

Using COVID-19 ICU Respiratory Training Simulators

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The coronavirus SARS-Cov-2 has led to an urgently worrying, universal problem: a lack in the amount of intensive care units (ICUs) with beds allowing patients to be put on mechanical ventilation and healthcare staff trained to operate these ventilators.

At the outset of the SARS-COV-2 pandemic, it had not been determined why respiratory support in the ICU was necessary in COVID-19 patients. Experts spoke about "COVID-19 pneumonia" and "non-ARDS" (non-acute respiratory distress syndrome) [1], leading to confusion amongst those operating ventilators. This confusion stemmed from the fact that established guidelines for mechanical ventilation did not address this novel kind of respiratory failure.

However, very soon after this confusion was identified, guidelines were proposed [2,3] to help practitioners. This article provides information on how to use TestChest to represent COVID-19 patients in order to practice using respiratory support devices and to alleviate the shortage of expertise in providing effective respiratory care in the ICU to COVID-19 patients.

TestChest: The autonomous lung simulator

TestChest is an autonomous lung simulator that provides two natural interfaces: the airway opening, to provide respiratory therapy and the ability to the measurement of lung function, and an artificial finger to measure oxygen saturation and plethysmography.

The trainer sets the disease and the breath pattern, and the trainee applies the respiratory therapy. TestChest will respond to the settings and therapy applied autonomously, without needing any further interaction from the trainer.

