**GORE® EXCLUDER®**
Infrarenal Device Family

# CONFIGURATIONS

*Together, improving life*

## GORE® EXCLUDER® AAA Endoprosthesis with C3® Delivery System

### Trunk-Ipsilateral Leg Endoprosthesis

| Catalogue number | Intended aortic inner diameter (mm) | Aortic endoprosthesis diameter (mm) | Intended iliac inner diameter (mm) | Iliac endoprosthesis diameter (mm) | Overall device length (cm) | Recommended sheath size (Fr) |
|---|---|---|---|---|---|---|
| RLT231212 | 19–21 | 23 | 10–11 | 12 | 12 | 16 |
| RLT231214 | 19–21 | 23 | 10–11 | 12 | 14 | 16 |
| RLT231216 | 19–21 | 23 | 10–11 | 12 | 16 | 16 |
| RLT231218 | 19–21 | 23 | 10–11 | 12 | 18 | 16 |
| RLT231412 | 19–21 | 23 | 12–13.5 | 14.5 | 12 | 16 |
| RLT231414 | 19–21 | 23 | 12–13.5 | 14.5 | 14 | 16 |
| RLT231416 | 19–21 | 23 | 12–13.5 | 14.5 | 16 | 16 |
| RLT231418 | 19–21 | 23 | 12–13.5 | 14.5 | 18 | 16 |
| RLT261212 | 22–23 | 26 | 10–11 | 12 | 12 | 16 |
| RLT261214 | 22–23 | 26 | 10–11 | 12 | 14 | 16 |
| RLT261216 | 22–23 | 26 | 10–11 | 12 | 16 | 16 |
| RLT261218 | 22–23 | 26 | 10–11 | 12 | 18 | 16 |
| RLT261412 | 22–23 | 26 | 12–13.5 | 14.5 | 12 | 16 |
| RLT261414 | 22–23 | 26 | 12–13.5 | 14.5 | 14 | 16 |
| RLT261416 | 22–23 | 26 | 12–13.5 | 14.5 | 16 | 16 |
| RLT261418 | 22–23 | 26 | 12–13.5 | 14.5 | 18 | 16 |
| RLT281212 | 24–26 | 28.5 | 10–11 | 12 | 12 | 18 |
| RLT281214 | 24–26 | 28.5 | 10–11 | 12 | 14 | 18 |
| RLT281216 | 24–26 | 28.5 | 10–11 | 12 | 16 | 18 |
| RLT281218 | 24–26 | 28.5 | 10–11 | 12 | 18 | 18 |
| RLT281412 | 24–26 | 28.5 | 12–13.5 | 14.5 | 12 | 18 |
| RLT281414 | 24–26 | 28.5 | 12–13.5 | 14.5 | 14 | 18 |
| RLT281416 | 24–26 | 28.5 | 12–13.5 | 14.5 | 16 | 18 |
| RLT281418 | 24–26 | 28.5 | 12–13.5 | 14.5 | 18 | 18 |
| RLT311413 | 27–29 | 31 | 12–13.5 | 14.5 | 13 | 18 |
| RLT311415 | 27–29 | 31 | 12–13.5 | 14.5 | 15 | 18 |
| RLT311417 | 27–29 | 31 | 12–13.5 | 14.5 | 17 | 18 |
| RLT351414 | 30–32 | 35 | 12–13.5 | 14.5 | 14 | 18 |
| RLT351416 | 30–32 | 35 | 12–13.5 | 14.5 | 16 | 18 |
| RLT351418 | 30–32 | 35 | 12–13.5 | 14.5 | 18 | 18 |

