第二十三届世界中医药大会
中医药服务贸易展参展通知

由世界中医药学会联合会主办的世界中医药大会是全球中医药领域规模大、参与广、层次高的学术盛会。至今已在中国（2004）、法国（2005）、加拿大、新加坡、中国澳门、澳大利亚（2009）、荷兰、英国、马来西亚、美国、俄罗斯、西班牙、新西兰、泰国、意大利、匈牙利、中国（2020 线上）、中国香港、巴西、菲律宾、法国（2024）、澳大利亚（2025）成功举办了 22 届。

第二十三届世界中医药大会拟于 2026 年 10 月 17 至 18 日在新加坡召开，由新加坡中医师公会承办。会议将组织多种形式的学术交流活动，为来自世界各地的中医药专家学者、政府官员、企业代表分享他们理论研究和临床经验、科研成果和新发明新创造提供平台。

活动期间，大会附设中医药服务贸易展，现就有关事宜通知如下：

一、组织结构

主办单位：世界中医药学会联合会

承办单位：新加坡中医师公会

二、时间及地点：

2026 年 10 月 17 日—18 日 （10 月 16 日大会报到）

新加坡 莱佛士城会议中心

世界中医药大会办公室（世界中医药学会联合会国际联络部）：王晶、李昕雪、杨柳、王婧雯
Office for World Congress of Chinese Medicine of WFCMS: Wang Jing, Li Xinxue, Yang Liu, Virginia Wang
E-mail: wccm@vip.163.com, Tel: +86-10-58650243/58650058/58650026/58650240

三、展位注册

本届大会展位报名采用网上会议平台，点击下方链接 https://wfcms.daanjiankang.com/2026，或微信扫描二维码进入报名网站。在“展商报名”板块进行报名。

四、展品须知

为确保展会合规，展商展览展品需符合《参展产品资质合规要求一览表》要求（后简称“一览表”，具体详见附件）。

五、参展费用

展位类型	尺寸	报名时间	价格 （人民币与美金按 1:7 汇率）
标准展位	3 米 x 3 米	2026 年 9 月 15 日及之前	28000 元（4000 美金）
		2026 年 9 月 16 日及之后	33600 元（4800 美金）
经济展位	3 米 x 2 米	2026 年 9 月 15 日及之前	22400 元（3200 美金）
		2026 年 9 月 16 日及之后	28000 元（4000 美金）
迷你展位	3 米 x 1 米	2026 年 9 月 15 日及之前	14000 元（2000 美金）
		2026 年 9 月 16 日及之后	16800 元（2400 美金）

- 以上价格包含展位面积、围板、1 张桌子、2 把椅子、1 个插座、基本照明，免 1-2 人参会注册费（标准展位和经济展位免 2 人注册费，迷你展位免 1 人注册费）。
- 展位具体信息请与大会组委会联络。鉴于各展位资源有限，若未能满足第一志愿，请服从组委会的统一统筹与协调。展位位置将严格按照款项到账的先后顺序进行分配，敬请知悉。