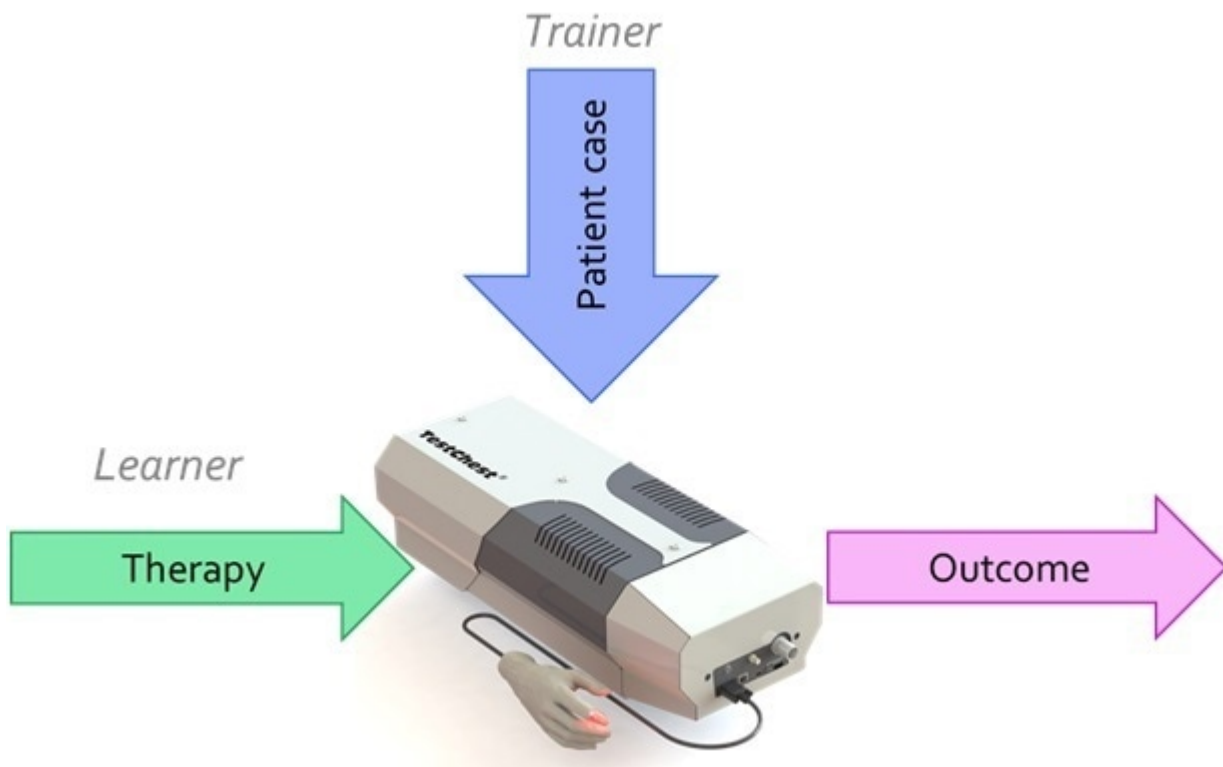


Figure 1. Autonomous lung simulator TestChest. The trainer sets a patient case and TestChest reacts autonomously to the therapy attempts of the trainee, without further interaction from the trainer. Outcomes are oxygen saturation, airway pressures, tidal volumes, respiratory rate, and other physiological responses. Image Credit: neosim AG

Method to set TestChest to simulate COVID-19 patients

Three successive scenarios are created to train operators to use mechanical ventilators for COVID-19 patients arriving in the ICU.

The overall goals of the simulation are to be able to practice the following:

- Selecting the appropriate ventilation mode
- Adjusting settings to achieve adequate SpO₂ and adequate end-tidal CO₂
- Applying lung-protective ventilation, including tidal volume (VT), plateau, positive end-expiratory pressure (PEEP).

The table below provides detailed information about setting the TestChest to achieve the different patient groups as described in [4], Type L and Type H.

Table 1. TestChest parameters set to provide COVID-19-like patient for training. Detailed explanation in [5]. Source: neosim AG

Parameter		COVID-19	COVID-19	COVID-19
		Type L: spontaneous breathing	Type L: paralysis, first hours	Type H: late stage (ARDS)
CW	ml/hPa	93	93	93
V'CO ₂	mlSTPD	250	197	250
P0.1	hPa/100 ms	8	0	1
Crs	ml/hPa	52	52	35
Raw	hPa/(L/s)@1 L/s	Rp5	Rp5	Rp5
f (spont)	/min	40	0	0
FRCmin	ml	1102	1102	1102
LowerInflection	hPa	1	1	12
UpperInflection	hPa	35	35	35
C1	ml/hPa	52	52	8
C3	ml/hPa	8	8	8
Pthreshold	hPa	80	80	35
Pcollapse	hPa	0	0	10
RCIh	s	0.5	0.5	0.5
Rcrecol	s	10	10	10
FRCpred	ml	1750	1750	1750
Tdelay	s	1	1	1
Vdaw	arbitrary	large	large	large
Cr	ml/hPa	2	2	15
RCcollapse	s	10	10	1
Pdiff	mmHg	400	400	300
QT	ml/min	6000	4000	6000
POPv@Pcardio=10 hPa%		50	50	50
POPv@Pcardio=20 hPa%		60	60	60
POPv@Pcardio=30 hPa%		70	70	70
Leak level	arbitrary	none	none	none
HeartRate	/min	90	70	80

It is important to note that the parameters of TestChest may need to be adapted to the rapidly evolving knowledge of COVID-19.

Scenario 1: COVID-19 Type L; respiratory failure and spontaneous breathing after intubation or return to spontaneous breathing.

Patients arrive in the hospital with a very high respiratory drive due to intense hypoxemia. They present a high dead space that can be explained by a reduction of hypoxic pulmonary vasoconstriction and the presence of multiple pulmonary microthrombosis. These patients have a high compliance and respond to FiO₂, some PEEP and anti-Trendelenburg position.

Ventilation challenges:

- Choosing the right mode
- Setting up the spontaneous breathing parameters such as trigger, e-cycling, PEEP, and FiO₂
- Setting alarms.

