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In order to define the relative sensitivities of end tidal CO₂ (ETCO₂), pulmonary artery pressure (PAP) and end tidal nitrogen analysis (ETN₂) in the detection of venous air embolism (VAE), the authors performed serial air injections in dogs.

Methods: Six dogs (weight 25.7+8.1 with pentobarbital, anesthetized were mechanically hyperventilated (VT:20ml.kg rate 15-20 bpm) and placed in a supine/200 head-up posture. Systemic and pulmonary artery pressure, ETCO, (Beckman LB2) and ETN, (Perkin-Elmer 1100 Series massspectrometer) were monitored continuously. An 18 gauge, two inch catheter was placed in the right external vein (EJV). After stabilization (45 min), 5 air injections (0.25, 0.5, 0.75, 1.0 and 1.5cc.kg $^{-1}$) were made via the EJV in random sequence at 30-Increases in PAP and 40 min intervals. , and decreases in ETCO. of 2mmHq, 0.04% and 0.2% respectively were considered positive responses. The time to maximum change and the time to return to baseline following VAE were noted.

Results: The results (expressed as % positive responses for each injectate volume) are shown in Table 1. The sensitivity of the three modalities was comparable. The times to maximum response for the 1.0 and 1.5 cc.kg⁻¹ injectates (Table 2) were not different for PAP and ETN2. Both were more (p<.005, paired t) ETCO2 rapid than although the range for the three was nar-ETN2 returned to baseline (Table 2) following VAE much more rapidly than did ETCO2 (p<.002) which was in turn slightly more rapid than PAP (p<.05).

Discussion: The results of this study indicate that the sensitivities of PAP, ETCO2 and ETN2 in the detection of VAE are very similar. While ETN2 enjoys the theoretical advantage of greater specificity for VAE and might therefore appear preferable, our study indicates two limitations. First, the instrumentation employed must be capable of reliably detecting ETN2 concentration changes on the order of 0.05%. This exceeds the capability of existing commercially available OR systems. Second, ETN2 returns to pre-VAE levels more rapidly than PAP and ETCO2 suggesting that it may not be a reliable indicator of the resolution of the physiologic disturbance caused by VAE. This observation may be the result of ai "held up" in the pulmonary vasculature proximal to the alveolus".

The specificity of ETN₂ analysis is attractive but ETN₂ analysis (with appropriately sensitive detection systems)

should probably be used in conjunction with rather than as an alternative to other monitoring modalities.

Reference: 1. Lechner et al. Quantitative recovery of expired nitrogen and nitrous oxide from venous gas emboli. Pflug Arch 397:225-231, 1983.

	Percent Positive Responses (n=6)				
Injectate Vol (cc/kg)	0.25	0.50	0.75	1.0	1.5
PAP	16.7	33.3	66.7	100	100
ETCO2	16.7	66.7	83.3	100	100
ETN ₂	16.7	50	83.3	100	100

Table 1. The frequency (expressed as percent) of positive responses of PAP, ETCO2 and ETN2 after intravenous injection of various volumes of air.

		Maximum Δ±SD	Time (mins) ± SD		
			To Max∆	To Baseline	
1.0cc.kg ⁻¹	PAP	7.33±4.5mmHg	1.35±1.5	19.7±3.5	
	ETCO2	0.77±.29%	2.10±1.1	15.3±2.1	
	ETN ₂	0.11±.06%	1.25±0.5	8.5±3.9	
1.5cc.kg ⁻¹	PAP	13.33±12.2mmHg	0.92±0.7	23.8±6.1	
	ETC02	0.88±.25%	1.85±0.7	19.4±6.0	
	ETN ₂	0.20±.16%	1.20±0.5	8.0±4.3	

Table 2. The mean maximum changes $(\overline{\Delta})$ in PAP, ETCO2 and ETN2, the time (in minutes) from air injection to maximum change (Max Δ), and the time from air injection to return to stable baseline after 1.0 and 1.5cc.kg⁻¹ air injections. Statistical comparisons in text.

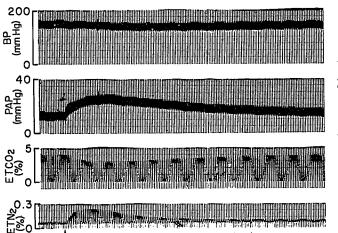


Figure. BP, PAP, $ETCO_2$, and ENT_2 response to injection of lcc/kg of air into the right ventricle of a dog.

1 MIN

INJECT AIR

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