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五、联系方式

● 主办方：

世界中医药大会办公室（国际联络部）

联系人：王晶、李昕雪、杨柳、王婧雯

电子邮箱：wccm@vip.163.com

电话：+86（10）58650243 / 0240 / 0026

● 承办方：

新加坡中医师公会

联系人：许文楷、邓碧莹

电子邮箱：assoc@singaporetcm.com

电话：+65 68774206/68774223

世界中医药学会联合会

2026 年 8 月

附件：

《参展产品资质合规要求一览表》

一、中国大陆展商资质要求

此部分适用于在中国大陆注册的企业及其生产或经营的产品。

展品类别	企业主体资质（门槛）	产品上市资质（一物一证）
药品(含中药饮片/成药)	1. 营业执照（三证合一）2. 药品生产许可证（厂家）或药品经营许可证（经销商）3. GMP/GSP 认证符合性检查结果	1. 药品注册证（国药准字批件）2. 药品检验报告书（近期）3. 说明书及标签备案
药材(含中药材/中药饮片/民族药材)	1. 营业执照（三证合一）2. 药品生产许可证（生产型企业）或药品经营许可证（经营型企业）3. GMP/GSP 认证符合性检查结果 4. 中药材种植/养殖备案证明（如适用）	1. 药品注册证（国药准字批件，中药饮片除外）2. 药材质量检验报告书（含农残、重金属等指标）3. 产地证明（中药材）4. 炮制规范备案（中药饮片）
医疗器械(含设备/耗材/试剂)	1. 营业执照 2. 生产/经营许可证（一类备案，二类备案，三类许可）	1. 医疗器械注册证（或备案凭证）2. 产品技术要求说明书 3. 若含软件，需提供软件版本号说明
药食同源产品(普通食品类)	1. 营业执照 2. 食品生产许可证（SC 证）或食品经营许可证	1. 第三方检测报告（近一年内，含重金属/微生物）2. 产品执行标准（企标或国标）3. 严禁宣称治疗功效
药食同源产品(保健食品类)	1. 营业执照 2. 食品生产/经营许可证	1. 保健食品注册证书（或备案凭证）2. 保健食品标志（蓝帽子）及批准文号
AI 智能体(医疗 AI/软件)	1. 营业执照（含软件开发/销售范围）2. 软件著作权证书	1. 若用于诊断/治疗：必须提供医疗器械注册证（通常含 AI 软件组件）2. 若仅为辅助/管理：需提供软件测试报告及功能说明书 3. 数据安全与隐私合规声明

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二、海外及港澳台地区展商资质要求

此部分适用于在中国大陆以外地区（含港澳台）注册的企业及其产品。

核心原则： 对于不符合第一部分“中国大陆展商资质要求”的海外展商，请提供贵国或地区相关监管机构颁发的、证明企业及产品合法上市销售的同等效力资质文件。

展品类别	企业主体资质（门槛）	产品上市资质（一物一证）
药品/药材	1. 企业注册证明（如商业登记证）2. 药品生产/经营许可证（由所在国/地区药监部门颁发，如美国 FDA、欧盟 EMA 等）	1. 药品上市许可证明（如美国 NDA/ANDA、欧盟 MA 等）2. 产品检验报告（由权威第三方实验室出具）3. 产品说明书及标签
医疗器械	1. 企业注册证明 2. 医疗器械生产/经营许可证（如美国 FDA 注册、欧盟 CE 证书等）	1. 医疗器械上市许可证明（如美国 FDA 510(k)/PMA、欧盟 CE 证书等）2. 产品技术规格说明书
食品/保健食品	1. 企业注册证明 2. 食品生产/经营许可证（如美国 FDA 注册、欧盟卫生证书等）	1. 产品成分分析报告（由权威第三方实验室出具）2. 产品执行标准 3. 严禁宣称治疗功效
AI 智能体(医疗 AI/软件)	1. 企业注册证明 2. 知识产权证明（如软件著作权、专利证书等）	1. 若用于诊断/治疗：必须提供所在国/地区的医疗器械上市许可（如美国 FDA、欧盟 CE-IVDR 等）2. 若仅为辅助/管理：需提供软件测试报告及功能说明书 3. 数据安全与隐私合规声明（如符合 GDPR 等）

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第二十三届世界中医药大会

The 23rd World Congress of Chinese Medicine

The 23rd World Congress of Chinese Medicine

Exhibition Notice for TCM Service & Trade Exhibition

Hosted by the World Federation of Chinese Medicine Societies (WFCMS), the World Congress of Chinese Medicine is a high-profile global academic event featuring large scale, extensive participation and top-tier standards in the field of TCM. To date, 22 sessions have been successfully held across the globe: China (2004), France (2005), Canada, Singapore, Macao of China, Australia (2009), the Netherlands, the United Kingdom, Malaysia, the United States, Russia, Spain, New Zealand, Thailand, Italy, Hungary, Online Event hosted by China (2020), Hong Kong of China, Brazil, the Philippines, France (2024), and Australia (2025).

