2013 Overall Compliance by Domain	Compliance
Filter Integrity Test	38.4%
Gloved Fingertip Sampling	51.4%
Compounding Facility Management: Airflows and Pressure Differential Monitoring	59.0%
Depyrogenation by Dry Heat	59.7%
Bacterial Endotoxin Testing	60.8%
Quality Management: General Viable Air and Surface Sampling Considerations	61.1%
Quality Management: Surface Sampling - A personnel metric	61.3%
Quality Management: Incubation	65.4%
Sterility Testing	66.1%
Quality Management: General	67.5%
Quality Management: Environmental Sampling Program	67.7%
Quality Management: Viable Air Sampling - A facility metric	68.4%
Low Risk Level CSPs with 12 Hour or Less BUD	69.9%
Hazardous Drug Compounding	73.2%
Compounding Facility Management: Cleaning and Disinfecting	73.3%
CSPs for Immediate Use	75.5%
Sterilization by Filtration	75.8%
General Facility Design	76.2%
Compounding Facility Management: Equipment Calibration	76.8%
Initial and Ongoing Training and Competency Measurement	77.8%
Sterilization by Dry Heat	78.3%
Hand Washing and Garbing	78.5%
Primary/Secondary Engineering Controls	82.2%
Personnel Media-Fill Challenge Testing	85.2%
Compounding Facility Management: Temperature and Humidity Monitoring	85.4%
Aseptic Technique	86.2%
Allergen Extracts as CSPs	86.4%
Quality Management: Patient/Caregiver Training	86.7%
Steam Sterilization	86.7%
Sterilization Methods	88.5%
Radiopharmaceuticals as CSPs	90.0%
Final Release Checks	90.3%
Beyond Use Dating	90.6%
Single and Multiple-Dose Vials	91.1%
Quality Management: Non-Viable Particle Testing	91.6%
Inventory Storage and Handling/Delivery of CSPs	94.2%
Grand Total	**77.2%**