Product Information

page 1 of 2

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Ultramid[®] B36 LN 01

Product description

® = registered trade mark of

BASF SE

Ultramid[®] B36 LN 01 is a polyamide 6 grade of intermediate to high viscosity that is well suited for the production of cast and blown film. Clarity and thermoformability are enhanced by the incorporation of nucleating and slip agent.

Specification		Test method	Unit	Value
	Relative Viscosity (RV) 1% [m/v] in 96% [m/m] sulfuric acid	According to ISO 307 (calculated by Huggins method)		3.49 - 3.71
	Viscosity Number (VN) 0,5% [m/v] in 96% [m/m] sulfuric acid	According to ISO 307	ml/g	210 - 226
	Moisture content	According to ISO 15512	% [m/m]	max. 0.06
	Extractables	According to ISO 6427- chips not ground/16h	% [m/m]	max. 0.6
	Lubricant	BASF method	(mg/kg)	250 - 550
	Nucleating agent	BASF method	(mg/kg)	250 - 550
	Film grade	BASF method		1 - 3

General properties		Test method	Unit	Typical value
	Melting point	According to ISO 3146	°C	220
	Density	According to ISO 1183	g/cm ³	1.12 - 1.15
	Bulk density		kg/m³	700
	Pellet size		mm	2 - 2.5
	Pellet shape			cylindrical
	Water absorption, 23°C/50% rh		%	2.6
	Water absorption, saturation in water 23°C		%	9.5

Supply form and storage	Ultramid [®] B36 LN 01 is supplied pre-dried and ready for processing in a varie- ty of moisture proof containers, such as boxes, bigbags (Asia) and bulk con- tainers. The material must be protected against moisture during storage. A storage time of 12 months should not be exceeded. Opened bags should be used up immediately in order to prevent moisture pickup.				
Food legislation	Ultramid [®] film grades (Ultramid [®] A, B, C, Flex F) comply with the current legis- lation on plastics in contact with food in several regions. If you need details on the food approval status of a particular Ultramid [®] grade, please contact BASF directly at plastics.safety@basf.com. We will be happy to provide you with an up-to-date declaration of conformity based on the cur-rent legal regulations.				
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Medical disclaimer	BASF has not developed or tested its plastics especially for the use in medi- cal devices (defined in risk classes I to III according to the European and US Medical Device legislation) and pharmaceutical applications. Therefore BASF makes no warranties, express or implied, concerning the suitability of any BASF plastics for use in any medical device and pharmaceutical applications. BASF does not supply its plastics for the manufacture of implants of any risk class. Please inform us in advance, if you intend to use BASF plastics in medical devices or pharmaceutical applications.				
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