

Pakro Kimya Dış Ticaret A.Ş.

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## Moplen HP602N

lyondellbasell

Polypropylene, Homopolymer

## **Product Description**

Moplen HP602N is a polyproylene homopolymer manufactured using the Spheripol process. It is known for its optimized balance of stiffness and toughness, and is particularly suitable for injection molding. Potential end use applications include housewares, toys, closures and other packaging items.

Product Characteristics					
Status Commercial:		I: Active			
Test Method used	ASTM	ASTM			
Availability	Asia-Pacific, Australia/NZ, Africa-Middle East				
Processing Methods	Injection Molding Good Color Stability, High Gloss , Homopolymer, Good Processability, Good Processing Stability, Good Stiffness , Good Surface Finish, Good Toughness, Low Warpage				
Features					
Typical Customer Applications		Caps & Closures, Housewares, Opaque Containers, Sports, Leisure and Toys			
Typical Properties		Method	Value	Unit	
Physical					
Melt Flow Rate (230°C/2.16kg)		ASTM D 1238	12	g/10 min	
Note: ASTM D1238L					
Density		ASTM D 1505	0.9	g/cm³	
Mechanical					
Flexural Modulus		ASTM D 790	16000	kg/cm²	
Tensile Strength @ Yield		ASTM D 638	350	kg/cm²	
Tensile Elongation @ Yield		ASTM D 638	12	%	
Impact					
Notched Izod Impact (23 °C)		ASTM D 256	3	kg-cm/cm	
Hardness					
Rockwell Hardness (R-Scale)		ASTM D 785	104		
Thermal					
Heat deflection temperature at 0.46 N/mm2		ASTM D 648	107	°C	

## Notes

Typical properties; not to be construed as specifications.

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- Equistar Chemicals, LP
- Basell Sales & Marketing Company B.V.
- Basell Asia Pacific Limited
- Basell International Trading FZE
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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:

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(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.

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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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