



## DECLARATION OF CONFORMITY

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**WE, DESIN LLC, UNDER OUR SOLE RESPONSIBILITY, HEREBY DECLARE:**

**DEVICE (MODEL):**

*Obi (IFD-500-030)*

**DESCRIPTION**

*Obi is a reusable robotic utensil intended to compensate for the function of a human arm during mealtime activities to restore functional eating status.*

*(Class I Medical Device, Active, Non-Sterile, Non-Measuring)*

*BASIC UDI-DI: 08691740001Obi34*

**CONFORMS WITH THE FOLLOWING STANDARDS:**

| NUMBER             | TITLE   |
|--------------------|---|
| ISO13485: 2016     | Medical Devices - Quality Management Systems - Requirements for Regulatory Purposes   |
| IEC60601-1: 2012   | Medical Electrical Equipment - General Requirements for Basic Safety and Essential Performance.   |
| NEN EN 15284: 2007 | Materials and Articles in Contact with Food Stuffs. Test Method for the Resistance to Microwave Heating of Ceramic, Glass, Glass-Ceramic or Plastic Cookware. |
| EN50581:2012       | Technical Documentation for the Assessment of Electrical and Electronic Products with Respect to the Restriction of Hazardous Substances                      |

**Note:** No Common Specification is applicable.

**AND IF APPLICABLE, CONFORMS WITH THE FOLLOWING DIRECTIVES:**

| NUMBER       | TITLE   |
|--------------|---|
| MDR 2017/745 | Medical Device Regulation                     |
| EC/1935/2004 | Food Contact Directive                        |
| 2014/30/EU   | Electromagnetic Compatibility (EMC) Directive |
| 2014/35/EU   | Low Voltage Directive                         |
| 2015/863     | RoHS3 Directive                               |
| 2012/19/EU   | WEEE2 Directive                               |
| 93/68/EEC    | CE Marking Directive                          |

**ORIGINATED, REVIEWED AND APPROVED BY:**

Jon Dekar, CEO, DESIN LLC

**SIGNATURE:** \_\_\_\_\_

**DATE (MM/DD/YYYY):** \_\_\_\_\_

*[Signature]*  
04/15/2025