
Gulf Pharmaceuticals, Julphar, the largest generic drug manufacturer in the UAE, chooses QAvalid™ to facilitate FDA compliance.

Clarmon Corporation announced today the completion of an agreement with Gulf Pharmaceutical Industries (Julphar). Clarmon will deliver QAvalid, an innovative and fully integrated compliance management solution, to help Julphar manage its FDA / GMP compliance processes in an easy and cost-effective manner.

QAvalid is fully integrated with Microsoft® Word and allows its users to work in a familiar way while providing all the benefits of a centrally controlled application. Linking related documents to specific FDA / GMP regulations and controlling content from creation and approval through to ongoing usage guarantees an automatic audit trail to prove full regulatory compliance.

QAvalid was initially developed for the pharmaceutical, biotechnology and medical device industries. However it has proven applicable to an array of other industries including Nuclear Decommissioning, Semiconductor Manufacturing, Civil Construction, Food and Beverages, Veterinarian Products, Cosmetics, Automotive, Aircraft Manufacturing and Telecommunications.

About Gulf Pharmaceutical Industries (Julphar)

Gulf Pharmaceutical Industries (Julphar) is the largest manufacturer for generic drugs in the UAE (United Arab Emirates). Julphar has seven manufacturing plants, five located in the UAE as well as one in Germany and one in Costa Rica.

Julphar has marketing authorizations for more than 477 pharmaceutical products and it is present on more than 54 outside markets worldwide.

Currently Julphar has two additional facilities under construction and will be using QAvalid during the construction and validation phase of these facilities. The facilities is planed to be Eurolex, GMP and FDA compliant, and will be among the region's most advanced solid dose and sterile Biotechnological production facilities.

For more information about Gulf Pharmaceutical Industries, please visit <http://www.julphar.com>.

About Clarmon Corporation

Clarmon Corporation provides integrated software applications for the management of structured content within a number of target industries. Most of Clarmon's clients to date have been in the Pharmaceutical, Biotechnology and Medical Device Industries. Through its QAvalid application Clarmon provides a powerful tool to allow companies to achieve regulatory compliance faster and easier. By removing tedious manual tasks for Quality Assurance,QAvalid™ drives costs down, and diminishes the risk of non-compliance. Clarmon Corporation is headquartered in London, with a development and consultancy team in Bucharest Romania and sales staff in Dubai and Singapore.

For more information about Clarmon Corporation, please visit <http://www.clarmon.com>.