NASCSA Conference 2021
Speaker Biographies

Haley Alder

Haley Alder serves as the administrator for the St. Louis County Prescription Drug Monitoring Program (PDMP). St. Louis County PDMP is the first and only locally-based PDMP, covering 85% of the state's population and 94% of healthcare providers. Haley has spear headed several system-level enhancements such as electronic health record integration, clinical alerts, NarxCare, and the expansion of interstate data sharing - increasing user-friendliness for providers and ultimately improving patient care. Haley brings a public health perspective to the program, having received a Master of Public Health from Saint Louis University prior to joining St. Louis County PDMP. Prior to that, she served as a Case Manager for clients with substance use disorders in Virginia. She believes public health encompasses the whole person and their environment, and that both health and racial equity must be at the forefront of any public health program. In her role she is also involved in developing and enhancing community partnerships and comprehensive strategies aimed to reduce the negative health outcomes and disparities associated with substance use, such as distributing Naloxone to community members in at no cost. The unique approach Haley brings to the PDMP has shaped the progress St. Louis County has made in an effort to bring a statewide program to Missouri.

Theo Antoniou

Theo Antoniou is the Chief Technology Officer at MDScripts during the past 12 years. In his role he is responsible for development of clinical dispensing features and pharmacy interfaces. He is responsible for the operation of claim adjudication interfaces, eprescribing interfaces, PDMP reporting, lab testing and drug pedigree interfaces. He is proficient in web application development, databases, communications and security.

Prior to MDScripts, Mr. Antoniou worked on real-time software development in communications systems at Motorola and access control systems at General Electric. Mr. Antoniou holds a Bachelor of Science degree in electrical and computer engineering from the University of Colorado.

Jenna Bluestein

Jenna Bluestein is a senior associate with Pew's substance use prevention and treatment initiative. In this role, she provides research support for state policy analysis and develops publications for policymakers on evidence-based treatment and overdose prevention for opioid use disorder. She also contributes to Pew's federal initiatives by compiling data for legislator education on medications for opioid use disorder. Before coming to Pew, she worked on physical and behavioral health integration in Medicaid at the National Academy for State Health Policy and supported local community health, preparedness, and response efforts at the National Association of County and City Health Officials. She holds a master's degree in public policy from the Johns Hopkins Bloomberg School of Public Health and a bachelor's degree from Wesleyan University.

Joe Bozenko

Joe Bozenko is a Senior Research Chemist with the DEA's Special Testing and Research Laboratory at Dulles, Virginia and a Scientific Advisor to DEA's Special Operations Division. He's been with the DEA for 22 years and investigates synthetic drug manufacturing around the world. Mr. Bozenko has processed some of the largest methamphetamine laboratories in the world, traveled extensively, and has authored and presented many reports and peer-reviewed scientific articles pertaining to the clandestine synthesis and analysis of controlled substances. Mr. Bozenko is closely involved with science-related officer safety and leads DEA's handheld instrumentation testing and evaluation. In addition to this, Mr. Bozenko has also been instrumental in the development of the DEA's High-Hazard Level 'A' Clandestine Laboratory Response Training Program. Mr. Bozenko is also charged with the specialized analysis of selected fentanyl, methamphetamine, and MDMA samples, both domestic and international, for intelligence purposes. Mr. Bozenko holds both a Baccalaureate and Master's Degrees in Chemistry and is an Adjunct Professor of Chemistry at Shepherd University. Mr. Bozenko also holds a patent, jointly with Harvard University, on the Archimedes Magnetic Levitation System.

Eric Brantley

Eric Brantley has 22 years of experience in the Pharmaceutical industry, with 19 years in Regulatory Compliance. Eric is currently an SOM Engagement Manager with IQVIA, and has been with the company for 2 ½ years. Prior to joining IQVIA, Eric managed SOM programs for both generic and brand pharmaceutical manufacturers, and spent 13 years with a major distributor involved in SOM and DEA Compliance. Eric has a passion for his work and engaging others on the topic of SOM. When not working, Eric enjoys listening to live jazz and cycling.

