EU DECLARATION OF CONFORMITY

Technical Documentation Name	senofilcon A with Light Filtering Additive contact lenses		
Version Number	2.0		
Product Identification	**Trade Name of Device** The following product listing includes Diagnostic, Revenue, and Kit Configurations:	**Device Name**	**Basic UDI-DI**
	ACUVUE® OASYS MAX 1-Day Contact Lenses	ACUVUE OASYS MAX 1-DAY	0733905a00011BK
	ACUVUE® OASYS MAX 1-Day MULTIFOCAL Contact Lenses		
Legal Manufacturer	Johnson & Johnson Vision Care, Inc. 7500 Centurion Parkway Jacksonville, Florida 32256 United States		
EU Authorised Representative	AMO Ireland Block B, Liffey Valley Office Campus, Quarryvale, Co. Dublin D22 X0Y3, Ireland		
Notified Body	BSI Group The Netherlands B.V. Say Building, John M. Keynesplein 9, 1066 EP Amsterdam Netherlands Phone : +31 (0)20 346 07 80 Notified Body number : 2797		
Intended Purpose	The ACUVUE® OASYS MAX 1-Day Contact Lenses are intended for Daily Wear for the optical correction of myopia (short-sightedness) and hyperopia (long-sightedness) in persons with healthy eyes that may have 1.00D or less of astigmatism. The ACUVUE® OASYS MAX 1-Day MULTIFOCAL Contact Lenses are intended for Daily Wear for the optical correction of myopia (short-sightedness) and hyperopia (long-sightedness) in presbyopic persons with healthy eyes that may have 0.75D or less of astigmatism. The contact lenses contain a UV blocker to help provide protection against transmission of harmful UV radiation to the cornea and into the eye.		
Classification	IIa		
Product Codes	Universal Product Codes (UPC) for the contact lenses are provided within the ERP (Enterprise Resource Planning) SAP system.		

GMDN Codes	47841, Soft corrective contact lens, daily-disposable
EMDN (CND) Code	Q021004010101, Contact lenses-Hydrogel, Daily Single-Use
Manufacturer's Single Registration Number (SRN)	Not Yet Available
Design, Manufacturing and Distribution Sites	This document is valid for all medical devices described originating from the following sites: Johnson & Johnson Vision Care, Inc. 7500 Centurion Parkway Jacksonville, Florida 32256 United States Johnson & Johnson Vision Care Ireland UC The National Technology Park Limerick V94 N732 Ireland

This Declaration of Conformity is issued under the sole responsibility of the manufacturer.

We, Johnson & Johnson Vision Care, Inc., hereby declare the above listed medical devices comply with Medical Device Regulation (MDR) 2017/745.

This declaration is made on the basis of MDR Certificate Number 732087, issued by above stated Notified Body, in accordance with the conformity assessment laid down in Annex IX of MDR 2017/745.

SIGNATURES

Place of Issue	**Refer to Manufacturer's Address above**

Victoria Brennand

Digitally signed by Victoria Brennand
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Reason: I attest to the accuracy and integrity of this document.
Date: 2022.06.02 13:06:29 -04'00'
Adobe Acrobat version: 2020.013.20064

Date	June 2, 2022

Victoria Brennand
Associate Director, Regulatory Affairs
Johnson & Johnson Vision Care, Inc.
Jacksonville, Florida 32256, USA

THOMAS WILKINSON JR

Digitally signed by THOMAS WILKINSON JR
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Date	June 2, 2022

Thomas Wilkinson
Director, Quality Systems, Quality Compliance
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