### Contralateral Leg Endoprosthesis

| Catalogue number | Intended iliac inner diameter (mm) | Iliac endoprosthesis diameter (mm) | Contralateral Leg length (cm) | Recommended sheath size (Fr) |
|---|---|---|---|---|
| PLC121000 | 10–11 | 12 | 10 | 12 |
| PLC121200 | 10–11 | 12 | 12 | 12 |
| PLC121400 | 10–11 | 12 | 14 | 12 |
| PLC141000 | 12–13.5 | 14.5 | 10 | 12 |
| PLC141200 | 12–13.5 | 14.5 | 12 | 12 |
| PLC141400 | 12–13.5 | 14.5 | 14 | 12 |
| PLC161000[a] | 13.5–14.5 | 16 | 9.5 | 12 |
| PLC161200[a] | 13.5–14.5 | 16 | 11.5 | 12 |
| PLC161400[a] | 13.5–14.5 | 16 | 13.5 | 12 |
| PLC181000[a] | 14.5–16.5 | 18 | 9.5 | 12 |
| PLC181200[a] | 14.5– 16.5 | 18 | 11.5 | 12 |
| PLC181400[a] | 14.5–16.5 | 18 | 13.5 | 12 |
| PLC201000[a] | 16.5–18.5 | 20 | 9.5 | 12 |
| PLC201200[a] | 16.5–18.5 | 20 | 11.5 | 12 |
| PLC201400[a] | 16.5–18.5 | 20 | 13.5 | 12 |
| PLC231000[a] | 18.5–21.5 | 23 | 10 | 14 |
| PLC231200[a] | 18.5–21.5 | 23 | 12 | 14 |
| PLC231400[a] | 18.5–21.5 | 23 | 14 | 14 |
| PLC271000[a] | 21.5–25.0 | 27 | 10 | 15 |
| PLC271200[a] | 21.5–25.0 | 27 | 12 | 15 |
| PLC271400[a] | 21.5–25.0 | 27 | 14 | 15 |

### Aortic Extender Endoprosthesis

| Catalogue number | Intended aortic inner diameter (mm) | Endoprosthesis diameter (mm) | Endoprosthesis length (cm) | Recommended sheath size (Fr) |
|---|---|---|---|---|
| PLA230300 | 19–21 | 23 | 3.3 | 16 |
| PLA260300 | 22–23 | 26 | 3.3 | 16 |
| PLA280300 | 24–26 | 28.5 | 3.3 | 16 |
| PLA320400 | 27–29 | 32 | 4.5 | 17 |
| PLA360400 | 30–32 | 36 | 4.5 | 18 |

### Iliac Extender Endoprosthesis

| Catalogue number | Intended iliac linner diameter (mm) | Endoprosthesis diameter (mm) | Endoprosthesis length (cm) | Recommended sheath size (Fr) |
|---|---|---|---|---|
| PLL161007 | 8–9 | 10 | 7 | 12 |
| PLL161207 | 10–11 | 12 | 7 | 12 |
| PLL161407 | 12–13.5 | 14.5 | 7 | 12 |

## GORE® EXCLUDER® Iliac Branch Endoprosthesis

### Iliac Branch Component

| Catalogue number | Intended external iliac vessel diameter (mm) | Diameter iliac end (mm) | Iliac branch component length (cm) | Recommended sheath size (Fr) |
|---|---|---|---|---|
| CEB231010H | 6.5–9 | 10 | 10 | 16 |
| CEB231210H | 10–11 | 12 | 10 | 16 |
| CEB231410H | 12–13.5 | 14.5 | 10 | 16 |

### Internal Iliac Component

| Catalogue number | Intended internal iliac vessel diameter (mm) | Diameter iliac end (mm) | Internal iliac component length (cm) | Recommended sheath size (Fr x cm) |
|---|---|---|---|---|
| HGB161007H | 6.5–9 | 10 | 7 | 12 x 45 |
| HGB161207H | 10–11 | 12 | 7 | 12 x 45 |
| HGB161407H | 12–13.5 | 14.5 | 7 | 12 x 45 |

## Aortic Accessories

### GORE® DRYSEAL Flex Introducer Sheath

| Catalogue number[b] | Sheath size (Fr) | Minimum sheath ID (mm) | Nominal sheath OD (mm) | Working length (cm) |
|---|---|---|---|---|
| GDSF1233 | 12 | 4.0 | 4.7 | 33 |
| GDSF1245 | 12 | 4.0 | 4.7 | 45 |
| GDSF1433 | 14 | 4.7 | 5.3 | 33 |
| GDSF1533 | 15 | 5.0 | 5.6 | 33 |
| GDSF1633 | 16 | 5.3 | 6.1 | 33 |
| GDSF1833 | 18 | 6.0 | 6.7 | 33 |

