

## Vibrasat® Pro ·

### Descripción:

Vibrasat Pro es la tecnología líder para liposucciones asistidas por potencia (PAL).

### Comodidad:

Su diseño fácil de usar elimina la fatiga incluso durante los procedimientos más largos, mientras que su tecnología maximiza los resultados del paciente.

Vibrasat Pro ofrece la forma más segura y suave de eliminar la grasa.

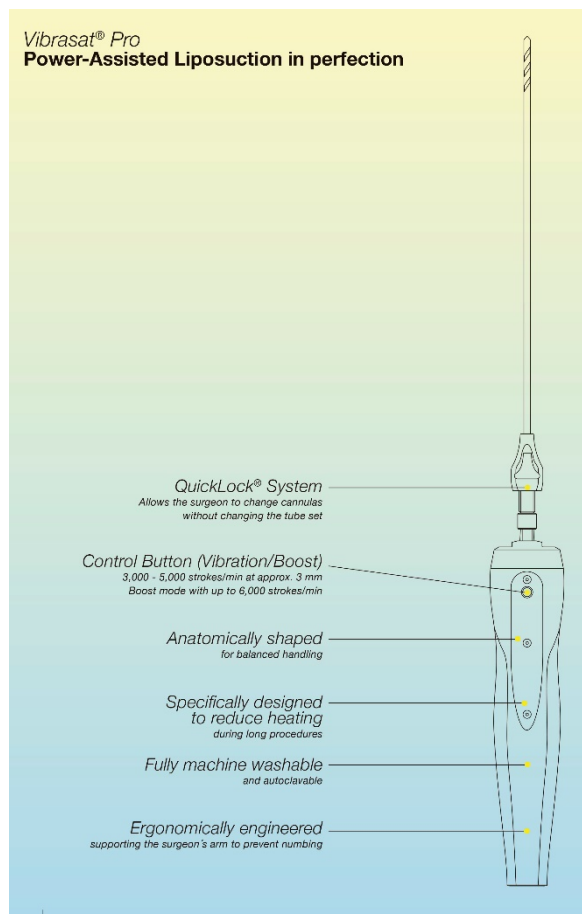
Se requiere menos esfuerzo para mover la cánula a través del tejido, asegurando un menor riesgo con resultados de máxima precisión.

Su empuñadura de forma anatómica lleva el punto de equilibrio justo con un peso bajo, lo que permite sostenerlo cómodamente durante cualquier período de tiempo.

El diseño permite un cambio de cánula rápido y fácil, proporcionando un flujo de trabajo óptimo y más eficiente.

### Aplicaciones:

- Liposucción de grasa.
- Liposucción HD · High Definition
- Infiltración TLA para volúmenes pequeños y grandes
- Infiltración de grasa autóloga en mamas y gluteos
- Tratamiento de Lipedema

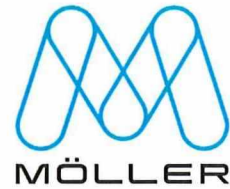


**CE0482**  
**Clase IIa**



### Datos Técnicos

Dimensiones pieza de mano (diámetro x largo)	52 x 300mm
Dimensiones Consola (Ancho x Alto x fondo)	170 x 90 x 205mm
Peso pieza de mano	0,7 Kg.
Peso consola	0,5 Kg.
Velocidad regulable (frecuencia de movimiento)	3.000 – 6.000 /min.
Longitud de movimiento aprox.	2,8mm
Fuente de alimentación	100 - 240 VCA, 50 - 60 Hz
Clase del dispositivo médico	IIa



**KONFORMITÄTSERKLÄRUNG / DECLARATION OF CONFORMITY**

Name und Adresse des Herstellers / **Möller Medical GmbH**  
Name and address of the manufacturer Wasserkuppenstraße 29-31  
36043 Fulda, Germany

Einmalige Registrierungsnummer / **DE-MF-000005100**  
Single Registration Number (SRN)

Wir tragen die alleinige Verantwortung für die Ausstellung dieser EU-Konformitätserklärung. / We are solely responsible for issuing this EU Declaration of Conformity.

