



redefine limits.

curebionics.com

Catalog



Cure Bionics



Rebuilding humans. Reinventing robots.

Hannibal Bionic Prosthetic Arm



The Hannibal hand Engineered to Evolve



CE
Certified



Realistic
Skin color



Lightweight
Comfort



Multi
Modes



+24 Hours
Battery Life



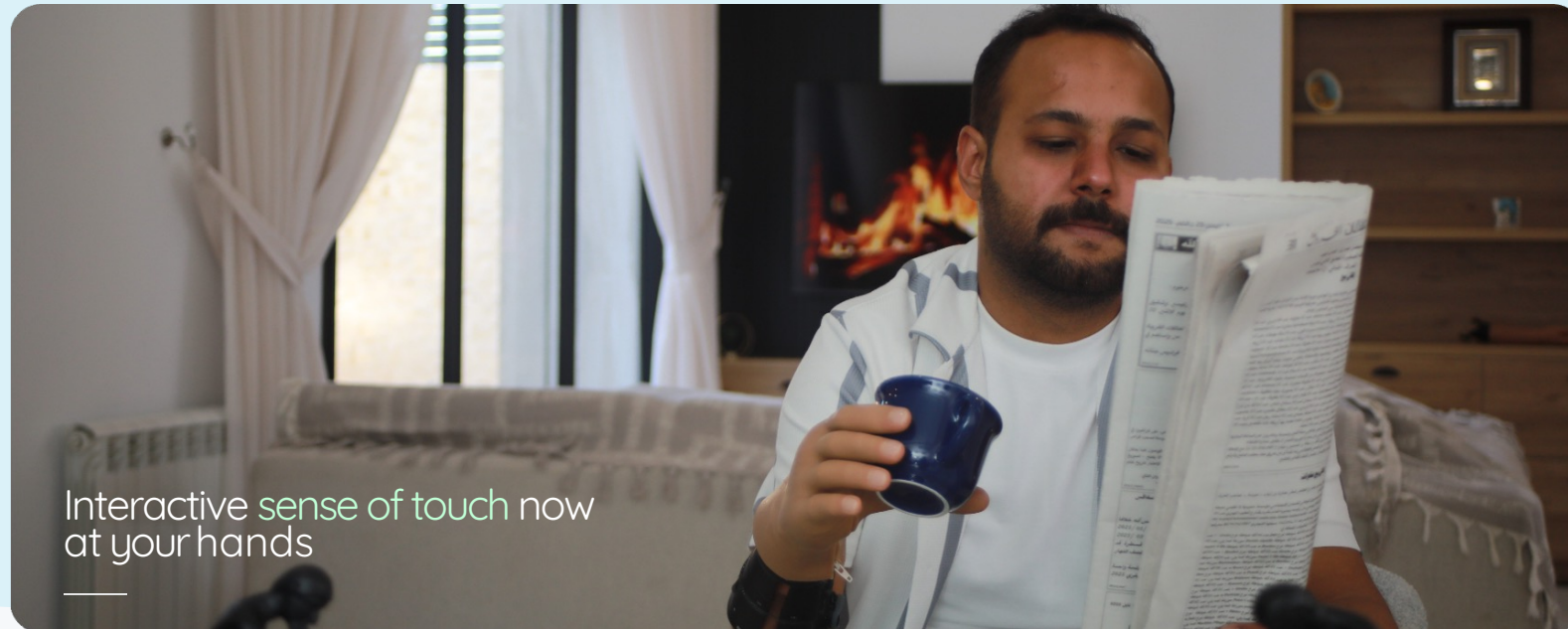
Freeze
Mode



Integration with our
Mobile App



Haptic
Feedback



Interactive **sense of touch** now
at your hands

Angel Aesthetic Prosthetic Arm





Flow Edition

Functional
Plug & Play
Terminal tools

Glow Silicone Parts



Hasdrubal Prosthetic Legs



Below-Knee Prosthetic



Above-Knee Prosthetic



Hip-Knee Prosthetic

Hasdrubal Prosthetic Sockets

Hasdrubal Socket
GEN 1

Fixed Socket: Provides a secure fit and strong support without the need for adjustments.



Hasdrubal Socket
GEN 2

Adjustable Socket: Offers greater flexibility, allowing size modifications for optimal comfort and better adaptation to physical changes.



MyoRehab Ecosystem

MyoRehab Ecosystem

MyoRehab Solutions

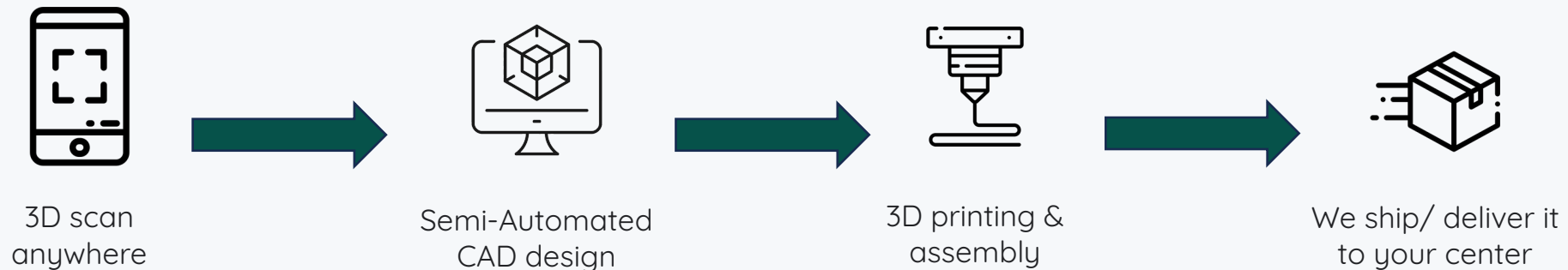
We offer an **innovative rehabilitation ecosystem solution** with a **smart Bluetooth devices** that links **muscle activity** to the **MyoLink app** and the **MyoController** for **interactive training** and **enhanced control**.



