AAPS PharmSci 360 - Poster Abstract Tracks (Subtracks) | Primary Topics | Keywords = (Review Groups)

There are four tracks covering all aspects of the pharmaceutical sciences. Three tracks are divided into two subtracks with Biomolecular and Chemical focuses.
The Preclinical, Clinical, and Translational Sciences track is not split into subtracks.

IMPORTANT NOTE: In the submission site the structure below, Track (Subtrack), Primary Topic, and Keywords, are referred to as Review Groups. Screeners, please select all Review Groups that apply to your expertise.

Review Group Selection Process

1. Select the Track-Subtrack that best fits your research. ***(see below)***

* Preclinical, Clinical, and Translational Sciences
* Bioanalytics
* Manufacturing and Analytical Characterization
* Formulation and Delivery

2. Select the Subtrack (Biomolecular or Chemical). Does not apply to the Preclinical, Clinical, and Translational Sciences track.

3. Select the Primary Topic that best fits your research.

4. From your Primary Topic, select the best Keywords for your research. If no listed Keywords fits your research, select 'Other'. (*Note: In some instances there are no Keywords.*)

TRACK (SUBTRACK)	PRIMARY TOPIC	KEYWORDS
Bioanalytics - Biomolecular	Analyte Stability	Ex Vivo
		In Vitro
		Solution
Bioanalytics - Biomolecular	Bioanalytical	New Applications of Existing Technology
		New Approaches
		New Technology
Bioanalytics - Biomolecular	Bioanalytical Innovations and Applications	n/a
Bioanalytics - Biomolecular	Bioanalytical Risk Assessment and Strategy	n/a
Bioanalytics - Biomolecular	Biology	Biomarkers
		Immunogenicity
Bioanalytics - Biomolecular	Biomarker Quantification	Biomarker/Pharmacodynamic Measurement
		Clinical Qualification
		Diagnostic Development (including companion diagnostics)
		Disease Heterogeneity Assessments
		Exosome
		Flow Cytometry Methods
		High Content Data Analysis
		Hybrid Methods (e.g. IP/LCMS)
		Imaging Methods
		Ligand Binding Assay (LBA) Methods
		Mass Spectrometry (LC-MS) Methods
		New Matrices
		New Modalities
		OMICs
		PCR Methods
		Preanalytical Variables
		Single-Cell-Based Biomarkers
		Target Engagement/Receptor Occupancy
		Vaccines
Bioanalytics - Biomolecular	Drug Quantification	Endogenous Homologs Quantification
		Flow Cytometry Methods
		Hybrid Methods (e.g. IP/LCMS)
		Imaging Methods
		Ligand Binding Assay (LBA) Methods
		Mass Spectrometry (LC-MS) Methods
		Other Methods/Techniques
		PCR Methods
		Post-marketing Commitment
		Surrogate Analyte
		Therapeutic Drug Monitoring
Bioanalytics - Biomolecular	Immunogenicity	Binding Antibody Methods
		Cell-Based Methodologies
		Immunogenicity Prediction
		Immunogenicity Risk Assessments
		Neutralizing Antibody Methods
Bioanalytics - Biomolecular	In Vivo and Ex Vivo Biotransformation	ADC Metabolism
		Evaluation of In Vivo Biotransformation
		Impact of Biotransformation on Immunogenicity
		Impact of Biotransformation on PK
		Molecule Variants Quantification Ex Vivo
Bioanalytics - Biomolecular	Life Cycle Management of Bioanalytical Methods	Collaboration with other Partners (Co-development)
		Data Management
		General Life Cycle Management
		Methods Transfer and CRO Management
Bioanalytics - Biomolecular	Novel Modalities	ADCs
		Alternative Scaffold
		CAR-T
		Cell-Based Therapy
		Encapsulated Drugs (Lipid, Nanoparticle, etc.)
		Multi-specific Antibodies
		Nanoparticle Based Modalities
		Oligos, RNAs, and Locked Nucleic Acids
		Other
		Viral Vectors

