

Technical Data Sheet

Moplen EP540V

Polypropylene, Impact Copolymer



Product Description

Moplen EP540V is a polypropylene impact copolymer, with very high melt flow, low warpage, low odor, and good stiffness-impact balance. Typical applications are component of compounds for moulding automotive interior and exterior parts like door panels and bumper fascia, direct injection moulding of multimedia packaging like DVD cases, and injection moulding of general housewares and thin wall containers.

Regulatory Status

For regulatory compliance information, see Moplen EP540V [Product Stewardship Bulletin \(PSB\) and Safety Data Sheet \(SDS\)](#).

Status	Commercial: Active
Availability	Africa-Middle East; Asia-Pacific; Australia and New Zealand
Application	Bumpers; Exterior Automotive Applications; Housewares; Interior Automotive Applications; Multi Media Packaging; Opaque Containers
Market	Automotive; Consumer Products; Rigid Packaging
Processing Method	Injection Molding
Attribute	High Flow; Impact Copolymer; Low to No Odor; Low Warpage

Typical Properties	Nominal Value	Units	Test Method
Physical			
Melt Flow Rate, (230 °C/2.16 kg)	110	g/10 min	ASTM D1238
Density	0.90	g/cm ³	ASTM D792
Mechanical			
Flexural Modulus	1600	MPa	ASTM D790
Tensile Strength at Yield	24	MPa	ASTM D638
Tensile Elongation at Yield	4	%	ASTM D638
Impact			
Notched Izod Impact Strength			
(23 °C)	40	J/m	ASTM D256
(-20 °C)	30	J/m	ASTM D256

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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Date: 7/25/2017

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Recipient Tracking #:
Request #: 907420

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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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