



**Current as of June 22, 2026**

**Short talks selected from proffered abstracts**

**PR001 ELF3-driven epigenetic reprogramming creates ERK pathway dependency in SERD-resistant ER+ breast cancer.** Na Zhang. Dana-Farber Cancer Institute, Boston, MA, United States.

**PR002 Dark kinase dependency screening reveals STK17A as a previously unrecognized targetable vulnerability in EGFR-altered glioblastoma.** Eric Spinetti. STARX Therapeutics Inc, Miami, FL, United States.

**PR003 Targeting the “undruggable” oncogene CCNE1 using a molecular glue degrader in CCNE1 amplified cancers.** Nina Ilic-Widlund. Monte Rosa Therapeutics, Inc., Boston, MA, United States.

**PR004 enFoldX: AI classification of AlphaFold3-derived structural ensembles enables T cell specificity prediction.** Jonathan Levine. MSKCC, NEW YORK, NY, United States.

**PR005 JAB-BX600, a first-in-class EGFR-directed antioEdy drug conjugate delivering a novel KRAS G12D inhibitor.** Andrea Wang-Gillam. Jacobio (US) Pharmaceuticals, Burlington, MA, United States.

**PR006 Patient-level prediction of oncology clinical trial outcomes using a calibrated pan-cancer foundation model for synthetic control arms and indirect treatment comparisons.** Samantha Liang. Unlearn.AI, Inc, San Francisco, CA, United States.

**PR007 Discovery of a Next-Generation KAT6A Inhibitor with an Improved Therapeutic Index and Reduced Hematotoxicity Risk.** Davide D'Alia. Qubit Pharmaceuticals, Paris, France.

**PR008 VBC106: A First-in-class FOLR1/MSLN Targeted Tri-specific Antibody Drug Conjugate (ADC) in Ovarian Cancer, Endometrial Cancer, Lung Adenocarcinoma and Beyond.** Jing Li. VelaVigo (Shanghai) Limited, Shanghai, China.

**PR009 DLL3 CAR-T and CAR-NK cell combination therapy promotes synergistic antitumor activity in small cell lung cancer.** Alan Bers. Dana-Farber Cancer Institute, Boston, MA, United States.

**PR010 Targeted degradation of cyclin T1 constrains transcription elongation and triggers tumour cell apoptosis.** Janice Wenzheng Neng. Department of Pharmacology, University of Cambridge, Cambridge CB2 1PD, UK Gurdon Institute, University of Cambridge, UK, Cambridge, United Kingdom.

**PR011 A first-in-class Pin1 degrader provides robust PKPD response and efficacy in PDAC models.** Morgan O'Shea. Larkspur Biosciences, Cambridge, MA, United States.

**PR012 GLIO-1 is a selective DHODH inhibitor that is effective in IDH-mutant gliomas and KDM6-mutated cancers.** Diana Shi. Harvard Medical School, Boston, United States.



## AACR DRUG DISCOVERY AND DEVELOPMENT (AACR D3)

July 21-24, 2026 | Boston, MA

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**PR013 CDK4/6 bifunctional degraders overcome CDK4/6i resistance in breast cancer and demonstrate antitumor efficacy in additional tumor types.** Maechen Wang. Memorial Sloan Kettering Cancer Center, New York, NY, United States.



**Poster Session A (To be presented on July 22 from 6:15-8:45 p.m.)**

**A001 A Murine Translational Platform for ADC-Associated ILD Supports Rational Linker Optimization and Development of ARK-101, a Low-Toxicity HER2 ADC.** Suresh Anaganti. NJ Bio, Inc., Princeton, NJ, United States.

**A002 Paired PDX-PDXO Models Serve an Integrated Preclinical Platform for High-Throughput Payload Screening and ADC Drug Development.** Ludovic Bourre. Crown Bioscience, San Diego, CA, United States.

**A003 Novel ATR inhibitor antibody conjugates with potent single payload antitumor activity for personalized medicine.** Martin Duplessis. DCx Biotherapeutics, Saint Laurent, QC, Canada.

**A004 Precision degrader-antibody conjugates (pDACs) with a SMARCA2/4 dual payload as targeted therapeutics for acute myeloid leukemia.** Ryan Holmes. Prelude Therapeutics, Wilmington, DE, United States.

**A005 VBC106: A First-in-class FOLR1/MSLN Targeted Tri-specific Antibody Drug Conjugate (ADC) in Ovarian Cancer, Endometrial Cancer, Lung Adenocarcinoma and Beyond.** Jing Li. VelaVigo (Shanghai) Limited, Shanghai, China.

**A006 VBC104: A First-in-class CD79b/CD20 Targeted Bi-specific Antibody Drug Conjugate (ADC) with Superior Efficacy vs 1st line CD79b ADC in DLBCL, RT, FL, MCL, MZL and Beyond.** Jing Li. VelaVigo (Shanghai) Limited, Shanghai, China.

**A007 Sacituzumab Govitecan as a Potential Strategy for Sensitizing Cisplatin-Resistant HNSCC to Senolytic Treatment.** Natalie Luffman. Virginia Commonwealth University, Richmond, VA, United States.

**A008 ERBB2 activating mutations define a delivery-permissive state enhancing trastuzumab deruxtecan activity in HER2-non-amplified breast cancer.** Sharanya Nag. Memorial Sloan Kettering Cancer Center, New York, New York, NY, United States.

**A009 Preclinical safety and efficacy QP101: A novel HER2-targeting dual payload ADC combining topoisomerase I (TOP1) and cyclin dependent kinase 7 (CDK7) inhibitors.** Kiyeon Nam. Qurient Co. Ltd., Seongnam, Korea, Rep.

**A010 Post-translational modifications of transferrin receptor are novel therapeutic targets for Antibody-Drug Conjugates in Small Cell Lung Cancer.** Amanda Riley. Fred Hutch Cancer Center, Seattle, WA, United States.

