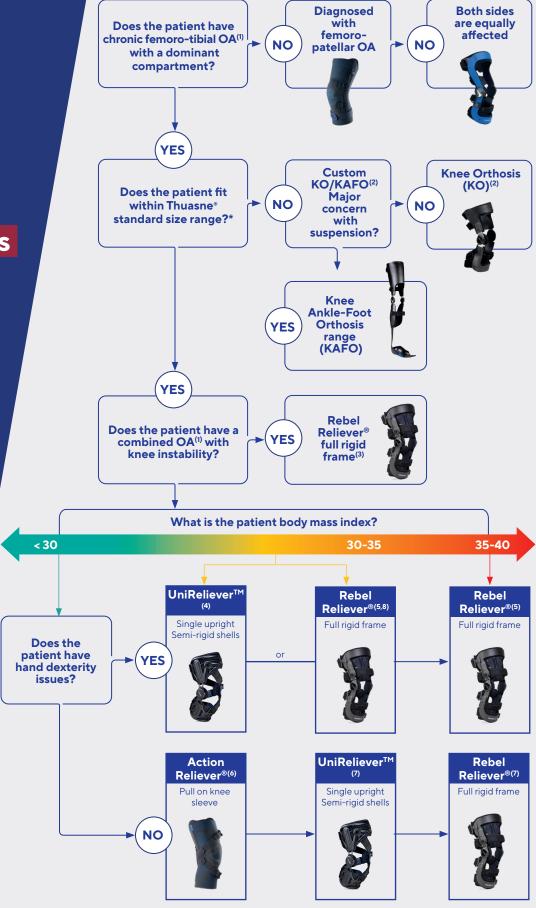


**Osteoarthritis** knee braces prescription guide

If the patient can be fitted with each of these products, please make sure to determine in **ACCORDANCE** with the patient, which product would suit the best their **EXPECTATIONS** and **NEEDS** (Discretion, Desired activities...)



thigh circumference = 71 cm/28" à: Osteoarthritis ): Knee Orthosis / KAFO: Knee-Ankle

Availability of these products might vary from a given country or region to another, as a result of specific local regulatory approval or clearance requirements for sale in such country or region. All the medical devices mentioned on this document are CE marked according to the Regulation 2017/745 on medical devices, they are also UKCA-marked according to S.I. 2002/618 regulation on medical devices. Medical devices mentioned in this document are CE class I. Please contact Thuasne® should you need any additional information on devices classification. Please carefully read the instructions for use, indications and contraindications of the products. The full list of medical indications, are available in the instructions for use. Last revision date: 03/2024. Ref.: 2402100.

Thuasne SAS – SIREN/RCS Nanterre 542 091186 – capital 1950 000 euros – 120, rue Marius Aufan 92300 Levallois-Perret (France)

## Patella Reliever® - Action Reliever® - Rebel Reliever®:

Thuasne 120, rue Marius Aufan 92300 Levallois-Perret France

## UniReliever<sup>™</sup> - SpryStep® KAFO - SpryStep® KO:

Thuasne Deutschland GmbH Im Steinkamp 12 30938 Burgwedel - Deutschland

Thuasne USA 4615 Shepard Street - Bakersfield, CA 93313 United-States

MedEnvoy Global BV Prinnses Margrietplantsoen 33, Suite 123 - 2595 AM The Hague The Netherlands

All products except Action Reliever® and Patella Reliever®:
UK Responsible Person (UKRP):
THUASNE UK Ltd
Unit 4 Orchard Business Centre
North Farm Road
Tunbridge Wells, TN2 3XF,
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