PCA

NERS CERTIFICATE

ACCURAGE



Certificate

ISO 13485: 2016

KAF GRUP SAĞLIK HİZ. İNŞ. SAN. VE TİC. LTD. ŞTİ.

Head Office: Atakent Mah. 221. Sk. Rota Office A Blok No: 3 A İç Kapı No: 83
Küçükçekmece /İstanbul /TURKEY
İstanbul Manufacturing Address: Hadımköy Mahallesi Deniz Kızı Sok. No: 4 / 5
Arnavutköy / İstanbul/TURKEY

This certificate shows that the medical devices - quality management system (ENISO 13485:2016) of the above company was approved by PCA Certification for the following scope, the validity of the certificate depends on the company's pass the annual surveillance audits and company's maintenance the related management system conditions according to international accreditation criteria

SCOPE

Manufacturing and sales of barrier cream wash gloves, washcap-shampoo cap, incontinence body wipes and perineal cloths; body and perineal are cleansing wipes, splints, fixers / flaster tapes and tube fixers, ultrasound ecg gel, eeg paste, alcohol-free foam hand sanitizer, oral care swab, oral care set swab, alkali, neutralizer and enzyme containing solution, stretcher paper roll

GROUP CODE

A

 Certificate No
 : TC-75264

 Registration Date
 : 24.03.2020

 Reissue Date
 : 29.03.2022

Expiry Date

Certificate Period : 3 Years (From the date of registration)

Exclusion : 4.1.5 / 4.1.6 / 7.3.7 / 7.5.3 / 7.5.4 /

7.5.9.2 / 7.5.10 / 8.3.4

: 23.03.2023

IAS ACCREDITED

Management Systems Certification Body

MSCB-103

PCA Certification Approval

PCA Sertifikasyon Hizmetleri Limited Şirkeli Orta Mah. Ordu Sk. İzpark C Blok No:26/23 Kartal / İSTANBUL Tel: +90 216 510 63 48-49 Pbx Faks: +90 216 517 63 49 www.pca-tr.com info@pca-tr.com

FR.86 Rev.4



KAF GRUP SAĞLIK HİZMETLERİ SANAYİ VE TİCARET LİMİTED ŞİRKETİ

MAIN ADDRESS: ATAKENT MAH.221.SOK ROTA OFFICE A BLOK NO:3A D:82-83 KÜÇÜKÇEKMECE İSTANBUL FACTORY ADDRESS: HADIMKÖY MAH.DENİZ KIZI SOK.NO:4/5 ARNAVUTKÖY İSTANBUL

ISO 9001:2015

PRODUCTION AND SALES OF STERILE AND NON-STERILE MEDICAL DEVICES AND CONSUMABLES (BODY CLEANER, HAND CLEANING WIPES, HAIR RESTRAINT ETC)

This certificate is valid for 3 years on condition that surveillance audits are carried out before the period of each year in accordance with the accreditation rules. With this certificate, the holder of the certificate has been audited on the address mentioned above by TRB Certification Body in accordance with the condition and requirements of the license contract and it has been registered that the company satisfies the conditions of the standard. This certificate is valid as long as the certificate holder follows TRB certification rules and the certificate is seen as valid on www.trb.com.de web site certificate inquiry section.

Certification Number

: DE-QC-1914

Date of Issue

: 15.02.2021

Expiry Date
Certification Period

: 14.02.2023 : 3 Years (2. Year)

Printing Number

: 01













KAF GRUP SAĞLIK HİZMETLERİ İNŞAAT SANAYİ VE TİCARET LİMİTED ŞİRKETİ

MAIN ADDRESS: ATAKENT MAH.221.SOK ROTA OFFICE A BLOK NO:3 A D:82-83 KÜÇÜKÇEKMECE İSTANBUL FACTORY ADDRESS: HADIMKÖY MAH.DENİZ KIZI SOK.NO:4/5 ARNAVUTKÖY İSTANBUL

GMP

PRODUCTION AND SALES OF STERILE AND NON-STERILE MEDICAL DEVICES

This certificate is valid for 3 years on condition that surveillance audits are carried out before the period of each year in accordance with the accreditation rules. With this certificate, the holder of the certificate has been audited on the address mentioned above by TRB Certification Body in accordance with the condition and requirements of the license contract and it has been registered that the company satisfies the conditions of the standard. This certificate is valid as long as the certificate holder follows TRB certification rules and the certificate is seen as valid on www.trb.com.de web site certificate inquiry section.

