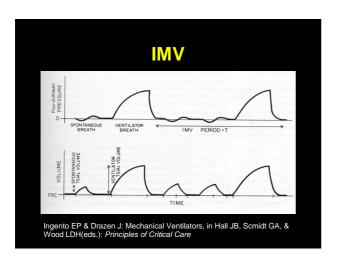
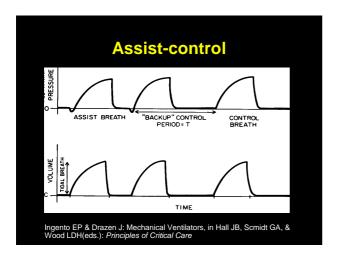
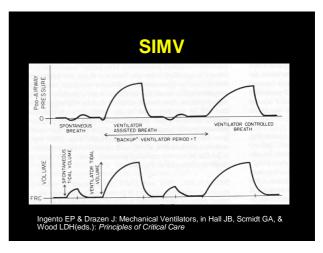


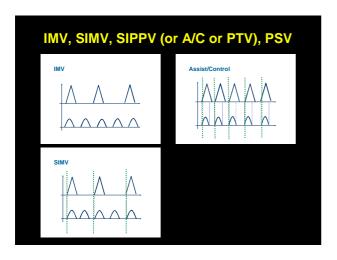
- every breath is fully supported by the ventilator - in classic control modes, patients were unable to breathe except at the controlled set rate - in newer control modes, machines may act in assist-control, with a minimum set rate and all triggered breaths above that rate also fully supported.

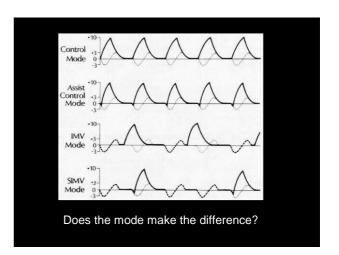


SIPPV (or PTV or A/C) and SIMV Terminology: Triggered ventilation can be divided into patient triggered (PTV), otherwise known as synchronous intermittent positive pressure ventilation (SIPPV) or assist control (A/C), the infant being able to trigger a positive pressure inflation with each breath, and synchronized intermittent mandatory ventilation (SIMV), the infant being able to trigger only a pre-set number of positive pressure inflations.



