

Our Validation Services

Industrial Vision Systems Ltd (IVS) provide validation services for their full range of vision system products and integration services to the pharmaceutical, medical devices, wound care, biotechnology and consumer healthcare manufacturing customers. These industries increasingly rely on automation so the quality of computer software is assuming greater significance in their operations. All vision systems in these industries must be validated in order to achieve Good Manufacturing Practice (GMP) and Good Laboratory Practice (GLP) - collectively referred to as GxP.

Our validation packages follow the standard International Society of Pharmaceutical Engineering (ISPE) and Good Automated Manufacturing Practices (GAMP5) guidelines; as well as assessment of validation requirements for 21 CFR Part 11. Our packages guarantee you peace of mind, our years of experience in installing and validating vision systems on pharmaceutical production lines and cells gives us in-depth knowledge of equipment functionality.

IVS validation engineers are trained to understand the specific needs of the industry and identify the vision solutions required for each customer. Each validation project is unique and requires a planned and targeted approach.



We assess the individual requirements of a project through a process of risk assessment. All necessary project and validation documentation is developed using our standard templates and formats, or your own as required.

By using original IVS validation services, manufacturers can ensure that their vision systems are tested and proven to the highest standards in a timely and cost effective manner, achieving fully validated and compliant vision solutions.







"Establishing documented evidence which provides a high degree of assurance that a specific process will consistently produce a product meeting its pre-determined specifications and quality attributes."

FDA Guidelines on General Practices of Process Validation.

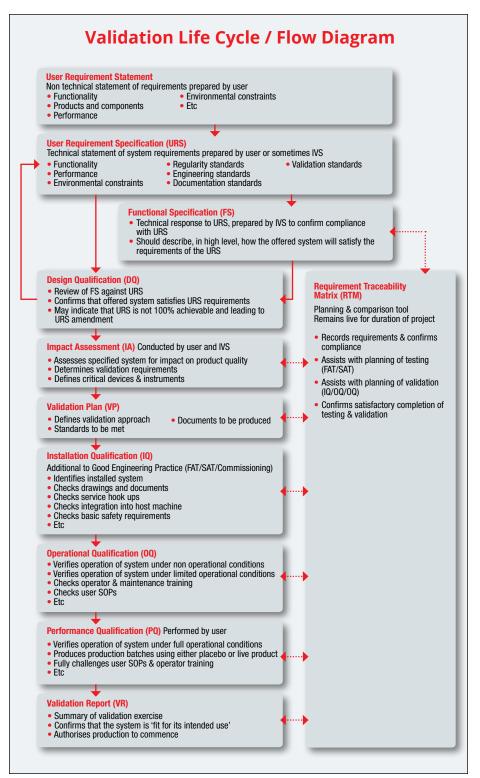
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Industrial Vision Systems Ltd have a defined lifecycle policy regarding our vision system validation as shown below. We have a well established Quality Management System (QMS) and have undergone formal audit supplier reviews from our blue-chip pharmaceutical customers.









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