The 23rd World Congress of Chinese Medicine is scheduled to take place in Singapore from October 17 to 18, 2026, organized by the Singapore Chinese Physicians' Association. The conference will host diverse academic exchange sessions, offering a platform for TCM experts, scholars, government officials and enterprise representatives worldwide to share theoretical research, clinical experience, scientific research outcomes, new inventions and innovations.

Concurrent with the conference, a TCM Service & Trade Exhibition will be held. Relevant arrangements are notified as follows:

I. Organizational Structure

Host: World Federation of Chinese Medicine Societies (WFCMS)

Organizer: Singapore Chinese Physicians' Association

II. Date & Venue

Conference Dates: October 17 – 18, 2026

Check-in Date: October 16, 2026

Venue: Raffles City Convention Centre, Singapore

III. Booth Registration

Booth application shall be completed via the official online conference platform. Applicants may access the registration page through the link: https://wfcms.daanjiankang.com/wfcms-m/#/meet-home, or scan the QR code below via WeChat, then submit applications under the "Exhibitor Registration" section.

IV. Exhibition Product Guidelines

To ensure exhibition compliance, all displayed products by exhibitors must meet the requirements specified in the Compliance Checklist for Exhibited Product Qualifications (hereinafter referred to as the Checklist, see attachment for full details).

世界中医药大会办公室（世界中医药学会联合会国际联络部）：王晶、李昕雪、杨柳、王婧雯
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V. Exhibition Fees

Booth Type	Dimensions	Registration Deadline	Price
Standard Booth	3m × 3m	On or before September 15, 2026	RMB 28,000 (USD 4,000)
Standard Booth	3m × 3m	On or after September 16, 2026	RMB 33,600 (USD 4,800)
Economy Booth	3m × 2m	On or before September 15, 2026	RMB 22,400 (USD 3,200)
Economy Booth	3m × 2m	On or after September 16, 2026	RMB 28,000 (USD 4,000)
Mini Booth	3m × 1m	On or before September 15, 2026	RMB 14,000 (USD 2,000)
Mini Booth	3m × 1m	On or after September 16, 2026	RMB 16,800 (USD 2,400)

- The above fees cover booth space, partition panels, 1 table, 2 chairs, 1 power socket and basic lighting. Registration fees for 1-2 conference participants are waived(standard and economy booths waive registration fees for 2 participants; mini boothwaives registration fee for 1 participant).
- For detailed booth information, please contact the conference organizing committee. Please be advised that due to the limited availability of exhibition booths, you may be required to accept alternative arrangements if your first preference is unavailable. Booth assignments will be strictly allocated on a first-come, first-served basis, determined by the chronological order of payment receipt.

VI. Contact Information

Host: Office of World Congress of Chinese Medicine (International Liaison Department of WFCMS)

Contact Persons: Wang Jing, Li Xinxue, Yang Liu, Wang Jingwen

E-mail: wccm@vip.163.com

Tel: +86 (10) 58650243 / 58650240 / 58650026

Organizer: Singapore Chinese Physicians Association

Contact Persons: Koh Boon Khai, Tin Pit Inn

E-mail: assoc@singaporetcm.com

Tel: +65 68774206 / 68774223

World Federation of Chinese Medicine Societies

August 2026

Attachment:

Compliance Checklist for Exhibited Product Qualifications

Part 1: Qualification Requirements for Mainland China Exhibitors

Applicable to enterprises registered in Chinese Mainland and their manufactured or distributed products.

