

hCG

Verify Reportable Ranges (Calibration Verification) every 6 months

Verify that the FastPack® IP System is accurate to the limits of the reportable range specified by Qualigen, Inc. by using the FastPack® IP hCG Method Verification Kit.

<p style="text-align: center;">Low</p> <p>Target <input data-bbox="311 443 539 499" type="text"/></p>	<p style="text-align: center;"><i>Write "Low Verifier", the lot number, and your initials on the peel-off FastPack® label and place it here</i></p>
<p style="text-align: center;">Mid</p> <p>Range <input data-bbox="300 863 539 919" type="text"/></p>	<p style="text-align: center;"><i>Write "Mid Verifier", the lot number, and your initials on the peel-off FastPack® label and place it here</i></p>
<p style="text-align: center;">High</p> <p>Target <input data-bbox="311 1283 539 1339" type="text"/></p>	<p style="text-align: center;"><i>Write "High Verifier", the lot number, and your initials on the peel-off FastPack® label and place it here</i></p>

The Low Verifier result must be < 1.8 mIU/mL and the High Verifier result must be > 1000.0 mIU/mL to accept the Manufacturer's Reportable Range. If either result is different than above, record the reportable range based on actual observed values. The mid value must be within the range designated on the range card. If it is not, repeat the test. If any value is out of range after repeating the test, contact Qualigen System Support.

- Accept Manufacturer Reportable Range: 1.8 mIU/mL to 1000.0 mIU/mL
- DO NOT Accept. Derived Reportable Range: _____