Laporan Praktek Kerja Profesi Apoteker PT. Taisho Pharmaceutical Indonesia Tbk. Jalan Raya Bogor Km 38. Periode17 Juni - 30 Agustus 2013 = Report of Pharmacist Internship Program at PT. Taisho Pharmaceutical Indonesia Tbk. Jalan Raya Bogor Km 38. 17 Juni - 30 Agustus 2013 Period

Bhata Bellinda, author

Abstrak

Pharmaceutical Industry is the place to meet the needs of public health by increasing the quantity and quality of the drug to be in production All drugs are in production and will be circulated to be guaranteed safety efficacy and quality To achieve this we need a guideline that covers all aspects of production and quality control in order to ensure the drug is made consistently meeting the requirements set out and in accordance with their intended use namely the Good Manufacturing Practice GMP All GMP pharmaceutical industry shall apply in all aspects and activities of a series of drug manufacturing Work Practice Pharmacist PKPA conducted during the period 17 June to 30 August 2013 in the PT Taisho Pharmaceutical Indonesia Tbk to provide supplies for prospective pharmacists to apply the knowledge they have learned during the course in a practical and direct the quality control of drugs in the Pharmaceutical Industry The PKPA activities aimed to compare the application of the provisions of the implementation of GMP in Pharmaceutical Industry particularly in PT Taisho Pharmaceutical Indonesia Tbk dan understand the duties and responsibilities of pharmacists in the Pharmaceutical Industry is mainly in charge of production quality assurance and quality control