CQAF Website User Guide

CQAF, 26 Nov 2017





Welcome to visit: www.cqaf.org

点击下方,快速链接到具体步骤



注意:

• 如果您在公司网络无法登录此网页,请联系您公司IT Service.





成为CQAF会员后,您将可以:

- 免费报名COAF 季度会
- 免费获得在网站向核心组提问题的机会
- 免费阅读COAF发表文章等,参与COAF项目

请按以下步骤完成会员申请



http://www.cqaf.org/index/membership





Projects

Initiatives

- 具体步骤:
 - 点击 网站主页面右上方 Registe 1. 注册页面 http://www.cqaf.org
 - 完成所有信息录入后点击Regist 2.

News & Publications

- 登录信息 •
- 个人信息 •

www.cqaf.org

China QA Forum

Home Page

- 工作单位信息 •
- 专业背景信息 •

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	Phase I unit Others, please specify Graph rest
	If Regulatory Agency
	Inspector Policy maker Others, please specify Single input Your Role(s) / Area(s) of Expertise
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	Dhanghai Beijing Other city, please specify Single input Quality Audit
	>Syears 2-Syears
	Quality Compliance Systems 2-5 years 42 years
r提交	Clinical (GCP)
	>5 years 2-5 years Pharmacovigilance / Drug Safety
	>5years 2-5years <2-5years
	Regulatory Inspection +5 years 2-5 years 42 years
	Education / Training
	□>5 years □ 2.5 years □ <2 years. Manufacturing (GMP)
	Systems 2-5 years 42 years Clinical Lab (GCLP)
	□ >5 years □ 2-5 years □ <2 years
	Laboratory (GLP) \$ 5 years 2-5 years 42 years
	Others, please specify Tingle loar
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	Register
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Personal Details					
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Job Title	Single input	*	Email Address	Single input	*
Organization Name	Single input	*			



- 请 📀 Google Chrome 浏览器进行信息注册或修改
- 请务必在网络好的环境下完成此步骤,否则您可能收不到相应短信验证码;
- 请牢记您的密码,这将用于您的后续会员登录;
- *是必填项
- 请务必确保信息正确,以便于CQAF告知您最新信息





3. 填写工作单位信息▼

Organizat	ion Details			
Organiz	ation type *			
	Pharma Company	CRO	Freelancer	Hospital
	Regulatory Agency	Others, please specify	Single input	
lf Hospi	tal			
	GCP office/CTI	Investigator	EC EC	Laboratory
	Phase I unit	Others, please specify	Single input	
If Regul	latory Agency			
	Inspector	Policy maker	Others, please specify	Single input





用户注册 用户缴费 会员登录

4. 填写专业背景信息►

注	意:		
•		Google Chrome	浏览器进行信息
	注册或	修改	
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请务必保持信息止确,因为这涉及	
到您是否能参与CQAF项目	

Your Role	(s)/Area(s) of Experti	se			
Working	g In: *				
	Shanghai		Beijing	Others, please specify	Single input
Quality	Audit *				
	>5 years		2-5 years	<2 years	
Quality	Compliance *				
	>5 years		2-5 years	<2 years	
Clinical	(GCP) *				
	>5 years		2-5 years	<2 years	
Pharma	acovigilance / Drug Sa	afety	*		
	>5 years		2-5 years	<2 years	
Regulat	tory Inspection *				
	>5 years		2-5 years	<2 years	
Educati	ion Training *				
	>5 years		2-5 years	<2 years	
Manufa	cturing (GMP) *				
	>5 years		2-5 years	<2 years	
Clinical	Lab (GCLP) *				
	>5 years		2-5 years	<2 years	
Laborat	tory (GLP) *				
	>5 years		2-5 years	<2 years	
Others,	please specify Single	e inpu	ıt		
	>5 years		2-5 years	<2 years	



填写专业背景信息▶ 4.

用户注册

勾选同意CQAF Vision 5. 等,后点击Register

I hereby to confirm that I agree with COAF Vision and Mission, CQAF Scope and Principle as in www.cqaf.org.



