

**ALFATER XL A50I 3EF0010**

(Last update: 08.09.2023)

**MOCOM**

Base Polymer	Thermoplastic Vulcanisate (PP+EPDM)
Colour	black
Service Temperature	max. range (depending on load, geometry, etc.) -40-125 °C
Special Features	easy flow, high surface quality, black fulfills weathering dry, hot (Kalahari, PV3929), black fulfills weathering moist, hot (Florida, PV3930), good weathering resistance, good chemical resistance, good ozone resistance, good hot air resistance, good hydrolysis resistant
Market Segment	Automotive, building and construction, do-it-yourself, electrical and electronic, sport and leisure, Furniture
Application Area	injection moulded parts
Typical Applications	seals, sealing lip, handles, soft touch elements, impact protection / shock absorber, protection caps, thin walled parts, profile, cable ducts, 2K components
Adhesion to	Polypropylene Homopolymer, Polypropylene Copolymer, Polyethylene, Thermoplastic Vulcanisate (PP+EPDM), Thermoplastic Olefin Elastomer, Thermoplastic Elastomer general

Pre-Drying Conditions	in a dry air (dessiccant) dryer 60-70 °C for 2-4 h max. moisture content <0,10 %
Processing Injection Moulding	melt temperature 190-220 °C mould temperature 10-60 °C injection speed middle - high back pressure (hydraulic) >3 bar holding pressure 50-70% of injection pressure
Storage	dry, protected from light not above 30°C tightly sealed do not stack the pallets

Properties	Value	Dimension	Test Norm
<b>Mechanical Properties</b>			
Tensile Stress at 100% Elongation (Type 2)	1.9	MPa	ISO 37
Tensile Strength (Type 2)	3.6	MPa	ISO 37
Tear Strength (Method Ba)	18	kN/m	ISO 34-1
Elongation at Break (Type 2)	385	%	ISO 37
Shore Hardness A (15s)	50	-	ISO 868
<b>Thermal Properties</b>			
Compression Set (22h / 100 °C / 25% / Type A)	42	%	ISO 815
Compression Set (22h / 70 °C / 25% / Type A)	37	%	ISO 815
Compression Set (72h / 22 °C / 25% / Type A)	21	%	ISO 815

## Technical Data Sheet

# Alfater XL<sup>®</sup>



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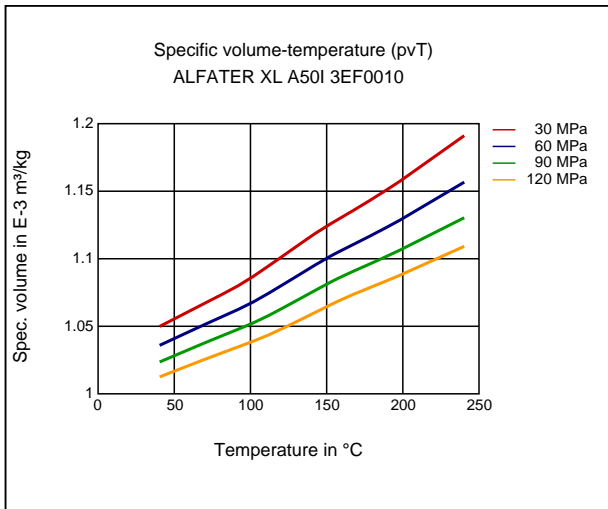
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Compression Set (72h / 125 °C / 25% / Type A)	62	%	ISO 815
<b>Electrical Properties</b>			
Tracking Resistance (CTI)	600	-	IEC 60112
<b>Rheological Properties</b>			
Spiral Flow (2 mm wall thickness)	55	cm	-
<b>Physical Properties</b>			
Density	950	kg/m <sup>3</sup>	ISO 1183
<b>Ageing Properties</b>			
Ozone resistance (method A)	passed	-	ISO 1431-1A

### Diagrams

#### Specific volume-temperature (pvT)



#### Liability Exclusion

These are guide values and not a specification. The test values mentioned are representative values only and not binding minimum or maximum figures. These test values have been determined on standardised test specimens and can be affected by pigmentation, mould design and processing conditions.

Any information given on the chemical and physical characteristics of our products, including, without limitation, technical advice on applications, whether verbally, in writing or by testing the product, is given to the best of our knowledge and in good faith and does not exempt the buyer from carrying out their own investigations and tests in order to ascertain the product's specific suitability for the purpose intended.

The buyer is solely responsible for confirming the suitability of the product for a particular application, its utilization and processing and must observe any applicable laws and government regulations. **NO EXPRESS OR IMPLIED RECOMMENDATION OR WARRANTY IS GIVEN WITH REGARD TO THE SUITABILITY OF THE PRODUCT FOR A PARTICULAR APPLICATION, SUCH AS, BUT NOT LIMITED TO, SAFETY-CRITICAL COMPONENTS OR SYSTEMS.**

**Healthcare uses:** the supply of any product by ALBIS for any medical, pharmaceutical or diagnostic application is conditional to an

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assessment by ALBIS in terms of compliance with ALBIS' internal risk management policy – even for products which are in general designated for use in Healthcare applications.

**Important:** irrespective of product type or designation, ALBIS does not recommend or support the use of any products it supplies which fall into the following medical, pharmaceutical or diagnostic application categories:

- risk class III applications according to EU directive 93/42/EEC
- any bodily implant application for greater than 30 days
- any critical component in any medical device that supports or sustains human life.

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