

Safripol a division of KAP Diversified Industrial (Pty) Ltd

# **Safrene**<sup>®</sup> (High Density PolyEthylene)

#### **Technical Data Sheet**

## M 7650

#### High Density Polyethylene for Blow Moulding and Extrusion Applications

Safrene® M 7650 High Density Polyethylene Resin is a bimodal high molecular mass grade suitable for extrusion blow moulding of containers greater than 20 litre capacities. It has a high rigidity and hardness with very good impact strength and excellent environmental stress-crack resistance properties.

Safrene® M 7650 High Density Polyethylene Resin is primarily designed for the blow moulding of large hollow articles but may be used for smaller mouldings where exceptional environmental stress-crack resistance and impact strength are required.

#### Typical Applications

- General purpose containers
- Sheet extrusion
- Optical fibre sheathing

PROPERTIES(1)	VALUE	UNIT	TEST METHOD
Physical Melt Flow Rate, 190 °C/ 5 kg Melt Flow Rate, 190 °C/ 21.6 kg Density <sup>(2)</sup>	0.35	g/10min	ISO 1133
	9.5	g/10min	ISO 1133
	0.953	g/cm <sup>3</sup>	ISO 1183
Mechanical (3) Hardness Shore D Tensile Yield <sup>(4)</sup> Ultimate Tensile <sup>(4)</sup> Ultimate Elongation <sup>(4)</sup> Flexural Modulus Environmental Stress-Crack Resistance <sup>(5)</sup>	62	Units	ISO 868
	26	MPa	ISO 527
	40	MPa	ISO 527
	>600	%	ISO 527
	1300	MPa	ISO 178
	>1000	Hours	ASTM D1693
Impact (3) Charpy Notched Impact Strength (23 °C) Charpy Notched Impact Strength (-30 °C)  Thermal Vicat Softening Point (10N) Vicat Softening Point (50N) Crystalline Melting Range	23	kJ/m²	ISO 179
	20	kJ/m²	ISO 179
	128	°C	ISO 306/A50
	80	°C	ISO 306/B50
	130-133	°C	ISO 3146

<sup>(1)</sup> Typical values; not to be construed as specification limits.

<sup>(2)</sup> Unannealed

Typical processing conditions	Blow Moulding	Extrusion
Feed Zone (°C)	180	180
Zone1 (°C)	190 - 200	190 - 200
Zone 2 (°C)	200 - 210	200 - 210
Zone 3 (°C	200	200
Die (°C)	200	200
Melt Temp (°C)	200	200

<sup>(3)</sup> Compression moulded samples

<sup>(4)</sup> Test speed 50 mm/min

<sup>(5)</sup> Bell test condition B, 50°C, 10% Igepal, F50.



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Product Stewardship	At Safripol*, protecting people and the environment will be a part of everything we do and every decision we make. Each employee has a primary responsibility in ensuring that our products and operations meet applicable government standards.  Our goal is to eliminate all injuries, prevent adverse environment and health impacts, reduce wastes and emissions and promote resource conservation at every stage of the life cycle of our products. The success of this rests with each and every individual involved with Safripol* products throughout the life cycle.		
Food Contact Compliance	Safrene® M 7650 High Density Polyethylene Resin should comply with Commission Regulation (EU) No 10/2011 and with U.S. FDA 21 CFR 177.1520(c)3.1c food contact regulations when used unmodified and processed according to good manufacturing practices for food contact applications. The purchaser remains responsible for determining whether the use complies with all relevant regulations.		
Customer Notice	Safripol* strongly encourages its customers to review both their manufacturing processes and their applications of Safripol* products from the standpoint of human health and environmental quality to ensure that Safripol* products are not used in ways for which they are not intended or tested. Safripol* personnel are available to answer your questions and to provide reasonable technical support. Safripol* product literature, including safety data sheets, should be consulted prior to use of Safripol* products. Current safety data sheets are available from Safripol*.		
Safripol Medical Application Policy	<ul> <li>Safripol* will not knowingly sell or sample any product or service (*Product*) into any commercial or developmental application that is intended for:</li> <li>a. permanent (Long term) contact with internal body fluids or internal body tissues. Long term is a use which exceeds 72 continuous hours.</li> <li>b. use in cardiac prosthetic devices regardless of the length of time involved; (Cardiac prosthetic devices include, but are not limited to, pacemaker leads and devices, artificial hearts, heart valves, intra-aortic balloons and control systems, and ventricular bypass assisted devices);</li> <li>c. use as a critical component in medical devices that support or sustain human life; or</li> <li>d. use specifically by pregnant women or in applications designed specifically to promote or interfere with human reproduction.</li> <li>Additionally, all Products intended for use in pharmaceutical applications, other than pharmaceutical packaging, must pass the current Pharmaceutical Liability Guidelines.</li> </ul>		
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Additional Information			
Our contact details			
		Gauteng: Tel: +27 (0) 11 575 4549	
2021 South Africa		<b>Kwa-Zulu Natal</b> : Tel: +27 (0) 31 450 4111	
Website: www.safripol.com Email: info@safripol.com		Email: info@safripol.com	
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