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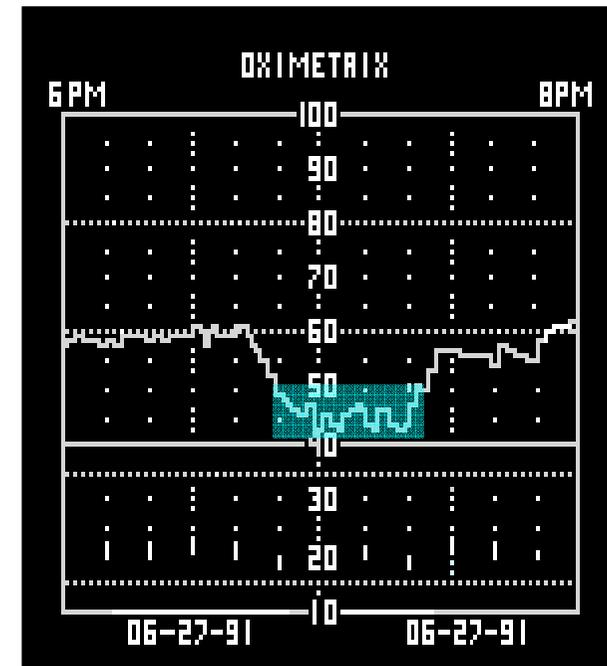
brain tissue PO_2 ($P_{br}O_2$) monitoring in
traumatic brain injury: past & future

why do we measure brain tissue PO_2
($P_{br}O_2$) ?

SjvO₂, desaturation episode

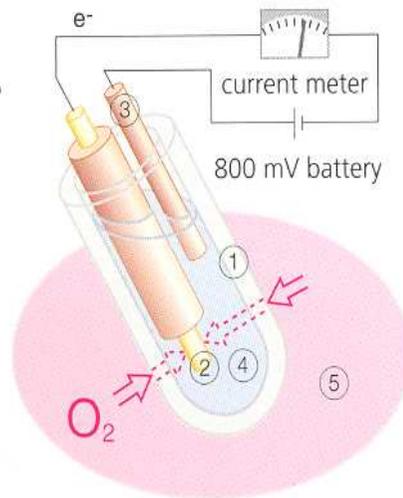
- Def.: SjvO₂ < 50%, t > 10 min

- increased morbidity & mortality
 - multiple DE: 90% „unfavourable outcome“
 - single DE: 74% „unfavourable outcome“
 - no DE: 55% „unfavourable outcome“



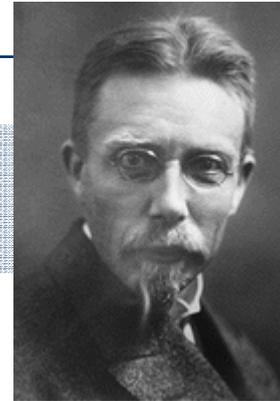
PbrO₂, Clark-type electrode

- 1 Polyethylene tube diffusion membrane
- 2 Polarographic Gold cathode
- 3 Polarographic Silver anode
- 4 Electrolyte filled cell
- 5 Tissue

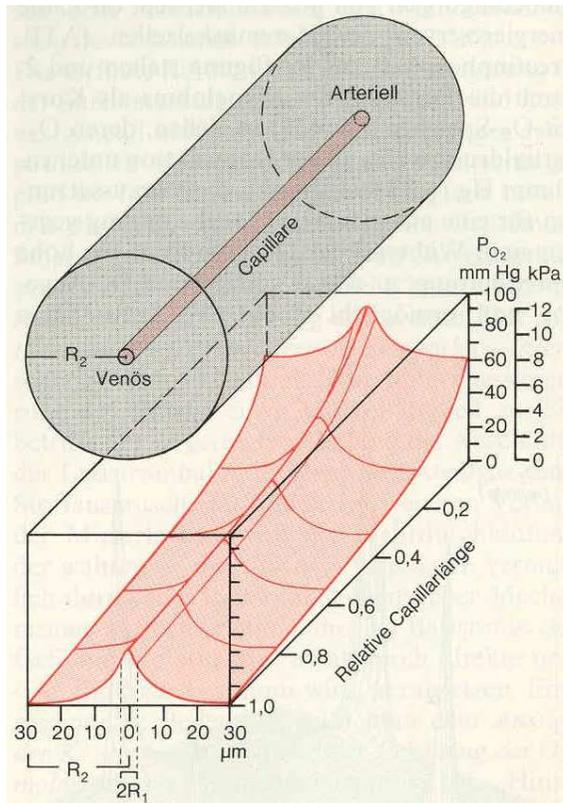




PO₂ in the Krogh cylinder



August Krogh (1874 – 1949)
Nobel Prize 1920



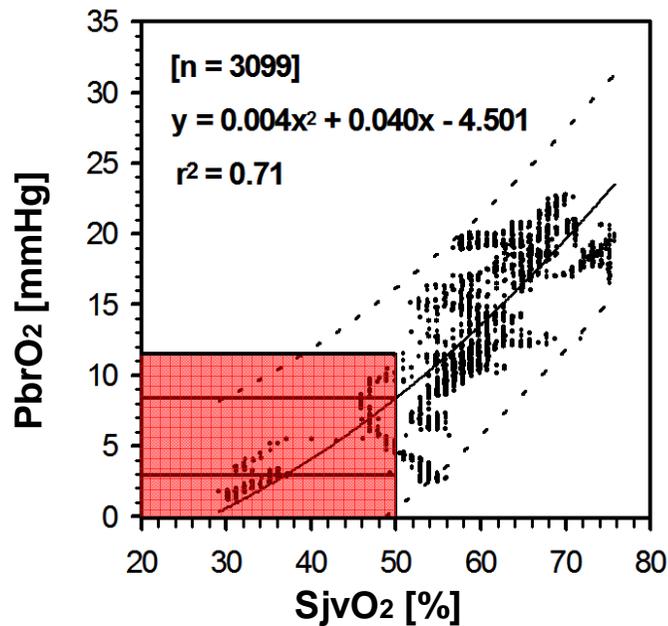
■ PO₂...
...determines O₂ diffusion distance

