EG-KONFORMITÄTSERKLÄRUNG · EC DECLARATION OF CONFORMITY
DÉCLARATION CE DE CONFORMITÉ · DICHIARAZIONE CE DI CONFORMITÀ

Name und Adresse des Herstellers: /
Name and address of the manufacturer: /
Nom et adresse du fabricant: /
Nome e indirizzo del fabbricante:

Gauke Healthcare Co.,Ltd

Chengnan Industrial Park, Tuanfeng County, HuangGang, Hubei 438800, China

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that /
Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che

das Medizinprodukt: /
the medical device: /
le dispositif médical: /
il dispositivo medico:

ART. NO.	NAME OF PRODUCTS
B49003	DIN13157 first aid kit-fillings
B49003-201	DIN13157 first aid kit with BOX 201 ABS, with wall bracket
B49003-200	DIN13157 first aid kit with BOX 200 PP
B49003-303	DIN13157 first aid kit with BOX 303 ABS,
B49004	DIN13169 first aid kit-fillings
B49004-304	DIN13169 first aid kit with BOX 304 ABS,
B49004-300	DIN13169 first aid kit with-BOX 300 ABS,

The content list of the first aid kits:

Adhesive tape DIN13019-A 5x2.5	Class I
Plaster set	Class I
First aid dressings bandages, Sterile,DIN13151-K 6x8cm	Class Is
First aid dressings bandages, Sterile,DIN13151-M 8x10cm	Class Is
First aid dressings bandages, Sterile,DIN13151-G 10x12cm	Class Is
Sterile Compress(Burn dressings) DIN13152-A 60x80cm	Class Is
Conforming Elastic bandages DIN61634 FB6, 6cmx4m	Class I
Conforming Elastic bandages DIN61634 FB8, 8cmx4m	Class I
First aid blanket	Class I
Sterile wound compress 10x10cm	Class Is
Triangular bandages DIN13168-D 96x96x136cm	Class I
Scissor DIN58279 A145	Class I
Disposable gloves DIN EN 455-1	Class I
Wet wipe for cleaning skin	Class I
Eye pad 55x75mm	Class Is
Instant Cold compress	Class IIa
Garbage Plastic Bag	
Fleece	Class I

der Klasse: /
of class: /
de la classe: /
di classe:

Class I, Class Is and Class IIa

nach Anhang IX der Richtlinie 93/42/EWG / according to annex IX of directive 93/42/EEC /
selon l'annexe IX de la directive 93/42/CEE / secondo l'allegato IX della direttiva 93/42/CEE

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 93/42/EWG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“. /

meets the provisions of the directive 93/42/EEC and its transpositions in national laws which apply to it. The declaration is valid in connection with the “final inspection report” of the device. /

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remplit toutes les exigences de la directive sur les dispositifs médicaux 93/42/CEE et de ses transpositions en droit national qui le concernent. La déclaration est valable si elle est associée au «rapport de l'inspection finale» du produit. /

soddisfa tutte le disposizioni della direttiva 93/42/CEE e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il "rapporto di ispezione finale" del prodotto.

Konformitätsbewertungsverfahren: /
Conformity assessment procedure: /
Procédure d'évaluation de la conformité: /
Procedura di valutazione della conformità:

Directive 93/42/EEC Annex VII and V

Registrier-Nr.: /
Registration No.: /
N°d'enregistrement: /
Numero di registrazione:

DD 60038309 0001

Benannte Stelle: /
Notified Body: /
Organisme notifié: /
Organismo notificato:

TÜV Rheinland LGA Products GmbH
Tillystraße 2
90431 Nürnberg
Deutschland
CE 0197

Huanggang,11.Jan.2017

Ort, Datum / Place, date /
Lieu, date / Luogo, data

Name und Funktion / Name and function /
Nom et fonction / Nome e funzione