Certification Number

: DE-GMP-051

Date of Issue Expiry Date : 15.01.2021 : 14.01.2023

Certification Period

: 3 Years (2. Year)

Printing Number

:01





A g out of General Manager



KAF GRUP SAĞLIK HİZMETLERİ SANAYİ VE TİCARET LİMİTED ŞİRKETİ

MAIN ADDRESS: ATAKENT MAH.221.SOK ROTA OFFICE A BLOK NO:3A D:82-83 KÜÇÜKÇEKMECE İSTANBUL FACTORY ADDRESS: HADIMKÖY MAH.DENİZ KIZI SOK.NO:4/5 ARNAVUTKÖY İSTANBUL

ISO 14001:2015

PRODUCTION AND SALES OF STERILE AND NON-STERILE MEDICAL DEVICES AND CONSUMABLES (BODY CLEANER, HAND CLEANING WIPES, HAIR RESTRAINT ETC)

This certificate is valid for 3 years on condition that surveillance audits are carried out before the period of each year in accordance with the accreditation rules. With this certificate, the holder of the certificate has been audited on the address mentioned above by TRB Certification Body in accordance with the condition and requirements of the license contract and it has been registered that the company satisfies the conditions of the standard. This certificate is valid as long as the certificate holder follows TRB certification rules and the certificate is seen as valid on www.trb.com.de web site certificate inquiry section.

Certification Number

: DE-EC-1914

Date of Issue

: 15.02.2021

Expiry Date

: 14.02.2023

Certification Period

: 3 Years (2. Year)

Printing Number

: 01













KAF GRUP SAĞLIK HİZMETLERİ SANAYİ VE TİCARET LİMİTED ŞİRKETİ

MAIN ADDRESS: ATAKENT MAH.221.SOK ROTA OFFICE A BLOK NO:3A D:82-83 KÜÇÜKÇEKMECE İSTANBUL FACTORY ADDRESS: HADIMKÖY MAH.DENİZ KIZI SOK.NO:4/5 ARNAVUTKÖY İSTANBUL

ISO 45001:2018

PRODUCTION AND SALES OF STERILE AND NON-STERILE MEDICAL DEVICES

This certificate is valid for 3 years on condition that surveillance audits are carried out before the period of each year in accordance with the accreditation rules. With this certificate, the holder of the certificate has been audited on the address mentioned above by TRB Certification Body in accordance with the condition and requirements of the license contract and it has been registered that the company satisfies the conditions of the standard. This certificate is valid as long as the certificate holder follows TRB certification rules and the certificate is seen as valid on www.trb.com.de web site certificate inquiry section.

Certification Number

: DE-SC-1914

Date of Issue Expiry Date :01.03.2021

Certification Period

: 28.02.2023

Printing Number

: 3 Years (2. Year)

: 0:









THE REPUBLIC OF TURKEY MINISTRY OF HEALTH TURKISH MEDICINES AND MEDICAL DEVICES AGENCY

Certificate No: 222489

Date of Issue: 26 April 2022

CERTIFICATE OF FREE SALE

To whom it may concern,

It is hereby certified that the products detailed in the attached schedule, which are manufactured by "KAF GRUP SAĞLIK HİZMETLERİ İNŞAAT SANAYİ VE TİCARET LİMİTED ŞİRKETİ" (Atakent Mh. 221 Sk. No:3/A KÜÇÜKÇEKMECE İSTANBUL), have been affixed with the CE mark in accordance with Medical Device Directives of the European Union (EU) and are freely sold in Turkey and EU.

This certificate is issued to be given to the relevant competent authorities of other countries and is valid for 36 months from the date of issue.

Yours sincerely,

Omer Faruk KURU

Head of Medical Devices

Registration and Coordination Department

This certificate consists of 2 page/s and 8 products. The products listed in the attached schedule are registered from the date of issuance of this certificate and information about the current status of these products is accessible through

https://utsuygulama.saglik.gov.tr/UTS/vatandas#/vatTibbiCihazListele.





Address: Sögütözü Mahallesi, 2176. Sokak No:5 06520 Çankaya/ANKARA Phone: +90 312 218 30 00 Fax: +90 312 218 34 60 https://www.titck.gov.tr

Date of Issue: 26 April 2022

PRODUCT SCHEDULE

#	Barkod	Brand	Label Name	Reference No / Version / Model	GMDN Code
1	8682022062215	ONECARE	BARRIER CREAM INCONTINENCE PERINEAL CLOTHS	KAFG02	46205
2	8682022062246	ONECARE	BARRIER CREAM INCONTINENCE PERINEAL CLOTHS	KAFG06	46205
3	8682022062208	ONECARE	BARRIER CREAM INCONTINENCE WIPES	KAFG01	46205
4	8682022062222	ONECARE	BODY CARE SET	KAF G04	46205
5	8682022062260	ONECARE	BARRIER CREAM INCONTINENCE WIPES	KAFG08	46205
6	8682022062253	ONECARE	BARRIER CREAM INCONTINENCE WIPES	KAFG07	46205
7	8682022062239	ONECARE	BARRIER CREAM INCONTINENCE PERINEAL CLOTHS	KAFG05	46205
8	8682079003155	ONECARE	WASH GLOVES WITH BARRIER CREAM	KAFG16	46205

End of product schedule.









