

ORFIBRACE™ NS 3.0 mm (1/12")

Thickness	mm (inches)	3.0 (1/8)	3.0 (1/8)
Perforation	% (type)	0 (non perfo)	1.0 (macro)

Thermoforming conditions

Optimum activation temperature (in water bath)	°C (°F)	65 (149)	65 (149)
Activation time (in water bath)	minutes	3 - 4	3 - 4
Transparent when activated		no	no
Working time	minutes	1 - 1 ½	1 - 1 ½
Hardening time	minutes	6 - 6 ½	5 ¾ - 6 ¼
Time to completion	minutes	24 - 25	22 - 23
Resistance to stretch		high	high
Drape		moderate	moderate
Memory (after 200 % elongation)		moderate	moderate
Maximum elongation when activated	%	2500	1800
Memory (after maximum elongation)		moderate	moderate
Sticks to itself when activated and wet		no	no
Sticks to itself when activated, after drying		temporarily	temporarily
Adhesion (velcro strip) using heat gun		no	no

Mechanical properties at 21°C

Flexural modulus	MPa	700	650
Elastic modulus	MPa	340	340
Tensile strength	MPa	17.0	13.0
Strain at break	%	no break	60

General properties

Density	g cm ⁻³	1.21	1.21
Hardness (shore D)		59	59
Surface feeling		smooth	smooth
Color		grey	grey
Odor		none	none
Fatigue	cycles	> 10000	> 10000
Biocompatible		yes	yes

INFORMATION

The hardening time indicates the time period during which the material remains flexible, but no longer mouldable.

The time to completion indicates the length of time until the orthosis is finished and can be worn by the patient.

The memory indicates the ability of the material to regain its original shape after reheating.

The flexural modulus indicates the resistance of the material to a force causing it to bend.

The elastic modulus defines the ratio of the applied tensile stress to the change in shape of the material.

The tensile strength is the pulling force required to break the material.

The strain at break is the length increase of the material when stretched until failure.

The hardness indicates the resistance of the material to compression.

Fatigue indicates the minimum number of stress cycles the material sustains when bending over 90 degrees without failure.

The biocompatibility is studied according the guidelines of the International Organization for Standardization 10993 – Biological Evaluation of Medical Devices:

- Primary skin irritation study.
- Delayed dermal contact sensitization study.
- Cytotoxicity study.

Note:

Although the information in this publication is believed to be accurate and reliable, the data shown are for guidance only. Orfit Industries gives no guarantees about the results and assumes no liability in connection with them. The properties reported here are intended primarily to facilitate comparison among Orfit products. Standard testing methods often allow alternative measuring methods. Therefore, data from other sheet manufacturers may not be directly comparable. For additional information, please contact Orfit Industries.



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Belgium T +32 (0)3 326 20 26
America T 516-935-8500

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