

## Analytical Scientist

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Company: Tekwissen UK

Location: Kent

Category: life-physical-and-social-science

### Overview:

TekWissen Group is a workforce management provider throughout the UK Europe and many other countries in the world. The below client is an American multinational pharmaceutical and biotechnology corporation.

**Job Title: Analytical Scientist**

**Location: kent sandwich CT139ND**

**Job Type: 12 Months**

### Job Description

The Materials Characterisation Team based within the Analytical Technology Function of clients Global Supply Technology and Engineering (GT&E) department are looking for a scientist to support the quantitation of small molecule impurities using liquid chromatography with mass spectrometry. This is to enable technical support for complex analytical and process investigations along with the assessment and support of on market products in the areas of nonconforming materials line extensions and second sourcing activities.

### SKILLS NEEDED:

Technical understanding of the development and validation of LCMS methods.

Appreciation of challenges associated with the analysis of trace impurities in pharmaceutical materials.

Strong working knowledge of quantitative chromatographic analysis and the

necessary laboratory skills required to achieve accurate precise and reliable data first time.

Ability to work independently and to document analytical results and conclusions through presentations and technical reports.

Oral and written communication skills with emphasis on clarity of communication.

Ability to follow scientific protocols / methods and departmental procedures with excellent attention to detail.

Highly motivated self directed and proactive.

Creative entrepreneurial attitude with appetite to work in a fast paced environment.

Ability to execute key priorities to timelines whilst effectively balancing multiple priorities.

Interpersonal skills and ability to work with personnel at all levels.

## **ROLE RESPONSIBILITIES:**

Responsibilities will include but are not limited to the following:

Supporting the LCM testing of drug product lots for the quantitation of small molecule impurities.

Designing studies to evaluate method suitability (linearity accuracy precision etc.).

Developing appropriate sample preparation methods in a range of different matrices.

Executing accelerated drug product stability studies including trending of impurities data.

Performing experiments in accordance with good laboratory practices (GLP) established business processes and protocols and applicable SOPs.

Identifying and investigating atypical analytical results.

Troubleshooting analytical instrumentation.

Documenting laboratory work in a detailed clear and timely manner in compliance with

GL Requirements.

Selecting appropriate presentation techniques for analytical datasets in order to deliver concise headline results to stakeholders in a timely manner.

### **QUALIFICATIONS:**

Bachelors Degree in relevant scientific field.

Practical analytical laboratory experience in the field of quantitative chromatographic analysis.

Desirable

Experience with LCMS and associated quantitative processing software.

Experience in the development / validation of LCMS methods for trace analysis.

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