

# Amy Mihalchik, Ph.D., DABT, RAC

SUPERVISING SCIENTIST

## CONTACT INFORMATION

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## PROFESSIONAL PROFILE

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Dr. Amy Mihalchik is a board-certified toxicologist with a Regulatory Affairs Certification (R.A.C.) from the Regulatory Affairs Professionals Society with expertise in human health risk assessment. Her work primarily focuses on integrating quantitative structure activity relationship (QSAR) model data, read-across analyses, and other weight-of-evidence approaches to support the safety of substances, including drug degradants, cosmetics, consumer product ingredients, and industrial chemicals. Dr. Mihalchik is experienced in interpreting QSAR output from numerous platforms commercially and freely available, including Lhasa DEREK, Leadscope Model Applier, VEGA, EPA T.E.S.T., and ToxTree, among others, and has contributed to working groups assessing the potential for using in silico tools to support drug degradant safety and carcinogenicity hazard assessments.

She also integrates results from literature-based risk assessments with client-generated data, regulatory agency documents, and corporate technical reports to address complex exposure and toxicity issues. To support workplace safety in the pharmaceutical manufacturing arena, Dr. Mihalchik derives occupational exposure limits (OELs) and acceptable daily intakes (ADIs) and develops permissible daily exposure limits (PDEs) and tolerable exposure values (TEs) for pharmaceutical compounds and devices, respectively. She also has expertise analyzing data from leachables and extractables, testing of medical devices, including familiarity with ISO guidelines, and ISO-10993 biocompatibility assessments of medical devices.

Her doctoral research at the National Institute for Occupational Safety and Health (NIOSH) focused on addressing potential toxicities associated with multi-walled carbon nanotubes using in vitro methods to support “safety by design” efforts.

Dr. Mihalchik is familiar with the body of regulatory documents issued by the FDA, ICH, ISO, and other entities, and she maintains current knowledge of toxicology and regulatory science through extensive commitment to continuing education courses, webinars/seminars, and active participation in professional societies. She has published multiple peer-reviewed book chapters and scientific articles, and has presented at scientific conferences since 2014.

## EDUCATION AND DEGREES EARNED

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- 2016 Doctor of Philosophy (Ph.D.)  
Pharmaceutical and Pharmacological Sciences (field: nanotoxicology)  
West Virginia University, Morgantown, WV
- 2011 Bachelor of Arts (B.A.)  
General Biology (minor: English)  
Washington & Jefferson College, Washington, PA

## AWARDS AND GRANTS

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- AAAS/Science Program for Excellence in Science  
Society of Toxicology Student Travel Award, March 2015  
Integrated Graduate Education and Research Training (IGERT) Fellow, 2014–2016  
WV NANOSafe Graduate Fellow, 2013–2014

## SELECTED PROFESSIONAL EXPERIENCE

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### *Toxicology Consultant*

- Contributed to risk assessments across sectors
- Prepared gap analyses in support of GRAS and GRASE dossiers.
  - Conducted targeted literature searches addressing key endpoints of toxicological relevance (e.g., reproductive and developmental toxicity, carcinogenesis, genotoxicity) in databases, including PubMed and Embase.
- Developed proficiency in gathering relevant toxicological data from databases—including ToxPlanet, EPA CompTox Dashboard, PubChem, and FDALabel—to support projects across sectors (e.g., consumer products, foods, pharmaceuticals).
- Utilized a variety of freely available (e.g., VEGA, EPA T.E.S.T., ToxTree) and commercial (Q)SAR platforms to address complex client queries regarding mutagenicity, genotoxicity, carcinogenicity, and target-organ effects.
- Assessed cosmetic ingredients using literature review and (Q)SAR techniques.
- Supported a litigation project by reviewing and summarizing available literature regarding the case.
- Reviewed and interpreted nonclinical study data intended to support pharmaceutical products and medical devices.
- Supported ISO 10993-compliant evaluations of medical devices, including:
  - Gap analysis of available biocompatibility data
  - Review of device components, extractables, and leachables, including derivation of TE values per ISO 10993-17

