QUALITY MANAGEMENT ENGINEER

A. Company Introduction:

B. Braun Vietnam, a subsidiary of B. Braun Melsungen AG Group - one of the world's leading manufacturers of medical devices and pharmaceutical products and services - now operates one of the largest medical complex in Vietnam and has become a prestigious trademark in Vietnam healthcare market. With more than 1,400 employees nationwide, we develop high quality “made in Vietnam” products, especially our infusion solutions, dialysis solutions and plastic medical instruments have widely been used and appreciated in international and domestic markets. Every service provided by B. Braun Vietnam incorporates the entirety of our knowledge and skills, the company's deep understanding of users' needs and extensive expertise. Our mission is to PROTECT and IMPROVE the health of people around the world. Let’s become a part of the B. Braun family and share your expertise.
www.bbraun.com.vn

B. Job function:

Be responsible to plan, conduct validation and follow-up re-validation in accordance with B. Braun's Validation SOPs, such as Design Qualification, Installation Qualification, Operational Qualification, Performance Qualification and Releasing Final Validation Report for all production processes in production facility.

This includes as followings:

- Create, monitor and follow the Master Validation Plan for production processes/equipment used for producing and quality controlling at the plants of BBV VN
- Co-ordinate with the QC, QM and Production to make a risk assessment then define and perform needed activities related to Qualification and Validation of equipment and processes according to Validation Master Plan or department's requests, as well as WHO GMP/ISO 9001/ ISO 13485 requirements.
- Monitor and co-ordinate with related persons to undertake quality engineering projects with guidance from the Superior and the Management in co-operation with

Production and Quality Control department. To ensure that the validation and calibration activities are undertaken objectively based on clear technical aspects.

- Assist other QM function for performing process audit arise due to corrective actions implementation.
- Plan and execute qualifications and validation computerized system validation, data integrity and other as request.
- Implement Corrective Action and Preventive Action, Risk assessment, Deviation Control.
- Prepare validation document to support drug registration activities (if any)
- Implement other quality issues whenever required.
- Ensure that the requirements of ISO, GMP, GSP. GLP as well as legal requirements are well implemented and maintained.
- Ensures that the audit programs are developed and executed effectively.
- Handles complaints system from clients and product regulatory affair processes.

C. Job requirements:

- Graduated from universities with technical engineering background
- Good command of English both in communicating & writing
- Willing to learn and do

D. Strive for more…

Motivated and competent employees are our most important assets. We are committed to invest in our people, through continuous career development, on-the-job training and professional qualifications. The opportunities are endless at B. Braun, as we are in a continuous growth phase. You can really drive your own career here and are trusted to do a fantastic job.
Whatever role you are in, you are in some way, shape or form contributing to protecting and improving the health of people around the world. Our team love that they genuinely make a difference!"

E. How to apply

Please send your updated CV to: recruitment.vn@bbraun.com

F. For more information, please visit:

www.bbraun.com.vn

www.facebook.com/bbraunvn

www.linkedin.com/in/bbraunvncareer/

Or contact our Hotline: +84 24 3357 1616 (Ext. 1129)

“Candidates are always welcome at B.Braun Vietnam. We are an equal opportunity employer and commit to ensure fairness and transparency during selection process as well as in your development later on with us”.