■ arterial O₂ content...
...is not determined by PO₂ but
...Hb and SaO₂

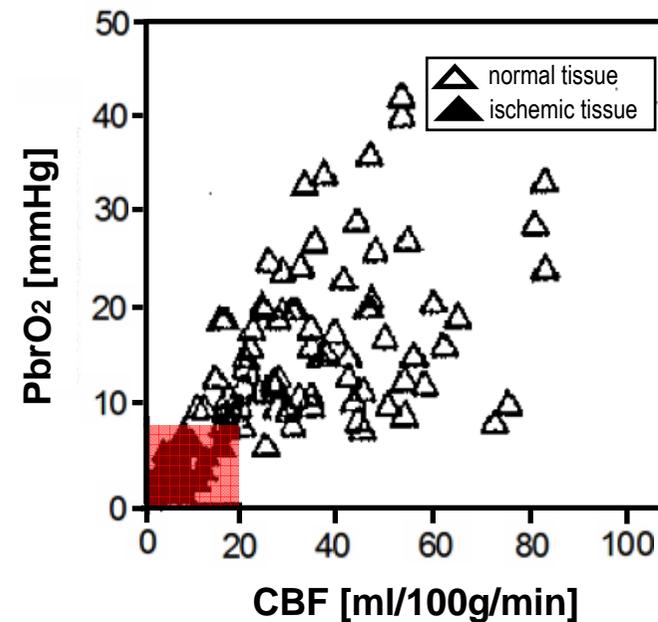
	PaO ₂ [mmHg]	PcapO ₂ [mmHg]	O ₂ diffusion distance [μm]
RA (FiO ₂ = 0,21)	100	90 - 30	64 - 36
NBHO (FiO ₂ = 1)	400	280 - 120	90 - 66
HBHO (3 bar)	2000	1400 - 600	247 - 135



critical threshold of PbrO₂ – S_{iv}O₂ & CBF



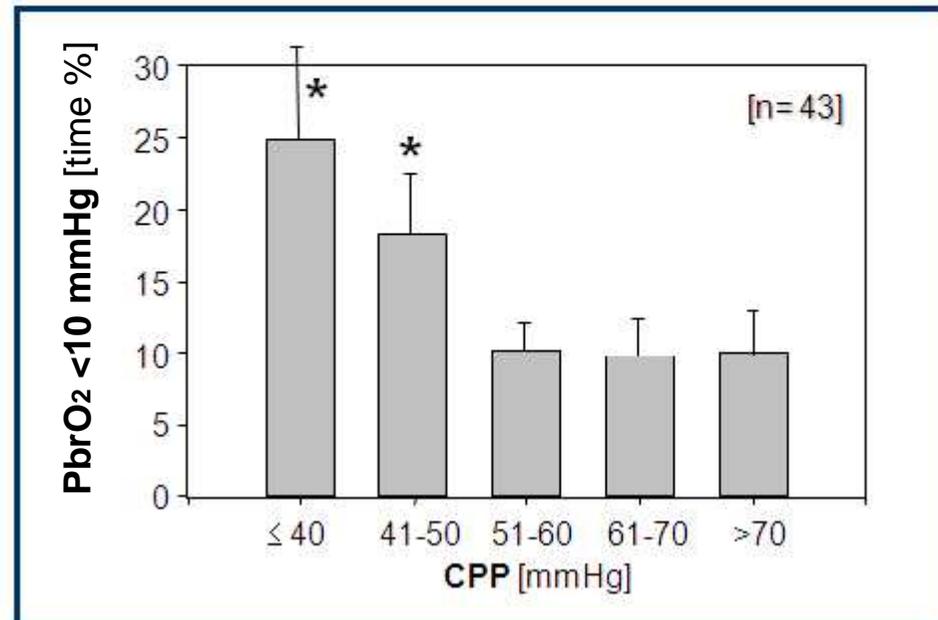
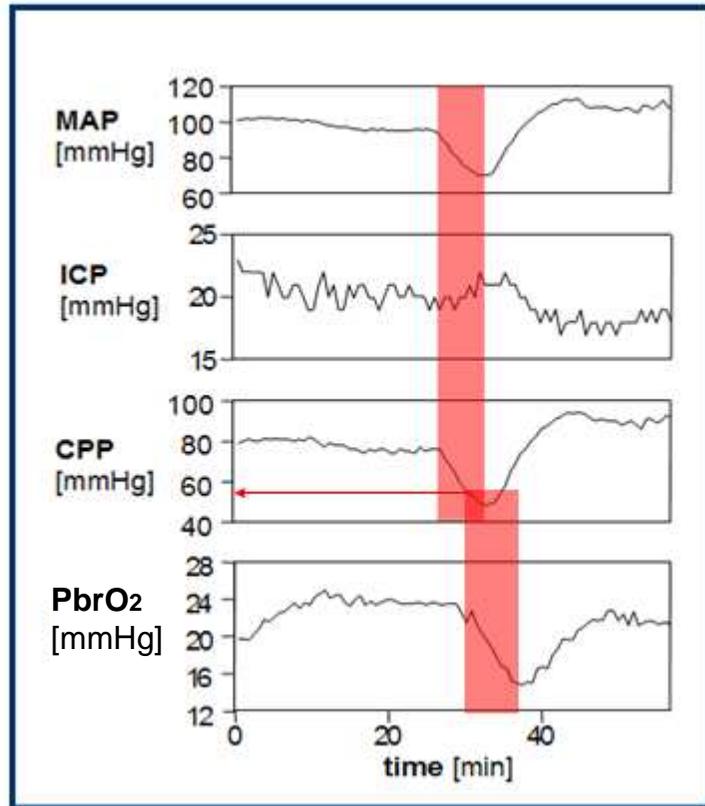
- TBI
- clinical data
- critical P_tiO₂ = 8.5 mmHg