TRACK (SUBTRACK)	PRIMARY TOPIC	KEYWORDS
Bioanalytics - Biomolecular	Pharmacology	Bioanalytical
		Biomarkers
Bioanalytics - Biomolecular	Reagents and Reference Standards	Characterization and Quality Control
		Life Cycle Management
		Stability
Bioanalytics - Biomolecular	Regulations (BMV/GLP/GCP/CLIA)	Biomarkers
		Drug (and Metabolites)
		GCP/GLP Compliance for Bioanalytical Labs
		General Topics
		ICH and Harmonization
		Immunogenicity and Risk Assessment
		Samples and Reagent Stability
Bioanalytics - Biomolecular	Samples and Laboratory Management	Bioanalytical Documentation and Reports
		Biorepositories
		Informed Consent
		Laboratory Information Management System (LIMS)
		Patient-Centric Sampling (microsampling and dried blood
		Post-Collection Sample Condition and Record Management
Bioanalytics - Biomolecular	Vaccines	Binding Antibody Methods
		Cell-Based Methodologies
		Correlates of Protection
		Neutralizing Antibody Methods
Bioanalytics - Chemical	Analyte Stability	Ex Vivo
		In Vitro
		Solution
Bioanalytics - Chemical	Bioanalytical Innovations and Applications	n/a
Bioanalytics - Chemical	Bioanalytical Risk Assessment and Strategy	n/a
Bioanalytics - Chemical	Biomarker Quantification	Biomarker/Pharmacodynamic Measurement
		Clinical Qualification
		Diagnostic Development (including companion diagnostics)
		Disease Heterogeneity Assessments
		Exosome
		Flow Cytometry Methods
		High Content Data Analysis
		Hybrid Methods (e.g. IP/LCMS)
		Imaging Methods
		Ligand Binding Assay (LBA) Methods
		Mass Spectrometry (LC-MS) Methods
		New Matrices
		New Modalities
		OMICs
		PCR Methods
		Preanalytical Variables
		Single-Cell-Based Biomarkers
		Target Engagement/Receptor Occupancy
Bioanalytics - Chemical	Drug Quantification	Endogenous Homologs Quantification
		Flow Cytometry Methods
		Hybrid Methods (e.g. IP/LCMS)
		Imaging Methods
		Ligand Binding Assay (LBA) Methods
		Mass Spectrometry (LC-MS) Methods
		Other Methods/Techniques
		PCR Methods
		Post-marketing Commitment
		Surrogate Analyte
		Therapeutic Drug Monitoring
Bioanalytics - Chemical	Immunogenicity	Binding Antibody Methods
		Cell-Based Methodologies
		Immunogenicity Prediction
		Immunogenicity Risk Assessments
		Neutralizing Antibody Methods
Bioanalytics - Chemical	In Vivo and Ex Vivo Biotransformation	ADC Metabolism
		CYP450 Assessment
		Evaluation of In Vivo Biotransformation
		Impact of Biotransformation on Immunogenicity
		Impact of Biotransformation on PK
		Metabolite Quantification
Bioanalytics - Chemical	Life Cycle Management of Bioanalytical Methods	Collaboration with other Partners (Co-development)
		Data Management
		General Life Cycle Management
		Methods Transfer and CRO Management
Bioanalytics - Chemical	Medicinal Chemistry	Structure Activity Relationship
Bioanalytics - Chemical	Novel Modalities	ADCs
		Alternative Scaffold
		CAR-T
		Cell-Based Therapy
		Encapsulated Drugs (Lipid, Nanoparticle, etc.)
		Multi-specific Antibodies
		Nanoparticle Based Modalities
		Oligos, RNAs, and Locked Nucleic Acids
		Other
		Viral Vectors
Bioanalytics - Chemical	Reagents and Reference Standards	Characterization and Quality Control
		Life Cycle Management
		Stability