**A011 Trastuzumab deruxtecan kills skin and uveal melanoma in vivo by HER2-independent payload release.** Elif Tanaydin. University of Gothenburg, Gothenburg, Sweden.



**A012 Preclinical Development of LGR4/5/6–FOLR1 Bispecific Antibody-Drug Conjugates for the Treatment of High-Grade Serous Ovarian Cancer.** Yukimatsu Toh. University of Texas, Houston, TX, United States.

**A013 Preclinical Development of a Humanized CD44v9-Directed Antibody-Drug Conjugate for Treatment-Refractory Breast Cancer.** Naoto Ueno. University of Hawai'i Cancer Center, Honolulu, HI, United States.

**A014 CAN020, a Trop2-targeting, dual-payload ADC from CanWell Pharma.** Shaoshan Wang. CanWell Pharma, Inc., Woburn, MA, United States.

**A015 JAB-BX600, a first-in-class EGFR-directed antitumor drug conjugate delivering a novel KRAS G12D inhibitor.** Andrea Wang-Gillam. Jacobio (US) Pharmaceuticals, Burlington, MA, United States.

**A016 High drug-to-antibody ratio (DAR) anti-TROP2 degrader-antibody conjugate (DAC) demonstrated poor solubility at high dose level leading to the fast clearance in mice.** Victor Yip. Genentech, South San Francisco, CA, United States.

**A017 A novel TROP2 targeting dual action bispecific tumor immunity enhancing ADC (TIE-ADC) that simultaneously eliminates cancer cells and immunosuppressor cells to overcome immunosuppression to enhance anti-tumor immune response in TNBC.** Shiva Bhowmik. TRIO Pharmaceuticals, Inc., San Diego, CA, United States.

**A018 A Novel CD33 and TRAIL-R2 Targeting Bispecific Antibody Sowing Selective Elimination of Leukemic Blasts With No Harm to Normal Myeloid and Other Immune Cells to Treat Relapsed/Refractory Acute Myeloid Leukemia.** Shiva Bhowmik. TRIO Pharmaceuticals, Inc., San Diego, CA, United States.

**A019 Targeted CNS Oncology Using Multispecific DNA-Based Therapeutics.** Emily Lin. Taipei Medical University and Hospital, Taipei, Taiwan.

**A020 Engineering developability of anti-CD3 scFv-based multispecific T-cell engagers.** Robert Pejchal. Adimab LLC, Lebanon, NH, United States.

**A021 Enhancing NK cell immunotherapy in non-small cell lung cancer using dual-antigen targeting Tri-Specific Killer Engagers (TriKEs/TriKE PACCs).** Rachel Steinmetz. University of Minnesota, Minneapolis, MN, United States.

**A022 Agentic AI-Driven Knowledge Graph Centrality with Reciprocal Rank Fusion for Interactive Discovery of Cancer Driver Genes and Pathways.** Dony Ang. Siloam Therapeutics, La Habra, CA, United States.

**A023 Using an agentic AI research platform for rare cancer drug discovery: Case of rhabdoid tumors.** Reed Bender. Lantern Pharma, Dallas, TX, United States.



**A024 FastBindRank, a novel, scalable method for high-fidelity virtual screening of ultra-large chemical libraries identifies novel HDAC11 inhibitors.** Jiawei Dai. Yale Cancer Center, New Haven, CT, United States.

**A025 Real-World Data–Driven Target Discovery in Advanced NSCLC.** Aaron Hardin. Guardant Health, Palo Alto, CA, United States.

**A026 In-silicoPhase III Clinical Trial of Avelumab Plus Axitinib Versus Sunitinib in Advanced Renal Cell Carcinoma Using Machine Learning Model Transfer Approach.** Elshad Hasanov. Division of Medical Oncology, Department of Internal Medicine, Pelotonia Institute for Immuno-Oncology, The Ohio State University Comprehensive Cancer Center, Columbus, OH, United States.

**A027 Artificial intelligence for detection of treatment-related adverse events using electronic health records in oncology: Focus on Sub-Saharan African (SSA) countries.** Elinda Kuhoga. Muhimbili University of Health and Allied Sciences, Dar-Es-Salaam, Tanzania.

**A028 enFoldX: AI classification of AlphaFold3-derived structural ensembles enables T cell specificity prediction.** Jonathan Levine. MSKCC, NEW YORK, NY, United States.

**A029 AI-Driven Network-Based Discovery on TCGA-PAAD Identifies KRAS-Axis Drug-Repurposing Targets for Pancreatic Cancer Interception.** Jianfu Li. Mayo Clinic, Jacksonville, FL, United States.

**A030 Patient-level prediction of oncology clinical trial outcomes using a calibrated pan-cancer foundation model for synthetic control arms and indirect treatment comparisons.** Samantha Liang. Unlearn.AI, Inc, San Francisco, CA, United States.

**A031 Large language models ensemble deciphers spatial proteogenomic landscapes to identify a novel trop2-cd47 co-targeting axis in non-small cell lung cancer.** Ellen McNeeley. University of Alabama at Birmingham, Birmingham, AL, United States.

**A032 Analysis of health records and genetics data to identify drug effects on cancer.** Rachel Melamed. UMass Lowell, Lowell, MA, United States.

**A033 Integratingcell-SELEX anddeeplearning method fordiscovery ofhigh-affinityaptamerstargetingcholangiocarcinoma.** Sourav Pal. Mayo Clinic, Jacksonville, FL, United States.

**A034 Repurposing adult clinical trials for pediatric target discovery using the CURE AI foundation model.** Neil Pfister. Numenos and the University of Alabama at Birmingham, Birmingham, AL, United States.

**A035 Diagnostic Accuracy of Virtual Programmed Death Ligand-1 Immunohistochemistry in Head and Neck Squamous Cell Carcinoma using Artificial Intelligence.** Charles Jeffrey Tan. Princess Margaret Cancer Center, Toronto, ON, Canada.



**A036 Smart Health Screening in Identification of Individuals at High Cancer Risk Using Artificial Intelligence.** Aida Yavari Kondori. Mashhad University of Medical Sciences, Mashhad, Iran, Islamic Rep.

