

Gentium S.p.A is a Biopharmaceutical Company whose mission is to discover, develop and provide to care-givers new therapies for diseases that currently have few or no treatment options, with the aim to improve the lives of patients with rare diseases and with high unmet medical needs, located in Villa Guardia near Como.

**DRUG SAFETY ASSOCIATE
(Rif. MI 2)**

- Assist in processing adverse event reports for clinical studies;
- Prepare expedited report, cover letters, handle follow up request;
- Maintain data base;
- Evaluate of adverse event reports to confirm regulatory reporting status; process adverse event including preparation of clinical narrative summaries, and coding adverse events, indication for use and concomitant medical conditions;

Requirements:

Degree in life Sciences

2/3 years experience in Pharmacovigilance

Good written and spoken English

Excellent computer skills (MS office)

Please send your curriculum to: humanresource@gentium.it