TRACK (SUBTRACK)	PRIMARY TOPIC	KEYWORDS
Bioanalytics - Chemical	Regulations (BMV/GLP/GCP/CLIA)	Biomarkers
		Drug (and Metabolites)
		GCP/GLP Compliance for Bioanalytical Labs
		General Topics
		ICH and Harmonization
		Immunogenicity and Risk Assessment
		Samples and Reagent Stability
Bioanalytics - Chemical	Samples and Laboratory Management	Bioanalytical Documentation and Reports
		Biorepositories
		Informed Consent
		Laboratory Information Management System (LIMS)
		Patient-Centric Sampling (microsampling and dried blood
		Post-Collection Sample Condition and Record Management
Formulation and Delivery - Biomolecular	Administration	DP Handling
		In-Use Compatibility
		Nasal/Pulmonary
		Ocular
		Other
		Otic
		Potent Modalities
		Sterility and Microbiology
		Strategies
		Transdermal
Formulation and Delivery - Biomolecular	Drug Delivery	Extended Release (Non-implant)
		Implants
		On Body Delivery Systems (OBDS)
		Other
		Other Routes of Administration - Ocular
		Other Routes of Administration - Other
		Other Routes of Administration - Otic
		Other Routes of Administration - Transdermal and Topical
Formulation and Delivery - Biomolecular	Drug Delivery, Devices, and Drug Device	Design Control
		Hardware
		Human Factor Engineering
		New Delivery Technologies
		Patient-Centric Development
		Software
Formulation and Delivery - Biomolecular	Formulation	Cell Therapy
		Free Oligonucleotide
		Gene Therapy
		Other
		Protein - Developability Assessment
		Protein - Excipients
		Protein - High-Throughput Screening
		Protein - Lyo
		Protein - Other
		Protein - Syringes
		Protein - Topics
		Vaccine/Tolerance Induction
Formulation and Delivery - Biomolecular	Pharmaceutics	Drug Delivery Novel Systems
		Drug Delivery Ocular
		Drug Delivery Transdermal
		Pharmaceutical Polymers
Formulation and Delivery - Biomolecular	Primary Packaging	Compatibility
		Container Closure Integrity
		Extractables/Leachables
		New Materials
Formulation and Delivery - Biomolecular	Regulatory Considerations	(Sub)visible Particles
		Accelerated Approval Pathways
		Bioequivalence
		Biosimilars
		Innovative Technologies
		Inspections and GMPs
		Large Market Developments
		New Regulations and Guidances
		Risk Assessment Implementation
		Smaller Market Developments
		Stability Requirements
Formulation and Delivery - Chemical	Biopharmaceutics	BCS, DCS
		Bioequivalence (also Regulatory)
		Comparability Assessments
		IVIVC
		Other
		Predictive Modeling
Formulation and Delivery - Chemical	Drug Delivery	Extended Release (Non-implant)
		Implants
		Nanoparticles
		Other
		Other Routes of Administration - Ocular
		Other Routes of Administration - Other
		Other Routes of Administration - Otic
		Other Routes of Administration - Transdermal and Topical
Formulation and Delivery - Chemical	Drug Delivery, Devices, and Drug Device	Design Control
		Hardware
		Human Factor Engineering