### GORE® Molding & Occlusion Balloon Catheter

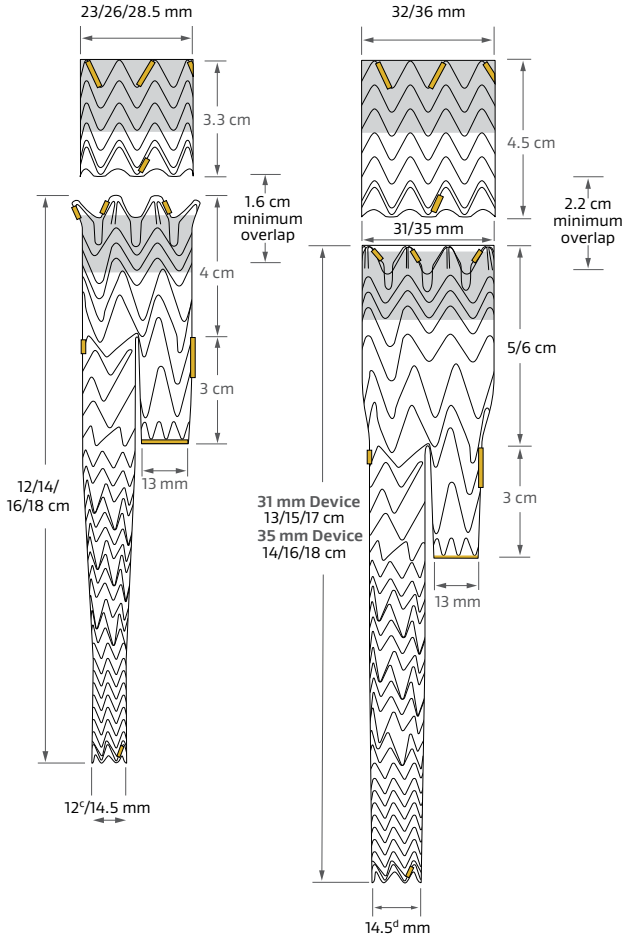
| Catalogue number | Size (Fr) | Balloon diameter (mm) | Catheter length (cm) |
|---|---|---|---|
| MOB37 | 10 | 10–37 | 90 |

[a] PLEASE NOTE: All large diameter Contralateral Leg Endoprostheses (16, 18, 20, 23, 27 mm) can be used as Iliac Extenders.

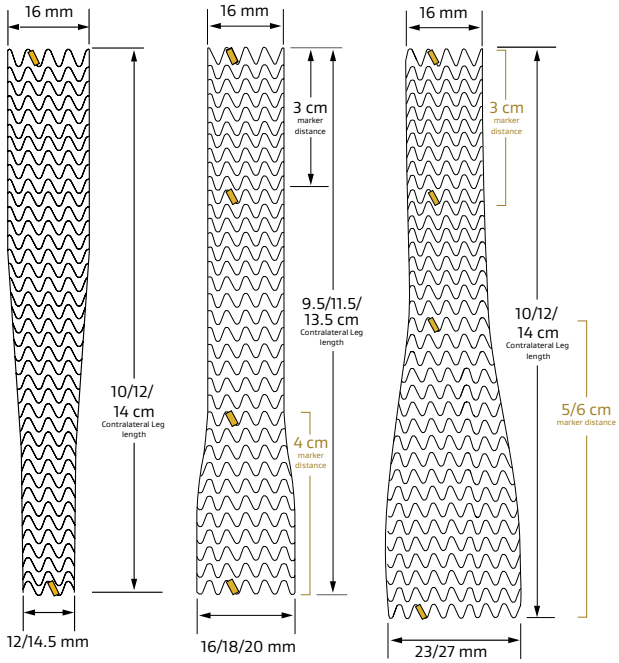
[b] GDSF catalogue numbers are not available in all regions; use DSF catalogue numbers instead where appropriate.

