



BED HEAD UNITS

MODELS

AURA



**Aura
Modular Bed Head Unit**

Providing efficiency to various areas

- AURA 100
- AURA 200
- AURA 300



WWW.TEDISELMEDICAL.COM

info@tedisel.com



TECHNICAL DATA

MODEL: AURA

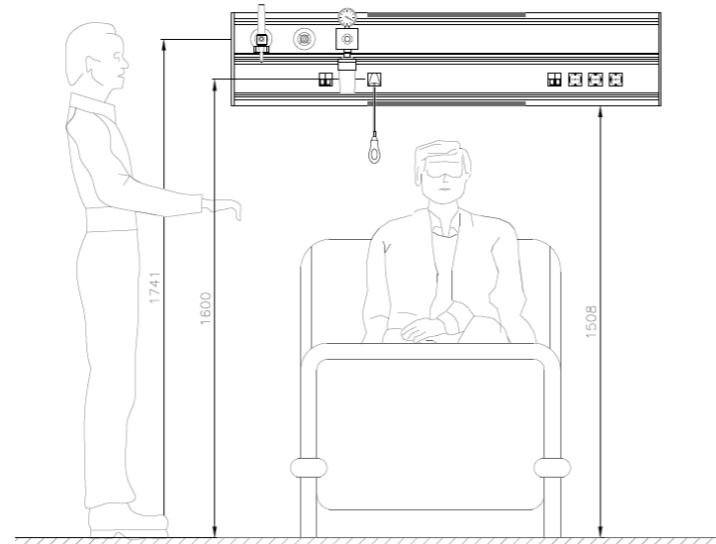
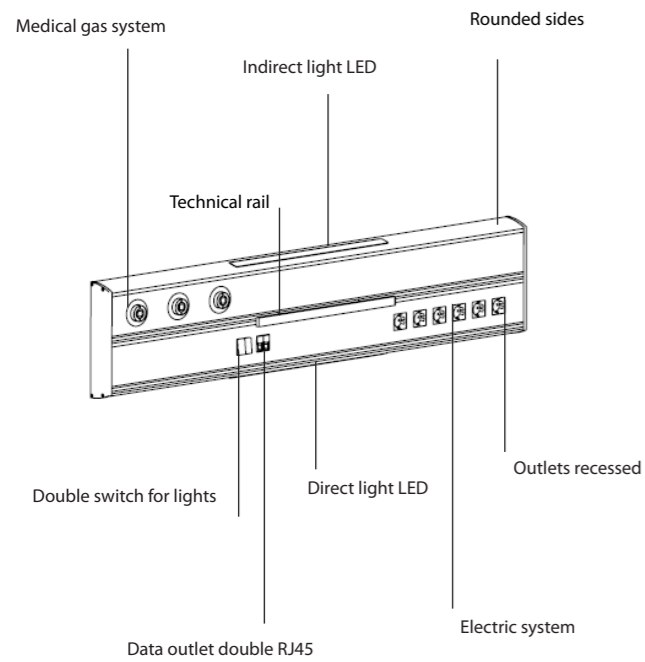


Profile	
Length	Other lenghts available depending on the project*
* Max. manufacturing length 3000mm. Possibility of installing consecutive sections for several beds.	
Technical rail	
Max. Load	25 kg x meter
Electrical System	
Rated voltage	230 Vca
Rated frequency	50Hz

Lighting System (optional)	
Direct LED light	10W - 20W (4500K)*
Indirect LED light	10W - 20W - 40W (4500K)*
Night light	1W

* Possibility to regulate lights (0-10V) and DALI option

Medical Gas System	
	O ₂ - Air - N ₂ O - CO ₂ - VAC - AGSS - Air 800 - N ₂ 800 - O ₂ /He - O ₂ /N ₂ O
Materials	
Structure	Aluminium with epoxy painting or anodized, HPL (upon request)
Technical rails	Stainless steel AISI 304 (standard) / Aluminium (optional)
Plastic diffuser	Extruded polycarbonate