Socket manufacturing service

We can build your prosthetic socket with just a 3D scan

But how?



We deliver your prosthetic anywhere, anytime—completely remotely.

Regulatory compliance

CE mark / ISO 13485 / ISO 9001

Our company has obtained the **CE** for medical devices, which confirms our commitment to adhering to European standards for quality and user safety.

	<small>Registered Office Address Bd Brand Whitlock 30 B-1200 Brussels Belgium Registered Address Bd Général Wahnis 53 B-1030 Brussels Belgium</small>	<small>www.obelis.net +32 2 732 59 54</small>
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CERTIFICATE OF MDR NOTIFICATION

Reference No.: MG 0568-2024 **Date:** 21/10/2024
Order No.: EU AM 0362-2024

This is to certify that, according to the Regulation (EU) 2017/745 we, here at Obelis s.a., have performed all notification duties and responsibilities as the European Authorized Representative (EC REP) of:

Name: Cure Bionics **Address:** Rue Abderrahman Ibn Abi Bakr, 231
4023, Sousse
Tunisia

AS STIPULATED AND DEMANDED BY THE AFOREMENTIONED REGULATION:

The manufacturer declares that the Class I device(s) comply(ies) with the Regulation including all general safety and performance requirements.
The manufacturer has provided Obelis s.a. with all the appropriate declarations as per the Regulation (EU) 2017/745 Article 52 requirements, including the EC Declaration of Conformity (according to Annex IV), confirming that their Class I medical device(s), as stipulated here below, is/are fulfilling the applicable requirements of the Regulation (EU) 2017/745.
The notification of the following medical device(s) has been completed by Obelis s.a. (EC REP) in compliance with the Regulation (EU) 2017/745 on the **14/10/2024**

MEDICAL DEVICE(S): PLEASE SEE ANNEX A - LIST OF DEVICES (1 Device 1 Page)

As of the **15/10/2024**, and provided that the manufacturer will continue complying with the hereabove mentioned requirements*, he therefore:

- Is required to affix the CE marking on this(ese) device(s);
- May place this(ese) device(s) on the European Union and EEA territory.


Mr. G. Elkayam CEO
Obelis sa

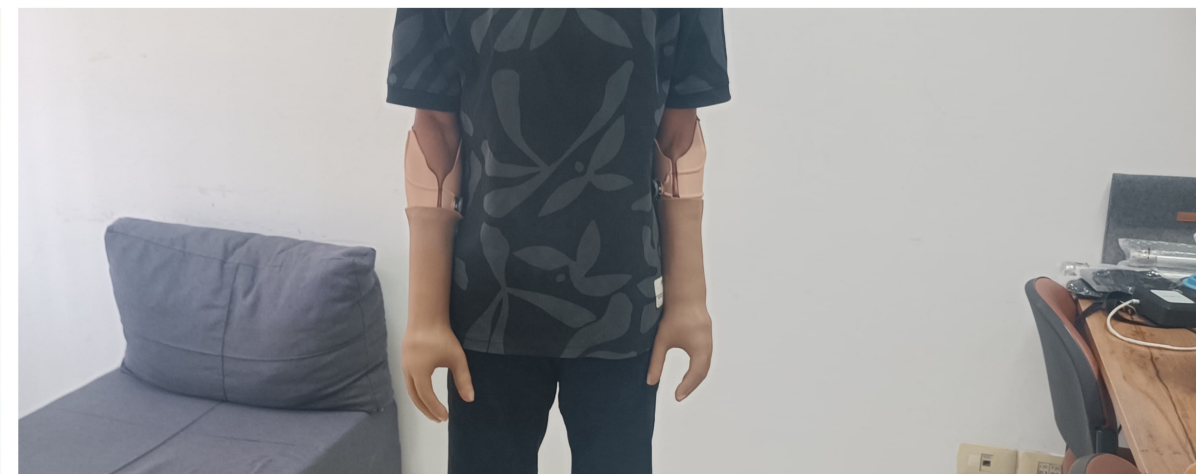
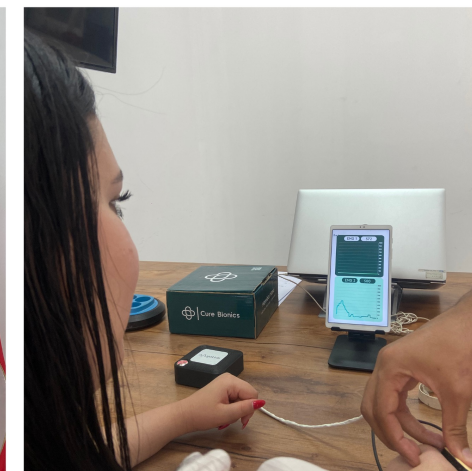


Obelis European Authorized Representative Center is member of the European Association of Authorized Representatives (E.A.A.R.).
Obelis s.a. is ISO 9001:2015 and ISO 13485:2016 certified.

*This certificate will become void automatically upon termination of the EAR agreement or removal of the products from EARMandate

V5 - ID: 00072820 DOC11.01.0004 - 12/09/2022

Some Happy Patients



Supporting organizations



Media coverage



Supported &
recognized
by

First and only certified prosthetic technology company in Africa and ME



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