- ischemia (MCA occlusion)
- experimental data (cats)
- critical P_tiO₂ = 8 mmHg



PbrO₂ vs. CPP



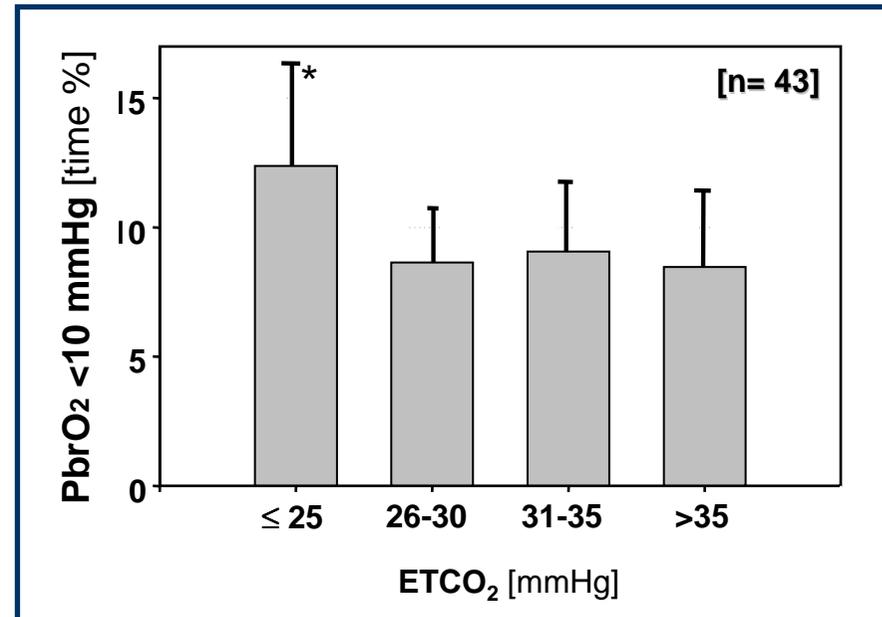
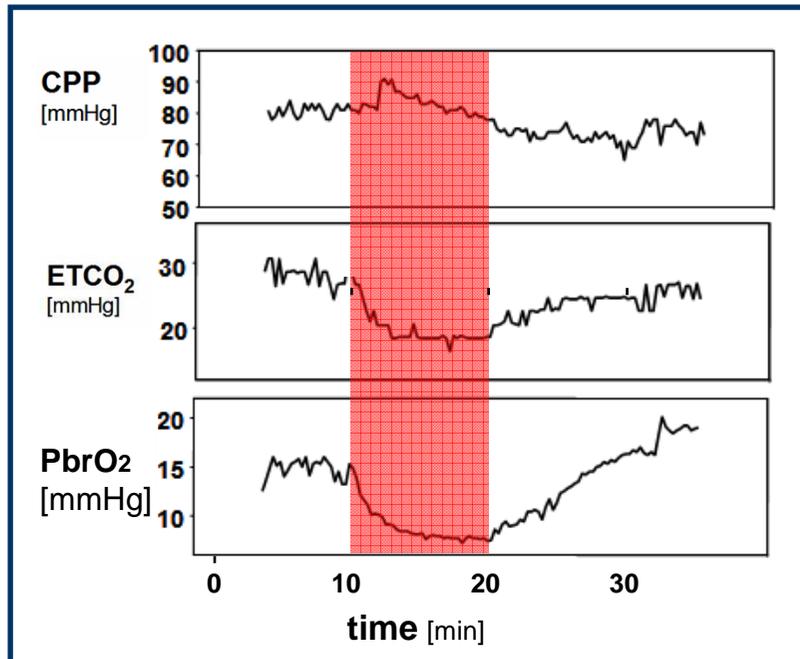
Kiening, *J Neurosurg*, 1996

