



Hostacom SB572C

Compounded Polyolefin

Product Description

Hostacom SB572C fractional melt flow, 1,100 MPa flexural modulus, precolored, extrusion grade polypropylene copolymer has balanced strength and processability.

Product Characteristics

Status	Commercial: Active
Test Method used	ASTM
Availability	North America, Asia-Pacific, Australia/NZ, Africa-Middle East, Latin America
Processing Methods	Extrusion Pipe Sheet and Semi Finished Products
Features	Copolymer, Low Flow , Good Melt Strength , Good Processability, High Strength
Typical Customer Applications	Panels & Profiles, Soft Profile & Sheets

Typical Properties	Method	Value	Unit
Physical			
Density -Specific Gravity	ASTM D 792	0.91	g/cm ³
Melt Flow Rate (230°C/2.16kg)	ASTM D 1238	0.6	g/10 min
Mechanical			
Flexural Modulus	ASTM D 790	1100	MPa
Tensile Strength @ Yield	ASTM D 638	23	MPa
Tensile Strength @ Break	ASTM D 638	34	MPa
Tensile Elongation @ Brk	ASTM D 638	650	%
Impact			
Notched Izod Impact	ASTM D 256		
(23 °C)		530	J/m
(-30 °C)		130	J/m
Additional Information			
Mold shrink, Linear-Flow	ASTM D 955		
Note: Please contact Basell for shrinkage recommendations.			

Additional Properties

Note: Mold shrinkage values are determined on laboratory injection molded 100 mm x 150 mm x 3.2 mm plaques and, as such, are not necessarily representative of actual field data. Since, for example, wall thickness, gate type and location, flow length and paint oven temperature affect final part dimensions, it is recommended that you contact your Basell representative before any tools are cut.

Notes

Typical properties; not to be construed as specifications.

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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