

Makroblend® UT235 M

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(PC+PET)-blend, 15% mineral filled, easy flow, low coefficient of linear thermal expansion, easy release, injection molding. Molded parts from UT235M having exceptional dimensional stability.

ISO Shortname

	Property	Test Condition	Unit	Standard	typical Value
D.					-
	eological properties Melt volume-flow rate	270 °C; 5 kg	cm ³ /10 min	ISO 1133	43
П	Molding shrinkage, parallel	150x105x3 mm; 270 °C / MT 70°C; 600 bar	%	b.o. ISO 2577	0.5 - 0.6
	Molding shrinkage, normal	150x105x3 mm; 270 °C / MT 70°C; 600 bar	%	b.o. ISO 2577	0.5 - 0.6
Me	echanical properties (23 °C/50 % r. h.)	.		<u>'</u>	<u> </u>
_	Tensile modulus	1 mm/min	MPa	ISO 527-1,-2	4500
1	Yield stress	5 mm/min	MPa	ISO 527-1,-2	68
1	Yield strain	5 mm/min	%	ISO 527-1,-2	3.5
3	Stress at break	5 mm/min	MPa	ISO 527-1,-2	67
3	Strain at break	5 mm/min	%	ISO 527-1,-2	4.0
İ	Flexural modulus	2 mm/min	MPa	ISO 178	4650
İ	Flexural strain at flexural strength	2 mm/min	%	ISO 178	5.0
İ	Flexural stress at 3.5 % strain	2 mm/min	MPa	ISO 178	110
İ	Flexural strength	2 mm/min	MPa	ISO 178	115
j	Charpy impact strength	23 °C	kJ/m²	ISO 179-1eU	85
j	Charpy impact strength	-30 °C	kJ/m²	ISO 179-1eU	85
į	Puncture maximum force	23 °C	N	ISO 6603-2	4500
j	Puncture energy	23 °C	J	ISO 6603-2	36
İ	Izod impact strength	23 °C	kJ/m²	ISO 180-U	85
İ	Izod impact strength	-30 °C	kJ/m²	ISO 180-U	85
'n	ermal properties			<u>, </u>	<u> </u>
_	Temperature of deflection under load	1.80 MPa	°C	ISO 75-1,-2	114
1	Temperature of deflection under load	0.45 MPa	°C	ISO 75-1,-2	128
İ	Vicat softening temperature	50 N; 120 °C/h	°C	ISO 306	139
7	Coefficient of linear thermal expansion, parallel	23 to 55 °C	10 ⁻⁴ /K	ISO 11359-1,-2	0.45
2	Coefficient of linear thermal expansion, transverse	23 to 55 °C	10 ⁻⁴ /K	ISO 11359-1,-2	0.45
1	Coefficient of linear thermal expansion, parallel	23 to 85 °C	10 ⁻⁴ /K	ISO 11359-1,-2	0.45
1	Coefficient of linear thermal expansion, transverse	23 to 85 °C	10 ⁻⁴ /K	ISO 11359-1,-2	0.48
)t	her properties (23 °C)		Ų.	<u>'</u>	'
	Water absorption (saturation value)	Water at 23 °C	%	ISO 62	0.4
	Water absorption (equilibrium value)	23 °C; 50 % r. h.	%	ISO 62	0.2
1	Density		kg/m³	ISO 1183-1	1340
j	Bulk density		g/cm³	ISO 60	0.75
j	Filler content	Method A	%	b.o. ISO 3451-1	15
re	ocessing conditions for test specimens				,
	Injection molding-Melt temperature		°C	ISO 294	270
	Injection molding-Mold temperature		°C	ISO 294	70
	Injection molding-Injection velocity		mm/s	ISO 294	200
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C These property characteristics are taken from the CAMPUS plastics data bank and are based on the international catalogue of basic data for plastics according to ISO 10350.

Impact properties: N = non-break, P = partial break, C = complete break



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Disclaimer

Information Impact properties

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Typical value

These values are typical values only. Unless explicitly agreed in written form, the do not constitute a binding material specification or warranted values. Values may be affected by the design of the mold/die, the processing conditions and coloring/pigmentation of the product. Unless specified to the contrary, the property values given have been established on standardized test specimens at room temperature.

General

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Disclaimer Non Medical Grade

This product is not designated for the manufacture of a medical device or of intermediate products for medical devices (1). [This product is also not designated for Food Contact (2), including drinking water, or cosmetic applications. If the intended use of the product is for the manufacture of a medical device or of intermediate products for medical devices, for Food Contact products or cosmetic applications Covestro must be contacted in advance to provide its agreement to sell such product for such purpose.] Nonetheless, any determination as to whether a product is appropriate for use in a medical device or intermediate products for medical devices, for Food Contact products or cosmetic applications must be made solely by the purchaser of the product without relying upon any representations by Covestro. 1) Please see the "Guidance on Use of Covestro Products in a Medical Application" document. 2) As defined in Commission Regulation (EU) 1935/2004.

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