peter Kranke, Frank J. Overdyk, David K. Whitaker, Federico Gordo, Albert Dahan and Andreas Hoeft

Patient monitoring on low acuity general hospital wards is currently based largely on intermittent observations and

appropriate management, thereby reducing the need for higher acuity care, reducing hospital lengths of stay and

costs and even, at times, improving survival. This degree of monitoring has thus far been commonly inappropriate for general hospital ward settings due to device costs and the need for staff expertise in interpretation. In this review, we discuss some developments to improve patient monitoring and thus detection of deterioration in low acuity general hospital wards.

Online 22 February 2018

Integrated multimodality monitoring:

Ventilation: capnography, impedance plethysmography

Oxygenation: SpO₂, ScO₂,

Hemodynamics: HR,