Michael Gross, Ph.D. RAC

Education and Training

- B.Sc., Chemistry, Philadelphia College of Pharmacy and Science
- Ph.D., Organic Chemistry, Temple University
- Post-doctoral Fellow, National Institute of Dental Research
- National Institutes of Health Grants Associate Program, Intern
- RAPS-RCAB, Regulatory Affairs Certified (RAC)

Industrial Experience

- Triton Biosciences-Shell Oil Company, Alameda, California, Director Regulatory Affairs
- Schering-Plough Research Institute, Kenilworth, New Jersey, Director Regulatory Affairs
- Becton Dickinson & Company, Franklin Lakes, New Jersey, Director Corporate Regulatory Affairs
- Aventis Behring, King of Prussia, Pennsylvania Vice-President World-Wide Compliance
- QLT Inc., Vancouver, British Columbia Vice-President, Regulatory Affairs
- Cardiome Pharma Corp. Vancouver, British Columbia Vice-President, Regulatory Affairs

Government Experience

- Health Scientist Administrator, National Heart Lung and Blood Institute, Bethesda, Maryland
- Food and Drug Administration, Bethesda, Maryland, Bureau of Biologics, Chemist

Committee Appointments

- Advisory Committee on Biotechnology-California State Assembly, Member
- Industrial Biotechnology Association (BIO), FDA Affairs Committee, Founder and Member
- Pennsylvania Biotechnology Association, Board of Directors, Member
- Pharmaceutical Manufacturers Association, Committee on Racemic Mixtures, Member
- United States Pharmacopea, Pharmaceutical Waters Expert Committee, Member
- Regulatory Affairs Professionals Society, Southern BC Chapter, Chair
- Parenteral Drug Association, Combination Products Interest Group, Leader
- NIH-NIAID Special Emphasis (Grant and Contract Review) Panel, Member
- PDA Foundation, Board Member