Kiening, *Neurol Res*, 1997

Bardt, *Acta Neurochir*, 1998



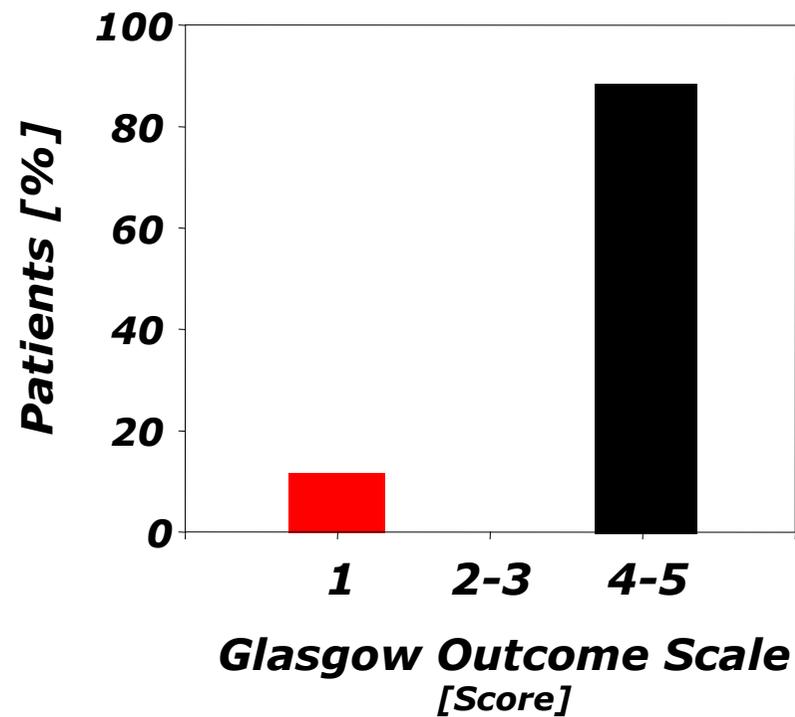
PbrO₂ vs. PCO₂



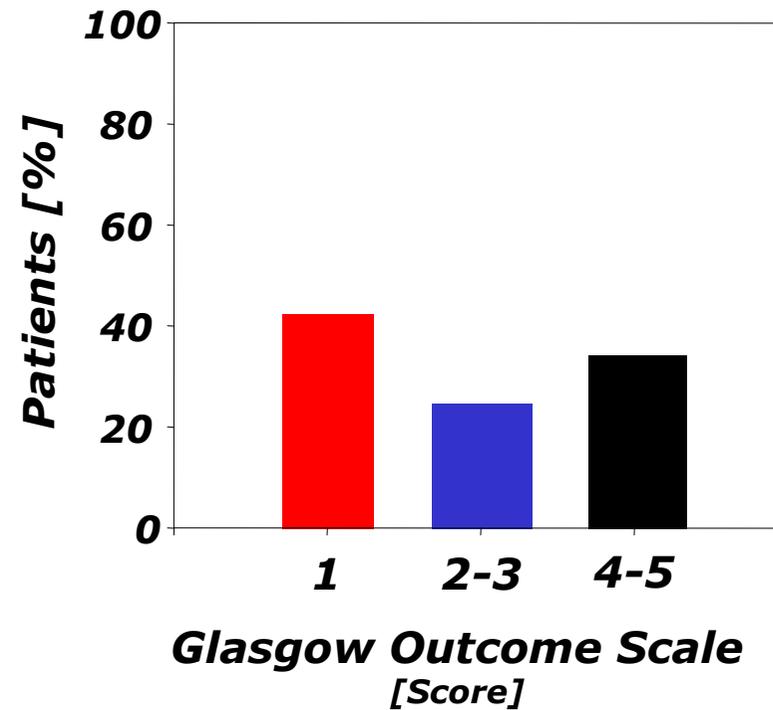


PbrO₂ impact on outcome

**PbrO₂ < 10 mmHg
< 30 Min [n=20]**



**PbrO₂ < 10 mmHg
> 30 Min [n=23]**



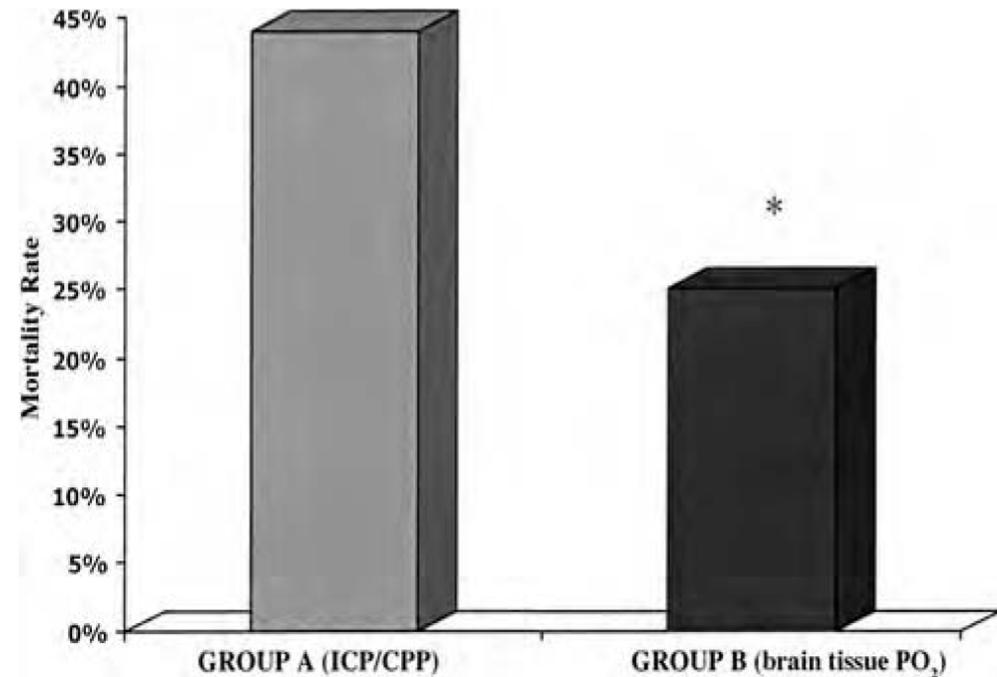
PbrO₂ based therapy

group A:

- $n = 25$
- $ICP < 20 \text{ mmHg} / CPP > 60 \text{ mmHg}$
- „Guidelines“

group B:

- $n = 28$
- + $PbrO_2 \geq 25 \text{ mmHg}$
- $FiO_2 \uparrow$ / decompressive surgery

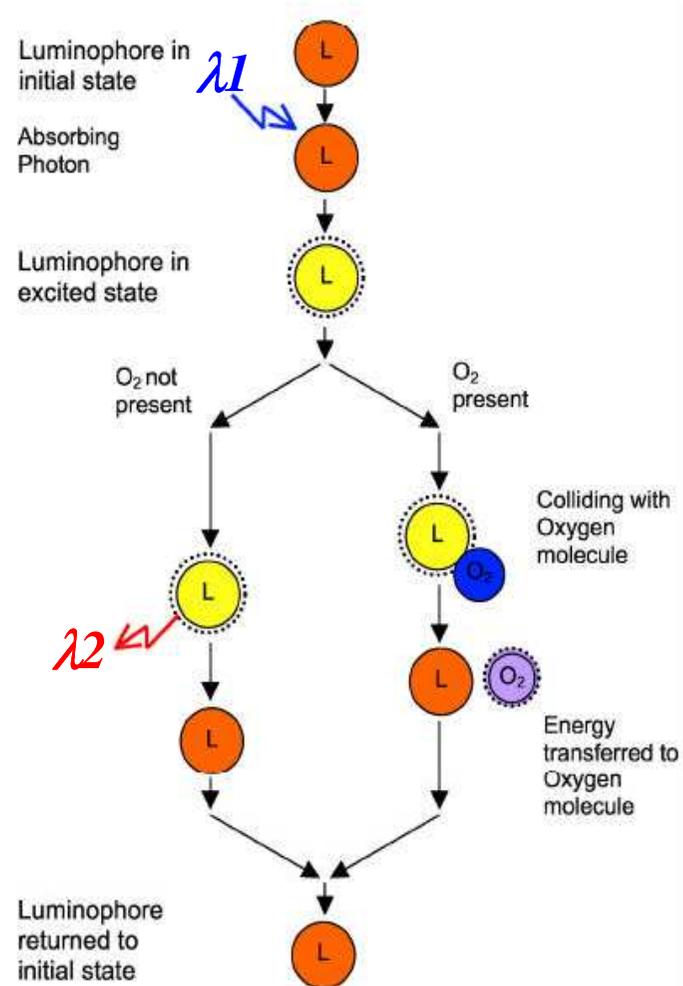




PbrO₂ measurement by oxygen quenching Neurovent

LICOX features

storage	fridge
consumes oxygen	yes
consumes electrolytes	yes
long-term monitoring	restricted