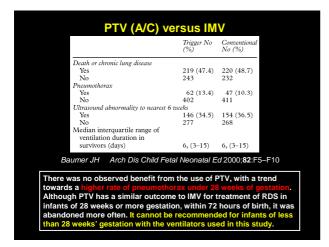




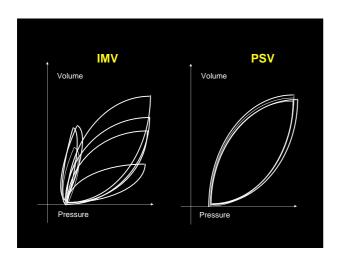


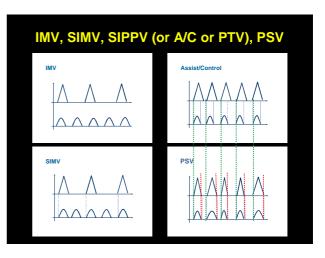
	BW <1000 gm		BW 1000-1499 gm		BW 1500-2000 gm		BW >2000 gm		All BW	
	IMV (n = 45)	SIMV (n = 45)	IMV (n = 44)	SIMV (n = 48)	IMV (n = 25)	SIMV (n = 27)	IMV (n = 46)	SIMV (n = 47)	IMV (n = 160)	SIMV (n = 167)
Duration of venti- lation (hr)	801	596	138	136	77	102	93	724	120	103
Median (95% CD	(499-1020)	(148-973)	(101-280)	(98-344)	(43-107)	(72-107)	(72-106)	(64-77)	(101-142)	(94-118)
Sedation required: No. (%)*	15 (43)	23 (60)	26 (59)	21 (44)	16 (64)	13 (48)	35 (78)	33 (73)	92 (58)	90 (54)
Total doses first 4 days	5.6 (4.5)	6.8 (5.4)	5.9 (4.4)	3.2 (3.1)*	7.3 (7.5)	4.7 (3.7)	6.6 (6.1)	6.2 (6.5)	6.3 (5.5)	5.4 (5.3)
At 1 hr: Pto ₂	0.48 (0.23)	0.52 (0.27)	0.61 (0.27)	0.53 (0.25)	0.59 (0.26)	0.48 (0.18)	0.70 (0.25)	0.71 (0.27)	0.60 (0.26)	0.57 (0.26)
MAP (cm H ₂ O)	7.4 (1.7)	7.6 (2.2)	9.1 (2.8)	8.3 (2.4)	8.8 (2.8)	7.7 (2.0)	9.2 (2.6)	8.6 (1.8)	8.6 (2.6)	8.1 (2.2)b
OI	6.4 (4.4)	8.3 (8.7)	10.3 (8.8)	6.8 (4.6)b	7.7 (7.3)	5.4 (3.3)	6.2 (5.0)	5.3 (4.3)	7.7 (6.7)	6.5 (5.8)
Paralysis: No. (%)	5 (11)	2(4)	2(5)	2(4)	1(4)	0	7 (15)	6 (13)	15 (10)	10 (6)
Air leak: No. (%)	10(22)	7 (15)	9 (20)	10(21)	3 (12)	1(4)	2(4)	0	24 (15)	18 (11)
IVH (grade III or IV): No. (%)	6 (13)	7 (16)	2 (5)	5 (10)	3 (12)	0	0	0	11 (7)	12 (7)
Death: No. (%) Oxygen	7 (15)	7 (16)	2 (5)	3 (6)	1 (4)	1 (4)	0	0	10 (6)	11 (7)
At 28 days: No. (%)†	32 (79)	29 (77)	22 (50)	25 (52)	7 (28)	7 (26)	0	1(2)	59 (39)	61 (39)
At 36 wk PCA: No. (%)†	28 (72)	18 (47) ^b	14 (32)	12 (25)	6 (24)	4 (15)	NA	NA	NA	NA
Results are mean (SE CI, Confidence innerv Significant difference ${}^{1}p = 0.02$. ${}^{3}p < 0.05$. *Excludes infants wh †Percentage of surviv	al; PCA, posso s between IM o received ann	conceptional in V and SIMV iconvulsant d	ige; NA, not a within BW gr	oups:	at leas	st as ef	: We fo ficacion y have BW-sp	us as c improv	onvent ed cert	ional ain

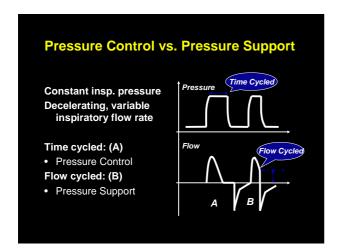
	Trigger No (%)	Conventional No (%)
Death or chronic lung disease		_
Yes	219 (47.4)	220 (48.7)
No	243	232
Pneumothorax		
Yes	62 (13.4)	47 (10.3)
No	402	411
Ultrasound abnormality to nearest 6	5 vveeks	
Yes	146 (34.5)	154 (36.5)
No	277	268
Median interquartile range of		
ventilation duration in		
survivors (days)	6, (3–15)	6, (3–15)
Baumer JH Arch Dis Child Fe V with inspiratory times of between the trigger at each inspiratory effort.	n 0.2 and 0.25	seconds, the ventilat
et to trigger at each inspiratory effort, I	backup rate of	35 breaths a minute.

