公司简介
Company Profile

本公司专业生产 6827 中医器械：针灸针、电针仪、掀针、皮内针、梅花针、火罐等。公司已有二十多年的生产经验，所有产品不仅具有传统的工艺技术同时也运用先进的生产技术和设备。

公司所生产产品不仅有国内的注册证和许可证，还获得欧盟的CE认证，并通过德国TUV的ISO13485质量体系认证，公司自主品牌“顺和”在国内外享有客户的一致好评。公司还与多家知名公司以“OEM”方式合作。

我们将竭尽全力为我们的客户提供优质的产品和具有竞争力的价格。我们真诚地欢迎您与我们联系以获取更多的信息。我们也期待不久的将来成为您可以信赖的供应商。

Our company is specialized in the production of 6827 Chinese medicine instruments: acupuncture needle, electronic acupuncture instrument, press needle, intradermal needle, seven star plum blossom needle and cupping jars etc.All the products not only have the traditional processing technology,but also adopt advanced testing equipment and quality inspection equipment .

The products produced by us not only own domestic registration certificates and licenses, also have the CE certification of EU,and passed the ISO13485 quality system certification issued by German TUV . Our company brand"Shunhe"enjoys good reputation at home and abroad market. What's more, our company have cooperated with several well-known companies in the "OEM" way.

We will try our best to provide our customers high quality products and competitive prices. We sincerely welcome you to contact us for more information. We also look forward to be your reliable supplier in the near future .

厂区图 The factory Image

苏州市华伦医疗用品有限公司
Suzhou Hualun Medical Appliance Co.,Ltd.

公司生产线一览图
General Chart of Production Line

1.生 产 Produce

公司现在多个机械自动化生产流水线车间，产量高、产品质量稳定

Now our company have multiple mechanical automation production line workshops with high output and stable product quality.

2.检验 Test

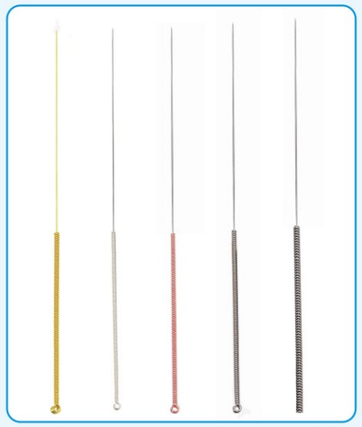
运用先进的投影技术检测产品针尖

Using advanced projection technology to detect needlepoint.

3.包装 Packaging

十万级净化车间保证产品质量

Hundreds of grade purification workshop to ensure products' quality



镀金针 镀银针 铜柄针 钢柄针 钢丝平柄针

4.针灸针部分手柄图
Image of acupuncture needles handle

针灸针规格表 Specification table of acupuncture needles

直径Diameter : 0.12 , 0.14 , 0.16 , 0.18 , 0.20 , 0.22 , 0.25 , 0.30 , 0.35.....

长度 Length : 7mm,9mm,13mm,25mm,40mm,50mm,60mm,75mm,100mm.....

特殊规格可根据客户要求订做

Special specifications can be customized according to customer requirements

File name：Declaration of Conformity

Document No：HL/CE-002-10

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:

Suzhou HuaLun Medical Appliance Co., Ltd.
Shengtian Economic Development Zone, HuangQiao
XiangCheng District, SuZhou, 215132 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 Acupuncture needles
Disposable Press needles

UMDNS-Code: 12-730

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

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. /

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 V

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

Certificate No.: DD 60140340 001
Issue date: 2019-07-09
Expiry date: 2024-05-27

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

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

Suzhou China 2020-12-28
Ort, Datum / Place, date /
Lieu, date / Luogo, data

MG ：俞瑞兴
Name und Funktion / Name and function /
Nom et fonction / Nome e funzione