**A037 Drug Development Using Machine Learning Approches: The Therapeutic Impact of Golnar on Inhibition of Tumor Growth in Colorectal Cancer.** Aida Yavari Kondori. Mashhad University of Medical Science, Mashhad, Iran, Islamic Rep.

**A038 Graph Neural Network Identification of Conserved Adaptive Resistance Circuits Across Glioblastoma Multiforme and KRAS-Mutant Lung Cancer.** Shivi Kumar. University of Pennsylvania, Dallas, TX, United States.

**A039 Physics-Informed Multi-Scale Modeling Identifies Redox–Membrane Stability Dependencies as a Therapeutic Vulnerability in Residual Glioblastoma Multiforme.** Shivi Kumar. University of Pennsylvania, Dallas, TX, United States.

**A040 Network Analysis and Deep Learning Identify Cascade Vulnerabilities in Glioblastoma Stem Cell Plasticity.** Darsh Dadhich. Hillsborough High School, Hillsborough, NJ, United States.

**A041 Glyco-Immune Checkpoint Activation in Inflammatory Breast Cancer Defines a Targetable Immune Evasion Program.** Ecem Kalemoglu. RUTGERS- Jersey City Medical Center, Jersey City, NJ, United States.

**A042 Identifying actionable dsRNA biomarkers from sense-antisense transcript pairs.** Otto Morris. Recursion Pharmaceuticals, London, United Kingdom.

**A043 The NF Target Hub: Connecting Transcriptomic Data to Targets in Neurofibromatosis.** Kara Quaid. The Children's Tumor Foundation, New York City, NY, United States.

**A044 Decoding the endometriosis immune–pain axis: A single-cell pipeline for neuroimmune hub discovery and drug prioritization.** Ariana Rahman. Mayo Clinic, Phoenix, AZ, United States.

**A045 In-silico based approach to characterize a group of small-molecule inhibitors of checkpoint CHK1, CHK2, WEE1 protein kinases for drug discovery.** Ashok Varma. Advanced Centre for Treatment, Research, and Education in Cancer, Navi Mumbai, Maharashtra, 410210, India Homi Bhabha National Institute, Anushaktinagar, Mumbai, Maharashtra, 400094, India, Navi Mumbai, India.

**A046 CLDN18.2 Expression in PDAC: KRAS Independence and Stage IV Prognostic Value.** Sacha El Khoury. The University of Texas MD Anderson Cancer Center, Houston, TX, United States.

**A047 Targeting mono-adv-ribosyltransferase 1 (ART1) suppresses tumor growth and oncogenic signaling in colorectal cancer.** Fnu Mimansa. Albert Einstein College of Medicine, Bronx, NY, United States.



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**A048 Oncolytic virotherapy as a novel therapeutic strategy in prostate cancer.** Ayodeji Ojetunde. Medserve-LUTH Cancer Centre, Lagos University Teaching Hospital, Lagos, Nigeria.

**A049 Rational grafting of p53TAD onto CTPR scaffolds generates potent p53-MDM2 antagonists.** Yawei Qi. University of Cambridge, Cambridge, United Kingdom.

**A050 Determining extracellular domain of EpCAM in the tumor niche for cancer diagnosis and prognosis of therapy response.** Han-Chung Wu. Institute of Cellular and Organismic Biology, Academia Sinica, Taiwan, Taiwan.

**A051 Optimization of Dual Combination [DOX][NIR] Nanomedicine: Effect of Size Tuning on Photophysical Behavior and Photothermal Efficiency for Enhanced Anti-Tumor Activity.** Fatima Abbas. University of Arkansas at Little Rock, little rock, AR, United States.

**A052 Sol-Moiety: A Generalizable Platform for Solubility-Driven Drug Delivery.** Arvin Karbasi. University of California, Los Angeles, Los Angeles, TX, United States.

**A053 Discovery and modular design of CDK12–DDB1 molecular glues for targeted Cyclin K degradation.** Andrii Monastyrskiy. Moffitt Cancer Center, Tampa, FL, United States.

**A054 Molecular glue discovery enabled by targeted degron display.** Zhe Zhuang. Stanford University, Stanford, CA, United States.

**A055 In situ intratumoral microdosing and spatial profiling enable rapid clinical evaluation of novel therapeutics and predict response in glioblastoma.** oliver jonas. Brigham and Women's Hospital, Boston, MA, United States.

**A056 Network-integrated clinical trial navigation with ctDNA profiling improves early-phase trial access: Final results from the PANNA-COTA study.** Anthony Joshua. Kinghorn Cancer Centre, St Vincents Hospital, Sydney, Australia.

**A057 Real-World Target Trial Emulation of FLAURA Trial Using InfinityAI: First-Line Osimertinib versus First-Generation EGFR Tyrosine Kinase Inhibitors in Advanced EGFR-Mutant NSCLC.** Abbas Shojaee. Guardant Health, Palo Alto, CA, United States.

**A058 Demographics, genomic testing, and JAK-inhibitor (JAKi) utilization in a novel, U.S. community oncology prospective registry for essential thrombocythemia (ET) and myelofibrosis (MF).** Christer Svedman. N-Power Medicine, Stockholm, Sweden.

**A059 Phase I study of the PKMYT1 inhibitor RP-6306 (lunresertib) in combination with carboplatin and paclitaxel for recurrent p53-abnormal ovarian and uterine cancer (GyneRep / OZUHN-019).** Laura Venegas. Princess Margaret Cancer Centre, Toronto, ON, Canada.



**A060 Targeting metabolic–DNA repair vulnerabilities in SDHB- and FH-deficient renal cell carcinoma with the novel alkylating agent KL-50.** Suparna Basu. Yale University, New Haven, CT, United States.

**A061 Application of CRISPR–Cas9 Technology in the Treatment of Chronic Lymphocytic Leukemia with Mutations P53 gene.** Manole Cojocar. Titu Maiorescu University of Bucharest, Faculty of Medicina, Bucharest, Romania.

