forward AM.

Ultrafuse® PP

Chemical resistant | biocompatible | low density

Extended TDS

Complete Technical Documentation and Testing Summary



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Technical Data Sheet

High-performance thermoplastic with low density, high elasticity and high resistance to fatigue.

Filament Properties		
Filament Diameter	1.75 mm	2.85 mm
Average diameter Tolerance	±0.050 mm	±0.1 mm
Average ovality	<0.050 mm	<0.050 mm
Available Spool size	750 g	750 g
Available colors	natural	

Spool Properties				
Spool size	750 g	2.0 kg	4.0 kg	8.0 kg
Outer diameter	200 mm	300 mm	350 mm	355 mm
Inner diameter	50.5 mm	51.5 mm	51.7 mm	36 mm
Width	55 mm	103 mm	103 mm	167 mm

Recommended 3D-Print processing parameters		Used for test specimens
Printer	FFF printer	Ultimaker S5
Nozzle Temperature ¹⁾	220 – 240°C	240 °C
Build Chamber Temperature	-	-
Bed Temperature	60 – 80°C	80°C
Bed Material	PP adhesive or tape	Glass + magigoo PP
Nozzle Diameter	≥ 0.4 mm	0.4 mm
Print Speed	20 - 50 mm/s	40 mm/s
Max Volumetric Speed ²⁾	8 mm³/s	//

Please check your standard and/or high speed print profile availability for an easy start at www.forward-am.com.

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Please contact us for further product information, like for example REACH, RoHS,

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Process materials in a well-ventilated room, or use professional extraction systems.

¹ Fast printing might require an additional increase of the nozzle temperature; the stated printing speed is based on current validations. As equipment and technology continues to evolve, it is possible that even higher printing speeds may be attainable in the future.

² Based on Bambu Lab X1C with a nozzle diameter of 0.4 mm



Further Recommendations

Drying recommendations to ensure printability and best mechanical properties³⁾

General Properties

60 °C in a hot air dryer or vacuum oven for 4 to 16 hours

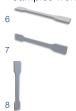
Average Values

deficial Froperties	Stariuaru	Average va	alues	
Filament Density ⁴⁾	ISO 1183-1	901 kg/m ³		
Tensile Properties ⁵⁾	Standard	Average Va	alues	
		XY- Direction ⁶⁾	XZ- Direction ⁷⁾	ZX- Direction ⁸⁾
Tensile strength ⁹⁾	ISO 527	15.5 MPa	-	9.0 MPa
Elongation at Break ⁹⁾	ISO 527	118.6 %	-	5.4 %
Young's Modulus ¹⁰⁾	ISO 527	541 MPa	-	435 MPa

Standard

Flexural Properties ^{6) 11)}	Standard	Average V	alues	
		XY- Direction	XZ- Direction	ZX- Direction
Flexural Strength	ISO 178	22.9 MPa	21.4 MPa	15.6 MPa
Flexural Modulus	ISO 178	575 MPa	494 MPa	380 MPa
Flexural Elongation at Break	ISO 178	9.4 %	8.8 %	7.9 %

⁵ Samples were conditioned in standard climate (23°C, 50% RH 72h)



⁹ Testing speed: 5 mm/min ¹⁰ Testing speed: 1 mm/min

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³ Please note: To ensure constant material properties the material should always be kept dry.

⁴ measured on filament

¹¹ Testing speed: 2 mm/min

Impact Properties ⁶⁾	Standard Average		alues		
		XY- Direction	XZ- Direction	ZX- Direction	
Impact Strength Charpy (notched)	ISO 179-2	5.3 kJ/m ²	8.3 kJ/m ²	2.5 kJ/m ²	
Impact Strength Charpy (unnotched)	ISO 179-2	41.8 kJ/m²	62.3 kJ/m²	13.6 kJ/m²	
Impact Strength Izod (notched)	ISO 180	5.3 kJ/m ²	10.6 kJ/m²	2.3 kJ/m ²	
Impact Strength Izod (unnotched)	ISO 180	37.7 kJ/m²	37.6 kJ/m²	11.6 kJ/m²	

Thermal Properties ⁶⁾	Standard	Average Values
HDT A at 1.8 MPa	ISO 75-2	49 °C
HDT B at 0.45 MPa	ISO 75-2	69 °C
Vicat softening point at 50 N	ISO 306	61 °C
Vicat softening point at 10 N	ISO 306	106 °C
Crystalization Temperature	ISO 11357-3	83 °C
Melting Temperature	ISO 11357-3	131 °C
Melt Volume-Flow Rate (MVR)	ISO 1133	6.7 cm ³ /10 min (230 °C, 2.16 kg)
Melt Mass-Flow Rate (MFR)	ISO 1133	4.8 g/10 min (X°C, X kg)

Hardness and Abrasion	Standard	Typical Values
Shore Hardness D (15s)	DIN ISO 7619-1	52

Biocompatibility	Standard	Typical Values
Cytotoxicity - Neutral Red	ISO 10993-5 (2009)	PASS
In Vivo Sensitization - Local Lymph Node Assay	ISO 10993-10 (2013)	PASS
In Vitro Skin Irritation	ISO 10993-10 (2013)	PASS

For the statement on Biocompatibility data see Chapter Biocompatibility.

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Biocompatibility



Biocompatibility product Information

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Product: Ultrafuse® PP

Revision: 12/3/2014 Version: 1.4

Contact:

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Ultrafuse® PP material is 3D printed as test specimens and <u>successfully passed</u> the requirements of the stated tests below:

- Cytotoxicity XTT Test Neutral red (ISO 10993-5:2009)
 The extract of the product Ultrafuse® PP resulted in a cell vitality of more than 70% in comparison to the negative control and can therefore be considered to be not cytotoxic.
- Skin Irritation Test (ISO10993-10:2013)
 All 10 volunteers exhibited no dermal changes in the test zone at 24h, 48h and 72h when exposed to Ultrafuse® PP.
- Skin Sensitisation Test Local Lymph Node Assay KretinoSens (ISO10993-10:2020)
 The extracts of the product Ultrafuse® PP resulted in an induction of the luciferase activity of less than 1.5 times compared to the DMSO control and is therefore assessed as non-sensitizing.

The performed biocompatibility tests were recorded on test specimen of the above referenced product to show compatibility of the material in general. The biocompatibility tests listed are not part of any continuous production protocol. The test assessments reflect only the test specimen and have to be retested on the final product. It remains the responsibility of the device manufacturers and/or endusers to determine the suitability of all printed parts for their respective application.

Disclaimer:

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