Claire Brennan

Claire Brennan is a Diversion Investigator for the Drug Enforcement Administration (DEA) who is currently the Chief of the Liaison Section for the Diversion Control Division in Arlington, VA. In this role, Ms. Brennan is responsible for the Liaison Section, ensuring that program objectives are met and acts as a liaison with DEA registrants and other partners in the prevention of diversion of controlled substances. Ms. Brennan has been employed with the DEA for 26 years and throughout her career has worked in the New Jersey, Seattle, and New England Division Offices as a Diversion Investigator, Diversion Group Supervisor, and Diversion Program Manager, respectively. Ms. Brennan holds a Bachelor of Science degree from the University of Scranton.

Kevin Borcher

Kevin Borcher is the Vice President, Pharmacy Informatics for Nebraska at CyncHealth. In his role, he is responsible for pharmacy systems, including the coordination of operations and enhancements of the PDMP and serving as the primary liaison between CyncHealth, professional associations, member organizations and stakeholders in relation to PDMP data collection and reporting, training, and publicity.

Prior to joining CyncHealth, Kevin served as a Pharmacy Informatics Coordinator for Nebraska Methodist Health System for more than 19 years. He also possesses extensive experience in clinical pharmacy, geriatrics, behavioral health, and hospital and community pharmacy. Kevin has a Doctor of Pharmacy degree from the University of Nebraska Medical Center, College of Pharmacy. He has served on the Nebraska Board of Health for over six years and is currently a member of the Nebraska Board of Pharmacy. Kevin is active in numerous organizations, and speaks to local, state, and national organizations and federal agencies on the activities surrounding CyncHealth and the PDMP.

Mark W. Caverly

Mark W. Caverly is a Senior Regulatory Consultant employed by IQVIA during the past 10 years. He specializes in assisting clients with the regulatory requirements of the Drug Enforcement Administration (DEA) and is responsible for working with clients to assure that potential regulatory issues are identified and remediated.

Mr. Caverly joined the company after a distinguished 31 year career with the U.S. Drug Enforcement Administration. In his last position with DEA, he was Chief of the Liaison and Policy Section, Diversion Control Division, at DEA Headquarters. In this capacity, Mr. Caverly was the primary point of contact to regulated industry, healthcare industry and associations and provided interpretations of federal law and regulations to Congressional staff, DEA registrants, and the general public. He represented DEA as a policy expert regarding compliance with Federal controlled substance and listed chemical laws and regulations. Mr. Caverly and his staff were also responsible for drafting and implementing new regulations.

Mr. Caverly's career at DEA also included 23 years of field experience. He served as a diversion investigator in the Miami, Boston, and Louisville divisions before becoming the Supervisory Diversion Investigator in the Louisville District Office.

Tony Garcia

Tony Garcia currently serves as the Executive Director for the South Texas High Intensity Drug Trafficking Area (HIDTA) program. The South Texas HIDTA is an alliance of over 87 federal, state and local law enforcement agencies working in close partnership to combat drug trafficking and related crime. Its area of operations includes fifteen Texas counties, thirteen of which are located between Del Rio and Brownsville, Texas on the border with Mexico. It also includes Bexar County and the San Antonio metropolitan area as well as Travis County and the Austin metropolitan area. Mr. Garcia also serves as the National Domestic Highway Enforcement (DHE) Chair for the HIDTA Program.

Mr. Garcia has been the South Texas HIDTA Executive Director since September 2008, having served as the Executive Director for the New Mexico HIDTA for eighteen months immediately prior to his present role. Mr. Garcia comes from a law enforcement background to include 31 years with the Texas Department of Public Safety (TXDPS). Mr. Garcia held numerous positions within TXDPS before he retired from that agency in 2007 as a Deputy Commander for the Narcotics Service. During his tenure at TXDPS, Mr. Garcia served on various committees, and was a member of the National Alliance of State Drug Enforcement Agencies, the International Association of Chiefs of Police and represented the state of Texas on the Border Governors' Security Commission. Mr. Garcia is a native Texan and was raised in the Rio Grande Valley. He attended Pan American University in Edinburg, Texas. Mr. Garcia is married and has two daughters and two grandchildren.

Jennifer Donnelly

Jennifer Donnelly is a Senior Epidemiologist and Assistant Director at the Kansas Board of Pharmacy. She is responsible for the oversight of the Kansas Prescription Drug Monitoring Program, K-TRACS, as well as being a lead epidemiologist for Kansas' Overdose Data to Action Cooperative agreement with Centers for Disease Control and Prevention. Jennifer came to Kansas in January 2019 after serving the state of Colorado for over 20 years at the Colorado Department of Public Health and Environment, HIV/STI and Viral Hepatitis Section. She has a Master of Public Health Degree with an Epidemiology emphasis from the Colorado School of Public Health. In May of 2019, Jennifer completed a fellowship at the Regional Institute for Health and Environmental Leadership. With over 20 years of public health experience, Jennifer understands the value of using data to guide program evaluation efforts and other policy decisions. She strives to tell stories through data visualization to promote understanding of complex public health issues.

