Comparative study of Trilogy200 and LTV1000 ventilators on a pediatric patient

Clinical report

Colleen Witt, MS, RAC

Clinical Research Associate, Philips Respironics Monroeville, Pennsylvania

William Truschel

Engineer, Philips Respironics Monroeville, Pennsylvania

Anandi Mahadevan

Engineer, Philips Respironics Monroeville, Pennsylvania

Abbreviations PC pressure control RR respiratory rate PS pressure support I-Time inspiratory time PEEP positive end expiratory pressure Vte tidal volume exhaled SIMV synchronized intermittent mandatory ventilation SpO, pulse oximetry FiO, oxygen concentration

LTV1000 and Trilogy200 ventilator settings		
Туре	Invasive via tracheostomy	
Mode	PC/SIMV	
Rate	18	
I-Time	0.6 sec	
PC	18 H ₂ O	
PS	10 cm H ₂ O	
PEEP	5 cm H ₂ O	
Flow Trigger Sensitivity	3 lpm	
Oxygen	32%	

Introduction

This clinical report is intended to compare the LTV1000 and the Trilogy200 ventilators in regards to triggering and patient/ventilator synchrony on a pediatric patient.

A 13 month-old female weighing 5.7 kg with a diagnosis of Cornelia de Lange syndrome requiring ventilatory support was observed at the Children's Hospital in New Brunswick, New Jersey. The primary goal of the study was to observe patient/ventilator synchrony and the response to the inspiratory efforts. A noninvasive cardiac output monitor (NICO) was used to collect comparison breath wave form data while the patient was breathing on her current ventilator (LTV1000) and Trilogy200. Additional data collected included exhaled tidal volume, respiratory rate and SpO2 to compare therapy efficacy while the patient was on both the LTV1000 and the Trilogy200 ventilators.

The infant was assisted by her current ventilator, an LTV1000, through a tracheostomy tube. The ventilator settings were: PC-SIMV mode, PC 18, PS 10, PEEP 5, RR 18, I-Time 0.6, and flow trigger sensitivity of 3 lpm.

During the study, the infant was switched from her current ventilator (LTV1000) to a Trilogy200 ventilator set to the same current prescription. Three liters per minute of low-flow oxygen was introduced into the Trilogy200 device to approximate the prescribed FiO_2 .

The infant was observed for two hours and spent approximately one hour on each of the ventilators in order to collect comparative data. The results of the study are contained herein.

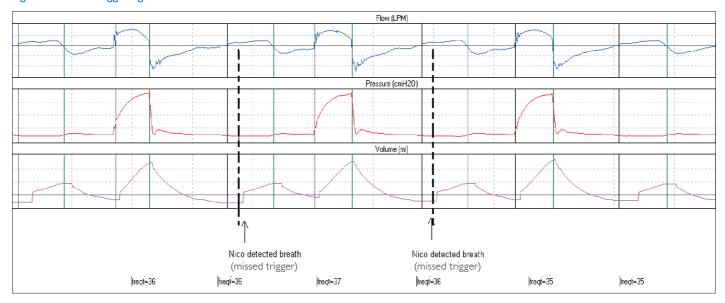


Observations

During the study, the NICO monitor was detecting breaths (see **Figure 1**); however the LTV1000 failed to detect the breath and as a result, did not trigger. The LTV1000 ventilator was not

synchronizing with the infant's effort and failed to trigger on every other breath (as shown in **Figure 1**).

Figure 1: Patient triggering while on the LTV1000



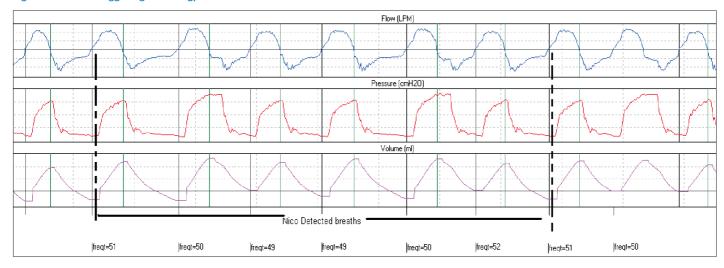
Throughout the study, the infant's SPO_2 level was maintained at 100%. There was no change in SpO_2 even when the ventilators were exchanged. The average Vte measured by the NICO while the infant was on the LTV1000 was 54 and the average breath rate was 45.

