We are a leading world supplier of integrated processing solutions for customers in the area of pharmaceuticals, food processing, feeds and fine chemicals. Our products and services stand for reliability and quality worldwide. About 3,000 employees work for Glatt at more than 15 companies and offices all over the world.

For our site in Ramsey, NJ (USA) we have an immediate opening for an:

**TECHNICAL SPECIALIST**

**Job description:**

- Create, manage and oversee the generation of technical documents in support of Pharmaceutical Development, Operations, Technical Operations, Process Validation, Cleaning Validation and Equipment Qualification activities for the Pharmaceutical Services Division. Should possess a pharmaceutical manufacturing/development and cGMP background in solid dosage form environment with preferred background in fluid bed processing.

**Preferred skills:**

- Minimum BA/BS or certificate in a technology related field and at least 5 years' development or manufacturing experience in a cGMP solid dosage manufacturing environment required.

- Must have the ability to understand the fundamental processing stages required to independently create the protocols, change controls, investigations, deviations and final reports required to conduct equipment qualification, process validation and cleaning validation activities.

- Strong computer skills are a necessity, with an ability to accurately convey information in both spoken and written form. Strong command of the English language is essential.

- Must meet deadlines and be detail oriented. Solid organizational skills are a necessity, with a demonstrated ability to organize and track several projects simultaneously. The ability to follow up with multiple departments is required.

Qualified candidates may apply to: Glatt Air Techniques Inc., Human Resources, 20 Spear Road, Ramsey, NJ 07446, USA

or via E-Mail to HR@glatt.com, www.glatt.com