

Director of Regulatory Affairs

Case Study #3448

Search Metrics:

11 Days until hired candidate was presented 4 Total candidates presented 53 Days until search completed

Client:

A 180 person biopharmaceutical company expecting 5 late phase drug candidates by 2015, engaging in the development and commercialization of therapeutic products based on RNA interference (RNAi) in the United States.

Objectives:

- I. Leading regulatory strategy to meet a milestone of being eligible for a fast track and orphan designation.
- II. Drive 3+ IND, NDA/sNDA submissions, develop future milestones and strategies.
- III. Set strategic direction for regulatory affairs and regulatory operations

Prestige Solution:

On site meeting to conduct full needs analysis
Implement the Prestige Performance Search methodology
Develop unique Performance Profile describing objectives of role
Selectively introduce profile directly to target audience