# GORE® EXCLUDER® AAA Endoprosthesis featuring C3® Delivery System

## Aortic Extender and Trunk-Ipsilateral Leg Endoprosthesis

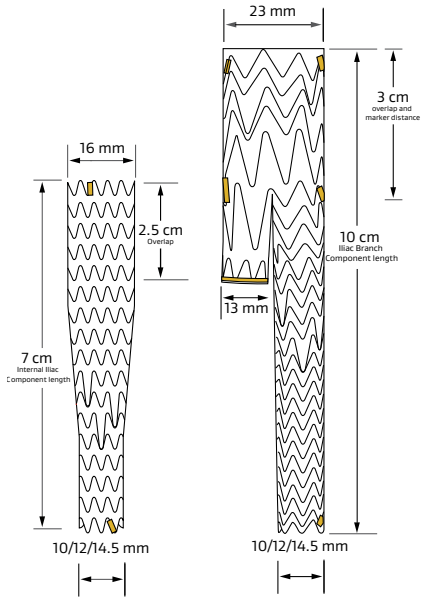



## Contralateral Leg Endoprosthesis[1,2]

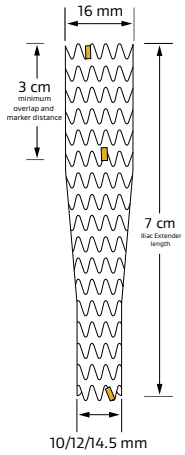



## GORE® EXCLUDER® Iliac Branch Endoprosthesis System

### Internal Iliac Component
### Iliac Branch Component




## Iliac Extender Endoprosthesis[3]




[c] 12 mm only available in select trunk sizes.
[d] 14.5 mm only available in select trunk sizes.

1. All large diameter Contralateral Leg Endoprostheses (16, 18, 20, 23, 27 mm) can be used as Iliac Extenders.
2. The 27 mm x 10 cm Contralateral Leg Endoprosthesis does not have a radiopaque marker 6 cm above the distal end.
3. Traditional sizes of Iliac Extender (10, 12, 14.5 mm) are not compatible with large diameter (16, 18, 20, 23, 27 mm) Contralateral Leg Endoprostheses.

## GORE® EXCLUDER® Conformable AAA Endoprosthesis with ACTIVE CONTROL System

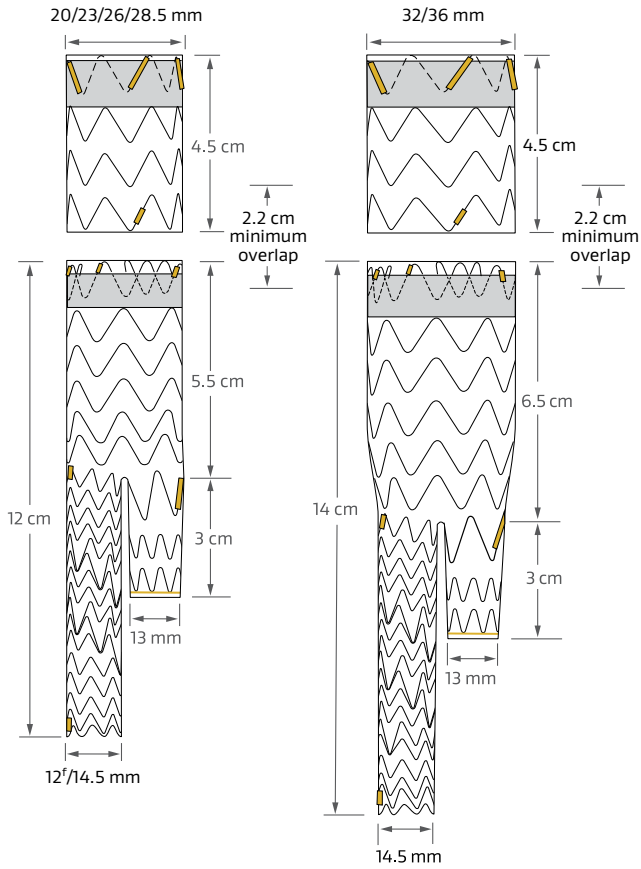
### Trunk-Ipsilateral Leg Endoprosthesis

| Catalogue number[e] | Intended aortic vessel diameter (mm) | Aortic endoprosthesis diameter (mm) | Intended iliac vessel diameter (mm) | Iliac endoprosthesis diameter (mm) | Overall device length (cm) | Recommended introducer sheath (Fr) |
|---|---|---|---|---|---|---|
| CXT201212H | 16–18 | 20 | 10–11 | 12 | 12 | 15 |
| CXT201412H | 16–18 | 20 | 12–13.5 | 14.5 | 12 | 15 |
| CXT231412H | 19–21 | 23 | 12–13.5 | 14.5 | 12 | 15 |
| CXT261412H | 22–23 | 26 | 12–13.5 | 14.5 | 12 | 16 |
| CXT281412H | 24–26 | 28.5 | 12–13.5 | 14.5 | 12 | 16 |
| CXT321414H | 27–29 | 32 | 12–13.5 | 14.5 | 14 | 18 |
| CXT361414H | 30–32 | 36 | 12–13.5 | 14.5 | 14 | 18 |