TRACK (SUBTRACK)	PRIMARY TOPIC	KEYWORDS
Formulation and Delivery - Chemical	Drug Delivery, Devices, and Drug Device	New Delivery Technologies
		Patient-Centric Development
		Software
Formulation and Delivery - Chemical	Formulation	Advanced Dissolution Testing
		Amorphous and Co-crystal Systems
		Bioavailability Enhancement
		Drug Substance Properties
		Excipients
		Fixed Dose Combinations
		Inhalation and Nasal
		Oral - Immediate Release
		Oral - Modified Release
		Parenterals
		Predictive Modeling
		Preformulation
		Special Populations
Formulation and Delivery - Chemical	Pharmaceutics	Drug Delivery Novel Systems
		Drug Delivery Ocular
		Drug Delivery Transdermal
		Pharmaceutical Polymers
Formulation and Delivery - Chemical	Primary Packaging	Compatibility
		Container Closure Integrity
		Extractables/Leachables
		New Materials
Formulation and Delivery - Chemical	Regulatory Considerations	(Sub)visible Particles
		Accelerated Approval Pathways
		Bioequivalence
		Biosimilars
		Innovative Technologies
		Inspections and GMPs
		Large Market Developments
		New Regulations and Guidances
		Patient Focused Drug Development Guidelines
		Risk Assessment Implementation
		Smaller Market Developments
		Stability Requirements
Manufacturing and Analytical Characterization - Biomolecular	Analytical	(Sub)visible Particles
		Combination Products
		Excipients
		Immunogenicity
		Impurities
		Modality Specific Methods - Cell Therapy
		Modality Specific Methods - Free Oligonucleotide
		Modality Specific Methods - Gene Therapy
		Modality Specific Methods - Other
		Modality Specific Methods - Protein
		Modality Specific Methods - Vaccine/Tolerance Induction
		New Technology
		Other
		Potency/Bioassay
Manufacturing and Analytical Characterization - Biomolecular	Automation	Computer Validation
		Other
Manufacturing and Analytical Characterization - Biomolecular	Biosimilar Manufacturing	Biosimilarity Assessment
		Other
		Patent Protection
Manufacturing and Analytical Characterization - Biomolecular	Drug Product Manufacturing and Development	Aseptic Technologies - Mixing, Sterilization, and Filling
		Cell Therapies
		Freezing and Thawing
		Immunogenicity and Critical Quality Attributes
		Lyophilization and Drying Technologies
		Manufacturing and Assembly of Drug/Device Combinations
		Manufacturing of Drug Delivery Systems
		Other
		Primary Packaging - Container Closure Integrity
		Primary Packaging - Other
		Primary Packaging - Syringes
		Primary Packaging - Vials
		Process Characterization and Optimization
		Protein Aggregation and Degradants
		Secondary Packaging
		Storage Considerations
		Vaccines
		Viral and Non-viral Vectors and Gene Therapy
		Visible and Subvisible Particles
		Visual Inspection
Manufacturing and Analytical Characterization - Biomolecular	Drug Substance Manufacturing and Development	API Packaging and Storage
		Cell Line Development
		Cell Therapies
		Clonality Assessments
		Expression Systems - Cellular and Cell-Free
		Genetic and Cell Line Engineering
		Mammalian Cell Culture
		Media Development
		Microbial/Yeast Fermentation
		Other

TRACK (SUBTRACK)	PRIMARY TOPIC	KEYWORDS
		Process Optimization and Intensification
		Protein Aggregation during Processing and Immunogenicity
		Purification and Virus Removal
		Vaccines
		Viral and Non-viral Vectors and Gene Therapy
		Virus Safety/Removal
Manufacturing and Analytical Characterization - Biomolecular	General Aspects and Strategies	Change Control
		CMO Management
		Drug Master Files
		Drug Substance and Drug Product Shipment
		Electronic Records
		Handling Control Substances (DEA)
		Inspections and GMP
		Lean Manufacturing/Six Sigma/Operational Excellence
		Life Cycle Management
		Manufacturing Economics
		Materials Management and Warehousing
		Other
		Regulatory Strategy
		Supply Chain
Manufacturing and Analytical Characterization - Biomolecular	Health, Safety, and Environment	Containment and Isolators
		High-Potent Drug Manufacturing
		OEL and PDE
		Other
Manufacturing and Analytical Characterization - Biomolecular	Innovative/Novel Processing Technologies and Concepts	For Use in Drug Product Manufacture
		For Use in Drug Substance Manufacture
		Other
Manufacturing and Analytical Characterization - Biomolecular	Integrated and Continuous Processing and Manufacturing	For Use in Drug Product Manufacture
		For Use in Drug Substance Manufacture
		Other
Manufacturing and Analytical Characterization - Biomolecular	Manufacture of Clinical Supplies	Blinding of Comparator Drugs
		Other
		Phase Appropriate GMP
		Speed to Patient
Manufacturing and Analytical Characterization - Biomolecular	Plant Engineering and Maintenance	Facility Design
		Legacy Facility Innovation/Renovation
		Media
		Media Fills
		Media/Buffer Preparation and Fluid Management
		Modeling and Scheduling Multiproduct Batch Plants
		Modular Manufacturing
		Other
		Plant Incident Investigations
Manufacturing and Analytical Characterization - Biomolecular	Process Design and Controls	Cleaning Validation
		Control of Impurity Formation
		In-Process Controls
		Other
		Process Analytical Technology and Parametric/Real-Time
		Process Modeling and Simulations
		Process Validation/Continuous Process Validation
		QbD and Assessment of Process Parameters
		Scale-Up/Process Transfers
		Statistical Process Controls and Six Sigma
		Use of Prior Knowledge and Risk-Based Approaches
Manufacturing and Analytical Characterization - Biomolecular	Single-Use and Disposable Systems	For Use in Drug Product Manufacture
		For Use in Drug Substance Manufacture
		Leachables and Extractables
		Other
Manufacturing and Analytical Characterization - Chemical	Analytical	Continuous/Real-Time Release
		Drug Release Measurement - Biorelevant Dissolution
		Drug Release Measurement - Cascade Impaction
		Drug Release Measurement - Dissolution
		Drug Release Measurement - Forms
		Drug Release Measurement - Other
		Excipients
		Impurities and Degradation - Forced Degradation
		Impurities and Degradation - Impurity Quantitation
		Impurities and Degradation - In Silico Predecision of Stability
		Impurities and Degradation - Other
		Method Development Strategies
		Model Maintenance
		New Analytical Technologies
		Other
		Physical Characterization Techniques
		Process Analytical Technology and Continuous Release
		Real Time Release testing
Manufacturing and Analytical Characterization - Chemical	Automation	Computer Validation
		Other
		Aseptic Technologies and Sterilization - Filling
		Aseptic Technologies and Sterilization - Filtration
		Aseptic Technologies and Sterilization - Mixing
		Aseptic Technologies and Sterilization - Other
		Bulk Packaging
		Freezing and Thawing
		Immunogenicity and Critical Quality Attributes

