Compliance Officer (Newtown, PA) – Review and analyze Manufacturing and Packaging Batch Records, Process Validation and Stability protocols and reports in compliance with 21 CFRs FDA guidelines; Review and inspect pharmaceutical production materials to ensure compliance with FDA, and cGMP standards and regulations; Maintain and control materials and stocks as per user department requirements in SAP and FIFO; Compile SOPs; Coordinate and facilitate FDA inspections; Verify and document consumption statements for accuracy.

Must have a Bachelor's Degree in Regulatory Affairs, Management, or Business Administration, plus 3 years experience in job offered or as Manager in pharmaceutical industry. Require skills and working knowledge of SOP, FDA regulatory requirements, cGMP, SAP, FIFO, FDA Inspections.

Job location: Newtown, PA. Submit résumé referencing job code VEA002 to HR, KVK-Tech, Inc., 110 Terry Drive, Suite 200, Newtown, PA 18940.