是必填项

Yo	our Role	(s)/Area(s) of Expertis	se				
	Workin	g In: *					
		Shanghai		Beijing		Others, please specify	Single input
	Quality	Audit *					
		>5 years		2-5 years		<2 years	
	Quality	Compliance *					
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	Clinical	(GCP) *					
		>5 years		2-5 years		<2 years	
	Pharma	acovigilance / Drug Sa	afety	*			
		>5 years		2-5 years		<2 years	
	Regula	tory Inspection *					
		>5 years		2-5 years		<2 years	
	Educat	ion Training *					
		>5 years		2-5 years		<2 years	
	Manufa	cturing (GMP) *					
		>5 years		2-5 years		<2 years	
	Clinical	Lab (GCLP) *					
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	Labora	tory (GLP) *					
		>5 years		2-5 years		<2 years	
	Others,	please specify Single	e inpu	ıt			
		>5 years		2-5 years		<2 years	

会员登录

用户缴费







具体步骤:

- 点击网站主页面右上方 Login 进入会员登录页面 <u>http://www.cqaf.org</u>
- 进入缴费页面,微信支付300元/年, 如需要发票请填写下方发票信息

注意: 缴费成功后 , 您会看到您的会员日的自动倒计时

公司名称(必填): 纳税人识别号(必填): 公司地址: 公司电话: 开户行: 开户账号: 发票名头(必填):如会员费,会议注册费,等

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	China QA Forum	
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PassWo	rd 🔒 Please input a password	
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	Log In	
	Forget PassWord? click here	
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> 用户注册 > 用户缴费 > 会员登录

具体步骤:

- 点击网站主页面右上 方 Login 进入会员登录 页面 <u>http://www.cqaf.org</u>
- 2. 录入手机号和您注册时 输入的密码

	(I)	
	China QA Forum	_
Phone	🔲 Please enter the phone	-
PassWord	Please input a password	
	Record the login information	
	Log In	
	Forget PassWord? click here	

会员登录后您将可以:

- 免费报名CQAF 季度会
- 免费获得在网站向核心组提问题的机会
- 免费阅读COAF发表文章等,参与COAF项目





请首先确保您已经完成上述会员申请流程,并能看到您的会员日倒计时,后续 您将定期收到季度会邀请信邮件

 请您按邀请信的指导,在网站注册报名季度会 <u>http://www.cqaf.org/active?active_type_id=1</u>





如何在网站提问

请首先确保您已经完成上述会员申请流程,并能看到您的会员日倒计时,

- 登录后,点击Q&A页面<u>http://www.cqaf.org/qa</u>
- 点击右上方 I ask 描述您的问题,并点击Submit

注意: COAF核心组将定期审核近期会员通过网站提出的问题,并将回复发布在网站和服务号

☆	i www.cqaf.or	r g /qa				Q
Chi	ina QA Forum				ALL V	Search In Site
	Home Page	News & Publications	Initiatives	Projects	Q&A	🌡 Amy Jiang [logout]
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	Q 众所周知,稚 吗?(2017-7-	瑄是申办方的责任(China GCF -12)	第六章第三十二条)和档	双利(China GCP第六章第三	王十九条),那么,只有申	申办方可以发起稽查
		5.周是否包括临床试验药品,还是	口活用工会业共日 2 /00	17.0.00.)		





请首先确保您已经完成上述会员申请流程,并能看到您的会员日倒计时! 您可以点击下方每个模块和子模块,以获得相应的资源





Home page (主页)

Home Page

您将看到:

- COAF基本介绍
- CQAF会员基本介绍
- CQAF联系信息
- 最新CQAF网站信息,包括
 - 最新发表文章,最新会议, 活动等



Projects

rest Notice

News & Publications

Welcome to CQAF

The China QA Forum (CQAF) was set up in May 2010 and is a non-profit professional organization aiming to promote GxP Quality Standards in the Healthcare Industry. We have set up a platform for quality professionals to share knowledge and best practices in the GxP area, and to communicate within the Forum and externally with regulatory authorities and the healthcare industry with regards to quality.

