

**ALTECH PA6 A 2065/500 GF65**

(Last update: 11.04.2024)

**MOCOM**

Base Polymer	Polyamide 6
Filler/Additive System	65 % glass fibres
Special Features	heat stabilised,easy release (demoulding),injection moulding grade,high stiffness
Market Segment	various,Automotive,Machinery,electrical and electronic
Application Area	electrical and electronic (E&E),electrical components

Pre-Drying Conditions	in a dry air (dessiccant) dryer <80 °C for 2-12 h max. moisture content <0,15 %
Processing Injection Moulding	melt temperature 270-290 °C mould temperature 80-100 °C
Storage	dry, protected from light

Properties	dry/cond.	Dimension	Test Norm
<b>Mechanical Properties</b>			
Flexural Modulus	21000 / -	MPa	ISO 178
Flexural Strength	320 / -	MPa	ISO 178
Tensile Modulus	22000 / -	MPa	ISO 527
Tensile Strength at Break	195 / -	MPa	ISO 527
Tensile Elongation at Break	2 / -	%	ISO 527
Impact Strength (Charpy, 23°C)	70 / -	kJ/m²	ISO 179/1eU
Impact Strength (Charpy, -40°C)	75 / -	kJ/m²	ISO 179/1eU
Notched Impact Strength (Charpy, 23°C)	13 / -	kJ/m²	ISO 179/1eA
Notched Impact Strength (Charpy, -40°C)	11 / -	kJ/m²	ISO 179/1eA
<b>Thermal Properties</b>			
HDT / A (1,8 MPa)	217 / *	°C	ISO 75-1/-2
DSC (Melt Point)	220 / *	°C	ISO 11357
<b>Rheological Properties</b>			
Shrinkage (lengthwise, 24h)	0.1 - 0.3	%	ISO 294-4
Shrinkage (lateral, 24h)	0.1 - 0.3	%	ISO 294-4
<b>Physical Properties</b>			
Density	1780 / -	kg/m³	ISO 1183

**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



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- 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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