

FLUENT[®]

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

2

TECHNOLOGY

DESIGNED TO INCREASE CLINICAL CONFIDENCE



ACCURATE FLUID DEFICIT READINGS

within +/- 50 mL
(1.69 oz)¹



ADVANCED PRESSURE CONTROL

maintains consistent
intrauterine distention²



FLOPAK™ TECHNOLOGY

helps manage fluid
use throughout the
procedure



IMPROVED VISUALISATION

for enhanced patient
and procedural
benefits

3

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.

¹When 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.