



## Associate Scientist/Principal Scientist Process Engineering P...

武田薬品工業株式会社での募集です。プラントエンジニア（電気）のご経験のある方...

### 募集職種

#### 人材紹介会社

株式会社ジェイ エイ シー リクルートメント

#### 採用企業名

武田薬品工業株式会社

#### 求人ID

1488722

#### 業種

医薬品

#### 雇用形態

正社員

#### 勤務地

神奈川県

#### 給与

550万円～900万円

#### 勤務時間

09:00～17:45

#### 休日・休暇

【有給休暇】有給休暇は入社後4ヶ月目から付与されます 初年度 12日 4か月目から 【休日】完全週休二日制 土日 祝日 年末...

#### 更新日

2024年08月01日 15:23

### 応募必要条件

#### キャリアレベル

中途経験者レベル

#### 英語レベル

ビジネス会話レベル

#### 日本語レベル

ネイティブ

#### 最終学歴

大学卒：学士号

#### 現在のビザ

日本での就労許可が必要です

### 募集要項

【求人No NJB2186975】

Job Description

Synthetic Molecule Process Development (SMPD) is responsible for the development of robust and cost effective processes for the manufacture of new small molecule pharmaceuticals along with methods for achieving and controlling high standards of purity and quality.

How you will contribute:

The successful candidate will be responsible for all aspects of reaction particle engineering including the development of scale down models for the study of unit operations as well as technical transfer to external contract manufacturing organizations. He/ she will have deep experience in using process analytical technologies (PAT) in combination with mathematical models (both statistical first principle) to enhance process understanding to effectively develop/ optimize/ scale up and troubleshoot processes. The Senior Engineer will have experience building scale down equipment and developing innovative advanced process control strategies for both batch and continuous processes.

He/ she will be recognized as a technical resource/expert within SMPD and across Pharmaceutical Sciences and utilize his/ her technical expertise to contribute across multiple projects and drive technical/scientific strategy. The Senior Engineer will be responsible for benchmarking current trends in research development and manufacturing technologies developing initiating and/or participating at a high level in projects that involve extraordinary well considered risks along with scientific/technical challenges as well as directing and managing outsourcing across a product platform as appropriate.

## スキル・資格

※英文レジュメが必要となります※

### Accountabilities:

Develops project or significant technical strategy and leverages technical skill (s) as a resource/expert within the department  
 Contributes significantly and independently to project work which may include multiple projects within functional area.  
 Plans and implements resolutions to technical problems/issues  
 Independently designs and executes experiments and reports results  
 Recommends justifies and implements technologies and innovations.  
 Owns a discipline/technical skill in its entirety and continues to develop expertise in other key technical skills.  
 Influences and supports initiatives related to driving scientific and technical improvement within function and potentially cross functionally.  
 Reviews interprets and communicates data cross functionally within pharmaceutical sciences and project teams  
 Conducts analysis of technical and conceptual risk and trends  
 Identifies process trends and defines/champions process strategy or use of novel technologies  
 Recognized as a technical expert and resource within function  
 Significant technical responsibility for a project area/technical program within the department and potentially across Pharmaceutical sciences  
 Identifies topics for initiatives and leads local/global initiatives on behalf of senior staff.  
 Defines appropriately complex/novel approaches and methodologies to solving outstanding technical challenges  
 Coordinates and leads technology transfer to internal or external manufacturing sites  
 Responsible for authoring relevant sections of regulatory documents validation plans reports and peer reviewed manuscripts.  
 Proactively identifies vendors and builds relationships to gain access to technologies as needed to deliver on pipeline goals.  
 Manages key vendor relationships across multiple projects as appropriate and proactively affects resolution of issues arising at vendors.  
 Represents Takeda and is an active member on pre competitive collaborations with academic and industrial partners.

### Education and Experience:

A Ph.D. degree with 3+ years of academic post doctoral or pharmaceutical industry experience; an MS degree with 9+ years of pharmaceutical industry experience; or a BS degree with 11+ years of pharmaceutical industry experience. Degrees in chemical engineering required.  
 Experience in the use of mathematical both statistical and first principle models as well as advanced process control systems preferred  
 Experience in building reaction kinetic models as well as process models preferred.  
 Experience in building laboratory and pilot plant equipment a plus  
 Experience in crystallization process development and scale up with an emphasis on form purity and particle size control a plus  
 Experience in the use and scale up of milling technologies (both dry and wet) for particle size control a plus  
 Experience in the use of process analytical technologies (FT IR NIR FBRM UV vis etc.) required  
 Experience in building chemometric models preferred  
 Experience in developing continuous processes a plus  
 Sound knowledge of current Good Manufacturing Practices (cGMP) preferred  
 Experience working in a pilot plant a plus  
 Previous experience with the use of contract facilities and managing technical transfers a plus  
 Experience in working in a multi disciplinary team environment  
 Proven scientific track record through presentations at scientific conferences and publication of peer reviewed manuscripts

### Knowledge and Skills:

Analytical and Problem Solving Skills Able to troubleshoot critical issues or problems using appropriate information and determine causes and possible solutions  
 Teamwork Ability to work well on global cross functional teams.  
 Communication Skills Able to express one's self clearly and concisely within team; documents issues and/or concerns concisely with colleagues; adjusts communication style as appropriate for the audience; timely and effectively communicates with senior management; technical writing skills to support authorship and approval of internal technical documents  
 Organization · Exercises good time management and prioritization skills to balance multiple project and departmental objectives  
 Technical Subject matter expertise in a specific scientific area or areas.  
 Knowledge Sharing Ability to capture knowledge within the organization; improves solutions processes and deliverables through use of information; improves information capital by contributing experience theories deliverables and models for others to use

Resource Management Project management skills; ability to manage one's time within individual departmental  
External Involvement · Demonstrated involvement in professional community evidenced by presentation of scientific posters or lectures at professional conferences or events. Interacts with external vendors for projects  
Leadership Skills · Develops and uses knowledge and interpersonal skills to appropriately influence and guide others towards the accomplishment of department/function goals and objectives.

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## 会社説明

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