John Gilbert

John A. Gilbert, Jr. counsels and advises clients on legal and regulatory issues involving controlled substances, prescription drugs, and precursor chemicals. His expertise extends to international, federal and state laws and regulations governing the scheduling, manufacturing, distribution, dispensing, import and export of controlled substances drugs and precursor chemicals. Mr. Gilbert has advised numerous companies at all levels of the drug supply chain on legal, regulatory and enforcement matters involving the Controlled Substances Act (CSA) and state laws governing controlled substances and precursor chemicals. Mr. Gilbert frequently conducts investigations and inspections related to compliance with federal and state laws and has handled numerous civil litigation matters involving violations of federal and state laws including actions before the DEA Office of Administrative Law Judges and civil actions initiated by U.S. Attorney Offices in federal court.

Mr. Gilbert has extensive experience in scheduling and regulation of controlled substances under the international drug control treaties and issues related to the United Nations Drug Control Program. He has advised and represented clients on matters related to the World Health Organization's Expert Committee on Drug Dependence, the International Narcotics Control Board and the U.N. Commission on Narcotic Drugs.

Mr. Gilbert also advises clients on compliance with federal and state requirements on licensing, pedigree, track and trace, and drug sampling requirements, including regulations associated with the Drug Quality and Security Act.

Before joining the firm in 1995, Mr., Gilbert was an attorney in the DEA's Office of Chief Counsel, Diversion/Regulatory Section. He also served as law clerk to the DEA's Chief Administrative Law Judge as part of the U.S. Department of Justice's Honors Program.

Sherry Green

For 28 years, Ms. Green has served as a national subject matter expert (SME) on drug, controlled substance, and health laws and policies. She specializes in prescription drug monitoring program (PMP) laws, drug abuse prevention and intervention policies, and multi-disciplinary alignment of laws and policies. Ms. Green served as the Associate Director of the President's Commission on Model State Drug Laws. She co-founded the Commission's non-profit successor, the National Alliance for Model State Drug Laws (NAMSDL), and served as its CEO and Executive Director for over 20 years.

After her tenure with NAMSDL, Ms. Green has continued her drug policy work by serving as a national SME for CDC, the Office of National Coordinator for Health Information Technology (ONC), the Agency for Health Care Research and Quality (AHRQ), and Congressional staff.
She has also advised the National Governors Association (NGA), the National Conference of State Legislatures (NCSL), the National Association of State Controlled Substances Authorities (NASCSA), and various health care entities.

Ms. Green has helped write and edit over 40 national model laws during her career, including a model PMP act, a model naloxone access act, and a model chemical control act. She was a member of the NASCSA task force updating the national model PMP Act and she serves on the Operations Subcommittee of the national PMP standards organization. Ms. Green trains and educates on PMPs, HIPAA (Health Insurance Portability and Accountability Act), controlled substance prescribing requirements and other key drug laws and policies for diverse stakeholders. Ms. Green has a Juris Doctor with Honors from George Washington University's National Law Center in Washington, D.C. and is a member of the D.C. Bar. She received a Bachelor's of Arts degree in Political Science-Economics from the University of Montana.

Eric Griffin

Eric Griffin - Is the Director of Compliance & Enforcement for the State of Ohio Board of Pharmacy. He oversees the daily operations of the Compliance & Enforcement Division, managing a staff of more than thirty investigators and support staff. He routinely directs major investigations, facilitates public outreach and training, and coordinates multi-agency initiatives. Eric started his law enforcement career at Delaware County Sheriff's Office where he served in numerous positions including Detective, Sergeant, Drug Task Commander, and Lieutenant. Eric has led and participated in a multitude of investigations at the State and Federal Level in the prosecution of Homicides and Major Drug Offenders to Health Care Professionals. As a Lieutenant, he supervised and oversaw the daily operations of the Detective Bureau, Drug Task Force and Administrative Investigations. Eric has more than twenty years of law enforcement experience and has attended countless training sessions across the country. Eric regularly presents on issues concerning drugs of abuse and diversion to law enforcement, administrative agencies, and the general public.

