Technical Data Sheet





ALTECH ASA A 1000/729 UV

(Last update: 02.12.2024)

MCOM

Base Polymer Acrylonitile/Styrene/Acrylate/Copolymer

Filler/Additive System light stabiliser
Colour grey,black,orange

Special Features UV stabilised,injection moulding grade

Typical Applications various, exterior parts

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

for 2-4 h

80 °C in an air circulating dryer

for 3-6 h

dependant on moisture content max. moisture content <0,02 %

Processing Injection Moulding melt temperature 240-280 °C

mould temperature 40-80 °C

Storage dry, protected from light

Properties	Value	Dimension	Test Norm
Mechanical Properties			
Flexural Modulus	2200	MPa	ISO 178
Flexural Stress (3.5% Strain)	60	MPa	ISO 178
Tensile Modulus	2100	MPa	ISO 527
Tensile Stress at Yield	41	MPa	ISO 527
Tensile Elongation at Yield	2.9	%	ISO 527
Tensile Elongation at Break	20	%	ISO 527
Impact Strength (Charpy, 23°C)	175	kJ/m²	ISO 179/1eU
Impact Strength (Charpy, -40°C)	60	kJ/m²	ISO 179/1eU
Notched Impact Strength (Charpy, 23°C)	12	kJ/m²	ISO 179/1eA
Notched Impact Strength (Charpy, -40°C)	2.5	kJ/m²	ISO 179/1eA
Ball Indentation Hardness H358/30	75	MPa	ISO 2039-1
Thermal Properties			
Vicat B50	95	°C	ISO 306
HDT / A (1,8 MPa)	85	°C	ISO 75-1/-2
Ball Indentation Temperature	75	°C	DIN EN
			60695-10-2
Rheological Properties			
Melt Index (MVR)	17	cm ³ /10min	ISO 1133
MVR temperature	220	°C	-
MVR load	10	kg	-
Shrinkage (24h)	0.6 - 0.8	%	ISO 294-4

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M∷COM

Physical Properties

Density	1080	kg/m³	ISO 1183
Flammability			
Flammability (1.5 mm)	HB	class	UL 94
Yellow Card available	yes	-	-
Glow Wire (GWFI, 650°C, 1.0mm)	passed	-	DIN EN 60695
Glow Wire (GWFI, 650°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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