

Webinar For Overview Program on Fabrication and Inspection Requirements for Bioprocess Equipment as per ASME BPE (Weekend Batch)

Date : 18th October 2020, Time: 10:00 AM to 1:00 PM



Course Features

Give overview on Bio-process equipment requirements for material, welding, dimensions, surface finish, qualification, examination, inspection and testing to achieve required sterility, cleanability as per ASME BPE Code.

Course Objectives

This course help delegate to understand requirements of Bio Process Equipment of Code and common industry practices for their acceptance criteria, qualifications, cleanability and sterilizability. It is very important to have knowledge on the methods to assure quality and code compliance to maintain hygienic.

Who should attend ?

- ♣ QA / QC, Production, Design, Engineers and Manager
- ♣ Representatives of Inspection agencies
- ♣ Representatives of Pressure Vessels Manufacturers who supply to Bio-Pharma Industry
- ♣ Representatives of Pharmaceutical Industry
- ♣ Representative of PMC in the field of manufacturing of biopharmaceuticals and sanitary equipment's



Course Contents

- ♣ Introduction and Contents of ASME BPE
- ♣ General Requirements
- ♣ Inspector Qualification requirement levels
- ♣ Sterility & Cleanability requirements
- ♣ Material selection and their Dimensional tolerance, Surface finish and surface defects
- ♣ Welding requirements - Material Joining, Joining Process, Qualification
- ♣ Weld inspection - Weld defects, Weld finish, Weld color, etc.
- ♣ Use of Inspection tools like Boroscope, Weld gauge etc.

Issue of Certificate

Certificate of successful attendance shall be issued to all the delegates who attend entire duration of the course.

Duration : 180 minutes (3 Hours)



Registration Fee : Rs. 1500 + 18 % GST
Per Participant

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