



## International Regulatory Affairs Project Manager

Location: Allegan, Michigan

### About Perrigo

Perrigo Company is a leading global healthcare supplier that develops, manufactures and distributes over-the-counter (OTC) and prescription pharmaceuticals (Rx), nutritional products, active pharmaceutical ingredients (API) and consumer products. Perrigo Company is the world's largest manufacturer of OTC pharmaceutical products for the store brand market with over \$2 billion in annual sales. The Company's primary markets and locations of manufacturing facilities are the United States, Israel, Mexico and the United Kingdom.

Perrigo produces more than 44 billion tablets annually, is the largest producer of liquid antacid and the second largest producer of acetaminophen and ibuprofen. Perrigo has more OTC ANDA approvals (61) than any other company (including branded companies).

### Job Description

At Perrigo, your job will mean more than any other job you've ever had. You will join a team that is as driven and excited as you are about making healthcare products more affordable around the globe. Our products improve our consumer's quality of life – let Perrigo improve the quality of your career.

As an International Regulatory Affairs Project Manager, you will be responsible for:

- Managing product registrations in countries outside of the US
- Providing registration support to Perrigo global affiliates
- Researching all regulatory alternatives and developing recommendations in response to new international marketing opportunities
- Developing labeling for registrations owned by Perrigo
- Responding promptly to internal and external customer inquiries

### Qualifications

Bachelor's Degree in a scientific or technical field is required. Experience with managing technical documentation is required. A minimum of three years experience in the pharmaceutical industry with demonstrated progression in scientific or regulatory affairs role is required. Three plus years of experience in international registrations or other Regulatory Affairs experience highly preferred. Knowledge of FDA drug product submissions (NDA/ANDA) and the Canadian regulatory system (including drugs and natural health products) highly preferred. Prior leadership experience is preferred. Excellent technical writing, verbal communication, and strong interpersonal skills required. Ability to effectively interact with government agencies and company personnel at all levels required.

**Qualified applicants are requested to apply online at [www.perrigo.jobs](http://www.perrigo.jobs), under 'US Careers.'**

Perrigo Company  
515 Eastern Avenue  
Allegan, MI 49010  
(269) 673-8451  
[www.perrigo.com](http://www.perrigo.com)

Perrigo Company is an Equal Opportunity Employer.

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