

Pakro Kimya Dış Ticaret A.Ş.

İstoç Ticaret Merkezi Aktem Plaza Kat:5 Daire:33 Bağcılar/İstanbul

lyondellbasell

Moplen HP400R

Polypropylene, Homopolymer

Product Description

Moplen HP400R is a homopolymer polypropylene used by customers for injection moulding applications. It exhibits a high fluidity combined with a good stiffness.

Moplen HP400R is suitable for food contact.

Product Characteristics					
Status Commerce		ial: Active			
Test Method used	IS	50			
Availability	E	Europe, Africa-Middle East			
Features	Н	High Flow , Good Stiffness			
Typical Customer Ap	plications F	urniture,	Housewares		
Typical Properties			Method	Value	Unit
Physical					
Density			ISO 1183	0.9	g/cm ³
Melt flow rate (MFR) (2	30°C/2.16kg)		ISO 1133	25	g/10 min
Melt volume flow rate	230°C/2.16kg)		ISO 1133	34	cm³/10min
Mechanical					
Tensile Modulus			ISO 527-1, - 2	1350	MPa
Tensile Stress at Yield			ISO 527-1, - 2	32	MPa
Tensile Strain at Break			ISO 527-1, - 2	>50	%
Tensile Strain at Yield			ISO 527-1, - 2	10	%
Impact					
Charpy unnotched imp	act strength		ISO 179		
(23 °C, Type 1, Edgew	ise)			105	kJ/m²
(0 °C, Type 1, Edgewis	e)			25	kJ/m²
Charpy notched impact A)	strength (23 °C, Type 1, Edgewise	e, Notch	ISO 179	2	kJ/m²
Hardness					
Ball indentation hardne	ess (H 358/30)		ISO 2039-1	70	MPa
Thermal					
Heat deflection temper	ature B (0.45 MPa) Unannealed		ISO 75B-1, - 2	90	°C
Vicat softening temperature A/50 ISO 306 154 °C					°C

Vicat softening temperature B/50

Notes

Typical properties; not to be construed as specifications.

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LyondellBasell markets this product through the following entities:

Equistar Chemicals, LP Basell Sales & Marketing Company B.V. Basell Asia Pacific Limited Basell International Trading FZE LyondellBasell Australia Pty Ltd

For the contact details of the LyondellBasell company selling this product in your country, please visit http://www.lyb.com/.

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This product(s) may not be used in:

(i) any U.S. FDA Class I, Health Canada Class I, and/or European Union Class I Medical Devices, without prior notification to Seller for each specific product and application; or

(ii) the manufacture of any of the following, without prior written approval by Seller for each specific product and application: (1) U.S. FDA Class II, Health Canada Class II or Class III, and/or European Union Class II Medical Devices; (2) film, overwrap and/or product packaging that is considered a part or component of one of the aforementioned Medical Devices; (3) 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; (4) tobacco related products and applications; (5) electronic cigarettes and similar devices; and (6) pressure pipe or fittings that are considered a part or component of a nuclear reactor.

(iii) Additionally, the product(s) may not be used in: (1) U.S. FDA Class III, Health Canada Class IV, and/or European Class III Medical Devices; (2) applications involving permanent implantation into the body; (3) life-sustaining medical applications; and (4) lead, asbestos or MTBE related applications.

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

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

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