

DENNIS J. NAAS, BS, PMP®**Degrees and Certifications:**

Bachelor of Science, Biology
Central Michigan University
Mt. Pleasant, Michigan
1979

Certified PMP® (Project Management Professional®)
2005

Professional Experience:**06/01/06 to Present**

ProDev Consulting Services, Ltd.
14244 Silver Ridge Road
Poway, CA 92064
V: 858-8836-2992
F: 866-300-6027
Email: dennis@prodevconsulting.com

Position: Sole Proprietor

Provision of expert, detailed and experienced regulatory, scientific and strategic preclinical and non-clinical toxicology consulting services to the pharmaceutical, biotechnology, chemical and agrochemical industries. Product development and commercialization support. Assistance in the areas of testing program and specific study design, protocol preparation, contract research organization evaluation and selection, study monitoring, GLP compliance and data and report review. Toxicology data summary preparation and review, manuscript preparation and review.

12/10/01 to 05/30/06

Pfizer Global Research and Development
Global Project Management
11085 Torreyana Road
San Diego, CA 92121

12/10/01 to 05/30/06

Position: Associate Director – Pfizer GPM
Global Project Manager

Responsible for providing strategic and operational leadership for multi-disciplinary drug development project teams and for the development and implementation of plans for drug projects. Experience with both development and integrated development/commercial teams. Management of the teams for time, cost and quality; coordination and assembly of the development plans. Identify and communicate best practices; establish and maintain an overall development schedule; manage progress versus plan and perform critical path analyses. Assist in development and tracking of an integrated project budget. Identification and quantification of risks associated with the project, including implementation of a formal risk management plan. Accountable for team communication, including project documents, meeting agendas, minutes and follow-up on action items. Management of team dynamics. Sole liaison for management of a Japanese in-license partner on a late-stage candidate, including contract closure.

7/1/97 to 12/9/01

AccuTox Consulting Services, Ltd.
3743 E. Hubbard Road
Midland, MI 48642

7/1/97 to 12/9/01

Position: Sole proprietor

Provision of expert, detailed and experienced regulatory, scientific and strategic preclinical and non-clinical toxicology consulting services to the chemical, pharmaceutical, agrochemical and biotechnology industries. Product development and commercialization support. Assistance in the areas of testing program and specific study design, protocol preparation, contract research organization evaluation and selection, study monitoring, GLP compliance and data and report review. Toxicology data summary preparation and review, manuscript preparation and review, computer system *beta* testing services.

5/1/84 - 6/27/97**WIL Research Laboratories, Inc.
Ashland, Ohio**

03/92 - 6/27/97

Position: Assistant Director of Toxicology

Direct and manage all technical and administrative aspects of general toxicology department (staff of 35). Preclinical testing program design and consultation. Responsible for scientific interpretation of multi-disciplinary results from directed studies. Study implementation and management. Recruitment, interview, hiring and performance evaluation of technical, supervisory and managerial staff in general and inhalation toxicology groups. Review, creation and implementation of new and revised technical procedures. Study costing, proposal and protocol preparation, study scheduling and coordination. General facility design and management. Routine communication with sponsor representatives, contractors and consultants.

9/88 - 03/92

Position: Senior Toxicologist

Direct all technical and administrative aspects of acute, subchronic and special toxicology studies. Study costing, proposal and protocol preparation, study scheduling and coordination. Oversee report preparation for all acute and special studies. Responsible for scientific interpretation of results from directed studies. Management of toxicology department. Involved in general facility management. Routine communication with sponsor representatives, contractors and consultants.

2/85 - 9/88

Position: Toxicologist - Acute, Subchronic and Special Toxicology

Direct all technical and administrative aspects of acute, subchronic and special toxicology studies. Study costing. Prepare proposals for procurement of contract work. Coordinate and schedule interdepartmental study activities. Prepared study protocols. Supervise preparation of reports. Responsible for scientific interpretation of study results. Development of standard operating procedures. Reviewed data for accuracy, consistency, completeness. Routine communication with sponsor representatives, contractors and consultants.

10/84 - 2/85

Position: Study Director - Acute Toxicology

Managed the operation of the in-life department. Coordinated activities of in-life department with support services groups. Trained employees. Prepared study protocols. Reviewed data for accuracy, consistency, completeness. Developed standard operating procedures. Verified test material preparation instructions, dose calculations and master computer protocols. Organized and formatted reports. Prepared interpretive summaries of study results. Responded to quality assurance audits. Routinely communicated with sponsor representatives.

5/84 - 10/84

Position: Section Head II - Acute Toxicology

Scheduled and directed the daily activities of technical personnel. Trained employees. Prepared study protocols as per with federal and international regulations. Reviewed data for accuracy, consistency and completeness. Developed standard operating procedures. Verified test material and dose calculations. Organized and formatted reports. Prepared interpretive summaries of study results. Reviewed tabulated data and reports. Routinely communicated with sponsor representatives.

6/1/79 - 4/20/84

**International Research and Development Corporation
Mattawan, Michigan**

9/16/82 - 4/30/84

Position: Unit Supervisor (Report Preparation)

Scheduled and directed the daily work activities of the unit. Reviewed raw data for accuracy, consistency, completeness and protocol adherence. Trained employees. Organized and formatted reports for presentation of methods, data and results. Prepared interpretive summaries of study results. Reviewed tabulated data and reports.

11/1/80 - 9/15/82

Position: Unit Supervisor (Subchronic Toxicology)

Scheduled and directed the daily work activities of the unit. Trained employees. Communicated with Sponsor's representative as directed. Prepared study record books and selected the animals to be used for studies.

6/1/79 - 10/30/80

Position: Research Associate (Acute Toxicology)

Responsible for the technical conduct of a wide variety of preclinical and nonclinical mammalian toxicology studies.