TRACK (SUBTRACK)	PRIMARY TOPIC	KEYWORDS
Manufacturing and Analytical Characterization - Chemical	Drug Product Manufacturing and Development	Liquids Manufacture - Oral and Topical Liquids
		Liquids Manufacture - Other
		Lyophilization and Drying Technologies
		Manufacturing and Assembly of Drug/Device Combinations
		Manufacturing of Aerosols and DPI
		Manufacturing of Drug Delivery Systems
		Other
		Primary Packaging - Blisters
		Primary Packaging - Bottles
		Primary Packaging - Container Closure Integrity
		Process Optimization
		Secondary Packaging
		Semi-solids Manufacture - Cremes
		Semi-solids Manufacture - Liposomes, Solid Lipid Nanoparticles
		Semi-solids Manufacture - Other
		Shipping Studies
		Solids Manufacture - Capsules
		Solids Manufacture - Drug Product Intermediates
		Solids Manufacture - Mini-tablets
		Solids Manufacture - Other
		Solids Manufacture - Powders
		Solids Manufacture - Tablets and Granules
		Storage Considerations
		Visual Inspection
Manufacturing and Analytical Characterization - Chemical	Drug Substance Manufacturing and Development	API Kilo Lab
		API Packaging and Storage
		Control of Impurity Formation
		Crystal Structure/Polymorph Screening
		Crystallization Development
		Filtration
		Genotoxic Impurities
		Immunogenicity and Critical Quality Attributes
		Milling and Micronization Technologies
		Other
		Particle Size Control
		Process Chromatography
		Process Optimization
		Purification
Manufacturing and Analytical Characterization - Chemical	General Aspects and Strategies	Change Control
		CMO Management
		Drug Master Files
		Drug Substance and Drug Product Shipment
		Electronic Records
		Handling Control Substances (DEA)
		Inspections and GMP
		Lean Manufacturing/Six Sigma/Operational Excellence
		Life Cycle Management
		Manufacturing Economics
		Materials Management and Warehousing
		Other
		Regulatory Strategy
		Supply Chain
Manufacturing and Analytical Characterization - Chemical	Generic Manufacturing	Other
		Patent Protection
		Pharmaceutical Equivalence Assessment
Manufacturing and Analytical Characterization - Chemical	Health, Safety, and Environment	Containment and Isolators
		Explosion Protection
		Green Chemistry
		High-Potent Drug Manufacturing
		OEL and PDE
		Other
		Solvent Recovery
Manufacturing and Analytical Characterization - Chemical	Innovative/Novel Processing Technologies and Concepts	For Use in Drug Product Manufacture
		For Use in Drug Substance Manufacture
		Other
Manufacturing and Analytical Characterization - Chemical	Integrated and Continuous Processing and Manufacturing	For Use in Drug Product Manufacture
		For Use in Drug Substance Manufacture
		Other
Manufacturing and Analytical Characterization - Chemical	Manufacture of Clinical Supplies	Blinding of Comparator Drugs
		Other
		Phase Appropriate GMP
Manufacturing and Analytical Characterization - Chemical	Plant Engineering, Equipment and Maintenance	Clean Media
		Facility Design
		Media
		Media Fills
		Modeling and Scheduling Multiproduct Batch Plants
		Modular Manufacturing
		Other
		Plant Incident Investigations
Manufacturing and Analytical Characterization - Chemical	Process Design and Controls	Cleaning Validation
		Control of Impurity Formation
		In-Process Controls
		Other
		Process Analytical Technology and Parametric/Real-Time
		Process Modeling and Simulations

