Unlocking the Power of Integrated Drug Development

Connecting the complex building blocks of development to accelerate and rescue critical drug development projects

Solutia Pharmaceutical Advisors works closely with senior managers in pharmaceutical and biotech companies to implement programs that address typical time-to-market issues, as well as the inherent risks of drug development. We integrate the work of industry experts to deliver real benefits to your business.

**R&D organizations are facing increasing pressures to deliver faster.**
Our approach to risk management in drug development provides our clients with the depth and breadth of insight to integrate complex activities. We help clients make better decisions using the collective expertise of a team of experts who have been responsible for bringing numerous drugs to market.

**Development operations must mitigate risks that steal time and resources.**
Solutia’s network of advisors has a proven record of understanding how to structure and make the critical decisions that enable speed in delivering new drugs to market. Our solutions tap into knowledge from all parts of the organization to support effective decision-making, enabling clients to meet their objectives for:

- Reducing drug development risk, time and expense
- Managing attrition in getting new drugs to market
- Ensuring timely achievement of critical launch or licensing milestones

Development teams often need additional critical skills or more hours in the day to make key decisions. We augment, coach and support client development teams to broaden their understanding and improve their project’s likelihood of success or help them make the right stop decisions, earlier.

Solutia Pharmaceutical Advisors can deliver real benefit to critical development programs through focused and integrated planning and execution support, reducing time to market, costs, and managing attrition.
Solutia Pharmaceutical Advisors’ proven experts bring you their years of experience with hands-on R&D investigation. They have worked on and lead development teams, managed departments and led global functions for many of the world’s leading pharmaceutical and biotechnology organizations. Our advisors span all areas of expertise required for successful drug development including:

- CMC , Chemical and Biopharmaceutical Process Development, Formulation, Analytical Development, Regulatory
- Pre-Clinical and Clinical Strategy & Planning, Pharmacology and Drug Safety
- Engineering and Site Strategy and Implementation
- Quality Systems
- Chemical and Bio-Manufacturing and Operations, Supply Chain & Launch Readiness

An integrated view across these disciplines can significantly shorten the development cycle, and save you significant avoidable costs and time by optimizing planning and mitigating risks. Things do go wrong in drug development and our proven experiences and perspectives can help you anticipate and avoid these pitfalls. When things appear to have “gone wrong” on a project, the application of this combined technical and managerial experience, can identify what’s really going on and rescue a critically important drug project.

Solutia Pharmaceutical Advisors structures our projects around the best individual expert or teams of advisors, bringing you the appropriate level of the right resources for your organization at the right time. Our teams help you quickly sort out your priorities, establish a realistic timeline and optimize your program on the basis of time, cost and scope. Our unique perspective comes from years of collective experience applied to your individual needs.

**Case Study – Managing risks in a jeopardized development program**

Our client acquired the rights to a drug candidate that had blockbuster potential. However, due to a specific finding in the clinical program, the development of this candidate was stalled, requiring a “new perspective” and strategy for resurrecting the program and getting it to market as quickly and inexpensively as possible. Solutia Pharmaceutical Advisors - through the application of a focused team of technical experts representing clinical, pre-clinical, drug metabolism, toxicology, and formulation expertise - were able to rapidly define the critical issues, develop an optimized strategy, and define an integrated plan that mitigated the technical constraints and managed the risk in the program.