

Technical Data Sheet

Moplen RP348SK

Polypropylene, Random Copolymer

**Product Description**

Moplen RP348SK is a nucleated and high fluity polypropylene random copolymer, with good processability, excellent clarity in low processing temperature, good mechanical property balance, high productivity with less void formation. Typical customer applications are thin walled articles (TWIM) for food and non food applications, transparent containers and boxes, houseware, DVD casing.

Regulatory Status

For regulatory compliance information, see Moplen RP348SK [Product Stewardship Bulletin \(PSB\) and Safety Data Sheet \(SDS\)](#).

Status	Commercial: Active
Availability	Africa-Middle East; Asia-Pacific
Application	Clear Containers; Housewares; Multi Media Packaging
Market	Consumer Products; Rigid Packaging
Processing Method	Injection Molding
Attribute	Good Chemical Resistance; Good Dimensional Stability; Good Impact Resistance; Good Processability; High Clarity; High Flow; Medium Rigidity; Random Copolymer

Typical Properties	Nominal Value	Units	Test Method
Physical			
Melt Flow Rate, (230 °C/2.16 kg)	35	g/10 min	ASTM D1238
Density	0.90	g/cm ³	ASTM D792
Mechanical			
Flexural Modulus	1050	MPa	ASTM D790
Tensile Strength at Yield	30	MPa	ASTM D638
Tensile Elongation at Yield	14	%	ASTM D638
Impact			
Notched Izod Impact Strength, (23 °C)	50	J/m	ASTM D256
Hardness			
Rockwell Hardness, (R-Scale)	93		ASTM D785
Thermal			
Deflection Temperature Under Load, (0.46 N/mm ²)	87	°C	ASTM D648
Optical			
Haze	18	%	ASTM D1003

Notes

These are typical property values not to be construed as specification limits.

Company Information

For further information regarding the LyondellBasell company, please visit <http://www.lyb.com/>.

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Users should review the applicable Safety Data Sheet before handling the product.

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.

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