



TEVA'S TOP PRIORITY IS ENSURING THE SAFETY AND EFFECTIVENESS OF OUR MEDICINES

As a leading pharmaceutical company, Teva's top priority is ensuring the safety and effectiveness of our medicines. Our commitment starts in the research laboratory and continues throughout the manufacturing process and supply chain where numerous safeguards are implemented to ensure the highest quality and reliability.

Every pharmaceutical company – whether it produces innovative brand-name, generic or biologic drugs – must adhere to the same manufacturing practices and strict quality control procedures necessary to manage a complex, global network of suppliers and distributors. In addition to complying with all government regulations, Teva has taken additional steps to further strengthen our supply chain and promote safety, including voluntary supplier audits.

Teva's manufacturing process specifically includes:

- **FDA Approval** – The FDA sets the highest standards for the production and distribution of both brand and generic medications. These regulations govern the sourcing, chemistry and manufacturing controls for each product. Teva works closely with the FDA and uses current Good Manufacturing Practices (cGMP).
- **Proven Partners** – In addition to relying on FDA inspection reports and cGMP certifications, Teva also works with trusted suppliers and conducts on-site facility audits to verify these vendors meet our exacting standards. We ensure that each manufacturing site has established proper controls for providing consistently high-quality products.
- **Rigorous Testing of Raw Materials** – We physically inspect each shipment and perform analytic tests using FDA-approved methods to verify compliance with pre-established acceptance criteria. We also retain a sample for future investigation, if necessary.
- **Ongoing Monitoring & Reporting** – Once our drugs are on pharmacy shelves, Teva tracks patient reports and monitors for trends. Our quality reporting system is designed to identify any potential problems and evaluate if further action is warranted. We also submit annual product reports to FDA.