**A062 Selective USP10 inhibition by GL-458 re-sensitizes platinum-resistant non-small cell lung cancer through DNA damage response disruption.** Sadaf Dorandish. WSU, Detroit, MI, United States.

**A063 Application of CRISPR–Cas9 Technology in the Treatment of Chronic Lymphocytic Leukemia with Mutations P53 gene.** Aurelian Udristioiu. Titu Maiorescu University of Bucharest, Faculty of Medical Assistance-Targu Jiu, Targu Jiu, Romania.

**A064 Application of CRISPR–Cas9 Technology in the Treatment of Chronic Lymphocytic Leukemia with Mutations P53 gene.** Aurelian Udristioiu. Titu Maiorescu University of Bucharest, Faculty of Medical Assistance-Targu Jiu, Targu Jiu, Romania.

**A065 The Role of miR-100 in lung cancer cell line: Relevance to cancer biology and potential therapy.** Basil Alotaibi. King Saud bin Abdulaziz University for Health Sciences (KSAU-HS), RIYADH, Saudi Arabia.

**A066 Methyl N-[5-(3'-iodobenzoyl)-1H-benzimidazol-2-yl]carbamate: A potent, safe, and theranostic approach to microtubule-targeted cancer therapy.** Janina Baranowska-Kortylewicz. University of Nebraska Medical Center, Omaha, NE, United States.

**A067 Combined Top1 and CHK Inhibitor Dual-Payload Strategy for Synergistic Reversal of Tumor Drug Resistance.** Tiejun Bing. ICE Biosciences InC., Beijing, China.

**A068 Multi-omics validated KRASi-resistant cell lines for combination and sequential drug treatment discovery.** Tiejun Bing. ICE Biosciences InC., Beijing, China.

**A069 The rational design of innovative dual COX-2/TP modulators as a multitarget strategy to stop cancer progression and metastasis.** Francesca Boccato. University of Turin, Turin, Italy.

**A070 Discovery of a Next-Generation KAT6A Inhibitor with an Improved Therapeutic Index and Reduced Hematotoxicity Risk.** Davide D'Alia. Qubit Pharmaceuticals, Paris, France.

**A071 Catalyzing TBXT as a therapeutic target through an integrated discovery pipeline.** Lee Dolat. Chordoma Foundation, Durham, NC, United States.

**A072 Structure-based drug design targeting DNA topoisomerase II for cancer therapy.** Xuesen Dong. University of British Columbia, Vancouver, BC, Canada.



**A073 ONM-421, a pH-responsive polymer-drug conjugate nanoparticle for the antigen-independent delivery of MMAE to solid tumors by exploiting the tumor acidic microenvironment.** Ruolan Han. OncoNano Medicine, Inc., Dallas, TX, United States.

**A074 Styrene maleic acid lipid particles (SMALPs) as chemotherapeutic nanocarriers: biophysical characterisation, cellular uptake, and intracellular drug delivery in cancer cells.** Jushta Jaiswal. University of Warwick, Coventry, United Kingdom.

**A075 The myokine irisin directly inhibits COX-2.** Wonchung Lim. Cheongju University, Cheongju, Korea, Rep.

**A076 Seeing the Drug and Seeing What It Is Doing: Integrated Spatial PK/PD Analysis for Hybrid and Next-Generation Oncology Therapeutics.** Mark Lim. AmberGen Inc., Billerica, MA, United States.

**A077 Multi-pathway inhibition and synergistic anticancer activity of novel biguanides in triple negative breast cancer models.** Julie Liu. UCLA, Los Angeles, CA, United States.

**A078 TP101: First-in-class cancer stem cell-targeted small molecule combination therapy for epithelial malignancies.** Cody McHale. Tract Bio, Charlotte, NC, United States.

**A079 Structure-guided discovery of first-in-class TRIB2 small-molecule inhibitors for therapy-resistant neuroendocrine prostate cancer.** JITENDER MONGA. Henry Ford Health System, Detroit, MI, United States.

**A080 Activation of hSSB2/1 by BCN077 Induces Tumor-Selective Mitotic Catastrophe While Protecting Irradiated Epithelium: A New Therapeutic Paradigm for Expanding the Cancer Treatment Window.** Andrew Norris. BCN Biosciences, Inglewood, CA, United States.

**A081 Multi-omics integration and network pharmacology identify shared transcriptomic signatures across breast cancer, hepatocellular carcinoma, and ovarian cancer: in silico discovery of a novel phytochemical lead.** Souhrid Sarkar. University of Glasgow, Glasgow, GA, United Kingdom.

**A082 AP3-guided biological SAR enables discovery of ACR-2316, a novel clinical-stage WEE1/PKMYT1 inhibitor designed for CDK1/2 and PLK1 pathway activation.** Lei Shi. Acrivon Therapeutics Inc., Watertown, MA, United States.

**A083 Integrated multi-omic analysis and CRISPR screening identify GPX2 as a critical vulnerability in high-risk lung adenocarcinoma.** Akul Singhania. Tempus AI, Inc., Chicago, IL, United States.

**A084 Cloperastine inhibits glycolysis and fatty acid metabolism that drives electron transport chain in tongue cancer.** Sanjeev Waghmare. Advanced Centre for Treatment Research and Education in Cancer, Navi Mumbai, India.



**A085 Opremorphin Induces Ferroptosis in Primary Human Glioblastoma Neurospheres.** Aaron Wasserman. Michigan State University College of Human Medicine, East Lansing, MI, United States.

**A086 AI-assisted identification and lab-based validation of novel hedgehog pathway inhibitors targeting the downstream GLI transcription effectors.** Kevin Williams. North Carolina Central University, Durham, NC, United States.

**A087 Calcitriol enhances chemotherapy response but fails to overcome established resistance in breast cancer models via VDR-dependent mechanisms.** Sirin Adham. Sultan Qaboos University, Muscat, Oman.