Larry Houck

Larry K. Houck provides counsel on regulatory and enforcement actions by the DEA. His career encompasses over 30 years of conducting investigations and negotiating on behalf of both the government and the industry. Mr. Houck focuses on controlled substances, prescription drugs, and regulated chemicals, helping clients navigate federal and state licensing, registration, and compliance issues.

Mr. Houck counsels clients throughout the registrant supply chain on administrative, civil, and criminal proceedings. In situations where clients face enforcement action, Mr. Houck has extensive understanding of the DEA's approach and priorities. He advises pharmaceutical and chemical companies on DEA inspections and audits. By working with clients to review business practices, he helps create the infrastructure to ensure compliant reporting, record keeping, and security.

Before joining Hyman, Phelps & McNamara in 2001, Mr. Houck worked as a DEA diversion investigator and policy staff coordinator. As a diversion investigator in the Washington, D.C. and Portland, Oregon, field offices, Mr. Houck conducted a full range of regulatory and criminal investigations and inspections of controlled substance and chemical registrants. While serving as a staff coordinator for the DEA's Office of Diversion Control's Liaison and Policy, he advised government officials and pharmaceutical and health care professionals on the Controlled Substance Act and its regulations. Mr. Houck drafted and implemented the DEA's controlled substance policies and regulations on diversion control issues that included pain management.

Janetta L. Iwanicki, MD

Dr. Janetta Iwanicki is the Chief Scientific Officer for Research and Consulting at Rocky Mountain Poison & Drug Safety—Denver Health and Hospital Authority in Denver, Colorado. She is an Assistant Professor of Emergency Medicine at the University of Colorado School of Medicine. Dr. Iwanicki is board-certified by both the American Board of Emergency Medicine and the American Board of Medical Toxicology, and teaches as an attending physician in both fields.
Dr. Iwanicki conducts her clinical research with the RADARS® System, and is the PI of an FDA BAA Grant focused on understanding prescription stimulant. Her work focuses on innovative ways to evaluate patterns and trajectories of use of prescription and non-prescription substances, develop novel survey tools and recruitment methodologies, and provide insights into regulatory and policy implications of her findings. She has published more than forty manuscripts and abstracts to better describe the patterns of drug use as well as the theoretical constructs vital for understanding these patterns, and has presented at multiple national and international meetings on these topics. She has authored ten book chapters in the fields of Medical

Toxicology and Emergency Medicine.

Alan McGill

Alan McGill - Is a Senior Supervisory Special Agent with the Pennsylvania Office of Attorney General and currently assigned to the Office of Public Engagement. Agent McGill has been a professional investigator for 30 years with 23 years in law enforcement. He began as a Private Investigator and then became a Freeport Police Officer for three years before joining the Pennsylvania Office of Attorney General where he spent the last 20 years with the Bureau of Narcotics Investigation and Drug Control. Agent McGill has served the Attorney General's Office as both a street supervisor and Drug Diversion Unit Supervisor, Undercover Narcotics Agent, Drug Task Force Coordinator, Drug Diversion Agent, Computer Voice Stress Analyst, PDMP BNI Statewide supervisor, Clandestine Laboratory Enforcement Program member and an "A" and "B" Technician installing wiretap equipment. Although Agent McGill's undercover experience is extensive, he is best known as a subject matter expert in Drug Diversion and Pharmaceutical Investigations. Agent McGill has personally conducted and/or supervised hundreds of diversion investigations at pharmacies, hospitals and nursing homes along with health and oversight duties such as inspections of pharmaceutical drugs and records. Additionally, Agent McGill has been conducting Drug Diversion presentations to clinicians at various hospitals, and medical associations throughout Pennsylvania and has been a national speaker on the topic. Additionally, Agent McGill is a guest lecturer at 7 Pennsylvania University clinician programs. Agent McGill has been training law enforcement on prescription drug investigations since 2012 at the OAG Agent Academy and Pennsylvania State Police Training facilities. In 2017 and 2018 Agent McGill authored an 8 hour Basic Drug Diversion Course and a Three Day Advanced Drug Diversion Course of which both are certified by the MPOETC (Municipal Police Officers Education and Training Commission) for 8 hours and 18 hours of continued education respectively. Agent McGill is an Executive Committee member of where he serves as Chair and chairs the Education Committee.