Summary for trainers:

Table 2. Source: neosim AG

Lung mechanics	Metabolism	Spontaneous breathing	Hemodynamics
<ul style="list-style-type: none"> • Crs = 52 ml/mbar. • Raw=5 (without tube), • No lower inflection point • UIP at 35 mbar • Not recruitable 	<ul style="list-style-type: none"> • V'CO₂ = 250 ml/min STPD 	<ul style="list-style-type: none"> • P_{insp}= -14 mbar, • T_i=1.4 sec, • f=40 /min 	<ul style="list-style-type: none"> • Pulseoximeter pleth will react to high inspiratory pressure (high intrathoracic pressure) • C.O.= 6 L/min

The hemodynamics are very basic and are only visible on pulseoximeter pleth curve.

Scenario 2: COVID-19 Type L; exacerbated respiratory failure after paralysis to avoid self-inflicted lung damage due to high respiratory drive.

If a patient’s respiratory drive is very high, they may be at risk from lung damage caused by uncontrolled inspiratory effort, severe lung injury, and asynchronies. Paralysis can help in these cases.

Ventilation challenges:

- Switching to an appropriate mode
- Setting the parameters like VT, rate, FiO₂, PEEP, to achieve adequate gas exchange and oxygen saturation
- Setting alarms to accommodate the new patient condition.

Summary for trainer:

Table 3. Source: neosim AG

Lung mechanics	Metabolism	Spontaneous breathing	Hemodynamics
<ul style="list-style-type: none"> • Crs = 52 ml/mbar. • Raw=5 (without tube), • No lower inflection point • UIP at 35 mbar • Not recruitable 	<ul style="list-style-type: none"> • V’CO₂ = 250 ml/min STPD 	<ul style="list-style-type: none"> • None 	<ul style="list-style-type: none"> • Pulseoximeter pleth will react to high inspiratory pressure (high intrathoracic pressure) • C.O.= 6 L/min

Paralysis will create a slight drop in V_{LEE} requiring PEEP to increase SpO₂.

Scenario 3: COVID-19 Type H; exacerbation respiratory failure low compliance ARDS stage

Unfortunately, some patients with COVID-19 can progress to a low

compliance ARDS.

Ventilation challenges are:

- The titration of FiO_2 and PEEP
- Reducing of VT to maintain 6 ml/kg, driving pressure <14 cmH₂O
- Not exceeding a plateau pressure of 30 cmH₂O (<28)
- Recruiting trials and evaluations
- Setting alarms to accommodate the new patient condition.

Summary for trainer:

Table 4. Source: neosim AG

Lung mechanics	Metabolism	Spontaneous breathing	Hemodynamics
<ul style="list-style-type: none"> • Crs = 35 ml/mbar, • Raw=5 (without tube), • LIP=8 mbar • UIP=35 mbar • Venous admixture (QsQt) about 37% • Recrutable with P_{insp} >35 mbar • Recruitment will take 10 seconds to achieve a QsQt reduction from 37% to 15% • Lung collapse if pressure (PEEP) falls below 10 mbar (suction) 	<ul style="list-style-type: none"> • $\dot{V}'\text{CO}_2 = 250$ ml/min STPD 	<ul style="list-style-type: none"> • None 	<ul style="list-style-type: none"> • Pulseoximeter pleth will react to high inspiratory pressure (high intrathoracic pressure) • C.O. = 6 L/min

Summary

TestChest provides mechanisms to simulate COVID-19 patients to facilitate the training of healthcare practitioners in operating mechanical ventilators. The TestChest's response is autonomous, so trainers do not need to enter any

outcome variables because they are created automatically in direct response to the trainee's actions.

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neosim is a Swiss company founded by experts with strong background in lung physiology and mechanical ventilation of intensive care patients. The mission of neosim is to bring high-fidelity physiology and pathophysiology to the patient simulator community.

For training and education of clinicians, especially respiratory therapists and intensive care professionals, neosim simulators create realistic breathing in health and disease. In contrast to other simulators, neosim's simulators can be treated with intensive care therapy methods and responds like a real human patient. The result manifests itself clinically and can be measured quantitatively with state-of-the-art monitoring in real-time.

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neosim AG



Address

Susenbühlstrasse 12
Chur
CH-7000
Switzerland

Phone: +41 (79) 254 0312



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