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:

Jiangsu Jichun Medical Devices Co.,Ltd
No. 98, Baiyang Bridge, Zhenglu Town, Tianning, Changzhou, Jiangsu 213111, 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:

Disposable Syringes
Models:

	SYRINGES WITHOUT NEEDLE	
Product code (REF)	**Fitting type**	**Capacity**
02071000090150	luer slip	1 ml/U100 (insulin)
02071000090180	luer slip	1 ml (tuberculin)
02073000090100	luer slip	2,5 ml
02075000090100	luer slip	5 ml
02076000090100	luer slip	10 ml
02075000090110	eccentric luer	5 ml
02076000090200	eccentric luer	10 ml
02077000090200	eccentric luer	20 ml
02078000090200	eccentric luer	30 ml
02079000090200	eccentric luer	50 ml
02071000090600	Luer lock	1 ml
02073000090600	luer lock	2,5 ml
02075000090600	luer lock	5 ml
02076000090600	luer lock	10 ml
02077000090600	luer lock	20 ml
02078000090600	luer lock	30 ml
02079000090600	luer lock	50 ml
02079005090800	catheter tip	50 ml
02079001090800	catheter tip	100 ml

	SYRINGES WITH NEEDLE		
Product code (REF)	**Fitting type**	**Capacity**	**Needle**
02071250090150	Central luer	U100 (insulin)	G25x5/8"
02071260090150	Central luer	U100 (insulin)	G26x1/2"
02071270090150	Central luer	U100 (insulin)	G27x1/2"
02071280090150	Central luer	U100 (insulin)	G28x1/2"
02071260090180	Central luer	1ml (tuberculin)	G26x1/2"
02071270090180	Central luer	1ml (tuberculin)	G27x1/2"
02073210090100	Central luer	2.5 ml	G21x1 ½"
02073220090100	Central luer	2.5 ml	G22x1 ½"
02073221090100	Central luer	2.5 ml	G22x1 ¼"
02073230090100	Central luer	2.5 ml	G23x1"
02075210090100	Central luer	5 ml	G21x1 ½"
02075220090100	Central luer	5 ml	G22x1 ½"
02075221090100	Central luer	5 ml	G22x1 ¼"
02075230090100	Central luer	5 ml	G23x1"
02075231090100	Central luer	5 ml	G23x1 ¼"

02076200090100	Central luer	10 ml	G20x1 ½"
02076210090100	Central luer	10 ml	G21x1 ½"
02076220090100	Central luer	10 ml	G22x1 ¼"
02076190090200	Eccentric luer	10 ml	G19x1 ½"
02076200090200	Eccentric luer	10 ml	G20x1 ½"
02076210090200	Eccentric luer	10 ml	G21x1 ½"
02077180090200	Eccentric luer	20 ml	G18x1 ½"
02077190090200	Eccentric luer	20 ml	G19x1 ½"
02077200090200	Eccentric luer	20 ml	G20x1 ½"
02077210090200	Eccentric luer	20 ml	G21x1 ½"

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

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. /

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.

EU Representative: Caretechion GmbH
Address: Niederrheinstr 71, 40474 Duesseldorf, Germany

Konformitätsbewertungsverfahren: / **Directive 93/42/EEC Annex V**
Conformity assessment procedure: /
Procédure d'évaluation de la conformité: /
Procedura di valutazione della conformità:

Registrier-Nr.: / **DD 60150044 0001**
Registration No.: /
N°d'enregistrement: /
Numero di registrazione:

Benannte Stelle: / **TÜV Rheinland LGA Products GmbH**
Notified Body: / **Tillystraße 2**
Organisme notifié: / **90431 Nürnberg**
Organismo notificato: **Deutschland**
CE 0197

Changzhou, 2020-06-23

2020-06-23

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

Pingnan Zhu , CEO

Pingnan Zhu

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