**A088 SLC39A6-targeted combination therapy to reverse CAF-induced resistance in HER2+ breast cancer using a microfluidic platform.** Zeina Habli. University of Pittsburgh, Pittsburgh, PA, United States.

**A089 ADT-1004: A mechanistically distinct pan-RAS inhibitor with the potential to escape resistance and mitigate on-target toxicities that limit efficacy and safety of other RAS inhibitors.** Gary Piazza. Auburn University, Auburn, AL, United States.

**A090 Therapy sequencing and capivasertib response in hormone receptor-positive, HER2-negative metastatic breast cancer (HR+/HER2- MBC).** Emanuelle Rizk. Beth Israel Deaconess Medical Center, Boston, MA, United States.

**A091 Overcoming Microenvironment-Mediated Chemoresistance in Acute Megakaryoblastic Leukemia through JAK Inhibition and 3D Niche Modeling.** Akira SHIMADA. Jichi Medical University, Shimotuke, Japan.

**A092 Early Prediction of Imatinib Resistance in Chronic Myeloid Leukemia: Development of an Ex Vivo Diagnostic Approach.** Nipanshi Tyagi. Jawaharlal Nehru University, New Delhi, India.

**Poster Session B (To be presented on July 23 from 6:15-8:45 p.m.)**

**B001 Preclinical evaluation of novel synthetic derivatives of tamoxifen in hormone receptor positive breast cancer growth and drug resistance.** SARAH CRAWFORD. Southern Connecticut State University, New Haven, CT, United States.

**B002 Dissecting cellular heterogeneity and altered cell states in castration resistant prostate cancer.** Julia Gattozzi. Stony Brook University, Stony Brook, NY, United States.

**B003 VEGFR3-AKT Signaling Drives Acquired Resistance to Darovasertib in GNAQ/GNA11-Mutant Uveal Melanoma.** Dongyu Han. University of Texas at El Paso, El Paso, TX, United States.



**B004 Analysis of proteomic changes in enzalutamide-resistant prostate cancer cells and identification of potential therapeutic targets for drug repurposing.** Chidi Molokwu. Bradford Institute of Health Research, Bradford, United Kingdom.

**B005 ADT-030, a novel PDE10 inhibitor that blocks RAS and  $\beta$ -catenin signaling with robust and durable antitumor activity and potential to mitigate resistance and on-target toxicities of monospecific RAS inhibitors.** Gary Piazza. Auburn University, Auburn, AL, United States.

**B006 Identification of possible biomarkers for acquired resistance against anti EGFR treatment regimen in a patient derived NSCLC model in vivo.** Julia Schueler. Charles River Discovery Germany GmbH, Freiburg, Germany.

**B007 ELF3-driven epigenetic reprogramming creates ERK pathway dependency in SERD-resistant ER+ breast cancer.** Na Zhang. Dana-Farber Cancer Institute, Boston, MA, United States.

**B008 The Sum of Its Parts: The Prevalence of Contribution of Effect through an Evaluation of Combination Clinical Trial Regimens of Immune Checkpoint Inhibitors and the VEGF Tyrosine Kinase Inhibitor Axitinib.** Jordan Weiss. Dana Farber Cancer Institute, Boston, MA, United States.

**B009 Machine Learning-Assisted Epigenetic Profiling Reveals SOX2 Promoter Methylation Associated with CDK1 Dysregulation and Therapeutic Targeting in Breast Cancer.** Bushra Khan. JAMIA MILLIA ISLAMIA, NEW DELHI, India.

**B010 Therapeutic targeting of PRC1 complexes to induce neuroblastoma differentiation.** Nathaniel Mabe. Purdue University, West Lafayette, IN, United States.

**B011 Single-molecule protein footprinting with Fiber-seq resolves coordinated chromatin states across regulatory domain.** Keith Maier. EpiCypher, Inc., Durham, NC, United States.

**B012 FFPE-CUT&Tag transcription profiling in patient biopsies reveals novel druggable pathways in lymphoma.** Morgan Oatley. EpiCypher, Inc., Durham, NC, United States.

**B013 KAT6A/B Inhibition Durably Reprograms Neuroblastoma Cell State to Enhance Retinoid Response and GD2-Directed Immunotherapy.** Nina Weichert-Leahey. Dana-Farber Cancer Institute, Boston, MA, United States.

**B014 Therapeutic Reversal of Tumor Cell Plasticity by EHF mRNA Delivery in Prostate Cancer.** Giuseppina Carbone. Institute of Oncology Research, Bellinzona, Switzerland.

**B015 RNA interference via siRNA-lipid conjugates is a safe and effective approach for MCL1-targeted therapy in triple-negative breast cancer.** Nina Cassidy. Vanderbilt University, Nashville, TN, United States.



**B016 Locally Injectable CASP Hydrogel for Sustained Co-Delivery of Temozolomide and an EGFR-Targeted Tandem Peptide siRNA System for Synergistic Glioblastoma Therapy.** Jordan Kinnitt. Clemson University, Clemson, SC, United States.

**B017 CLD-401, a Systemic Gene Medicine for In Situ IL-15 Superagonist Delivery: Driving NK and  $\gamma\delta$ T Cell-Mediated Tumor Killing.** Duong Nguyen. Calidi Biotherapeutics, San Diego, CA, United States.

**B018 CLD-501, an In Situ T Cell Engager Targeting TROP2: Reprogramming Solid Tumors to Generate Their Own Immunotherapy.** Eric Poma. Calidi Biotherapeutics, SAN DIEGO, CA, United States.

**B019 Application of CRISPR-Cas9 technology in the treatment of chronic lymphocytic leukemia with mutations P53 gene.** Aurelian Udristioiu. in Titu Maiorescu University of Bucharest, Faculty of Medical Assistance-Targu Jiu, Targu Jiu, Romania.

**B020 Immune, signal transduction, endocrine, cell growth and death, and neural signaling pathways are associated with the paclitaxel excipient polyoxyl 35 castor oil in PBMCs: implications for neurotoxic adverse side effects of chemotherapy.** Esther Chavez-Iglesias. University of California San Francisco, San Francisco, CA, United States.

