HALKEY | ROBERTS®

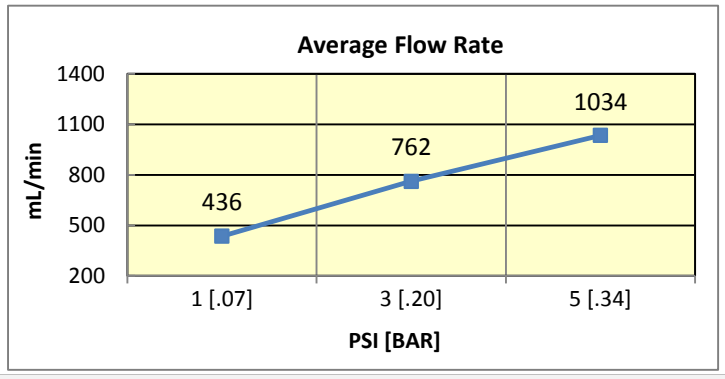
251 MALE LUER VALVE

MALE LUER 6.6MM TUBE END VALVE

GENERAL CHARACTERISTICS

- Halkey-Roberts Male Luer Valves are designed to attach directly to tubing during their production. These valves can be used to access a mating female luer connector or luer activated valve.
- All materials are Gamma resistant, ISO 10993 compliant, DEHP-free and not made with natural rubber latex.
- The Male Luer Valve is available in polycarbonate.
- Available in clear, red, blue, green and yellow
- Tubing port is designed for 6.6mm (.260 inch) O.D. tubing.
- Produced under GMP: Halkey-Roberts is an ISO 9001-2008, ISO 13485-2003 and FDA registered manufacturing facility.
- The Male Luer Valve series is a medical component: bulk, non-sterile, for manufacturing processing or repacking only.
- Customer is responsible for the Qualification/Verification of the HR® medical component in their final device application.
- Luer fittings are compatible with International Standard ISO 594.

MALE LUER VALVE WITH 6.6MM TUBE END
PART NUMBERS:
251001004, 251001004R, 251001004B, 251001004G, 251001004Y



PERFORMANCE CHARACTERISTICS

- Priming volume (without tubing): 0.171 ml

Flow Rate Averages

- Flow Rate @ 1 psi: 436 ml/minute (31,200 ml/hr @ 30 inch height)
- Flow Rate @ 3 psi: 762 ml/minute
- Flow Rate @ 5 psi: 1034 ml/minute

MATERIALS

- Swabable Stem: Clear Silicone
- Luer Valve Body:
  - Clear Polycarbonate
  - Clear Polycarbonate with Red Tint
  - Clear Polycarbonate with Blue Tint
  - Clear Polycarbonate with Green Tint
  - Clear Polycarbonate with Yellow Tint
- Luer: Clear Polypropylene
- Retainer: Clear Polypropylene
- Tube End Body: Clear Polycarbonate

PACKAGING AND SHIPPING

- Valves are bulk packaged, double bagged in clean, closed polybags
- Shipping container is clearly labeled with HR® part number, lot number and quantity

POTENTIAL STERILIZATION METHOD

- ETO and Gamma, based on raw material manufacturer's data

Important: All HR® Medical Components are shipped bulk, non-sterile, and are single patient use medical device components requiring further processing (e.g. assembly, packaging, sterilization) before clinical use. The buyer is responsible for determining effects of processing/multiple usage on these components, the appropriateness of the component in the final application, and pre/post shelf life.

December 2016