TRACK (SUBTRACK)	PRIMARY TOPIC	KEYWORDS
Manufacturing and Analytical Characterization - Chemical	Process Design and Controls	Process Validation
		QbD and Assessment of Process Parameters
		Robustness/CPV Continuous Process Verification
		Scale-Up/Process Transfers
		Statistical Process Controls and Six Sigma
		Use of Prior Knowledge and Risk-Based Approaches
Manufacturing and Analytical Characterization - Chemical	Single-Use and Disposable Systems	For Use in Drug Product Manufacture
		For Use in Drug Substance Manufacture
		Leachables and Extractables
		Other
Preclinical, Clinical, and Translational Sciences	ADME	Biotransformation (In Vitro and In Vivo)
		High Throughput Assays
		Metabolite Identification and Characterization
		Metabolizing Enzymes Biotransformation (In Vitro and In Vivo)
		Novel Drug Modality/Novel Drug Delivery
		Optimization of Protein Design
		Other
		Payload-Linker Identification/Optimization
		Permeability, Distribution, and Site of Action
		Pharmacokinetics
		Screening Tools for Candidate Optimization
		Transporters Biotransformation (In Vitro and In Vivo)
Preclinical, Clinical, and Translational Sciences	Biology	Cell Therapy CAR-T, STEM Cell, Other
		Cellular and Molecular Pathways
		Gene Therapy
		In Silico
		In Vitro
		In Vivo
		Metabolizing Enzymes
		Omics
		Protein Binding
		Protein/Gene Engineering and Expression
		Receptor/Target Interactions
		Target Identification
		Transporter
Preclinical, Clinical, and Translational Sciences	Biomarkers	Clinical Qualification
		Disease Heterogeneity Assessments
		Endpoints (Surrogate , Primary, Secondary, Exploratory)
		High Content Data Analysis
		Novel Biomarkers
		Observational/Epidemiology Studies
		Patient Selection
Preclinical, Clinical, and Translational Sciences	Biostatistical Methodologies	Bayesian Methods
		Clinical Study Design
		Other
		Regulatory Recommendations
		Statistical Analysis Models
		Statistical Reporting
		Tools/Software
Preclinical, Clinical, and Translational Sciences	Clinical Trials	Designs and Methodology (Including QdB and RBM)
		Dosing Strategies
		Drug Interaction Studies
		Ethics in Clinical Trials
		Ex-US and Multi-national Studies
		Monitoring
		Other
		Patient Stratification
		Real World Evidence
		Regulatory Guidance
		Special Population Studies
Preclinical, Clinical, and Translational Sciences	Immunogenicity	Clinical Relevance
		Factor in Species Selection
		Immunogenicity Risk Assessment
		Impact on Exposure
		Impact on Safety Assessment
		Integrated Summary of Immunogenicity
		Post-Marketing Surveillance and REMS
Preclinical, Clinical, and Translational Sciences	Medicinal Chemistry	Combinatorial Chemistry
		Fragment Based Design
		In Silico Based Design
		Natural Products
		Novel Drug Modality
		Peptides and other Large Molecules
		Rational Drug Design
		Small Molecules
		Synthetic Chemistry
Preclinical, Clinical, and Translational Sciences	Modalities	ADCs
		Alternative Scaffold
		Bispecifics, Trispecifics and Higher Order Complexities
		CAR-T and Other Modified Cell Lines
		Cell-Based Therapy
		Encapsulated Drugs (Lipid, Nanoparticle, and Viral Vectors)
		Multispecific Antibodies
		Nanoparticle-Based
		Oligos, RNAs, and Locked Nucleic Acids