Machelle Neal

Machelle Neal holds the position of Principal, Regulatory Consulting Services for IQVIA U.S. Compliance Solutions Group (formerly BuzzeoPDMA). Ms. Neal currently manages and oversees the US Compliance Regulatory Consulting Group and provides subject matter expertise to the industry and the health care community in the areas of the Controlled Substance Act, Prescription Drug Marketing Act and more specifically, state pharmacy and wholesale distribution regulatory issues.

With over 28 years of pharmacy operational and regulatory compliance experience in the healthcare industry, Ms. Neal assists clients with the development and maintenance of programs for pharmacy regulatory compliance, diversion surveillance and prevention, state and federal pharmacy and wholesale distribution licensing and regulatory determinations, and provides regulatory and operational training specific to these areas. Prior to this position, Ms. Neal was Compliance Director of State and Compliance Services, where she was responsible for the regulatory review and auditing functions of our consulting staff, and for the management of our regulatory research, state survey, and licensing departments. Previously, she held the position of Director, State Liaison for the company where she managed the company's federal and state regulatory research programs and established ongoing professional relationships with the federal and state offices, providing clients with a resource as liaison to federal and state agencies.

Prior to joining BuzzeoPDMA, Ms. Neal was employed as Compliance Manager, Regulatory Compliance by Omnicare, Inc., the nation's largest professional pharmacy related consulting and data management services for long-term care. Ms. Neal was responsible for the development of the company's regulatory compliance program and was responsible for the day-to-day regulatory oversight of approximately 230 institutional pharmacies nationwide. Prior to her position as Manager, Regulatory Compliance, Ms. Neal held the position of Manager, Corporate Communications with Omnicare where she was responsible for the management of the company's shareholder services, sales and marketing development, implementation, and training resources.

Ms. Neal graduated Cum Laude from Cincinnati State College, Cincinnati, Ohio.

Ralph Orr

Ralph A. Orr is the Director for the Virginia Prescription Monitoring Program (PMP) at the Virginia Department of Health Professions. Since 2005 Mr. Orr has seen the program go statewide, implement web-based 24/7 access, implement interoperability with 41 other programs nationwide as well as integration with 5,000 facilities across Virginia. Mr. Orr previously served as Deputy Executive Director of the Virginia Board of Pharmacy. Prior to joining the Department of Health Professions in 2002, Mr. Orr served 22 years in the United States Army. Mr. Orr was awarded the Virginia Governor's Award for Agency "Star" in 2009 and is a member of the Order of Military Medical Merit. Mr. Orr has served in various positions for the National Association of State Controlled Substances Authorities including President and Committee Chair.

Sidney Seal

Sidney "Sid" Seal, R.Ph., Oxford, MS., B.S. Pharmacy 1985, University of Mississippi School of Pharmacy; 1985-2011, Super D/USA Drugs, Pharmacist, PIC, District Supervisory positions; 2011-present, Mississippi Board of Pharmacy, Sr. Compliance Agent, Pharmacy regulatory/Drug Diversion/Fraud cases with MS Bureau of Narcotics, Medicaid, MS AGO, DEA, and FBI; 2014-2017, Secretary MS Chapter of NADDI (Nat'l Assn of Drug Diversion Investigators), Mississippi Opioid and Addiction Town Hall Meetings, Speaker/Presenter, Avid college sports fan, music aficionado, and collector of signed first addition books; Proud husband, father, and grandfather.

Mary-Lou Schoonover

Mary-Lou Schoonover currently serves as the Customer Due Diligence and State Licensing Manager for Par Pharmaceuticals, Inc., an Endo owned company. She is responsible for the licensing of all the Par/Endo entities as well as the DEA Suspicious Order Monitoring/Customer Due Diligence program. Mary-Lou has over 20 years of combined experience between PDMA and DEA compliance in the pharmaceutical industry. She has work for MedPro Systems, Allergan (formerly Actavis), Reckitt Benckiser and lastly, the United States Pharmacopeia. Considered a subject matter expert in State Licensing, PDMA and DEA Compliance, Mary-Lou attributes her wealth of knowledge and success to the many folks she has worked with. She has a Bachelor of Science in Business Administration and Management from Centenary University and; is a graduate of the National Academy of Paralegal Studies. Mary-Lou is a Certified Corporate Paralegal. She has served as an industry volunteer on several committees at NASCSA for many years. Mary-Lou has also presented at NASCSA and other industry conferences. She is a member of or affiliated with NASCSA, NADDI and IQVIA. In her spare time, Mary-Lou volunteers for the Center for Prevention in Sussex County, NJ.