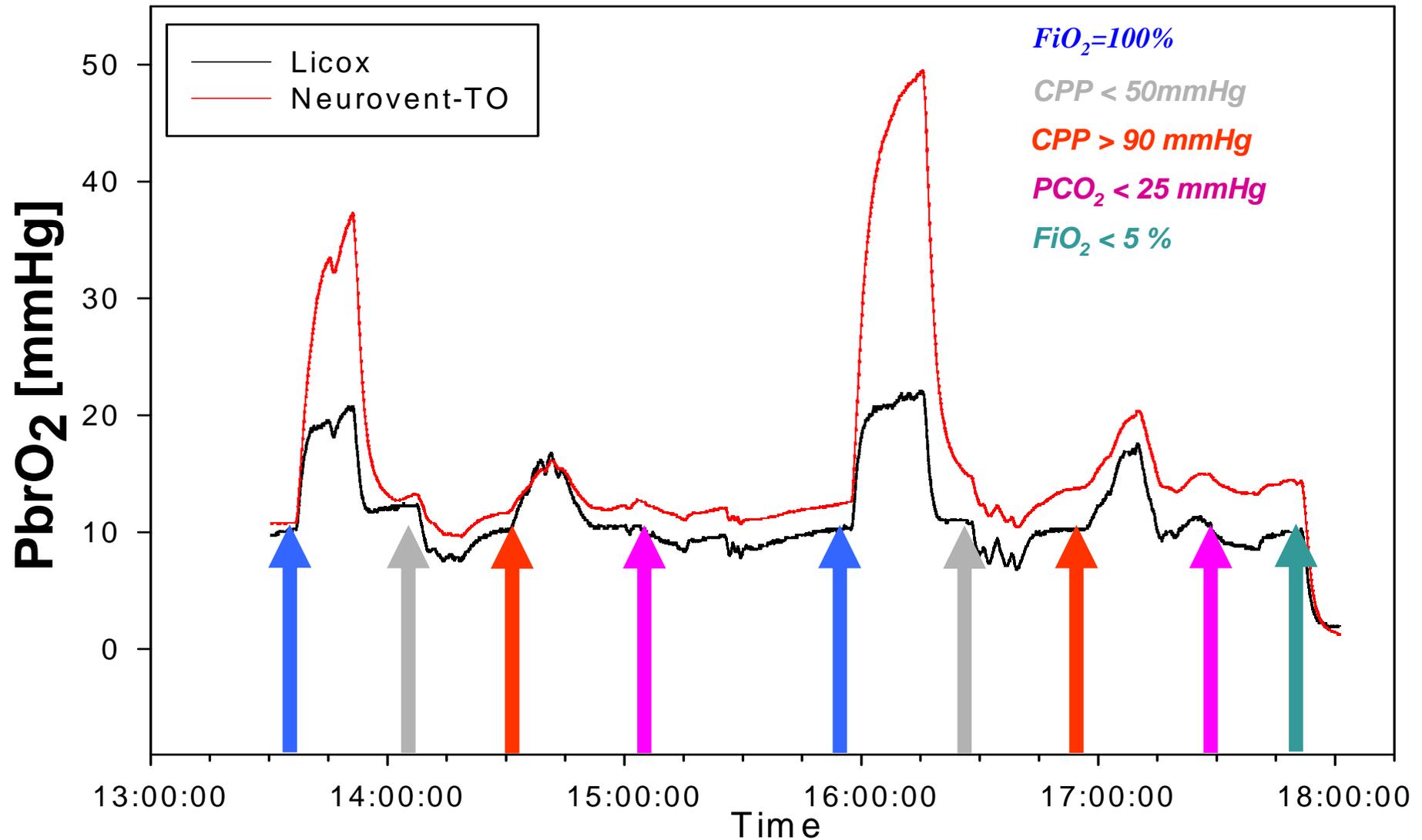


LICOX vs. Neurovent-TO / -PTO

	<i>LICOX</i>	<i>Neurovent-TO</i>	<i>Neurovent-PTO</i>
storage	fridge	room air	room air
consumes oxygen	yes	no	no
consumes electrolytes	yes	no	no
long-term monitoring	restricted	yes	yes
in-built thermocouple	no	yes	yes
in-built ICP catheter	no	no	yes
diameter	1.5 F / 0,49 mm	3 Fr / 1 mm	5 Fr / 1.65 mm
O₂ uptake area	13 mm ²	13 mm ²	22 mm ²
T_{90/35°C}	70 s	< 200 s	<200 s