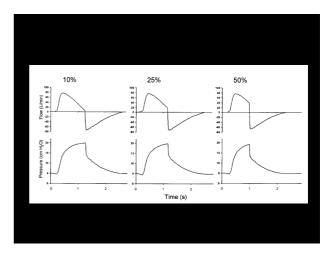


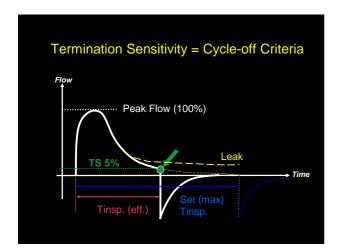
Synchronized mechanical ventilation for respiratory suppor in newborn infants (Review)									
Comparison 02. PTV / SIMV	<u> </u>								
Outcome title	No. of No. of studies participants		Statistical method	Effect size					
01 Death	5	1729	Relative Risk (Fixed) 95% CI	1.19 [0.95, 1.49]					
02 Airleaks	6	1769	Relative Risk (Fixed) 95% CI	1.03 [0.80, 1.34]					
03 Duration of ventilation (hours)	4	1402	Weighted Mean Difference (Fixed) 95% CI	-34.78 [-62.11, -7.44]					
04 Extubation failure	4	1056	Relative Risk (Fixed) 95% CI	0.93 [0.68, 1.28]					
05 Severe IVH	5	1729	Relative Risk (Fixed) 95% CI	1.03 [0.74, 1.43]					
06 CLD (oxygen dependency at 28 days)	4	805	Relative Risk (Fixed) 95% CI	0.91 [0.75, 1.12]					
07 CLD (oxygen dependent at 36 weeks PCA)	2	1310	Relative Risk (Fixed) 95% CI	0.90 [0.75, 1.08]					
Comparison 03. PTV vs SIMV									
Outcome title	No. of studies	No. of participants	Statistical method	Effect size					
01 Duration of weaning (hours)	3	120	Weighted Mean Difference (Fixed) 95% CI	-42.38 [-94.35, 9.60]					
02 Weaning failure	3	120	Relative Risk (Fixed) 95% CI	0.78 [0.31, 1.93]					
03 Extubation failure	3	120	Relative Risk (Fixed) 95% CI	1.00 [0.37, 2.67]					
04 Air leaks	3	120	Relative Risk (Fixed) 95% CI	0.80 [0.23, 2.83]					