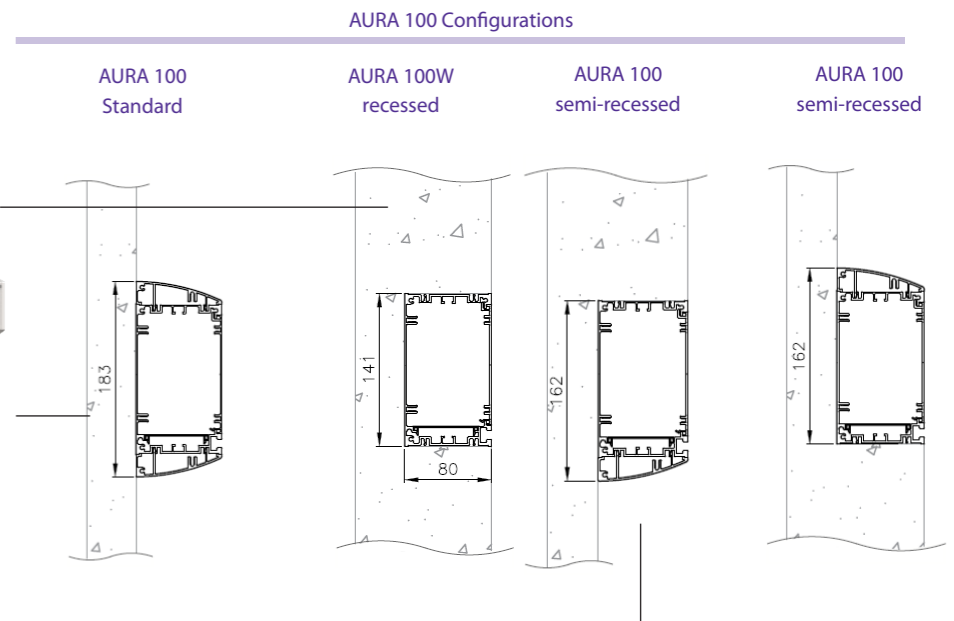
MODEL: AURA
VERSION: 100



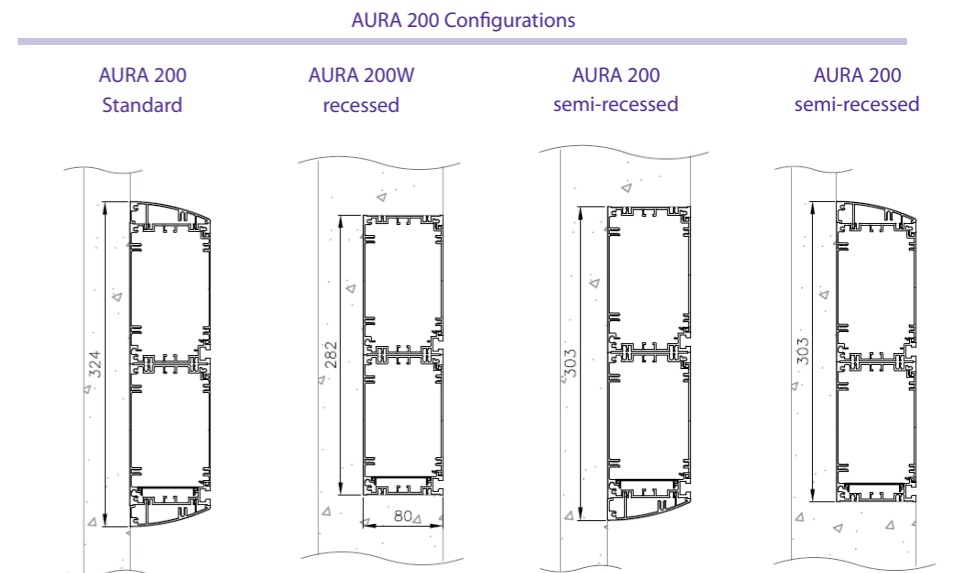
Wall Recessed

Surface Mounted

Rounded sides and geometric lines for optimal cleaning



MODEL: AURA
VERSION: 200

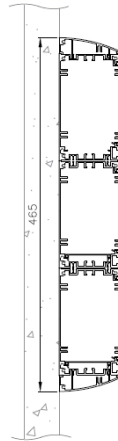


MODEL: AURA
VERSION: 300

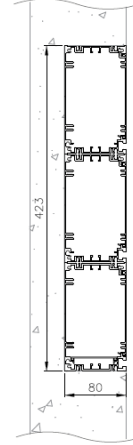


AURA 300 Configurations

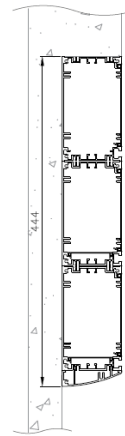
AURA 300
Standard



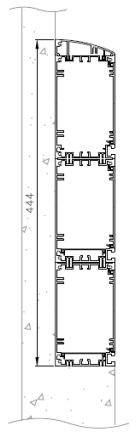
AURA 300W
recessed



AURA 300
semi-recessed



AURA 300
semi-recessed



Classification

Directive 93/42/EEC

CLASS II B

EC Conformity

Tedisel complies with the provisions of Directive 93/42/EEC (medical devices), ISO 11197 (medical supply units) and IEC 60601-1 (medical electrical equipment. Part1: general requirements for basic safety and essential performance).

Compatibility with other medical devices

Tedisel products may be equipped with medical devices from other manufacturers. Please follow the instructions provided by the manufacturers of this equipment for proper installation.

CE 0197

TüV Rheinland LGA Products GmbH

QUALITY

Tedisel Management System and all our products are certified with quality seal ISO 13485 and CE marking granted by notified body TÜV RHEINLAND LGA Products GmbH.

Tedisel, as medical device manufacturer, has the medical device manufacturer license number 6205-PS, granted by the Spanish Agency for Medicines and Health Products.

All medical devices developed and manufactured in Tedisel, are regulated by the European Directive 93/42/EEC. All equipment has accreditation and certification of electrical safety and electromagnetic compatibility standards ISO 11197 and EN 60601.