



# REPSOL HEALTHCARE HPP08G

HPP08G is a polypropylene homopolymer with a medium-low fluidity intended for injection moulding applications that require fair impact resistance balanced with the typical stiffness of a homopolymer. Articles manufactured with this grade have excellent chemical resistance, are easily decorated and can accept different colouring systems.

This grade has been produced with a Phthalate Free Catalytic system.

## Typical Applications

- ✓ Pharmaceutical packaging and healthcare applications
- ✓ Caps & Closures
- ✓ Pouches

Recommended melt temperature range from 190 to 250°C. Processing conditions should be optimised for each production line. Physical blends with other materials might cause incompatibilities

PROPERTIES	VALUE	UNIT	TEST METHOD
<b>General</b> Melt Flow Rate (230 °C; 2.16 kg) Density at 23°C Melting temperature	8 905 164	g/10' kg/m <sup>3</sup> °C	ISO 1133 ISO 1183 Internal (DSC)
<b>Mechanicals</b> Flexural Modulus Charpy Impact Strength Notched 23°C	1500 4	MPa kJ/m <sup>2</sup>	ISO 178 ISO 179
<b>Thermal</b> Heat Deflection Temperature 0.45MPa	82	°C	ISO 75

For further information, please contact our Technical Service and Development Laboratory or our Customer Care Service.

HPP08G complies in composition with EP 3.1.3, EP 3.1.6 and USP.

## STORAGE

HPP08G should be stored in a dry atmosphere, on a paved, drained and not flooded area, at temperatures under 60°C and protected from UV radiation. Storage under inappropriate conditions could initiate degradation processes which may have a negative influence on the processability and the properties of the transformed product.

\*This product(s) may not be used in:

- (i) any U.S. FDA Class I and/or European Union Class I Medical Devices (Non-invasive devices), without prior notification to Seller for each specific product and application
- (ii) the manufacture of any of the following, without prior written approval by Seller for each specific product and application: U.S. FDA Class II and/or European Union Class II Medical Devices:
  - Category IIa: Invasive devices with limited risk: e.g. syringes, lancets, insulin pens.
  - Category IIb: Invasive devices with higher risk: e.g. pouches for dialysis processes.
- (iii) in U.S. FDA Class III, and/or European Class III Medical Devices; Category III: Very high risk devices: long-term (> 29 days) or permanent implants, long term (> 29 days) applications in direct contact with any body part or any body fluid.

\*Repsol makes no warranties, express or implied, which extend beyond the description contained herein. Nothing herein shall constitute any warranty of merchantability or fitness for a particular purpose

\*Repsol accepts no liability from the use of Repsol products in conjunction with other materials.

\* Before using a product sold by Repsol, users should make their own independent determination that the product is safe, lawful and technically suitable for the intended use.

May 2017