**B021 A novel Fc $\gamma$ R+ cDC1 activation state drives therapeutic efficacy of TIGIT and PD-L1 co-blockade.** Ellen Duong. Genentech, South San Francisco, CA, United States.

**B022 Menin Inhibition and Leukemia Immunogenicity: A Novel Epigenetic Mechanism to Sensitize KMT2A-Rearranged Infant Leukemia to T-Cell-Mediated Cytotoxicity.** Aru Narendran. University of Calgary, Calgary, AB, Canada.

**B023 Mechanistic effects of Terfezia boudieri and its nanoformulation on cell cycle regulation and apoptosis in breast cancer models.** Roua Nouh. Biotechnology department, School of science and Engineering- The American University in Cairo, Cairo, Egypt.

**B024 Cisplatin primes an interferon-driven CD8<sup>+</sup> central memory T cell program to enable oncolytic virotherapy in gastrointestinal cancer malignancies.** Joseph Parker. Mayo Clinic Florida, Jacksonville, FL, United States.

**B025 Temporal Substate Flow and Multimodal Prediction Reveal Druggable Branch Points of EGFR TKI Resistance in NSCLC.** Michael Previte. Element Biosciences, San Diego, CA, United States.

**B026 PRISM mapping of the drug modality landscape: A systematic 900 cancer cell line reference of diverse therapeutic approaches.** Matthew Rees. Broad Institute of MIT & Harvard, Cambridge, MA, United States.

**B027 CRISPR screens identify thioredoxin reductase 1 as a target for inducing disulfidptosis in IDH1-mutant glioma models.** Dean Singleton. The University of Auckland, Auckland, New Zealand.



**B028 Identification of novel compounds that mediate transcriptional repression of estrogen receptor alpha gene ESR1 in breast cancer.** Jayalakshmi Sridhar. Xavier University of Louisiana, New Orleans, LA, United States.

**B029 Structures of human telomerase with BIBR1532 reveal novel mechanism of inhibition.** Yuqing Wang. UCLA, Los Angeles, CA, United States.

**B030 Fentanyl engages a GPCR-Neu1 signaling axis with divergent effects on pancreatic cancer cell behavior.** Abdulrahman Yaish. Queen's University, Kingston, ON, Canada.

**B031 MetAP2 inhibition by evexomostat (SDX-7320) decreases EZH2 and c-Myc and significantly prolongs survival in enzalutamide-resistant and neuroendocrine prostate cancer models.** Peter Cornelius. SynDevRx, Inc, Cambridge, MA, United States.

**B032 A High-Throughput Microfluidic Platform with Lab Automation and AI-Driven Morphological Profiling for Mechanism-of-Action Prediction of Anticancer Compounds.** Jenna McCormack. Xellar Biosystems, Boston, MA, United States.

**B033 Understanding the use and interpretation of duration of response endpoints in non-small cell lung cancer.** Brittany McKelvey. LUNgevity Foundation, Bethesda, MD, United States.

**B034 A case of myxoid variant of angiomatoid fibrous histiocytoma treated with tocilizumab before surgical resection.** Beatriz Mella Soares Pessoa. University of Connecticut, Farmington, CT, United States.

**B035 Restoring topoisomerase I inhibitor sensitivity in pediatric rhabdomyosarcoma using a pH-sensitive peptide-exatecan conjugate.** Shahyan Rehman. Yale School of Medicine, New Haven, CT, United States.

**B036 Allele Specific Characteristics of KRASQ61 Mutated Gastrointestinal Malignancies and Clinical Outcomes and Transcriptomic Features.** Sacha El Khoury. The University of Texas MD Anderson Cancer Center, Houston, TX, United States.

**B037 Convergent DNA damage and chromatin instability signatures as pharmacogenomic enrichment biomarkers for first-line Gemcitabine-Cisplatin ± Durvalumab in advanced biliary tract cancer: Beyond TMB/MSI.** Moonho Kim. University of Ulsan College of Medicine, Gangneung Asan Hospital, Gangneung, Korea, Rep.

**B038 Butyrylcholinesterase in uveal melanoma: Functional insights into a druggable prognostic biomarker.** Keyi Lyu. University of Nebraska Medical Center, Omaha, NE, United States.

**B039 Targeted Degradation of PD-L1 by Selective ER Translocation Inhibitors (SERTIs).** Thomas Bell. ASERTI Pharmaceutical, Reno, NV, United States.



**B040 Hold-to-Kill: RIPTAC Expanding Induced Proximity from Target Engagement to Therapeutic Window.** Tiejun Bing. ICE Biosciences Inc., Beijing, China.

**B041 PROTAC-mediated degradation of TACC3 sensitizes tumors to taxanes and induces apoptosis in FGFR3–TACC3 fusion cancers.** Eugen Dhimolea. Albert Einstein College of Medicine, New York City, NY, United States.

**B042 Clinical development and mechanism of action of LRK-4189, a first-in-class degrader of the lipid kinase PIP4K2C.** Krista Goodman. Larkspur Biosciences, Belmont, MA, United States.

**B043 Integrated organoid screening approaches enable preclinical assessment of protein degrader efficacy and target engagement.** Marten Hornsveld. Crown Bioscience, Leiden, Netherlands.

**B044 Targeting the “undruggable” oncogene CCNE1 using a molecular glue degrader in CCNE1 amplified cancers.** Nina Ilic-Widlund. Monte Rosa Therapeutics, Inc., Boston, MA, United States.

**B045 Targeted degradation of cyclin T1 constrains transcription elongation and triggers tumour cell apoptosis.** Janice Wenzheng Neng. Department of Pharmacology, University of Cambridge, Cambridge CB2 1PD, UK | Gurdon Institute, University of Cambridge, UK, Cambridge, United Kingdom.