Produktname/n / Product name/s **Vibrasat Pro Console REF. 00003921**  
**Vibrasat Pro Wand REF. 00003922**

Basis-UDI-DI / Basic UDI-DI **426027717MM.003WD**

Klassifizierung / Classification **Ila**  
Nach Anhang VIII der Verordnung 2017/745 / acc. to annex VIII of regulation 2017/745

Wir versichern, dass das hier erfasste Produkt der VERORDNUNG (EU) 2017/745 und den weiteren einschlägigen Rechtsvorschriften der EU entspricht. / We assure that the product covered here complies with REGULATION (EU) 2017/745 and other relevant EU legislation.

Gemeinsame Spezifikationen (GS), für die die Konformität erklärt wird / Common Specifications (CS) for which conformity is declared **Not available**

Konformitätsbewertungsverfahren / **MDR 2017/745**  
Conformity assessment procedure Annex IX Chapter 1

Zertifikatsnummer / Certificate No. **7295GB448220906**

Generische Produktgruppe / Generic Product Group **Z12040214 Liposuction Units**

Konformitätsbewertungsstelle (falls beigezogen) / **MedCert GmbH CE 0482**  
Notified Body (if consulted) Pilatuspool 2  
20355 Hamburg

**Fulda, 02. January 2023**

Ort, Datum / Place, date

**Andreas Bacher, Managing Director**

Name und Funktion / Name and function

Gültig bis 22. April 2026 / Valid until 22. April 2026

CONFIDENTIAL

## EU Quality Management System Certificate

### The Notified Body

**MEDCERT Zertifizierungs- und Prüfungsgesellschaft für die Medizin GmbH**  
**Pilatuspool 2 – 20355 Hamburg – Germany**

herewith certifies that the company

**Möller Medical GmbH**  
**Wasserkuppenstrasse 29-31**  
**36043 Fulda**  
**Germany**

**SRN: DE-MF-000005100**

with locations listed in the appendix

has introduced, applies and maintains a quality management system for the medical devices/groups of medical devices listed in the appendix.

The compliance of this quality management system to the requirements of the **Regulation (EU) 2017/745 on medical devices** was verified by assessment according to:

### Annex IX Chapter I

Any applicable limitations of this certification for certain medical devices are included in the appendix. This certification is subject to surveillance by MEDCERT.

**Effective date:** 2021-10-19

**Expiry date:** 2026-04-22

Final assessment report No.: 7295IA09F  
Procedure No.: QS – 7295  
Certificate No.: 7295GB448211019

Preceding certificate No.: —  
Preceding certificate date: —  
Identification of changes: —

Hamburg, 2021-10-19

MEDCERT Certification Body  
Lorenz Runge

The certificate is only valid when provided entirely with all of its pages.  
To verify the validity of this certificate, contact [info@medcert.de](mailto:info@medcert.de).

MEDCERT Notified Body Identification Number: 0482



## Appendix of EU Quality Management System Certificate

Procedure No.: QS – 7295

Certificate No.: 7295GB448211019

### Locations included in the scope of certificate

**Wasserkuppenstrasse 29-31  
36043 Fulda  
Germany**

This appendix is integral part of the above-referenced certificate.  
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**Appendix of EU Quality Management System Certificate**

Procedure No.: QS – 7295

Certificate No.: 7295GB448211019

**Class IIa medical devices**

Category	EMDN code	Medical devices/groups of medical devices
<b>MDA 0306</b>	<b>Z12040214</b>	Liposuction units
	<b>A018099</b>	Needles - other accessories
<b>MDN 1202</b>	<b>A010201</b>	Needles and kits - histologic and cytologic biopsy of small tissues
	<b>A018003</b>	Needle introducers
	<b>A019011</b>	Needles - bone infusion and vertebroplastic
	<b>Z12040214</b>	Liposuction units

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**Appendix of EU Quality Management System Certificate**

Procedure No.: QS – 7295

Certificate No.: 7295GB448211019

**Class IIb medical devices**

Category	EMDN code	Medical devices/groups of medical devices
<b>MDA 0306</b>	<b>Z12040214</b>	Liposuction units Intended purpose: Use for medical indications, including those accompanied by a change in fatty tissue, and for aesthetic body contouring / Administer tumescent local anaesthesia, other aqueous infusion solutions, as well as endogenous subcutaneous tissue and its components, into the body

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