### ***Regulatory Toxicology Consultant***

Performed risk assessments pertaining to pharmaceutical agents and medical devices:

- Provided literature-based risk assessments based on client data, regulatory agency documents and technical reports, and scientific literature
- Derived occupational exposure limits (OELs) and acceptable daily intakes (ADIs) to support workplace safety and appropriate cleaning carry-over limits in pharmaceuticals manufacturing
- Applied standard risk assessment methods to develop permissible daily exposure limits for pharmaceutical compounds and tolerable exposure values for medical devices
- Analyzed data derived from leaching and extraction testing of medical devices in compliance with ISO guidelines
- Performed ISO-10993 biocompatibility assessments of medical devices
- Determined F-values used to support child-proof packaging of pharmaceuticals.

Used Derek Nexus (knowledge-based), Leadscope Model Applier (statistics-based), and ToxTree software to predict mutagenicity and other endpoints associated with compounds in the absence of adequate data.

Provided expert review of QSAR output.

Identified appropriate surrogate compounds for assessment using read-across methods.

### ***Guest Researcher/Worksite Student***

Worked as a guest researcher for NIOSH:

- Studied effects of pristine multi-walled carbon nanotubes (MWCNT) and nitrogen-doped MWCNT (ND-MWCNT) at occupationally relevant doses on human lung small-airway epithelial cells
  - Conducted pioneering research on the impact of MWCNT physicochemical properties on the pulmonary system in a public health molecular toxicology lab
  - Provided an interface for translational interdisciplinary research between government and academic biomedical scientists and material scientists to address technical issues surrounding MWCNT safety
  - Determined that MWCNT and ND-MWCNT affect reactive oxygen species production, cell cycle progression, and cell signaling processes in lung cells.
- Studied effects of pristine MWCNT and ND-MWCNT on normal lung fibroblasts:
  - Investigated mechanisms of MWCNT-induced pulmonary fibrosis in an *in vitro* system using lung epithelial and fibroblast cells
  - Determined that MWCNT and ND-MWCNT induce production of mRNA for several genes that are key to pulmonary fibrosis.
- Gained experience in using and developing *in vitro* co-culture systems:
  - Used lung epithelial-endothelial co-culture to address toxicity of pristine MWCNT
  - Developed lung epithelial-fibroblast co-culture system to study mechanism of MWCNT and ND-MWCNT-induced pulmonary fibrosis.
- Also used various other techniques, including Western blotting, immunoprecipitation, confocal microscopy, ELISA, flow cytometry, cell cycle analysis, cell proliferation analysis, RNA extraction, and qRT-PCR.

### Doctoral Research

Completed research rotations in three laboratories at West Virginia University:

- Assisted on a project to assess the toxicity and potential therapeutic uses of nanoparticles in the Department of Orthopedics
- Became proficient in cell culture and basic molecular biology techniques through a breast cancer-focused research project
- Assisted on a project to identify specific molecular recognition elements (MREs) to detect malignant prostate cancer cells using systematic evolution of ligands by exponential enrichment (SELEX).

### PROFESSIONAL AFFILIATIONS

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American Association for the Advancement of Science — Member

American College of Toxicology — Associate Member

Society of Toxicology — Member, Associate Member

Association of Inhalation Toxicologists — Member

North Carolina Society of Toxicology Chapter — Member

Rho Chi Academic Honor Society in Pharmacy — Member

### MANUSCRIPTS

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Lea IA, Buerger AN, Feifarek D, **Mihalchik A**, Heintz MM, Haws LC, Nyambego H, Goyak K, Palermo C, Borghoff SJ. 2025. Evaluation of the endocrine disrupting potential of Di-isononyl phthalate. *Curr Res Toxicol* 8:100220; doi: 10.1016/j.crttox.2025.100220.

Lea IA, Feifarek D, **Mihalchik A**, Heintz M, Haws L, Nyambego H, Goyak K, Palermo C, Borghoff SJ. 2025. Evaluation of the endocrine disrupting potential of Di-isodecyl phthalate. *Curr Res Toxicol* 8:1002221; doi: 10.1016/j.crttox.2025.100221.

