



# CERTIFICATE



This is to certify that the company

## **SprintRay Inc.**

2710 Media Center Drive, Suite #100A  
Los Angeles, CA 90065  
United States of America

with the organizational units/sites as listed in the annex

has implemented and maintains a **Quality Management System**.

Scope of certification:

Design, manufacture and distribution of non-active, non-implantable, non-sterile dental materials for use in the area of digital dentistry.

**-AUS(a), BRA, CND, JPN, USA (a,b,c,d)**

Through an audit, documented in a report, performed by DQS Medizinprodukte GmbH, it was verified that the management system fulfills the requirements of the following standard:

## **ISO 13485 : 2016**

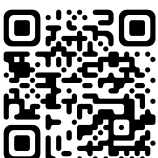
including applicable country-specific regulatory requirements, as indicated below the scope (full references of abbreviations are listed in the annex)

|                              |                  |
|------------------------------|------------------|
| Certificate registration no. | 31622718 MDSAP16 |
| Certificate unique ID        | 1000308909       |
| Effective date               | 2026-09-08       |
| Expiry date                  | 2029-09-07       |
| Frankfurt am Main            | 2026-09-08       |



## **DQS Medizinprodukte GmbH**

Sven Hoffmann  
Managing Director





**Annex to certificate**  
**Certificate registration No.: 31622718 MDSAP16**  
**Certificate unique ID: 1000308909**  
**Effective date: 2026-09-08**

## **SprintRay Inc.**

2710 Media Center Drive, Suite #100A  
Los Angeles, CA 90065  
United States of America

### **Audited site**

**31637665**

#### **SprintRay Inc.**

2710 Media Center Drive, Suite #100A  
Los Angeles, CA 90065  
United States of America

### **REPs FEI No.: site scope and country-specific requirements**

Administrative Activities and Head Quarters  
Functions.

**-AUS (a), BRA, CND, JPN, USA (a,b,c,d)**

**REPs FEI No.: F009064**

**31635806**

#### **SprintRay Inc.**

2705 Media Center Drive, Suite #100A  
Los Angeles, CA 90065  
United States of America

Customer Service, Supply Chain, Warehouse,  
Human Resources, Fulfillment, Distribution,  
Manufacturing and QC.

**-AUS (a), BRA, CND, JPN, USA (a,b,c,d)**

**REPs FEI No.: F009064**

**31622719**

#### **SprintRay Inc.**

17601 Sampson Ln  
Huntington Beach, CA 92647  
United States of America

Design, Manufacturing, Quality Assurance, and  
Regulatory Affairs.

**-AUS (a), BRA, CND, JPN, USA (a,b,c,d)**

**REPs FEI No.: F009064**



**Annex to certificate**  
**Certificate registration No.: 31622718 MDSAP16**  
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United States of America

### **Full references of country-specific requirements of MDSAP participating Regulatory Authorities**

| <b>Abbreviation</b> | <b>Jurisdiction</b> | <b>Reference</b>                                                                                                                                                                                                                         |
|---------------------|---------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| AUS                 | Australia           | (a) Therapeutic Goods (Medical Devices) Regulations 2002, Schedule 3, Part 1 – Full Quality Assurance Procedure<br>(b) Therapeutic Goods (Medical Devices) Regulations 2002, Schedule 3, Part 4 – Production Quality Assurance Procedure |
| BRA                 | Brazil              | RDC ANVISA n. 665/2022<br>RDC ANVISA n. 551/2021<br>RDC ANVISA n. 67/2009                                                                                                                                                                |
| CND                 | Canada              | Medical Devices Regulations – Part 1- SOR 98/282<br>Medical Devices Regulations – Part 1.1 – SOR 98/282 (as applicable)                                                                                                                  |
| JPN                 | Japan               | MHLW Ministerial Ordinance 169, Article 4 to Article 68<br>Japan PMD Act (as applicable)                                                                                                                                                 |
| USA                 | United States       | (a) 21 CFR Part 803<br>(b) 21 CFR Part 806<br>(c) 21 CFR Part 807<br>(d) 21 CFR Part 820<br>(e) 21 CFR Part 821                                                                                                                          |