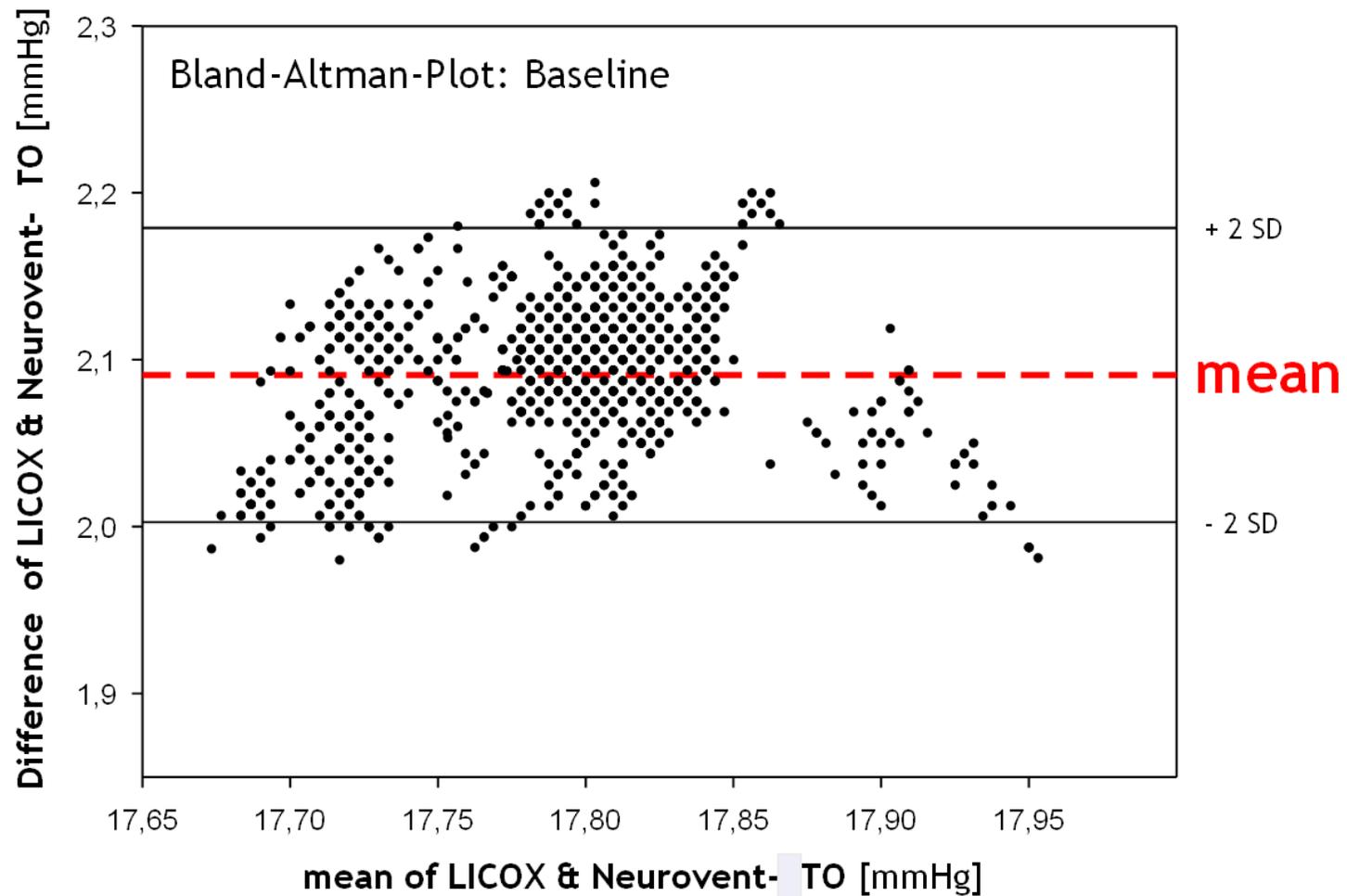


LICOX vs. Neurovent-TO (swine model; n = 9) - Protocol



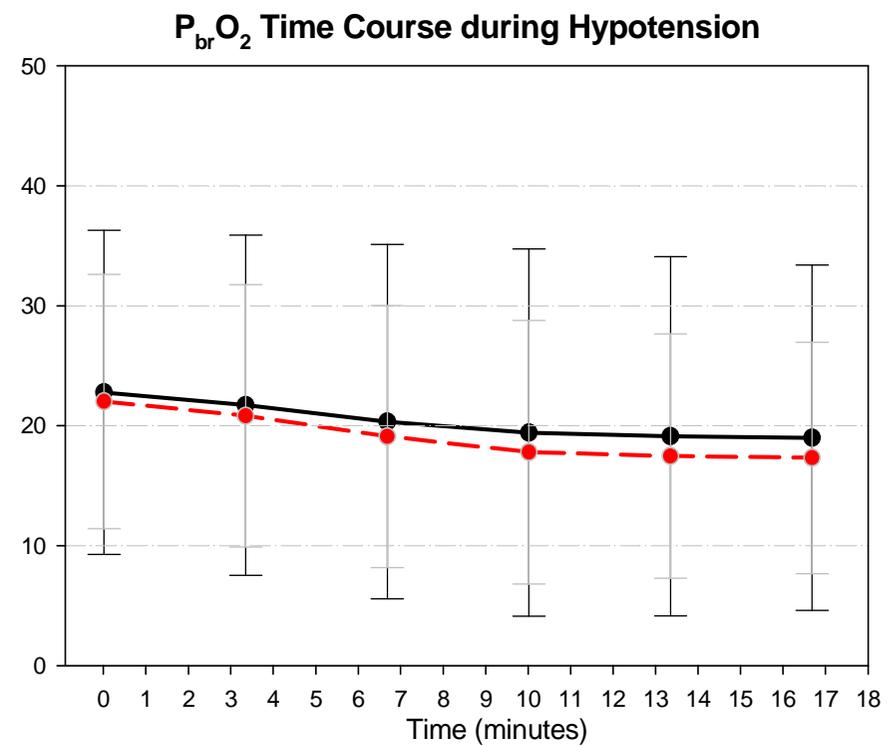
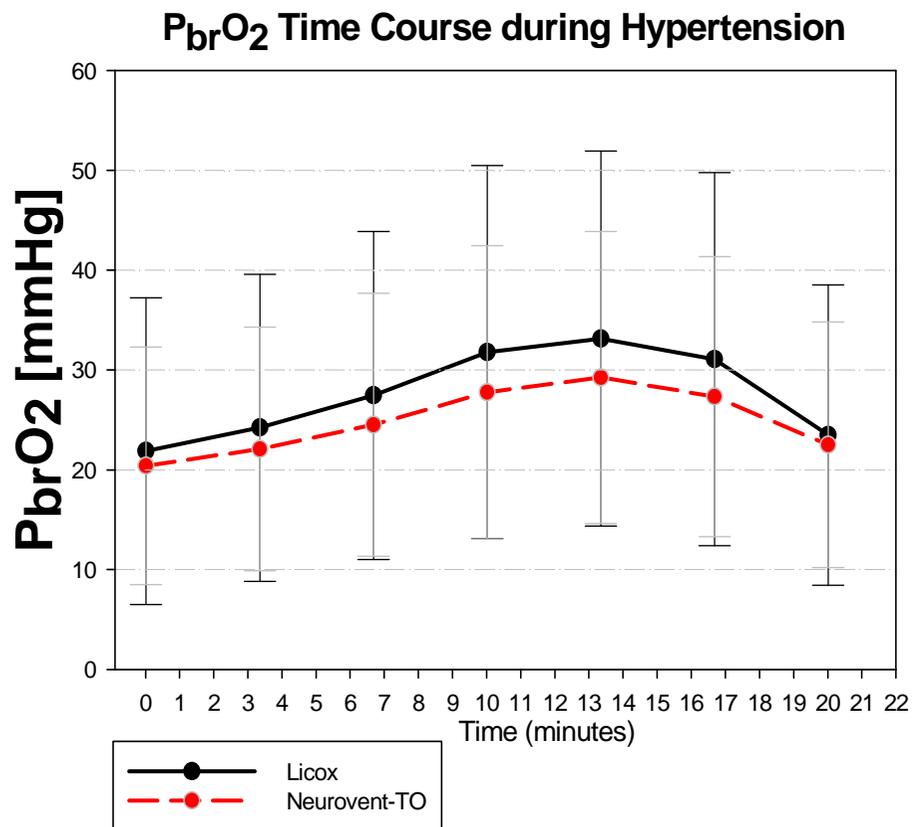