Atakent Mahallesi 221 Sk. No:3A Rota Office A Blok Kat 14 D:83 Küçükçekmece/İstanbul/Türkiye

EU Medical Device Regulation 2017/745

Declaration of Conformity

Manufacturer Name(*)	KAF GRUP SAĞLIK HİZMETLERİ İNŞ. SAN. VE TİC. LTD. ŞTİ			
Manufacturer Address(*)	Atakent Mah. 221 Sk. No:3A Rota Office A Blok Kat:14 D:83 Küçükcekmece/Istanbul/Turkey			
Manufacturer Individual Identity No.				
If the product is produced by someone else by the manufacturer, the Manufacturer's Name and Address (* if any)				
Name of the product(*)	Barrier Cream Incontinence Wipes Barrier Cream Incontinence Perineal Cloths Wash Gloves With Barrier Cream			
	Product Name Barrier Cream Incontinence Wipes (8 pcs)	Catalog No KAF G49		
	Barrier Cream Incontinence Wipes (12 pcs)	KAF G01		
	Barrier Cream Incontinence Wipes (25 pcs)	KAF G08		
Catalog/Reference No.(*)	Barrier Cream Incontinence Wipes (50 pcs)	KAF G07		
110.(")	Barrier Cream Incontinence Wipes (100 pcs)	KAF G50		
	Barrier Cream Incontinence Perineal Cloths (8 pcs)	KAF G51		
	Barrier Cream Incontinence Perineal Cloths (12 pcs)	KAF G06		
	Barrier Cream Incontinence Perineal Cloths (25 pcs)	KAF G02		



Atakent Mahallesi 221 Sk. No:3A Rota Office A Blok Kat 14 D:83

Küçükçekmece/İstanbul/Türkiye

	Barrier Cream Incontinence Perineal Cloths (50 pcs)		KAF G05	
	Barrier Cream Incontinence Perineal Cloths (100 pcs)		KAF G52	
	Wash Gloves With Barrier Cream		KAF G16	
Purpose of usage(*)	ONECARE Barrier Creamy Incontinence Wipes and Perineal Cloths are used to create a barrier layer to protect the incontinence-related skin of patients with special needs, such as those with reduced mobility, bedridden and intensive care patients, by cleaning urine and feces contact from external factors. It prevents skin damage that may occur due to pH change and irritation in the body. ONECARE Wash Gloves with Barrier Cream are used to create a barrier layer to protect the incontinence-related skin of patients with special needs, such as those with reduced mobility, bedridden and intensive care patients, by cleaning urine and feces contact from external factors. It prevents skin damage that may occur due to pH change and irritation in the body.			
Basic UDI-DI(*)	Catalog No KAF G49 KAF G01 KAF G08 KAF G07 KAF G50 KAF G51 KAF G06 KAF G05 KAF G52 KAF G16	Basic UDI-DI 868256095H 868256095H 868256095H 868256095H 868256095H 868256095H 868256095H 868256095H 868256095H 868256095H 868256095H	CAFG01JN CAFG08K4 CAFG07K2 CAFG50K3 CAFG51K5 CAFG06JY CAFG02JQ CAFG05JW CAFG52K7	
Product Classification / Classification Rule(*)	Class 1			
GMDN Code(*)	46205			
EMDN Code (*After activation)	Z120113			
Conformity Assessment Procedure(*)	ANNEX-IV (Annex II & III) ANNEX-IX (CHAPTER I & III)		Declaration of conformity Quality management system Technical Documentation Mod.	
			Type Examination	tation Mod.



Atakent Mahallesi 221 Sk. No:3A Rota Office A Blok Kat 14 D:83

Küçükçekmece/İstanbul/Türkiye

(Additions executed in	ANNEX-XI (PART A) Production Quality Assurance			
the product evaluation are marked)	ANNEX-XI (PART	ΓB) Product Ve	erification	
Notified Body Name and Number (**)				
EU Certificate No and Description Start/Effective date (**)				
Other EU Legislation / Common	Harmonized Standards			
Specifications / Harmonized	EN ISO 13485:2016	EN ISO 10993-5: 2009	EN ISO 10993-1: 2020	
Standards to which the product complies	EN ISO 15223-1: 2021	EN ISO 10993-10: 2021	EN ISO 20417: 2021	

(*) Sections beginning with are required.

