

Adflex Q 302 B

Advanced Polyolefin

Product Description

Adflex Q 302 B is a reactor TPO (thermoplastic polyolefin) manufactured using LyondellBasell's proprietary *Catalloy* process technology. It is an innovative material that resists gas fading and brings a warm and sensual feel, creating a truly unique sensation.

Part of a family of polymers with a soft, velvety texture, *Adflex* Q 302 B resin enables packagers to give their products a distinct edge over conventional plastics. Bottles blow molded from *Adflex* Q 302 B resin conveys an upscale, quality image that enhances point of purchase appeal. The unique texture is also ideal for applications requiring a no-slip surface, such as shower soaps and lotions.

For regulatory compliance information, see the Adflex Q 302 B Product Stewardship Bulletin (PSB).

Product Characteristics					
Status	Commercial: A	ctive			
Test Method used	ISO	ISO			
Availability	North America, Asia-Pacific, Australia/NZ, Africa-Middle East, Latin America				
Processing Methods	Extrusion Blow	Extrusion Blow Molding, Injection Blow Molding			
Features	Stress Crackin Heat Resistand	l Resistance, Higl g Resistance), Ga ce , Good Punctu erial, High Streng	as-fading F re Resista	Resistant, High nce ,	
Typical Customer Applications	Blow Moulding	Applications, Spo	orts, Leisu	re and Toys	
Typical Properties		Method	Value	Unit	
Physical					
Density (Method A)		ISO 1183	0.88	g/cm³	
Melt flow rate (MFR) (230 °C/ 2.16 kg)		ISO 1133	0.9	g/10 min	
Mechanical					
Tensile Stress at Break (°C)		ISO 527-1, -2	9.5	MPa	
Tensile Stress at Yield (23 °C)		ISO 527-1, -2	8.5	MPa	
Tensile Strain at Break		ISO 527-1, -2	>500	%	
Tensile Strain at Yield		ISO 527-1, -2	37	%	
Flexural modulus (23 °C)		ISO 178	350	MPa	
Impact					
Notched izod impact strength (23 °C, Type 1, Notch A)		ISO 180	47	kJ/m²	
Hardness					
Shore hardness D		ISO 868/ASTM D 2240	47		

Note: 15 seconds

Notes

Typical properties; not to be construed as specifications.

${f \mathbb{S}}$ LyondellBasell Industries Holdings, B.V. 2014
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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.

(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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