LICOX vs. Neurovent-TO (swine model; n = 9) – Baseline





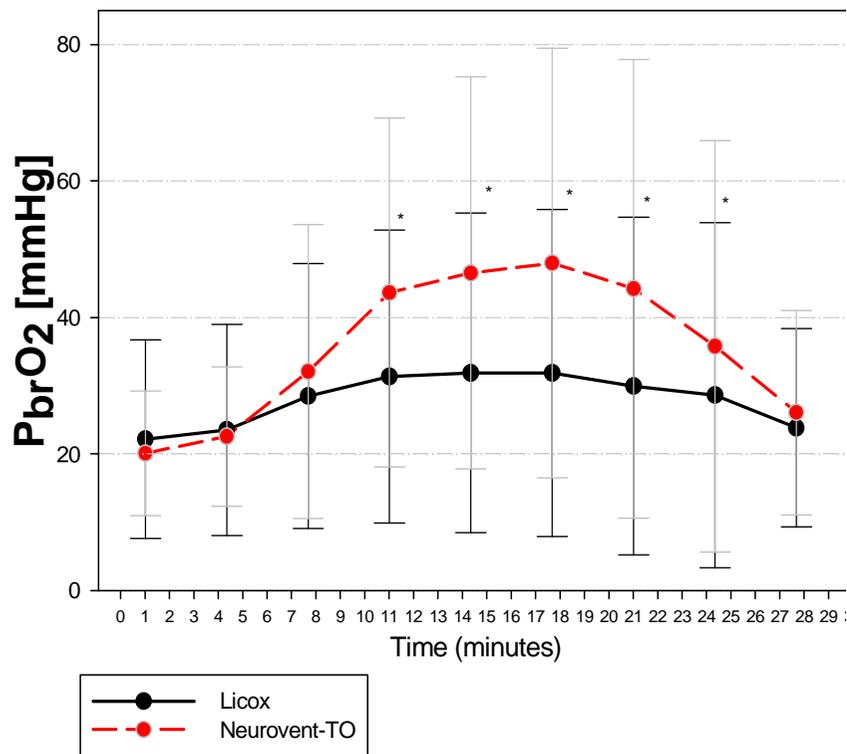
LICOX vs. Neurovent-TO (swine model; n = 9) – Hyper-/Hypotension



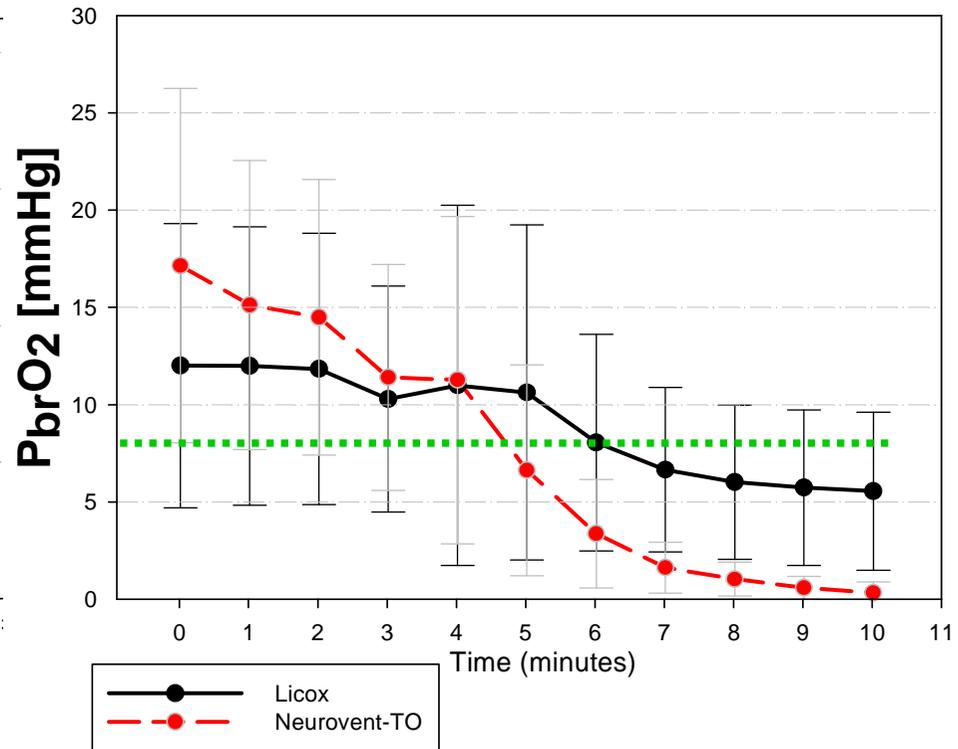


LICOX vs. Neurovent-TO (swine model; n = 9) – Hyper-/Hypooxygen.

P_{br}O₂ Time Course during Hyperoxygenation



P_{br}O₂ Time Course during Hypoxia



▪ significant differences in response to hyper-/hypooxygenation
⇒ probe-specific thresholds for critically low PbrO₂-values needed ?



PbrO₂: Summary

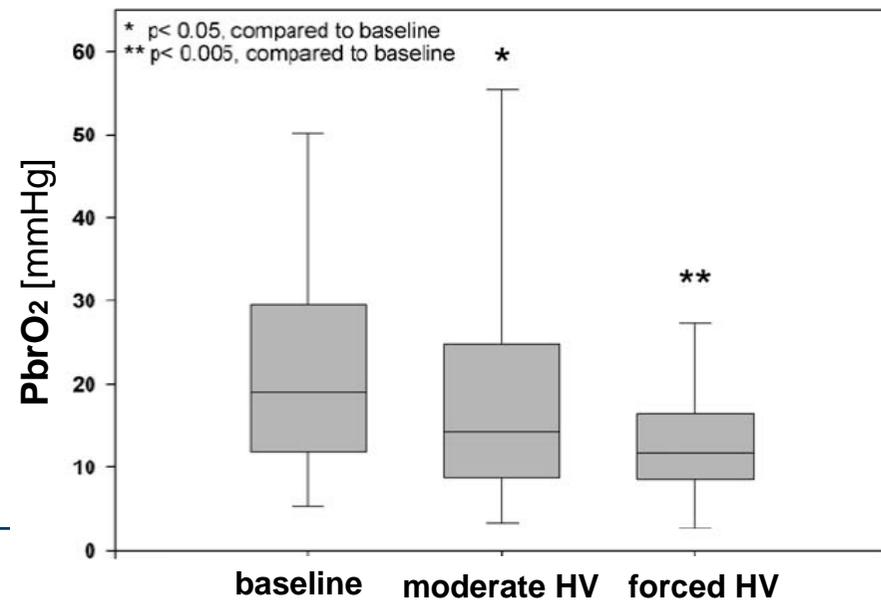
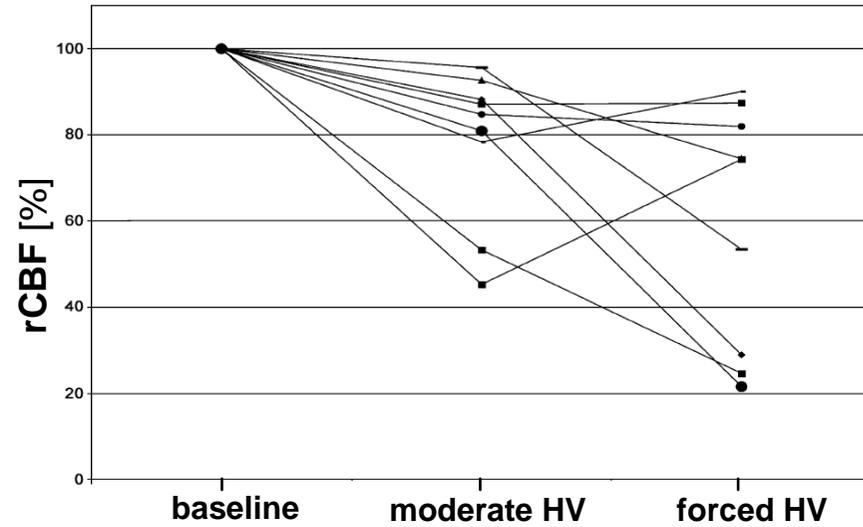
»the wish was father to the thought«
SjvO₂, Clark-type PbrO₂ electrode