### Aortic Extender Endoprosthesis

| Catalogue number[e] | Intended aortic inner diameter (mm) | Endoprosthesis diameter (mm) | Endoprosthesis length (cm) | Recommended sheath size (Fr) |
|---|---|---|---|---|
| CXA200005H | 16–18 | 20 | 4.5 | 15 |
| CXA230005H | 19–21 | 23 | 4.5 | 15 |
| CXA260005H | 22–23 | 26 | 4.5 | 15 |
| CXA280005H | 24–26 | 28.5 | 4.5 | 16 |
| CXA320005H | 27–29 | 32 | 4.5 | 18 |
| CXA360005H | 30–32 | 36 | 4.5 | 18 |

### Aortic Extender and Trunk-Ipsilateral Leg Endoprosthesis




[e] Currently available commercially.
[f] 12 mm only available in the 20 mm GORE® EXCLUDER® Conformable AAA Endoprosthesis.

Consult Instructions for Use
**eifu.goremedical.com**

Refer to *Instructions for Use* at eifu.goremedical.com for a complete description of all applicable indications, warnings, precautions and contraindications for the markets where this product is available. ℞ Only

**GORE® DRYSEAL Flex Introducer Sheath**
**INDICATIONS FOR USE:** The GORE® DrySeal Flex Introducer Sheath is intended to be inserted in the vasculature to provide a conduit for the insertion of endovascular devices while minimizing blood loss associated with such insertions.
**CONTRAINDICATIONS:** There are no known contraindications for this device.

**GORE® EXCLUDER® AAA Endoprosthesis**
**INDICATIONS FOR USE:** Trunk-Ipsilateral Leg and Contralateral Leg Endoprosthesis. The GORE® EXCLUDER® AAA Endoprosthesis is intended to exclude the aneurysm from the blood circulation in patients diagnosed with infrarenal abdominal aortic aneurysm (AAA) disease and who have appropriate anatomy as described below: Adequate iliac/femoral access; Infrarenal aortic neck treatment diameter range of 19–32 mm and a minimum aortic neck length of 15 mm; Proximal aortic neck angulation ≤ 60°; Iliac artery treatment diameter range of 8–25 mm and iliac distal vessel seal zone length of at least 10 mm. Aortic Extender and Iliac Extender Endoprosthesis. The Aortic and Iliac Extender Endoprostheses are intended to be used after deployment of the GORE® EXCLUDER® AAA Endoprosthesis. These extensions are intended to be used when additional length and/or sealing for aneurysmal exclusion is desired.
**CONTRAINDICATIONS:** The GORE® EXCLUDER® AAA Endoprosthesis is contraindicated in: patients with known sensitivities or allergies to the device materials; patients with a systemic infection who may be at increased risk of endovascular graft infection.

**GORE® EXCLUDER® Conformable AAA Endoprosthesis**
**INDICATIONS FOR USE:** Trunk-Ipsilateral Leg and Contralateral Leg Endoprosthesis. The GORE® EXCLUDER® Conformable AAA Endoprosthesis is intended to exclude the aneurysm from the blood circulation in patients diagnosed with infrarenal abdominal aortic aneurysm (AAA) disease and who have appropriate anatomy as described below: Adequate iliac/femoral access; Infrarenal aortic neck treatment diameter range of 16-32 mm and a minimum aortic neck length of 10 mm; Proximal aortic neck angulation ≤ 90°; Iliac artery treatment diameter range of 8-25 mm and iliac distal vessel seal zone length of at least 10 mm. Aortic Extender Endoprosthesis and Iliac Extender Endoprosthesis Components are intended to be used after deployment of the GORE® EXCLUDER® Conformable AAA Endoprosthesis. These extensions are intended to be used when additional length and/or sealing for aneurysmal exclusion is desired.
**CONTRAINDICATIONS:** The GORE® EXCLUDER® Conformable AAA Endoprosthesis is contraindicated in: patients with known sensitivities or allergies to the device materials. All components of the GORE® EXCLUDER® Conformable Endoprosthesis contain expanded polytetrafluoroethylene (ePTFE), fluorinated ethylene propylene (FEP), nitinol (nickel-titanium alloy) and gold. Patients with systemic infection who may be at increased risk of endovascular graft infection.