**Mihalchik AL**, Choksi NY, Roe AL, Wisser M, Whitaker K, Seibert D, Deore M, Pavlick L, Wikoff DS. 2024. Safety evaluation of 8 drug degradants present in over-the-counter cough and cold medications. *Regul Toxicol Pharmacol* 149:105621; doi: 10.1016/j.yrtph.2024.105621.

Tice RR, Bassan A, Amberg A, Anger LT, Beal MA, Bellion P... **Mihalchik-Burhans AL**, et al. 2021. *In silico* approaches in carcinogenicity hazard assessment: Current status and future needs. *Comp Toxicol* 20:100191 (Special issue: The in silico toxicology protocols initiative).

Sisler JD, Li R, McKinney W, Mercer RR, Ji Z, Xia T, Wang X, Shaffer J, Orandle M, **Mihalchik AL**, Battelli L. 2016. Differential pulmonary effects of CoO and La<sub>2</sub>O<sub>3</sub> metal oxide nanoparticle responses during aerosolized inhalation in mice. *Particle Fibre Toxicol* 13(1):42.

Sisler JD, Pirela SV, Shaffer J, **Mihalchik AL**, Chisholm WP, Andrew ME, Schwegler-Berry D, Castranova V, Demokritou P, Qian Y. 2016. Toxicological assessment of CoO and La<sub>2</sub>O<sub>3</sub> metal oxide nanoparticles in human small airway epithelial cells. *Toxicol Sci* 13(1):42.

**Mihalchik AL**, Ding W, Porter D, McLoughlin C, Schwegler-Berry D, Sisler JD, Stefaniak A, Snyder-Talkington BN, Cruz-Silva R, Terrones M, Tsuruoka S, Endo M, Castranova V, Qian Y. 2015. Effects of nitrogen-doped multi-walled carbon nanotubes compared to pristine multi-walled carbon nanotubes on human small airway epithelial cells. *Toxicology* 333:25–36.

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**BOOK CHAPTERS**

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**Mihalchik-Burhans AL**, Rogers EN. 2019. Considerations for leachables and extractables testing. In: Gad S (ed), Integrated Safety and Risk Assessment for Medical Devices and Combination Products, pp. 239–263. Springer.

**Mihalchik-Burhans AL**, Sullivan DW. 2019. Bridging issues of route. In: Gad S (ed), Integrated Safety and Risk Assessment for Medical Devices and Combination Products, pp. 273–297. Springer.

**Mihalchik AL**, Rogers EN. 2018. Classes of compounds with GI tract toxicity. Chapter 13 in: Gad SC (ed), Toxicology of the Gastrointestinal Tract. CRC Press.

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**ABSTRACTS AND PRESENTATIONS**

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**Mihalchik A**. Strategies to evaluate over-the-counter drug degradants. Abstract 1252, Society of Toxicology 64<sup>th</sup> Annual Meeting, Orlando, FL, March 2025.

Brown L, McMillan DA, Urban JD, **Mihalchik AL**. A tiered approach for assessing the safety of polymeric ingredients in cosmetics and personal care products. Poster presented at Society of Toxicology 62<sup>nd</sup> Annual Meeting, Nashville, TN, March 2023.

**Mihalchik AL**, Choksi NY, Wood ML. Toward best practices for read-across in evaluation of drug impurities, extractable, and leachable compounds. Poster presented at Society of Toxicology 62<sup>nd</sup> Annual Meeting, Nashville, TN, March 2023.

**Mihalchik AL**, Choksi NY, Lea I, Wood ML. Modern strategies to evaluate drug impurities. Session presented at Society of Toxicology 62<sup>nd</sup> Annual Meeting, Nashville, TN, March 2023.

**Mihalchik A**, Wood M. Considerations for standardization and derivation of pediatric and neonatal tolerable exposure limits for extractable and leachable compounds from medical devices. Poster presented at Society of Toxicology 61<sup>st</sup> Annual Meeting, San Diego, CA, March 2022.

