

Technical Data Sheet

Moplen EP300M

Polypropylene, Impact Copolymer



Product Description

Moplen EP300M is a heterophasic copolymer for injection moulding . The product offers an excellent balance of stiffness, impact strength and processability. Typical applications fo *Moplen* EP300M are thin walled packaging, housewares and automotive parts.

Regulatory Status

For regulatory compliance information, see *Moplen* EP300M [Product Stewardship Bulletin \(PSB\) and Safety Data Sheet \(SDS\)](#).

Status	Commercial: Active
Availability	Africa-Middle East; Europe
Application	Crates; Housewares; Industrial
Market	Automotive; Compounding; Consumer Products; Rigid Packaging
Processing Method	Compounding; Injection Molding
Attribute	Good Impact Resistance; Good Processability; Good Stiffness

Typical Properties	Nominal Value	Units	Test Method
Physical			
Melt Flow Rate, (230 °C/2.16 kg)	7	g/10 min	ISO 1133-1
Density	0.89	g/cm ³	ISO 1183-1
Mechanical			
Flexural Modulus	1000	MPa	ISO 178
Tensile Stress at Break	17.5	MPa	ISO 527-1, -2
Tensile Stress at Yield	21.5	MPa	ISO 527-1, -2
Tensile Strain at Break	600	%	ISO 527-1, -2
Tensile Strain at Yield	7	%	ISO 527-1, -2
Impact			
Charpy Impact Strength - Notched			
(23 °C, Type 1, Edgewise, Notch A)	45	kJ/m ²	ISO 179
(0 °C)	9	kJ/m ²	ISO 179
(-20 °C, Type 1, Edgewise, Notch A)	8	kJ/m ²	ISO 179
Thermal			
Vicat Softening Temperature, (A50)	143	°C	ISO 306
Deflection Temperature Under Load, (0.45 MPa, Unannealed)	77	°C	ISO 75B-1, -2
DSC Melting Point	163	°C	DSC

Notes

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

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Recipient Tracking #:
Request #: 907891

Processing Techniques

Specific recommendations for resin type and processing conditions can only be made when the end use, required properties and fabrication equipment are known.

Company Information

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

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- (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.

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