After approximately 30 minutes of collecting data from the LTV1000 ventilator, the infant was switched to the Trilogy200 ventilator. The Trilogy200 device was set to the same settings as LTV1000. The

Trilogy200 ventilator was able to detect all the breaths that the NICO monitor detected as a breath and the vent was triggering in synchronization with the patient to provide therapy to the child.

Figure 2 demonstrates the triggering efficacy of the Trilogy200 ventilator. Breath delivery is in synchronization with the infant with no missed triggers detected.

Figure 2: Patient triggering on Trilogy200



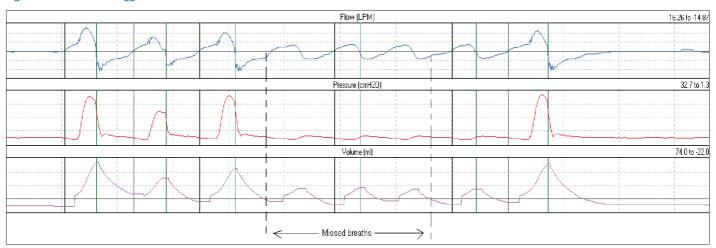
The average Vte while the infant was using the Trilogy200 device was 57 and the average breath rate was 50, similar to that of the LTV1000 ventilator. After one hour on the Trilogy200 ventilator, the infant was switched back to the existing ventilator and data

was recorded for another half hour to observe triggering and synchronization with the LTV1000 while the infant was awake. (The infant was asleep on the LTV1000 ventilator during the first part of the study.)

Observations

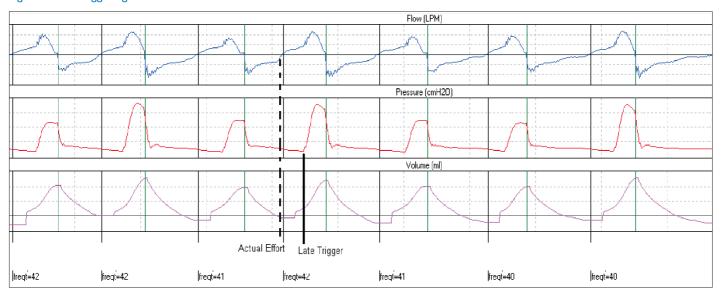
As illustrated in **Figure 3**, the NICO independent monitor detected breaths while the LTV1000 did not. This resulted in frequent missed breaths.

Figure 3: LTV missed triggers while on the LTV1000



When triggering did improve on the LTV1000 device, it was late as illustrated in **Figure 4**. The average Vte while on LTV1000 when the infant was awake was 54 ml.

Figure 4: Late triggering on the LTV1000 ventilator



Conclusion

Comparative data analysis between the two ventilators showed that patient/ventilator synchrony and response to the infant's inspiratory efforts was excellent when using Trilogy200.

Comparison statistics

Parameters	Trilogy200	LTV1000
Vte	57	54*
Respiratory rate	50	45
Triggering	No missed triggers**	Late or missed triggers***
SpO ₂	100%	100%

^{*}The Vte measurement on LTV1000 was consistently changing on a breath-by-breath basis with an average value of approximately 54. During the course of the study, the Trilogy200 device maintained a constant Vte of 57. SpO₂ measurements were similar for the two devices as indicated by performance for this parameter.

^{**}No missed triggers and synchronous with patient effort for the T200.

^{***}Late triggering detected and not synchronous with patient effort for the LTV1000.

Respironics and Trilogy are trademarks of Respironics Inc. and its affiliates. All rights reserved.

Please visit http://trilogy200.respironics.com



 $\ensuremath{@}$ 2010 Koninklijke Philips Electronics N.V.All rights are reserved.

Philips Healthcare reserves the right to make changes in specifications and/or to discontinue any product at any time without notice or obligation and will not be liable for any consequences resulting from the use of this publication.

 $\label{eq:caution:def} \textbf{CAUTION:} \textbf{US} \textbf{ federal law restricts these devices to sale by or on the order of a physician.}$

Geyer KKB 10/14/10 MCI 4103529 PN 1077255

Philips Healthcare is part of Royal Philips Electronics

Philips Respironics Asia Pacific +55 6882 5282 Philips Respironics Australia +61 (2) 9666 4444 Home Healthcare Solutions International +33 1 47 28 30 82 Philips Respironics United Kingdom +44 800 1300 845 Philips Respironics 1010 Murry Ridge Lane Murrysville, PA 15668

Customer Service +1 724 387 4000 800 345 6443 (toll free, US only) www.philips.com/respironics