**B046 A first-in-class Pin1 degrader provides robust PKPD response and efficacy in PDAC models.** Morgan O'Shea. Larkspur Biosciences, Cambridge, MA, United States.

**B047 dCBP-30: A rationally designed, orally bioavailable, near-linkerless PROTAC for potent CBP/p300 degradation.** Praveen Tiwari. Krantz Family Center for Cancer Research, Massachusetts General Hospital, Department of Medicine, Harvard Medical School, Boston, MA, United States.

**B048 CDK4/6 bifunctional degraders overcome CDK4/6i resistance in breast cancer and demonstrate antitumor efficacy in additional tumor types.** Maechen Wang. Memorial Sloan Kettering Cancer Center, New York, NY, United States.

**B049 Discovery of a small-molecule degrader targeting the intracellular immune checkpoint CISH.** Hao Xie. Mayo Clinic, Rochester, MN, United States.

**B050 Novel, conformational target discovery in TKI-resistant non-small cell lung cancer.** Faraz Choudhary. Immuto Scientific, Cambridge, MA, United States.

**B051 Targeted proteomics for parallel ADC biomarker screening: a pilot feasibility study within an Australian precision oncology program.** Nicholas McNamee. ProCan - Children's Medical Research Institute, University of Sydney, Sydney, NSW, Australia.

**B052 The Kinase Library: A global atlas of the human protein kinome and its applications in drug discovery.** Tomer Yaron-Barir. Dana-Farber Cancer Institute, Boston, MA, United States.



**B053 [18F]Fluoroglutamine PET radiotracer as a noninvasive biomarker of TP53-driven serine synthesis pathway activity in breast cancer.** Darshi Shah. Stony Brook Medicine, Stony Brook, NY, United States.

**B054 Alpha-Emitter vs. Beta-Emitter PSMA-Targeted Radioligand Therapy in Metastatic Castration-Resistant Prostate Cancer - A Systematic Review of Meta-Analyses.** Hina Zubair. Guthrie Robert Packer Hospital, Sayre, PA, United States.

**B055 Targeting LSD1 for anti-PD1 therapy resistance mechanism in Head and Neck Cancer for future therapy.** Manish Bais. Boston University, Boston, MA, United States.

**B056 DLL3 CAR-T and CAR-NK cell combination therapy promotes synergistic antitumor activity in small cell lung cancer.** Alan Bers. Dana-Farber Cancer Institute, Boston, MA, United States.

**B057 Evaluation of immune checkpoint inhibitors and targeted therapies ex vivo in live tumor fragments from core biopsies and resections accurately predicts response in the clinic.** Sean Caenepeel. Elephas Biosciences, Madison, WI, United States.

**B058 IMNN-001, a novel immune-oncology therapeutic delivering IL-12 to ovarian tumor and its microenvironment, facilitating immune reprogramming and survival benefit in the clinic.** Douglas Faller. IMUNON, Lawrenceville, NJ, United States.

**B059 Discovery of a “preparative survival checkpoint (signal-1 prime)” in B cells hijacked by cancer to confer chemotherapy resistance.** Jaewoong Lee. Korea university, Seoul, Korea, Rep.

**B060 Multimodal engineering of human mucosal-associated invariant T (MAIT) cells for enhanced cancer immunotherapy in liver cancer.** Yan-Ruide Li. University of California, Los Angeles, Los Angeles, CA, United States.

**B061 Harnessing stem cell-engineered CAR-NKT cells for universal low-toxicity autoimmune therapy.** Yan-Ruide Li. University of California, Los Angeles, Los Angeles, CA, United States.

**B062 Gene-transfer-resistant and biocontained synthetic organisms for discovery and controlled drug delivery.** Akos Nyerges. Harvard Medical School, Boston, MA, United States.

**B063 A gene-editable human regulatory T-cell platform reveals USP22 as a regulator of FOXP3 stability and IFN- $\gamma$  production.** Ajita Singh. Sai Life Sciences, Watertown, MA, United States.

**B064 Changes in tumormyeloid cell composition in patients with advanced pancreatic adenocarcinoma treated with lenvatinib plus pembrolizumab in second-line or later therapy.** Brandon Smaglo. UT MD Anderson Cancer Center, Houston, TX, United States.

**B065 RSK is a driver of immune suppression and tumor progression in basal-like breast cancer.** Ashley Spirrisson. Vanderbilt University, Nashville, TN, United States.



**B066 Human Tumor-on-Chip Platform for Mechanistic Investigation of Immune Exclusion and Immunotherapy Response in the Tumor Microenvironment.** Yu-Chieh Yuan. Xellar Biosystems, Boston, MA, United States.

**B067 Boosting immunogenic tumour cell death via nanotherapeutic targeting of the Stanniocalcin 1 phagocytosis checkpoint for enhanced cancer immunotherapy.** Jianqin Lu. Dept. of Pharmacology/Toxicology, R. Ken Coit College of Pharmacy The University of Arizona, TUCSON, AZ, United States.

**B068 A sphingolipid-derived paclitaxel nanovesicle enhances efficacy of combination therapies in triple-negative breast cancer and pancreatic cancer.** Jianqin Lu. Dept. of Pharmacology/Toxicology, R. Ken Coit College of Pharmacy The University of Arizona, TUCSON, AZ, United States.

**B069 Pharmacodynamic evidence of Wnt/ $\beta$ -catenin inhibition by tegavivint, a TBL1-directed transcriptional modulator, in advanced hepatocellular carcinoma: a phase 1 study.** Aundrietta Duncan. Iterion Therapeutics, Houston, TX, United States.

**B070 ATX-295, a Novel KIF18A Inhibitor, as a Therapeutic in Chromosomally Instable Tumors.** Laura Ghisolfi. Accent Therapeutics, Lexington, MA, United States.

**B071 TR $\beta$  Agonists as a First-in-Class Oncology Therapeutic Strategy: Broad Anti-Tumor Activity with Proof-of-Concept in Breast Cancer.** Noelle Gillis. University of Vermont, Burlington, VT, United States.

