









Advanced technology that pushes back on the challenges you face

The F&P 950 System was designed to resolve many of the pain points you may have experienced while administering care in the past or that you are facing now with your present device. **How many of the issues listed below are familiar to you?**



Circuit condensate



Damaged or lost probes



Complex to set up



Unclear alarm display



Limited temperature adjustability

By seeking to address these frustrations, the F&P 950 System frees you to concentrate your time more fully on patient care.

Discover how on the following pages.

The F&P 950 System paves the way to **greater patient care**



Circuit Technology:

A three-pronged approach
that tackles condensate

AirSpiral™

Wires have been embedded along the inspiratory limb wall for heating consistency, and to deliver essential feedback from the patient-end temperature sensor.

Insulating air pockets further protect the humidified gas from the ambient conditions, preventing it from losing heat and moisture.

Thermadapt™

Different heating zones along the neonatal inspiratory limb enable the breathing tube to adapt to changing ambient environments, such as those found in incubators and infant warmers. This technology also removes the need for an unheated extension.

Evaqua[™] technology

The permeable circuit membrane allows humidity to diffuse out of the expiratory limb before it condenses into liquid, reducing condensate in the expiratory limb and expiratory filter.



Probe Design: Facilitates ease of use and setup

The once-separated temperature and flow probes have been integrated into the consumables and hardware. This eliminates the need for the probes to be cleaned between patient use and the risk of these valuable components being damaged or misplaced.





Fewer Assembly Connections*:

Easy, three-step setup

The F&P 950 System is intuitive to set up and requires fewer assembly steps. This has been demonstrated to reduce user errors, even among those who have no experience with the product.



Interactive Display: Intuitive alarms

The interactive color touch screen reduces alarm confusion. The animations and text-based alarms provide users with clear guidance on when to react and how to resolve issues.



Comfort Settings: Temperature adjustability

Three separate modes are available for adult applications: Invasive, Mask and Optiflow. When set to Mask or Optiflow mode, the temperature can be adjusted to support patient comfort and encourage compliance.

What do users have to say

about the F&P 950 System?



could use the F&P 950 System without errors.¹



had to empty condensate during their shift, compared with 59% for their previous system.²



understand an alarm.3

Mode found it easy to

Four commonly asked questions

about the F&P 950 System

How does the F&P 950 System differ from the F&P 850 System?

The F&P 950 System has an advanced internal control system and new technology that improves performance.

Features include:

New circuits designed to minimize condensate
 Enhanced color touch screen and visual alarms for greater ease of use
 An easy and intuitive setup procedure





Can the temperature be adjusted?

Yes, the temperature can be adjusted when the adult circuit is connected in Mask and Optiflow modes. However, neonatal and invasive modes are always set to 37 °C.

The available comfort settings are:

Mode	Default	Medium	Low
Invasive	37 °C	-	-
Mask	31 °C	29 °C	27 °C
Optiflow	37 °C	35 °C	33 °(
Neonatal	37 °C	-	-



What therapies does the F&P 950 System support?

Adult patients:

Invasive, noninvasive/CPAP, nasal high flow

Neonatal patients:

Invasive, noninvasive/CPAP, nasal high flow, high-frequency ventilation

Note: When a breathing circuit is attached, the system automatically detects whether this is an adult or neonatal model and activates

the correct mode.

What is involved in managing the probes for the F&P 950 System?

Probes are either covered by protective SensiDomes™ or are disposed of with the single-use circuit. The SensiDomes provide a microbial barrier, which removes the need for disinfection or sterilization between patients.

Individual components may vary from those shown

^{*} Compared with the F&P 850 System

^{1.} Numbers presented represent findings from the F&P 950 Humidification System Clinical Usability Assessment – Product Validation (Adult).

^{2.} Numbers presented represent findings from the F&P 950 Usability Validation Report (Adult).

^{3.} Numbers presented represent findings from the F&P 950 Humidification System Clinical Usability Assessment – Product Validation (Neonatal).

All assessments and reports were conducted by Fisher & Paykel Healthcare.



For more information or to register for a trial, please contact your local Fisher & Paykel Healthcare representative or visit our website: www.fphcare.com/950



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