Linda DeGraw

Director, Contracts Specialist
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Linda has dedicated more than twenty-five years to drafting, negotiating and managing specialized contracts in the biopharmaceutical industry.

Her vast experience includes agreements for preclinical and clinical studies, research collaborations, master services and consulting, and other agreements relating to the development, manufacturing and testing of drug compounds and medical devices, and advice regarding contract management for clinical studies. Linda contributes a valuable expertise to Faber, by focusing much of her work on agreements for large multi-center international clinical studies.

EDUCATION
Fairleigh Dickinson University, M.B.A.
Thomas Edison State University, B.S., Business Administration
RKD Paralegal Studies, Inc., Paralegal Certificate

Experience

Consultant to Legal Division

Warner Chilcott (now Allergan)

- Drafted and negotiated various agreements relating to the pharmaceutical industry

Manager, Pharmaceutical Licensing & Business Development

Warner-Lambert Company (now Pfizer)

- Evaluated licensing opportunities
- Assisted in structuring business deals

Legal Advisor, Corporate Legal Division

- Responsible for a broad range of agreements relating to research and drug development activities

Client Work

- Multiple clinical trials sponsored by IDEAYA Biosciences, Inc., including multinational phase 2/3 trials involving sites in North America, Europe and APAC
- Multiple clinical trials sponsored by Zenas Biopharma (USA) LLC, including multinational phase 3 trials involving sites in North America, South America, Europe and Asia
- Phase 2/3 gene therapy clinical trial sponsored by bluebird bio, Inc., involving approximately 12 subjects across 5 sites in Europe and the US
- Multiple clinical trials sponsored by BeiGene, including a multinational phase 3 trial involving approximately 467 subjects across 150 sites in 20 countries located in Asia, Australia, Europe, Canada and the US
- Multiple clinical trials sponsored by Tesaro, Inc., including a multinational phase 3 trial involving approximately 960 subjects across 225 sites in 25 countries located in Asia, Europe, Canada and the US

Community Involvement

- Life Science Law Professionals, member
- Model Agreements & Guidelines International (MAGI), member
- Women in the Enterprise of Science & Technology (WEST), member