

Centurion OZIL® HP / Active Sentry HP, LF-SG-146-STED

DECLARATION OF CONFORMITY			
(check all additional conformity route(s) based on EU MDD Article 11 requirements for the device class and specifics)			
Annex II (3) <input checked="" type="checkbox"/> Annex II (4) <input type="checkbox"/>	Annex III <input type="checkbox"/> Annex IV <input type="checkbox"/>	Annex V <input type="checkbox"/> Annex VI <input type="checkbox"/>	Annex VII <input type="checkbox"/>
<p>Technical File Number: LF-SG-146-STED            Device Trade Name: Centurion OZIL® HP / Active Sentry HP            Supersedes (Date): 27-APR-2020</p> <p>Manufacturer: Alcon Laboratories, Incorporated      Authorized Representative in the European Community:            Address: 6201 South Freeway, Fort Worth, TX      Alcon Laboratories Belgium            76134-2099, USA      Address: Lichterveld 3, 2870 Puurs-Sint-Amands, Belgium</p>			
Medical Device (Trade Name)	GMDN Code	Catalogue Number	Class
ASSY, Ship, Centurion OZIL HP	34900	8065751761	IIb
ASSY, Ship, Active Sentry HP	34900	8065752914	IIb
<p>The device(s) covered by this declaration are in conformity with the European Medical Devices Directive 93/42/EEC as well as other, relevant Union legislation that make provisions for the issuing of a declaration of conformity as listed.            Alcon Laboratories, Incorporated. hereby declares under its sole responsibility that the listed device(s) and Quality Systems conform(s) to:</p> <p style="text-align: center;"><i>EU MDD 93/42/EEC as amended</i></p> <p>This Declaration is applicable to all products listed and released after the Date of Issuance of this Declaration of Conformity, and until a revised Declaration of Conformity is issued.</p> <p>Notified Body Information: Applicable <input checked="" type="checkbox"/> Not Applicable <input type="checkbox"/></p> <p>Conformity Assessment Certificate Number(s): G1 020895 0393</p> <p>Conformity Certificate Validity Period: Valid from 05-FEB-2021 Valid until 26-MAY-2024</p> <p>Notified Body: TÜV SÜD Product Service GmbH</p> <p>Identification number: 0123</p> <p>Address: Ridlerstraße 65, 80339 Munich, Germany</p>			
Place of Issue: Lake Forest, CA, USA	Date of Issue: 25-APR-2021	Name/Title/Function/Date: Kim Regis/Sr. Director/GRA Surgical Instrumentation	