(**) The conformity assessment is mandatory for products made by the notified body.

As a company KAF GRUP SAĞLIK HİZMETLERİ İNŞ. SAN. TİC. LTD. ŞTİ, we declare under our sole responsibility that the devices covered by this declaration comply with the Regulation (EU) 2017/745 of the European Parliament and of the Council on Medical Devices and that the requirements specified in the Regulation are fulfilled for these devices.

Signature Date and Place: 05.08.2022

Effective Date (if applicable):

Signatory

() 7 70.0

Gökmen Aytin

Mission

General Manager

[Signature and Seal/Stamp]



Atakent Mahallesi 221 Sk. No:3A Rota Office A Blok Kat:14 D:83 Küçükçekmece/Istanbul/Turkey

EU Medical Device Regulation 2017/745

Declaration of Conformity

Manufacturer Name(*)	KAF GRUP SAĞLIK HİZMETLERİ İNŞ. SAN. VE TİC. LTD. ŞTİ			
Manufacturer Address(*)	Atakent Mah. 221 Sk. No:3A Rota Office A Blok Kat:14 D:83 Küçükcekmece/Istanbul/Turkey			
Manufacturer Individual Identity No.		*		
If the product is produced by someone else by the manufacturer, the Manufacturer's Name and Address (* if any)				
Authorized European Representative	Anxietas Ug. Industriestrasse 43, 50389 Berzdorf Köln/GERMANY Köln HRB: 106071 info@anxietas.de			
Name of the product(*)	WANCARE ULTRASOUND ECG GEL			
Catalog/Reference . No.(*)	Name of the Product WANCARE ULTRASOUND ECG GEL 1000ml WANCARE ULTRASOUND ECG GEL 500 ml WANCARE ULTRASOUND ECG GEL 250ml WANCARE ULTRASOUND ECG GEL 5 lt	Catalog No KAF G31 KAF G31-1 KAF G31-2 KAF G31-3		
Purpose of usage(*)	It cuts the air between the skin and the probe in all kinds of Ultrasonography, Droppler, EKG, Exercise Test applications, and ensures that the ultrasonography waves come to the device screen more clearly and uninterruptedly.			
Basic UDI-DI(*)	KAF G31 8682079003KAI KAF G31-1 868207900KAF KAF G31-2 868207900KAF KAF G31-3 868207900KAF	G31-1N4 G31-2N6		
Product Classification / Classification Rule(*)	Class 1			



SAĞLIK HİZMETLERİ

KAF GRUP SAĞLIK HİZMETLERİ İNŞ. SAN. TİC. LTD. ŞTİ

Atakent Mahallesi 221 Sk. No:3A Rota Office A Blok Kat:14 D:83 Küçükçekmece/Istanbul/Turkey

GMDN Code(*)	15321			
EMDN Code (*After activation)	A108002			
Conformity		II & III) Declaration	n of conformity	
Assessment Procedure(*)	ANNEX-IX (CHAP	TFR I &	anagement system	
	☐ ANNEX-IX (PART	II) Technical	Documentation Mod.	
(Additions executed in	☐ ANNEX-X	Type Exar	Type Examination	
the product	ANNEX-XI (PART	(A) Production	Production Quality Assurance	
evaluation are marked)	ANNEX-XI (PART	B) Product V	Product Verification	
Notified Body Name and Number (**)				
EU Certificate No and Description Start/Effective date (**)				
Other EU Legislation / Common	Harmonized Standards			
Specifications / Harmonized	EN ISO 13485:2016	EN ISO 10993-5: 2009	EN ISO 10993-1: 2020	
Standards to which the product complies	EN ISO 15223-1: 2021	EN ISO 10993-10: 2013	EN ISO 20417: 2021	

(*) Sections beginning with are required.

(**) The conformity assessment is mandatory for products made by the notified body.

As a company KAF GRUP SAĞLIK HİZMETLERİ İNŞ. SAN. TİC. LTD. ŞTİ, we declare under our sole responsibility that the devices covered by this declaration comply with the Regulation (EU) 2017/745 of the European Parliament and of the Council on Medical Devices and that the requirements specified in the Regulation are fulfilled for these devices.

Signature Date and Place: 27.07.2022

Effective Date (if applicable):

Signatory

Gökmen AYTIN

Mission

General Manager

[Signature and Seal/Stamp] KAT GRU

Shent Mr. 221 SK. No.3H. Rota diffice Blok D 3 K. Yokmede 1 STANBUL 0812 07 42 V. Fax 02 2 47 42 01 Holkal J. J. 86 053 3864