Initiatives

"Quality Research and Quality Life"

- China Quality Assurance Forum (CQAF)

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Q&A



Home Page News & Publications Initiatives Projects Q&A

News & Publications (新闻和文章)

您将看到:

- CQAF网站新闻
- COAF发表文章:
 - 如就某个专业领域的COAF建 议
- CQAF简讯,
 - 如季度会简讯





Strategy on Management of Self-inspection

During the recent China QA Forum Face-to-Face meeng on 25 May, a workshop was organized to discuss how to manage self-inspecon in China in reflecon of CFDA self-inspecon requirements issued on 22 Jul 2015. Below is the summary of the discussion: Note: Self-inspecon in this arcle refers to sponsor's acvies to idenfy and report issues occurred in the study to CFDA via subming the self-inspecon report for studies that have been submied to CFDA but pending for the agency's approval. ...

2017-06-29 13:41:36



CQAF 2017 Annual Meeting Newsletter

In May 25th , 2017, China Quality Assurance Forum (referred to as CQAF) Annual Meeting was held in Roche Shanghai meeting room. Total 78 members from 40 organizations, including pharmaceutical companies, clinical trial institutions and CROs participated in this conference. This Annual Conference focused on the topics around Inspection Perspective in Electronic Source Data, Self-Assessment Strategy and Approach for CFDA PreApproval Inspection Readiness, and CQAF Projects Updates & Awards....

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您将看到:

CQAF会议信息

CQAF培训

CQAF录播

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Home Page News & Publications Projects Q&A Initiatives Q ALL V Search In Site China QA Forun Initiatives (活动) Home Page **News & Publications** Initiatives Projects Q&A & Jackey Zhang [logout] Meeting Training Living • 如季度会,年会 Meeting 如季度会,或某个专题 more> 2017 Q3 Quarterly Meeting on 11 Sep. 2017 Quarterly Meeting .. 2017-07-11 16:25:53

Training

Doentised from Descriptions.com



GCP Inspection Management Workshop如何迎接GCP现场核查研讨班

2017-07-26 17:24:45

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more>



Projects (项目)

您将看到:

• CQAF目前项目介绍,项 目负责人

Home Page

正在更新中



Projects



CQAF Website

Project Manager: Amy Jiang



US FDA Warning Letter Translation Project

Project Manager: Heidi Liu







Q&A (问答)

您可以看到

- 所有会员的提问和核心组的回答
 - 点击New,优先看到近期的Q&A
 - 点击Host,优先看到点击率最高的Q&A

China QA Forum Home Page News & Publications Initiatives Projects Q&A & Amy Jiang [logout] All New Hot I Ask I Ask I Ask I Ask	☆ ③ www.cqaf	.org/qa				ର୍
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众所周知,稽直是申办方的责任(China GCP第六章第三十二条)和权利(China GCP第六章第三十九条),那么,只有申办方可以发起稽直 吗?(2017-7-12)			第六章第三十二条)和杨	₹利(China GCP第六章第三日	十九条),那么,只有	同申办方可以发起稽查



注意:

请 💽

如何修改网站注册信息

具体步骤:

- 1. 点击 网站主页面右上方 您的姓名进入 个人信息中 心
- 点击头像下Change,可以 2. 修改头像
- 3. 点击Edit More,可以修改 注册信息

Google Chrome 浏览器进行信息注册或修改 请务必确保信息正确,以便于CQAF告知您最新信息

	My Information	
2	Name: Lanjing Zhang E-mail: Zijtxj77@aliyun.com	
	Membership Remaining 233 Days Time: Price: ¥300/year	
	Favorite News & Publications	All
	Favorite Q&A	AI





如何及时获得CQAF最新信息





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	文章	;	活动	≡ 会员	
)		



China QA Forum

中国质量保证论坛(CQAF)是成立于 2010年的非营利组织,致力于推动医 药行业内的GxP质量标准。

如您有任何问题,请联系:<u>chinaqaforum@gmail.com</u>