



ALTECH PP-H A 2020/554.01 CP FR GF20

(Last update: 11.01.2024)

MCOM

Base Polymer Polypropylene Homopolymer

Filler/Additive System 20 % glass fibres

Special Features halogen-free flame retardant, chemically coupled, heat stabilised, flame

retardant

Market Segment Automotive, electrical and electronic Typical Applications housings, fixing elements, bezels

Pre-Drying Conditions in a dry air (dessiccant) dryer 80-120 °C

for 2-3 h

in an air circulating dryer 80-120 °C

for 2-4 h

Processing Injection Moulding melt temperature 200-240 °C

mould temperature 20-70 °C

Storage dry, protected from light

Properties	Value	Dimension	Test Norm
Mechanical Properties			
Flexural Modulus	5500	MPa	ISO 178
Flexural Strength	115	MPa	ISO 178
Tensile Modulus	6200	MPa	ISO 527
Tensile Strength at Break	70	MPa	ISO 527
Tensile Elongation at Break	3.4	%	ISO 527
Impact Strength (Charpy, 23°C)	38	kJ/m²	ISO 179/1eU
Impact Strength (Charpy, -40°C)	30	kJ/m²	ISO 179/1eU
Notched Impact Strength (Charpy, 23°C)	8.5	kJ/m²	ISO 179/1eA
Notched Impact Strength (Charpy, -40°C)	6.5	kJ/m²	ISO 179/1eA
Thermal Properties			
Vicat B50	135	°C	ISO 306
HDT / A (1,8 MPa)	149	°C	ISO 75-1/-2
DSC (Melt Point)	166	°C	ISO 11357
Rheological Properties			
Melt Index (MVR)	5	cm ³ /10min	ISO 1133
MVR temperature	230	°C	-
MVR load	2.16	kg	-
Shrinkage (lengthwise, 24h)	0.2 - 0.4	%	ISO 294-4
Shrinkage (lateral, 24h)	0.4 - 0.6	%	ISO 294-4
Physical Properties			
Density	1200	kg/m³	ISO 1183

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Technical Data Sheet





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Flammability

Flammability (0.75 mm)	V-2	class	UL 94
Flammability (1.5 mm)	V-0	class	UL 94
Glow Wire (GWFI, 750°C, 1.0mm)	passed	-	DIN EN 60695
Glow Wire (GWFI, 750°C, 2.0mm)	passed	-	DIN EN 60695

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.

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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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