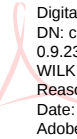




EU DECLARATION OF CONFORMITY

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 DN: c=US, o=JNJ, ou=Subscribers, 0.9.2342.19200300.100.1.1=1039331, cn=Victoria Brennand 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 DN: c=US, o=JNJ, ou=Subscribers, 0.9.2342.19200300.100.1.1=175047, cn=THOMAS WILKINSON JR Reason: I attest to the accuracy and integrity of this document Date: 2022.06.02 13:56:56 -04'00' Adobe Acrobat Reader version: 2020.013.20064	Date	June 2, 2022	
Thomas Wilkinson Director, Quality Systems, Quality Compliance Johnson & Johnson Vision Care, Inc. Jacksonville, Florida 32256, USA			