



Regulatory & Clinical Practice

Validant has an in-depth knowledge of the Medical Device, Pharmaceutical and Biologics sector and an extensive network of senior level professionals with the niche skill sets that are in high demand in the industry. We combine this market knowledge with our proven technical search capabilities to deliver a comprehensive, high quality search solution for our clients.



Key Focus Areas

Validant helps companies comply with the vast and ever-changing regulatory standards. We help clients efficiently move products past clinical trials, through approvals and into the marketplace.

Leadership

Validant has a strong history of recruitment and consulting in Clinical and Regulatory Affairs. We offer a unique combination of FDA, biologics, drug and medical device industry experience. We have successfully assisted our clients with US and International submissions, critical post-market surveillance initiatives, placement of Medical Directors and Executive Leadership.

Expertise

Our network and capabilities includes recruitment, consulting and executive search. We have a strong track record across multiple therapeutic areas including Oncology, Cardiovascular, Metabolic Disorders, Immunology, Ophthalmology, Infectious Disease, Dermatology and Neurology.

Partnering

Whether your organization is in need of occasional support from an experienced industry professional or is looking to ramp up for key projects, Validant is ready to deliver the expert talent you need. We invite you to learn more about our solutions and how we can become your trusted resource provider.

Key Areas of Focus:

- Clinical Operations
- Regulatory Affairs
- Clinical Drug Development
- Medical Affairs
- Drug Safety
- Clinical Data Management
- Biostatistics/Statistical Programming

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