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## DaelimPoly PP UE332K **Block copolymer**

## **Product Description**



DaelimPoly UE332K is the polypropylene blockcopolymermanufactured by Ulsan PPunder thelicenseof Lyondellbasell using the Spheripol process. This productisparticularlysuitableforinjectionmoldingoftoy, crates. DaelimPoly UE332KresinmeetstheFDArequirementsintheCodeof Federal Regulations in 21 CFR 177.1520 for food contact.

Features Optimized balance of stiffness and toughness / High impact strength at low temperature / High Stiffness / Low warpage

Market Consumer products, Industrial & Pipe

Application Toy / Crate / Housewares

ASTM Data			
Typical Properties	Nominal Value	Units	Test Method
Melt Flow Rate (230°C,2.16kg)	5	g/10 min	ASTM D1238L
Density	0.9	g/cm3	ASTM D1505
Flexural Modulus	11500	kg/cm2	ASTM D790
Tensile Strength at Yield	250	kg/cm2	ASTM D638
Elongation at Yield	9	%	ASTM D638
Izod Impact Strength (23°C)	15	kg <sup>f</sup> cm/cm	ASTM D256
Izod Impact Strength (-20°C)	7	kg <sup>f</sup> cm/cm	ASTM D256
Rockwell Hardness	95	R-Scale	ASTM D785
Vicat Softening Point	150	°C	ASTM D1525
HDT (0.46 N/mm²)	88	°C	ASTM D648

<sup>1)</sup> The above values are typical property values for reference only not be construed as specification limits.

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4) The use of this product(s) is strictly prohibited in

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- ii. applications involving permanent implantation into the body:
- iii. life-sustaining medical applications; or
- iv. lead, asbestos or MTBE related applications.

Users are solely liable for any injuries or damages resulting from any use of this product(s) in the above categories and Seller shall have no liability whatsoever.

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- i. U.S. FDA Class I, Health Canada Class I, and/or European Union Class I medicaldevices;
- ii. U.S. FDA Class II, Health Canada Class II or Class III, and/or European Union Class II Medical Devices;
- iii. film, overwrap and/or product packaging that is considered a part or component of one of the aforementioned Medical Devices:
- iv. packaging in direct contact with an active pharmaceutical ingredient and/or dosage form that is intended for inhalation, injection, intravenous, nasal, ophthalmic (eye), digestive, or topical (skin) administration;
- v. tobacco related products and applications;
- vi. electronic cigarettes and similar devices; or
- vii. pressure pipe or fittings that are considered a part or component of a nuclear reactor
  - \* All references to U.S. FDA, Health Canada, and European Union regulations include another country's equivalent regulatory classification.
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