Product Category	Mandatory Enterprise Qualifications	Product Market Access Certifications (One Certificate per Product)
Pharmaceuticals (Including TCM Decoction Pieces & Proprietary Chinese Medicines)	1. Unified Social Credit Business License 2. Drug Manufacturing License (for manufacturers) or Drug Distribution License (for distributors) 3. GMP/GSP compliance inspection report	1. Drug Registration Certificate (National Medicine Approval Number) 2. Recent product testing report 3. Filed product insert and label
Medicinal Materials (Including Chinese Medicinal Herbs, TCM Decoction Pieces & Ethnic Medicinal Materials)	1. Unified Social Credit Business License 2. Drug Manufacturing License (for manufacturers) or Drug Distribution License (for distributors) 3. GMP/GSP compliance inspection report 4. Cultivation/breeding filing certificate for applicable herbal materials	1. Drug Registration Certificate (National Medicine Approval Number, not required for TCM decoction pieces) 2. Herbal quality test report (covering pesticide residue, heavy metal indicators, etc.) 3. Origin certificate for Chinese medicinal herbs 4. Filed processing specifications for TCM decoction pieces

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Product Category	Mandatory Enterprise Qualifications	Product Market Access Certifications (One Certificate per Product)
Medical Devices (Including Equipment, Consumables & Reagents)	1. Business License 2. Manufacturing/Distribution License (Class I filing, Class II filing, Class III license)	1. Medical Device Registration Certificate / Filing Document 2. Product technical requirements and instructions for use 3. Software version statement for software-integrated devices
Food Homologous TCM Products (Ordinary Food Grade)	1. Business License 2. Food Production License (SC Certificate) or Food Distribution License	1. Third-party testing report issued within the past year (covering heavy metals, microorganisms) 2. Product implementation standard (national standard or enterprise standard) 3. Medical efficacy claims are strictly prohibited
Food Homologous TCM Products (Health Food Grade)	1. Business License 2. Food Production/Distribution License	1. Health Food Registration Certificate / Filing Document 2. Official health food logo (Blue Hat) and approval number
AI Medical Systems (Medical AI & Software)	1. Business License with software development/sales business scope 2. Software Copyright Registration Certificate	1. Medical Device Registration Certificate required if used for diagnosis/treatment (including AI software components) 2. Software test report and functional manual for auxiliary/administrative use only 3. Data security and privacy compliance statement

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Part 2: Qualification Requirements for Regions outside Chinese Mainland Exhibitors

Applicable to enterprises registered outside Chinese Mainland (including Hong Kong, Macao and Taiwan regions).

Core Principle: For exhibitors unable to meet the qualification standards for Mainland China exhibitors stipulated in Part 1, equivalent legal enterprise and product market access documents issued by regulatory authorities of their home country/region shall be provided.

Product Category	Mandatory Enterprise Qualifications	Product Market Access Certifications (One Certificate per Product)
Pharmaceuticals & Medicinal Materials	1. Enterprise registration certificate (e.g., Business Registration Certificate) 2. Drug Manufacturing/Distribution License issued by local drug regulatory authorities (e.g., US FDA, EU EMA)	1. Marketing Authorization Certificate (e.g., US NDA/ANDA, EU MA) 2. Test report issued by an accredited third-party laboratory 3. Product insert and label
Medical Devices	1. Enterprise registration certificate 2. Medical Device Manufacturing/Distribution License (e.g., US FDA registration, EU CE Certificate)	1. Medical Device Marketing Authorization (e.g., US FDA 510(k)/PMA, EU CE Certificate) 2. Product technical specification manual
Food & Health Food	1. Enterprise registration certificate 2. Food Manufacturing/Distribution License (e.g., US FDA registration, EU health certificate)	1. Third-party ingredient analysis report 2. Product implementation standard 3. Medical efficacy claims are strictly prohibited
AI Medical Systems (Medical AI &	1. Enterprise registration certificate 2. Intellectual property certificates (e.g., software copyright, patent certificate)	1. Local medical device marketing authorization required if used for diagnosis/treatment (e.g., US FDA, EU

Product Category	Mandatory Enterprise Qualifications	Product Market Access Certifications (One Certificate per Product)
Software)		CE-IVDR) 2. Software test report and functional manual for auxiliary/administrative use only 3. Data security and privacy compliance statement (compliant with GDPR and relevant local regulations)

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