



## Purell RP375RT

Developmental Grade - Gamma ray sterilizable syringes

### Features

- Superior transparency
- Good balance of mechanical and optical properties after sterilization
- Gamma ray sterilizable grade
- Compliance with USP class VI
- DMF listing as reference number 36412

### Typical applications

- Disposable syringes
- Diagnostic equipment
- Centrifuge tubes
- Vials/ Cryovials
- Pipette tips
- PCR tubes

PP Resin Properties (a)	Value	ASTM METHOD(b)
Melt flow rate (230°C / 2.16 kg), g/10min	25	D1238
Density, g/cm <sup>3</sup>	0.90	D792B
Tensile strength at yield, MPa	30	D638
Elongation at yield, %	14	D638
Flexural modulus, MPa	1200	D790A
Notched izod impact strength at 23°C, J/m	45	D256A
Deflection temperature, at 455 kPa, °C	90	D648
Haze(%), 0.8 mm Injection molded plaque	10.5	D1003

(a) Values shown are averages and are not to be considered as specifications.

(b) ASTM test methods are the latest under the Society’s current procedures. All molded specimens are prepared by injection molding.

Note: Due to the fact that different regulations in each country set different details of compliance, users of *Purell* RP375RT are recommended to undertake their own investigation of the requirements and comply with each regulation set forth, for instance, in applicable local F&DA requirements. Ultimately the users must make their own determination that their use of *Purell* RP375RT is safe, lawful and technically suitable in their intended applications.

This product(s) may not be used in the manufacture of any of the following, without prior written approval by Seller for each specific product and application:

- (i) U.S. FDA Class I or II Medical Devices; Health Canada Class I, II or III Medical Devices; European Union Class I or II Medical Devices;
- (ii) film, overwrap and/or product packaging that is considered a part or component of one of the aforementioned medical devices;
- (iii) packaging in direct contact with a pharmaceutical active ingredient and/or dosage form that is intended for inhalation, injection, intravenous, nasal, ophthalmic (eye), digestive, or topical (skin) administration;
- (iv) tobacco related products and applications, electronic cigarettes and similar devices.
- (v) safety components in automotive applications, for example: air bags, air bag unit housings and covers, seat belt mechanisms, brake systems, pedals and pedal supports, steering systems.

The product(s) may not be used in:

- (i) U.S. FDA Class III Medical Devices; Health Canada Class IV Medical Devices; European Class III Medical Devices;
- (ii) applications involving permanent implantation into the body;
- (iii) life-sustaining medical applications.

All references to U.S. FDA, Health Canada, and European Union regulations include another country’s equivalent regulatory classification.

### Storage

The resin should be stored in a dry location with good housekeeping practices during storage, transferring and handling. Process enclosures and adequate ventilation should be used to avoid excessive dust accumulation. Resin should be protected from direct sunlight, temperatures above 40°C and high atmospheric humidity during storage. Higher storage temperatures may reduce the storage time. The container should be kept closed to prevent contamination. For the additional recommended storage conditions, please refer to SDS.

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HMC Polymers is certified according to ISO 9001 and 14001  
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### The purpose of this document is only for technical support of the use of the product.

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