abbvie

Declaration of Conformity to Saudi Food and Drug Authority Medical Devices Interim Regulation

Manufacturer Identification Number assigned by the SFDA: M00980

I hereby declare that the medical device identified below complies with the MEDICAL DEVICES INTERIM REGULATION and its associated Implementing Rules and have been authorized by the SFDA to be placed on the KSA market.

The Authorised Representative is Cigalah Healthcare Company LLC, PO Box 19435, Jeddah, 21435, Kingdom of Saudi Arabia.

Import by: CIGALAH Healthcare Company LLC

#	Medical Device Trade Name1	Quantity	Serial Number/ Batch Number	Medical Device Listing National Registry Number (issued by MDMA system)2 or MDMA Application Number	Medical Device Listing National Registry Number (issued by MDNR system)3
1	CoolAdvantageCard,CD 2,Indirect,24 Cycles	17		A36ME0000000370SFDAA00004	
2	CoolAdhesive Pad, 8 Pack Carton	26	L202518204	A17ME0000000370SFDAA00004	
3	Kit, 8-Pk Gel Trap, CA Family	26	A202310-02	A40ME0000000370SFDAA00004	
4	Gasket, 24-Pack, CoolAdvantage	10	S202310-01	A38ME0000000370SFDAA00004	
5	Zeltiq Pretreatment Skin Wipe - Qty 24	17	U202408-01	A25ME0000000370SFDAA00004	
6	CoolAdhesive Pad, 8 Pack Carton	20	L202518204	A17ME0000000370SFDAA00004	
7	CoolAdhesive Plus Pad, 8 Pack Carton	6	L202517604	A18ME0000000370SFDAA00004	
8	Syringe, CoolGel Mini, 24-Pack Carton	2	U202503-01	A26ME0000000370SFDAA00004	

Note: Can be attached as a list

Authorised Signatory (on behalf of the manufacturer):

Name: Derek Carrick

Position: Customer Logistics Planner

Date: 20 August 2025

20/08/2025

Signature: X Derek Carrick

Derek Carrick
Supply Chain Analyst
Signed by: CARRIDX2

20/08/2025

AbbVie Logistics B.V.
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