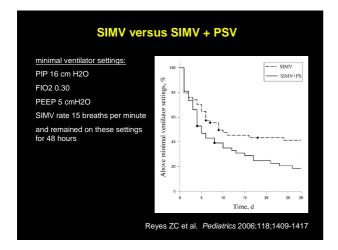


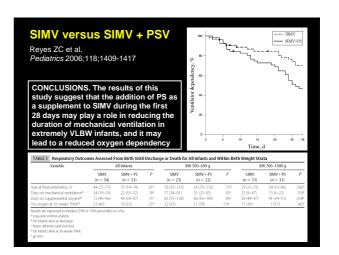


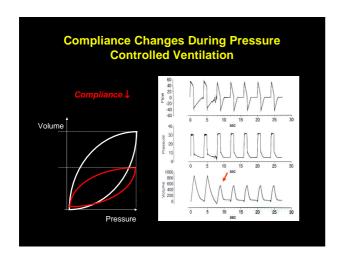


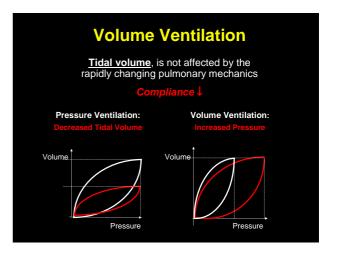


PSV improves respiratory function in VLBW infants when compared to SIMV TABLE 2—Results Concerning Spontaneous Respiratory Rate (RR), Tidal Volume $\{V_T\}$, Minute Volume $\{V_{min}\}$, and Mean Airway Pressure (MAP) During Four Ventilation Phases 1 First ventilation cycle Second ventilation cycle 3: SIMV (4 hr) 1: SIMV (4 hr) 2: PSV (4 hr) Test phase RR b/min: mean (SE) V_T ml/kg: mean (SE) V_{min} ml/kg: mean (SE) MAP cm H₂O: mean (SE) 43.63 (3.54) 6.88 (0.27) 349.0 (24.44) 7.73 (0.34) 56.08 (3.87) 53.05 (4.23) 43.18 (3.96) 6.96 (0.25) 388.0 (24.81) 7.56 (0.29) 7.05 (0.40) 7.06 (0.41) ¹Results are reported as mean and SE in parentheses. A significant difference was observed by ANOVA for repeated measures (RR, P=0.000000; $V_{\rm Tr}$, P=0.000000; $V_{\rm min}$, P=0.000000; MAP, P=0.018694). The Newman-Keuls post hoc test showed the following differences: spontaneous respiratory rate (RR) test phase 1 vs. 2, P=0.000124; test phase 3 vs. 4, P=0.000123; tical volume (Vr), test phase 1 vs. 2, P=0.000124; test phase 3 vs. 4, P=0.000123; test phase 1 vs. 3, P=0.000123; test phase 1 vs. 2, P=0.000123; test phase 1 vs. 2, P=0.000123; test phase 3 vs. 4, P=0.000123; mean airway pressure (MAP) test phase 1 vs. 2, P=0.0033747; test phase 3 vs. 4, P=0.001233; mean airway pressure (MAP) test phase 1 vs. 2, P=0.003747; test phase 3 vs. 4, P=0.0012438. Migliori C et al. Pediatr Pulmonol. 2003;35:364-367









Combination "Dual Control" Modes

Combination or "dual control" modes combine features of pressure and volume targeting to accomplish ventilatory objectives which might remain unmet by either used independently.

Combination modes are pressure targeted

Partial support is generally provided by pressure support

Full support is provided by Pressure Control

Combination "Dual Control" Modes

Volume Assured Pressure Support (Pressure Augmentation)

Volume Support (Variable Pressure Support)

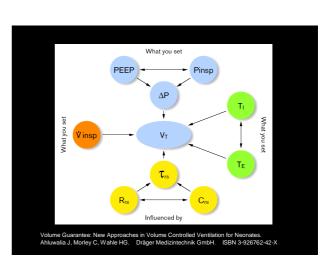
Pressure Regulated Volume Control (Variable Pressure Control, or Autoflow)

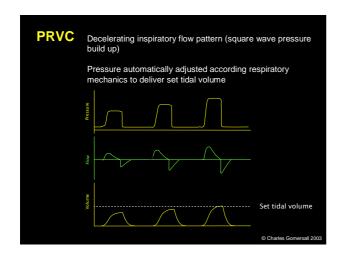
Airway Pressure Release (Bi-Level, Bi-PAP)

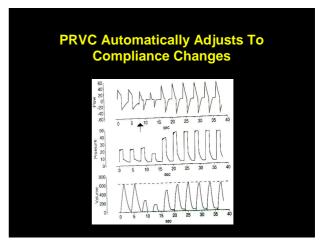
PRVC (Pressure regulated volume control)

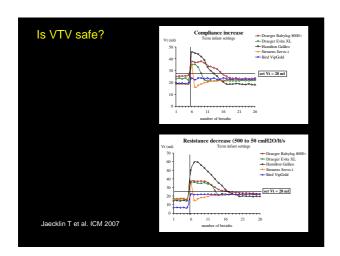
A control mode, which delivers a set tidal volume with each breath at the lowest possible peak pressure.

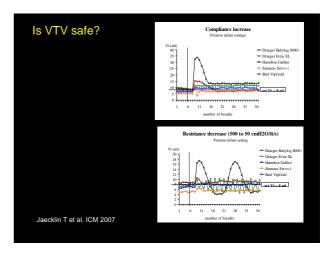
Delivers the breath with a decelerating flow pattern that is thought to be less injurious to the lung..... "the guided hand".

