

## Adflex KS021P

### Advanced Polyolefin

#### Product Description

Adflex KS021P is a reactor TPO (thermoplastic polyolefin) manufactured using LyondellBasell's proprietary *Catalloy* process technology. It is suitable for the extrusion and calendering of soft film and sheet, for the impact modification of polypropylene, as well as monolayer and multilayer air quenched blown films.

It is also used by our customers as high cold impact resistance material for automotive color-matched interior trim applications. Key characteristics are flexibility and low temperature impact resistance.

The grade is available in natural pellet form.

For regulatory compliance information see *Adflex* KS021P Product Stewardship Bulletin (PSB).

#### Product Characteristics

<b>Status</b>	Commercial: Active
<b>Test Method used</b>	ISO
<b>Availability</b>	North America, Asia-Pacific, Australia/NZ, Africa-Middle East, Latin America
<b>Processing Methods</b>	Extrusion Flat-die, Blown Film, Calendering, Extrusion Thermoforming
<b>Features</b>	Good Chemical Resistance, High ESCR (Environmental Stress Cracking Resistance), Good Flexibility, Low Temperature Impact Resistance, Good shape retention
<b>Typical Customer Applications</b>	Film, Interior Applications, Sports, Leisure and Toys

Typical Properties	Method	Value	Unit
<b>Physical</b>			
Density (Method A %)	ISO 1183	0.88	g/cm <sup>3</sup>
Melt flow rate (MFR) (230 °C/ 2.16 kg)	ISO 1133	0.9	g/10 min
<b>Mechanical</b>			
Tear Strength	ASTM D 624	85 N/mm	
<i>Note: (Graves, Die C, 50mm/min) - Load/Width @ Max Load</i>			
Tensile Stress at Break (23 °C, 50 mm/min)	ISO 527-1, -2	13	MPa
Tensile Stress at Yield (23 °C, 50 mm/min)	ISO 527-1, -2	8	MPa
Tensile Strain at Break (23 °C, 50 mm/min)	ISO 527-1, -2	>800	%
Tensile Strain at Yield (23 °C, 50 mm/min)	ISO 527-1, -2	38	%
Flexural modulus (23 °C, 2 mm/min, Chord)	ISO 178	300	MPa
<b>Impact</b>			
Notched izod impact strength (23 °C, Type 1, Notch A)	ISO 180	52	kJ/m <sup>2</sup>
<i>Note: Failure Mode: Partial</i>			
(-40 °C, Type 1, Notch A)		86	kJ/m <sup>2</sup>
<i>Note: Failure Mode: Partial</i>			
Multiaxial Impact Strength	ASTM D3763		
(Energy@ Max Load +23°C, 6.6m/s, 3.2mm plq; Failure Mode: Ductile)		17	J
(Energy@ Max Load -40°C, 6.6m/s, 3.2mm plq; Failure Mode: Ductile)		23	J

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**Hardness**

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Shore hardness (Shore D)	ISO 868	38
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Note: 15 seconds

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**Thermal**

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Melting temperature	161	°C
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Note: (ISO 11357-3)

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Heat deflection temperature B (0.45 MPa) Unannealed	ISO 75B-1, -2	48	°C
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Heat deflection temperature A (1.80 MPa) Unannealed	ISO 75A-1, -2	38	°C
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Vicat softening temperature (A50 (50°C/h 10N))	ISO 306	67	°C
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**Additional Information**

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Mold shrinkage	ISO 294-4
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Note: Please contact Equistar for shrinkage recommendations

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**Additional Properties**

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**AUTOMOTIVE SPECIFICATIONS:**

Chrysler MS-DB590 CPN 3900

Ford WSS M4D777-A8

GM GMW15072-170020

**Notes**

Typical properties; not to be construed as specifications.

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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 Safety Data Sheet before handling the product.

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