TRACK (SUBTRACK)	PRIMARY TOPIC	KEYWORDS
		Other
		PROTACs and Similar Molecular Glues
		Small Molecules
		Vaccines
Preclinical, Clinical, and Translational Sciences	Modeling and Simulation	Absorption Modeling
		Advanced Modeling Approaches
		Big Data and Artificial Intelligence
		Comparator Modeling
		Dose Project/Selection and Justification
		Drug interactions
		Imaging Based Approach
		In Vivo-In Vitro Correlation (IVIVC) Modeling
		Maternal/Fetal PK Model
		Other
		Other Special Populations
		Pediatric Model
		PK/PD Modeling
		Population PK Modeling
		Quantitative Systems Pharmacology (QSP)
		Resources, Tools and Software
		Standardization and Regulatory Advancement
		Translational Modeling
Preclinical, Clinical, and Translational Sciences	OMICS	Genomics
		Metabolomics
		Other
		Proteomics
Preclinical, Clinical, and Translational Sciences	Pharmaceutics	Drug Transport and Transporters
		Modeling and Simulation: New Approaches
		Molecular Biopharmaceutics
		Novel Drug Modality
		Pharmacokinetics
Preclinical, Clinical, and Translational Sciences	Pharmacology	Behavioral Pharmacology
		Cell Death and Differentiation
		DNA Damage and Repair
		Drug Abuse
		Drug-Drug Interactions
		Epigenetics and Epigenetic Therapy
		Experimental Therapeutics
		Gene Therapy
		Immunotherapy and Immunopharmacology
		Natural Products
		Neuropharmacology
		Omics (genomics, metabolomics, epigenomics, proteomics)
		Oncology (hematologic malignancies)
		Oncology (solid tumors)
		Orphan Drugs and Rare Diseases
		Pharmacokinetics
		Quantitative Pharmacology
		Signal Transduction
		Systems Pharmacology
		Toxicology
		Vaccines
Preclinical, Clinical, and Translational Sciences	Regulatory Guidance/Submissions	Clinical Study Reports (CSRs)
		Clinical Trial Protocols
		Data Standards and Management (Including CDISC)
		Harmonization and Guidance Documents
		Health Authority Interactions/Meetings
		Labeling
		Other
		Regulatory Filings and Submissions
		Safety
Preclinical, Clinical, and Translational Sciences	Safety	Cellular and Molecular Toxicology
		De-risking Strategies
		Genetic Toxicity
		Immuno-toxicity
		IND Enabling Studies
		Mechanistic Toxicity
		Other
		Screening Toxicity Studies
Preclinical, Clinical, and Translational Sciences	Translation	Drug-Drug Interaction (DDI)
		Formulation Selection
		Human PK and Dose Projections
		Other
		Patient Selection
		PK/PD
		Safety
Preclinical, Clinical, and Translational Sciences	Type of Human Studies	Adaptive and Accelerated Designs
		Bioequivalence
		Biosimilars
		Diseased Patient Population Studies
		Drug-Drug Interaction (DDI)
		First-Time-in-Human (FTIH)
		Food Effect
		Geriatric
		Organ Impairment

TRACK (SUBTRACK)	PRIMARY TOPIC	KEYWORDS
		Other
		Other Special Populations
		Pediatric
		Pregnant Women
		Radio-Labeled Mass Balance, Microdose and ADME
		Registrational Studies and Strategies
		Relative and Absolute Bioavailability
		Thorough QT/QTc (TQT)