Rogers EN, **Mihalchik AL**, Gad SC. Comparison and analysis of discrepancies among commonly used Cramer decision tree methods in Toxtree software. Abstract 1517, Annual Meeting, Society of Toxicology 60<sup>th</sup> Annual Meeting, Virtual, 2020.

**Mihalchik A** (contributor). In vitro and alternative models for regulatory submission. Podcast presentation available from American College of Toxicology, December 2018.

**Mihalchik-Burhans A**. And you want this by when? Making the successful transition from the bench to consulting. Oral presentation at the American College of Toxicology 39th Annual Meeting, West Palm Beach, FL, November 2018.

**Mihalchik A** (contributor). ACT annual meeting in West Palm Beach: Highlights and preview. Podcast presentation available from American College of Toxicology, October 2018.

**Mihalchik AL**, Rogers EN, Gad-McDonald SE, Sullivan DW Jr, Gad SC. Development of dermal permissible daily exposure (PDE) levels for elemental impurities by utilizing data from alternative routes. Poster presented at the Society of Toxicology 57th Annual Meeting, San Antonio, TX, March 2018.

**Mihalchik AL**. Effects of nitrogen-doped and pristine multi-walled carbon nanotubes in human bronchial epithelial cells and lung fibroblasts. Presented at Society of Toxicology 55th Annual Meeting, New Orleans, LA, March 2016.

**Mihalchik AL**. Addressing toxicity: In vitro caveats to the understanding of nanoparticle research and human health. Oral talk presented at the REN@WVU-NEEP Symposium, Morgantown, WV, October 2015.

**Mihalchik AL**. Nanoparticles: it's the little things in life that count. Oral talk presented at Davis & Elkins College in Elkins, WV, April 2015.

**Mihalchik AL**, Ding W, McLoughlin CE, et al. Effects of pristine and nitrogen-doped multiwalled carbon nanotubes (ND-MWCNT) on reactive oxygen species (ROS) and cell cycle progression. Presented at Society of Toxicology 54th Annual Meeting, San Diego, CA, March 2015.

**Mihalchik AL**, Ding W, Porter D, McLoughlin C, Schwegler-Berry D, Sisler JD, Stefaniak A, Snyder-Talkington BN, Cruz-Silva R, Terrones M, Tsuruoka S, Endo M, Castranova V, Qian Y. Effects of pristine and nitrogen-doped multiwalled carbon nanotubes (ND-MWCNT) on reactive oxygen species (ROS) and cell cycle progression. Poster session presented at the Society of Toxicology 54th Annual Meeting, San Diego, CA, March 2015.

**Mihalchik AL**, Ding W, Porter D, McLoughlin C, Schwegler-Berry D, Sisler JD, Stefaniak A, Snyder-Talkington BN, Cruz-Silva R, Terrones M, Tsuruoka S, Endo M, Castranova V, Qian Y. Effects of nitrogen-doped multi-walled carbon nanotubes compared to pristine multi-walled carbon nanotubes on human small airway epithelial cells. Poster session presented at the West Virginia University E.J. Van Liere Memorial Convocation & HSC Research Day, Morgantown, WV, February 2015.

**Mihalchik AL**, McLoughlin C, Schwegler-Berry D, Farcas M, Shvedova A, Porter D, Tsuruoka S, Endo M, Castranova V, Qian Y. Nitrogen-doped multi-walled carbon nanotube-induced effects in human small airway epithelial cells. Poster session presented at the West Virginia University Bench to Bedside: Translational Pharmacy Meeting, Morgantown, WV, June 2014.

**Mihalchik AL**, Porter D, Castranova V, Tsuruoka S, Endo M, Qian Y. Effects of MWCNT and nitrogen-doped MWCNT in lung epithelial cells. Presented at Society of Toxicology 53rd Annual Meeting, Phoenix, AZ, March 2014.

**Mihalchik AL**, McLoughlin C, Schwegler-Berry D, Farcas M, Shvedova A, Porter D, Tsuruoka S, Endo M, Castranova V, Qian Y. Nitrogen-doped multi-walled carbon nanotube-induced effects in human small airway epithelial cells. Poster session presented at the NIOSH Intramural Science Meeting, Morgantown, WV, July 2014.