MARK KESSLER, PH.D.

Over 20 years professional experience in the pharmaceutical industry. Have an expertise in biopharmaceutical drug development and manufacturing and sterile drug product manufacturing, as well as extensive experience in manufacturing facility design and commissioning, QC laboratory design, oral solid dosage form manufacturing and medical device manufacturing. In Israel, worked at Biotechnology General and Kamada and am currently at Teva Pharmaceuticals, serving increasingly senior roles in the areas of quality assurance, quality control and regulatory affairs. Most recent experience at Teva includes responsibility for quality operations for Teva plants and CMOs located in Israel, Europe, U.S. and Mexico that are producing biopharmaceutical APIs, LVPs, medical devices and sterile drug products.