**B072 SHY-ONC6, a first-in-class oral inhibitor of the proteasome 19S regulatory particle ATPases, drives tumor regression across solid and hematological xenograft models.** Yaron Hadari. SHY Therapeutics, LLC, Valhalla, NY, United States.

**B073 Rationale and Development of First-in-class Lysosome Disrupting Agent for Solid Tumours.** Anthony Joshua. Princess Margaret Cancer Centre, Toronto, ON, Canada.

**B074 Evaluation of a small molecule, AP-232, as a splicing factor inhibitor of U2 auxiliary factor 1 (U2AF1) in leukemia cells.** Mona Kazemi Sabzvar. The University of Tennessee Health Science Center, Memphis, TN, United States.

**B075 Selective targeting of inosine monophosphate dehydrogenase-2 impairs brain metastatic potential while preserving immune cell function.** Agata Kieliszek. McMaster University, Hamilton, ON, Canada.

**B076 Novel artesunate trimer derivatives as antitumor therapies for hepatocellular carcinoma.** Xiaohua Kong. The University of Texas at El Paso, EL PASO, TX, United States.



**B077 EMI-1725 Demonstrates Broad Antagonist Activity Against Drug-Resistant Androgen Receptor Mutations and Tumor Growth Suppression in Preclinical Models of Metastatic Castration-Resistant Prostate Cancer.** Steven Kregel. Loyola University Chicago, Maywood, IL, United States.

**B078 A brain-penetrant tumor-selective EGFR inhibitor for glioblastoma.** Sarah Lee. UCLA, Los Angeles, CA, United States.

**B079 Small-molecule inhibition of CFBF::MYH11 in chemoresistant inv(16) acute myeloid leukemia.** Thuy-An Nguyen. University of Virginia, Charlottesville, VA, United States.

**B080 Targeting tumor metabolism in breast cancer using LDHA inhibition to enhance chemotherapy response.** Khushboo Pathania. UIPS, Panjab Univeristy and University of Nottingham, Chandigarh and Nottingham, UK, India.

**B081 Lysyl oxidase inhibitor LXG6403 creates ferroptosis vulnerability in triple-negative breast cancer.** Ozge Saatci. Medical University of South Carolina, Charleston, United States.

**B082 Lysyl oxidase inhibitor LXG6403 effectively suppresses tumor progression and overcomes therapeutic resistance in clear cell renal cell carcinoma.** Ozgur Sahin. Medical University of South Carolina, Charleston, SC, United States.

**B083 GLIO-1 is a selective DHODH inhibitor that is effective in IDH-mutant gliomas and KDM6-mutated cancers.** Diana Shi. Harvard Medical School, Boston, United States.

**B084 Dark kinase dependency screening reveals STK17A as a previously unrecognized targetable vulnerability in EGFR-altered glioblastoma.** Eric Spinetti. STARX Therapeutics Inc, Miami, FL, United States.

**B085 Selective oral STK17A inhibitor UMF814A targets a previously unrecognized dark kinase vulnerability in spliceosome-mutant MDS and AML.** Eric Spinetti. STARX Therapeutics Inc, Miami, FL, United States.

**B086 Aspartyl/asparaginyl beta-hydroxylase (ASPH) blockade abrogates doxorubicin induced chemoresistance in chondrosarcoma.** Richard Terek. Brown University, Providence, RI, United States.

**B087 Dual MET/AXL inhibition suppresses hepatocellular carcinoma through tumour cell-intrinsic and microenvironment-dependent mechanisms.** Chenjing Zhu. Jiangsu Cancer Hospital, Nanjing, China.

**B088 A cryptic alternative bound state of HDAC6 drives mutation- and ligand-dependent remodeling of inhibitor interaction.** Paul Fong. Duke University, Durham, NC, United States.

**B089 Targeting metabolic vulnerabilities in cancer through structure-guided discovery of novel malic enzyme inhibitors.** Ben Krinkel. University of Auckland, New Zealand, New Zealand.



**B090 Label-free Raman molecular histology enables patient-tissue metabolic stratification for target selection in lung adenocarcinoma.** Katherine Zhao. SuperVision Medicine Co.,Ltd; Columbia University, Wuxi, China.

**B091 Synthetic Lethality of G6PD Deficiency and Asparaginase for Colorectal Cancer Therapy.** Alejandro Gutierrez. St. Jude Children's Research Hospital, Memphis, TN, United States.

**B092 MBD4 LOF defines a synthetic lethal therapeutic state targetable by ATR inhibition rather than PARP in high-grade serous ovarian cancer.** Fahad Kiani. CrisPRO.ai, Brooklyn, NY, United States.

**B093 Combinatorial CRISPR screening reveals synthetic lethal and suppressor interactions in RTK signaling and glycosylation networks.** Subin Kim. The University of Texas MD Anderson Cancer Center, Houston, TX, United States.

**B094 Suppressing ERG-driven oncogenesis by mRNA-based delivery of an intracellular mini-body targeting a unique protein methylation site in ERG fusion-positive prostate cancer.** Carlo Catapano. Institute of Oncology Research, Bellinzona, Switzerland.

**B095 Targeted inhibition of SLC2A5 prevents tumour growth and metastasis.** Jody Groenendyk. University of Alberta, Edmonton, AB, Canada.

**B096 Targeting NONO as a therapeutic strategy for metastatic castration-resistant prostate cancer (mCRPC).** Gaelle Mercenne. Talus Bio, Seattle, WA, United States.

**B097 Microfluidics-enabled asymmetric lipid vesicles for targeted delivery of RNA-based cancer vaccines in colorectal cancer.** Sabiruddin Mirza. School of Engineering and Applied Science, Harvard University, Cambridge, United States.

**B098 Ad5.F35-GUCY2C-PADRE vaccine safely induces dose-dependent immune responses associated with reduced recurrences in patients with GUCY2C-expressing GI cancers at high risk of relapse.** Adam Snook. Thomas Jefferson University, Philadelphia, PA, United States.