Jason Slavoski

Jason Slavoski joined the Delaware Office of Controlled Substances in September of 2017 as the Delaware PMP Administrator. He holds a Doctor of Pharmacy degree from Wilkes University and is a licensed pharmacist in Delaware, Maryland, and Pennsylvania.
Prior to becoming the Delaware PMP Administrator, Jason served in a management role for several pharmacy retailers. Jason also served as a consultant for RxWiki and as a Medical Science Liaison for a company specializing in pharmacogenomics. Jason is responsible for the management and daily operation of Delaware's PMP program. He sits on several advisory boards and committees in the State of Delaware and nationally. This is Jason's third year attending the annual NASCSA conference and first time running for a position on the Executive Committee.

Noreen S. Valentine

Noreen S. Valentine is the Chief of the Policy Section for the Drug Enforcement Administration's Diversion Control Division. In this role, Ms. Valentine's primary responsibility is to review all correspondence drafted as a result of receiving inquiries via letter, email, conference calls, and/or telephonic contact from DEA registrants, DEA personnel, Federal, state, and local authorities, and the public at large regarding DEA regulations.

Ms. Valentine has an employment history with the administration that spans 30 years. Ms. Valentine formerly held positions as a Group Assistant, Forfeiture Specialist, Operations Assistant, and Paralegal Specialist before embarking on her career as a Diversion Investigator. Ms. Valentine became a Diversion Investigator in August of 2003 and was assigned to the Washington Field Division, Washington, D.C. As a Diversion Investigator, Ms. Valentine has directed or been involved in regulatory, criminal, and complaint investigations, involving both pharmaceuticals and chemicals.
Ms. Valentine also served as a recruiter for 10 years and was detailed to the division's Financial Investigations Group. In 2013, Ms. Valentine was promoted to the position of Supervisory Diversion Investigator, and was assigned to the Policy Section for approximately three years.
In 2016, Ms. Valentine was selected to serve as a Group Supervisor of a Diversion Regulatory Group in the Philadelphia Field Division, where she was responsible for registrant population in the Eastern and Middle Districts of Pennsylvania and the state of Delaware. In this capacity, Ms. Valentine supervised seven Diversion Investigators, overseeing and assisting with a multitude of investigations.
Ms. Valentine received a Bachelor of Arts in Sociology from Saint Leo University.

Joshua S. Vinciguerra

Joshua S. Vinciguerra is the Director of the Bureau of Narcotic Enforcement at the New York State Department of Health. He is a former federal prosecutor in the U.S. Attorney's Office for the Northern District of New York. He has also served as an Assistant Attorney General in the New York State Attorney General's Criminal Enforcement and Financial Crimes Bureau, and as an Assistant District Attorney in the New York County District Attorney's Office under Mr. Robert Morgenthau, where he prosecuted narcotics and weapons crimes in the Office of Special Narcotics, and white-collar crime in the Frauds Bureau. He clerked in the New York State Supreme Court, Appellate Division, Fourth Department, in Rochester, New York. He is a volunteer firefighter.

Haley Winans

Haley Winans is the Specialist for the Michigan Automated Prescription System (MAPS), within the Bureau of Professional Licensing at the State of Michigan. Haley administers MAPS in partnership with Bamboo Health, formerly known as Appriss Health. Specifically, she oversees projects, grants and enhancements of MAPS, including the transition from a homegrown system to Appriss Health's PMP AWARxE software platform, implementing NarxCare, OpenBeds, and statewide integrations. She generates and reviews controlled substance prescribing and dispensing reports and assists practitioners, pharmacists, law enforcement officials, benefit plan managers with registration and data submission to MAPS. As a part of the Enforcement Division, Haley is the point-person for analytics of MAPS, and assists in reviewing data to open overprescribing, overdispensing, and drug diversion cases. As follow-up, she works closely with the Investigators of these cases, as well as the Assistant Attorney General's who litigate these cases by understanding the data and testifying at hearings if required. Haley has been with the State of Michigan for the past 10 years, 9 of which have focused on MAPS. Prior to joining the State of Michigan, she worked as a pharmacy technician at Meijer Pharmacy. Haley graduated from the University of Michigan-Flint with a Bachelor's in Health Care Administration.