

Quality Control Interview Questions & Answers

Change controls are of two types i.e., major and minor.

Q11. Corrective action & preventive action?

Corrective action: An action taken to eliminate the cause of the existing deviation, incident or problem in order to prevent its recurrence (occurring again).

Preventive action: An action taken to eliminate the cause of potential deviation, incident or problem in order to prevent its occurrence (an incident or event).

Q12. What is chromatography?

Chromatography is an analytical technique commonly used for separating a mixture of chemical substances into its individual components, so that the individual components can be thoroughly analyzed.

Chromatography is a laboratory technique for the separation of a mixture. The mixture is dissolved in a fluid called the mobile phase, which carries it through a structure holding another material called the stationary phase and the separation is based on differential partitioning between the mobile and stationary phases.

Q13. What is difference between Stationary Phase and Mobile Phase?

The key difference between stationary and mobile phase is that stationary phase does not move with the sample whereas mobile phase moves with the sample. Stationary phase and mobile phase are two important terms in chromatography, which is a technique of separation and identification of the components in a mixture.

Q14. What is column in chromatography?

A Chromatography column is a device used in chromatography for the separation of chemical compounds. A chromatography column contains the stationary phase, allowing the mobile phase to pass through it. The columns are mostly made of borosilicate glass, acrylic glass or stainless steel.

Q15. Which gas is used in Gas Chromatography?

In GC Nitrogen, Helium and Hydrogen are considered to be suitable carrier gases but Helium is most widely used due to safety concerns associated with hydrogen and also the fact that nitrogen is much less efficient.

Q16. What is HPLC in chemistry?

High performance liquid chromatography (HPLC) is a technique in analytical chemistry which is used to separate, identify, and evaluate each component in a mixture.

Q17. What is system suitability?

Before start of analysis of the Chromatographic system like HPLC & GC system suitability has to perform to know that the system is working properly or to know the performance.

System suitability criteria may include such factors as plate count, tailing, retention, and/or resolution and the above factors are most important as they indicate system specificity, precision, and column stability.

Quality Control Pharma Interview Question Answer

Jianjun Gao



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Pharma Interview Questions and Answers Abhishek Chouhan, Pharma Interview Questions and Answers This book contain all the information that will help you crack any Pharmaceutical interview as well as Questions and Answers This book is suitable for Production Quality assurance Quality control Regulatory affairs Research and development product development and Pharmacovigilance etc GRAB YOUR DREAM JOB IN PHARMA: INTERVIEW QUESTIONS & ANSWERS PATHAN AZHER KHAN,2024-05-06 A QUICK INTERVIEW REVISION BOOK Grab Your Dream Job in Pharma Interview Questions Answers for Drug Regulatory Affairs Scientific Research Writing Research and Development Pharma QA QC Production Pharmacovigilance Clinical Research Clinical Data Management Pharmaceutical Marketing List of companies in India QR Codes 100 Pharma Business ideas Overview This comprehensive questionnaire with answers written by industry experts educators and professionals is designed to bridge the gap between HR and candidates by offering common interview questions specific to pharmacovigilance Thus it enhances jobseeker s preparation and confidence The author aims to revolutionize the healthcare and pharmaceutical and research industries by equipping professionals with the knowledge and skills they need to ace their interviews jobs As the pharmaceutical and healthcare industry continues to evolve and expand there is a growing demand for professionals with specialized knowledge and skills in such areas We have gone the extra mile to develop specialized tools and support in this book such as career guidance exclusively for job seekers Our vision is to empower job seekers and professionals like you to take charge of their careers by providing them with the necessary market knowledge Key Features A trusted companion for job seekers with authentic data and references Pharmacovigilance Technical Interview Q A Everything a Candidate Needs in One Place Updated with Current Affairs 100 New Pharma Business Ideas Useful for Pharmacy Medicine and other healthcare sectors competitive exams Learn Technical Skills to get hired

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Frequently Asked Questions Prajjual MAKHAIK,2019-04-03 PHARMACEUTICAL INDUSTRY INTERVIEW FREQUENTLY ASKED QUESTIONS1 What is an SOP A Standard Operating Procedure SOP is a certain type of document that describes in a step by step outline form how to perform a particular task or operation Everyone in a company must follow the same procedures to assure that tasks are performed consistently and correctly Most companies have a wide variety of SOPs that describe how to do different tasks In many companies technicians and operators are trained in how to follow individual SOPs and their training record specifies which SOPs they are trained on and are authorized to use 2 What is 21 CFR part 11 Title

21 CFR Part 11 of the Code of Federal Regulations deals with the Food and Drug Administration FDA guidelines on electronic records and electronic signatures in the United States Part 11 as it is commonly called defines the criteria under which electronic records and electronic signatures are considered to be trustworthy reliable and equivalent to paper records

3 What are user Requirements User Requirements Specification describes what users require from the System User Requirement specifications are written early in the validation process typically before the system is created It is written by the System Owner and End Users with input from Quality Assurance Requirements outlined in the URS are usually tested in the Performance Qualification User Requirements Specifications are not intended to be a technical document readers with only a general knowledge of the system should be able to understand the requirements outlined in the URS

4 What is a validation plan Validation Plans define the scope and goals of a validation project Validation plans are written before a validation project and are specific to a single validation project Validation Plans can include Deliverables Documents to be generated during the validation process Resources Departments Personnel to participate in the validation project Time Line for completing the validation project

The Chemical Engineer ,2006 *Pharmacy Management* Shane P. Desselle,David P. Zgarrick,2005 Editors Desselle and Zgarrick have brought together 33 contributed chapters in their endeavor to prepare pharmacy students for the realities of managing a practice After a section on why it s important to study management in pharmacy school coverage includes the various ins and outs of managing oneself understanding stress operations people

Oil, Paint and Drug Reporter and New York Druggists' Price Current ,1919 Vols include the proceedings some summarized some official stenographic reports of the National Wholesale Druggists Association called 18 1882 Western Wholesale Druggists Association and of other similar organizations

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