

# FLUENT<sup>®</sup>

## fluid management

**SIMPLIFIED PROCEDURES WITH GREATER  
CONTROL FOR HYSTEROSCOPY**

- **Simplified** set-up and operation
- **Advanced technology** with intuitive user interface for increased clinical confidence
- **Time-saving** can cut set-up time by 50%



# Help your facility operate more efficiently

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## SIMPLICITY

REDUCED TUBING AND CONNECTION REQUIREMENTS



### FLOPAK™ CARTRIDGES

snap into place for simplified setup



### INTUITIVE TOUCHSCREEN

automatically guides setup and operation



### SINGLE-WASTE-BAG DESIGN

eliminates the need for multiple canisters

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## TECHNOLOGY

DESIGNED TO INCREASE CLINICAL CONFIDENCE



### ACCURATE FLUID DEFICIT READINGS

within +/- 50 mL (1.69 oz)<sup>1</sup>



### ADVANCED PRESSURE CONTROL

maintains consistent intrauterine distention<sup>2</sup>



### FLOPAK™ TECHNOLOGY

helps manage fluid use throughout the procedure



### IMPROVED VISUALISATION

for enhanced patient and procedural benefits

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## TIME SAVING

STREAMLINE PROCEDURES FOR GREATER CONTROL



**SETUP TIME**  
greatly reduced



**ALL-IN-ONE PROCEDURE KITS**  
give nurses all the components they need in one pack



**LESS EQUIPMENT REQUIRED**  
with integrated MyoSure® controller to help control OR space

**Fluent can be used with our MyoSure portfolio**

## Visit [FluentbyHologic.com](https://www.fluentbyhologic.com) for more details

### IMPORTANT SAFETY INFORMATION

The Fluent fluid management system is intended to provide liquid distention of the uterus during diagnostic and operative hysteroscopy, and to monitor the volume differential between the irrigation fluid flowing into and out of the uterus while providing drive, control, and suction for hysteroscopic morcellators.

The Fluent fluid management system may not be used to introduce fluids into the uterus when hysteroscopy is contraindicated. The system should not be used to remove pathologies from pregnant patients or patients exhibiting pelvic infection, cervical malignancies, or previously diagnosed endometrial cancer. For detailed benefit and risk information, including contraindications relative to endometrial ablation, please consult the Instructions For Use.

<sup>1</sup>When operating with a MyoSure device in a bench test environment in a uterine model (N = 20).

REFERENCES: 1. Hologic, Inc. Data on file, bench testing. DTP-00737. 2. Hologic, Inc. Data on file, bench testing. DTP-00591.