**GORE® EXCLUDER® Iliac Branch Endoprosthesis (IBE)**
**INDICATIONS FOR USE:** Iliac Branch and Internal Iliac Components. The GORE® EXCLUDER® Iliac Branch Endoprosthesis (IBE) is intended to be used with the GORE® EXCLUDER® AAA Endoprosthesis to isolate the common iliac artery from systemic blood flow and preserve blood flow in the external iliac and internal iliac arteries in patients with a common iliac or aortoiliac aneurysm, who have appropriate anatomy, including: Adequate iliac/femoral access; Minimum common iliac diameter of 17 mm at the proximal implantation zone of the IBE; External Iliac artery treatment diameter range of 6.5–25 mm and seal zone length of at least 10 mm; Internal iliac artery treatment diameter range of 6.5–13.5 mm and seal zone length of at least 10 mm; Adequate length from the lowest major renal artery to the internal iliac artery to accommodate the total endoprosthesis length, calculated by adding the minimum lengths of required components, taking into account appropriate overlaps between components. GORE® EXCLUDER® Components used in conjunction with GORE® EXCLUDER® Iliac Branch Endoprosthesis: Trunk-Ipsilateral Leg Component. The Trunk-Ipsilateral Leg is intended to provide proximal seal and fixation for the endovascular repair of the aneurysm. For more information on the Trunk-Ipsilateral Leg Component indications for use and deployment, see the GORE® EXCLUDER® AAA Endoprosthesis *Instructions for Use*. Contralateral Leg Endoprosthesis Component. The Contralateral Leg Endoprosthesis is intended to bridge the GORE® EXCLUDER® Device Trunk-Ipsilateral Component to the GORE® EXCLUDER® Iliac Branch Endoprosthesis following deployment of the GORE® EXCLUDER® Iliac Branch Endoprosthesis. Additionally, the Contralateral Leg Endoprosthesis is intended to be used for distal extension of the Iliac Branch Component in the external iliac artery. The Iliac Branch Component can treat external iliac artery diameters up to 13.5 mm. This ability to extend the Iliac Branch Component distally with any Contralateral Leg Endoprosthesis expands the external iliac artery treatment range up to 25 mm. For more information on the Trunk-Ipsilateral Leg and Contralateral Leg Endoprosthesis Component indications for use and deployment, see the GORE® EXCLUDER® AAA Endoprosthesis *Instructions for Use*. Aortic Extender and Iliac Extender Components. The Aortic and Iliac Extender Components can be used after deployment of the GORE® EXCLUDER® Iliac Branch and GORE® EXCLUDER® AAA Endoprostheses. These extensions are used when additional length and/or sealing for aneurysmal exclusion is desired. For more information on Aortic Extender and Iliac Extender indications for use and deployment, see the GORE® EXCLUDER® AAA Endoprosthesis *Instructions for Use*.
**CONTRAINDICATIONS:** The GORE® EXCLUDER® Iliac Branch Endoprosthesis is contraindicated in: patients with known sensitivities or allergies to the device materials. All components of the GORE® EXCLUDER® Iliac Branch Endoprosthesis, the GORE® EXCLUDER® AAA Endoprosthesis contains ePTFE, FEP, nitinol (nickel-titanium alloy) and gold. Patients with a systemic infection who may be at increased risk of endovascular graft infection.

**GORE® Molding & Occlusion Balloon Catheter (MOB)**
**INDICATIONS FOR USE:** The GORE® Molding & Occlusion Balloon Catheter is intended for temporary occlusion of large diameter vessels or to assist the expansion of self-expanding endovascular prostheses (stent grafts).
**CONTRAINDICATIONS:** The GORE® Molding & Occlusion Balloon Catheter is contraindicated in patients who: are contraindicated to contrast media or anticoagulants; have an arterial entry site that cannot accommodate a 10 Fr introducer sheath; are minors; are pregnant.

Products listed may not be available in all markets.


24AR1205-EN03 SEPTEMBER 2025

**W. L. Gore & Associates, Inc.**
goremedical.com

**Asia Pacific** +65 67332882 **Australia/New Zealand** 1800 680 424 **Europe** 00800 6334 4673
**United States** Flagstaff, AZ 86004 800 437 8181 928 779 2771