»Rome wasn't built in a day«
15 years from research to clinical routine

»new broom sweeps clean, but not always the same street«
Neurovent-TO /-PTO



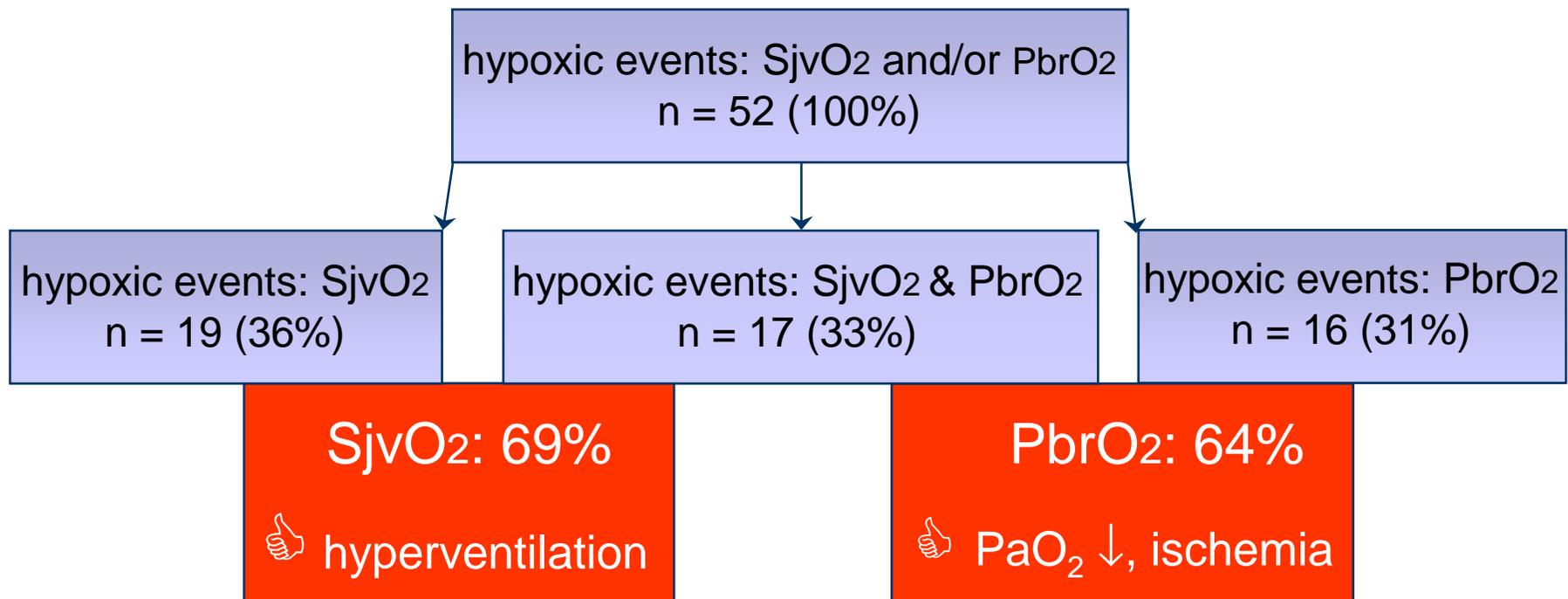
PbrO₂ vs. PCO₂ – “one size doesn’t fit all”





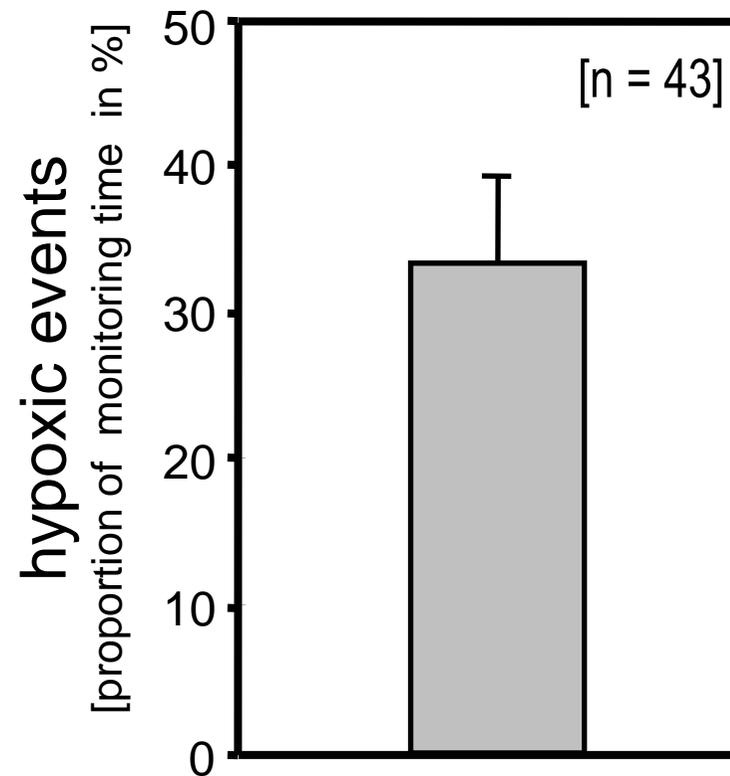
PbrO₂ vs. SjvO₂ (clinical data, TBD)

- n_{pat.}: 58, simultaneous SjvO₂ & PbrO₂ monitoring
- hypoxic thresholds used: SjvO₂: 50% | PtiO₂: 8 mmHg





hypoxic events despite “non-critical” ICP, CPP & PCO₂





critical threshold of $PbrO_2$ – microdialysis (clinical data, TBD)