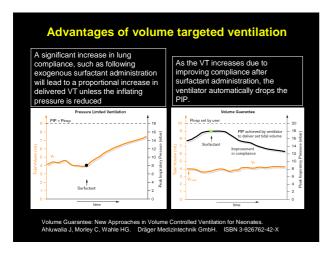




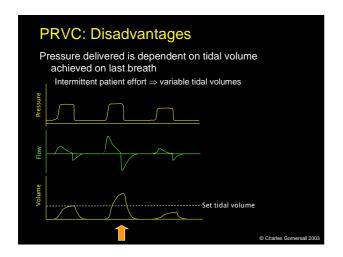


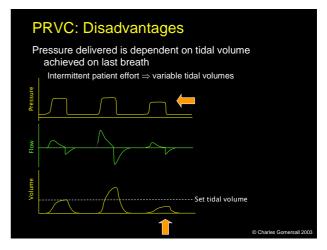


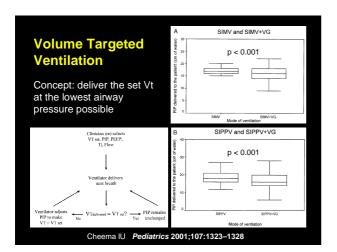


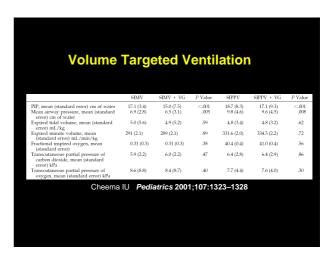


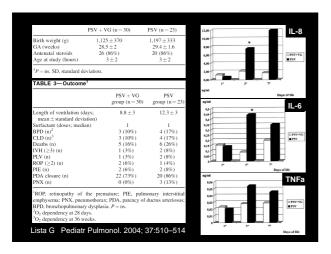
PRCV: Advantages Decelerating inspiratory flow pattern Pressure automatically adjusted for changes in compliance and resistance within a set range Tidal volume guaranteed Limits volutrauma Prevents hypoventilation









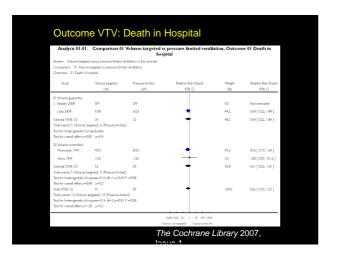


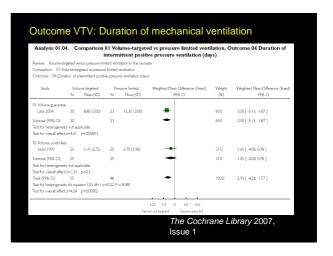
Methods:

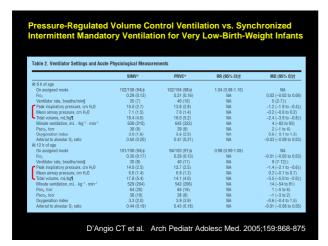
PSV group: The weaning strategy consisted of reducing the pressure support level progressively over time, so that the work of breathing was shifted from ventilator to the patient.

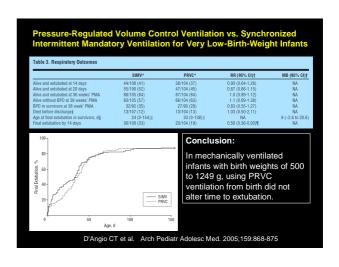
PSV-VG group: Weaning was a more automatic process once appropriate levels of Vt had been established.

Similar blood gas goals (e.g., pH>7.25; pO2, 50–75 mmHg; pCO2, 40–65 mmHg) were achieved during weaning from mechanical ventilation in both groups.









Lung-protective ventilation strategies in neonatology: What do we know—What do we need to know?

Anton H. van Kaam, MD, PhD; Peter C. Rimensberger, MD

Crit Care Med 2007; 35:925–931

A total of 24 RCTs and 3 systematic reviews comparing various CMV modes and settings and 2 RCTs investigating permissive hypercapnia reported no differences in mortality or bronchopulmonary dysplasia.