



Poster Session A

Thursday, May 9

12:30-2:30 pm

- A01 YAP1 drives ependymoma-like tumour formation in the brain.** Noreen Eder, The Francis Crick Institute, London, United Kingdom.
- A02 YAP1 opposes differentiation in mesenchymal tumors.** T.S. Karin Eisinger-Mathason, University of Pennsylvania, Philadelphia, PA.
- A03 Generation of primary sarcoma mouse models through CRISPR/Cas9 mediated activation of Yap1.** Jianguo Huang, Duke University, Durham, NC.
- A04 Genetic and pharmacologic inhibition of HES1 reduces YAP1 expression, impairing rhabdomyosarcoma cell growth.** Alexander Kovach, Duke University, Durham, NC.
- A05 RAS signaling promotes ERMS cell viability via sustaining TAZ expression and protein stability.** Liz (Yi-Tzu) Lin, Duke University Medical Center, Durham, NC.
- A06, PR03 YAP/TAZ requirement in mesenchyme-originated intestinal hamartomatous polyposis.** Junhao Mao, University of Massachusetts Medical School, Worcester, MA.
- A07 Role of YAP/TEAD and YAP/Smad signaling pathways in osteosarcoma tumour growth and lung metastasis dissemination.** Sarah Morice, INSERM, Nantes, France.
- A08 Loss of non-canonical Hippo signaling in fusion-positive alveolar rhabdomyosarcoma increases invasiveness and a dedifferentiated phenotype associated with metastasis.** Kristianne Oristian, Duke University Medical Center, Durham, NC.
- A09 Targeting Hippo-dependent and Hippo-independent regulation of the YAP1 oncoprotein in childhood rhabdomyosarcoma.** Katherine Slemmons, Children's Hospital Los Angeles, Los Angeles, CA.
- A10 Taz regulates aging of hematopoietic stem cells.** Anna Mura-Meszaros, Leibniz Institute on Aging, Jena, Germany.
- A11 Reawakening the regenerative potential of mammalian Müller glial cells to restore sight.** Ross Poche, Baylor College of Medicine, Houston, TX.

- A12 YAP and cancer stem cells in basal-like breast cancer.** Hazel Quinn, MDC Berlin, Berlin, Germany.
- A13 Yap activity in bile ducts, but not in hepatocytes, is required for normal liver regeneration.** Elisabeth Verboven, VIB - KULeuven, Leuven, Belgium.
- A14, PR10 Hippo Regulates Intestinal Regeneration By Inducing Revival Stem Cells.** Jeff Wrana, Lunenfeld-Tanenbaum Research Institute, Toronto, Ontario, Canada.
- A15 A balance of yki/sd activator and e2f1/sd repressor complexes controls cell survival and affects organ size.** Peng Zhang, Huntsman Cancer Institute, University of Utah, Salt Lake City, Utah.
- A16 The α -Arrestin ARRDC3 functions as a metastasis suppressor by regulating GPCR activation of the Hippo pathway.** Aleena Arakaki, UC San Diego, San Diego, CA.
- A17 The Tyrosine Phosphatase SHP2 regulates YAPY357 Phosphorylation, Sub-cellular Localization, and Transcriptional Co-Activity in Cholangiocarcinoma.** EeLN Buckarma, Mayo Clinic, Rochester, MN.
- A18 Location, location, location: Avenues to regulating Hippo.** Philamer C Calses, Genentech Inc., South San Francisco, CA.
- A19 Regulation of glioblastoma tumor growth and stem cell properties through G α ;12 and tissue factor, upstream and downstream players in YAP signaling.** Olga Chaim, UCSD, La Jolla, California.
- A20 G α q controls the Hippo Pathway through MOB1 tyrosine phosphorylation by FAK.** Xiaodong Feng, Moores Cancer Center, University of California, San Diego, La Jolla, California.
- A21, PR02 Spatial resets modulate YAP-dependent transcription.** Matt Franklin, Stanford University, Stanford, CA.
- A22, PR04 Integrin-mediated mechano-transduction controls HER2 oncogenic signaling and activation of YAP in breast cancer.** Filippo Giancotti, UT MD Anderson Cancer Center, Houston, TX.
- A23 Verteporfin as a new treatment paradigm for platinum-resistant ovarian cancer cells.** Radhika Gogoi, Geisinger Clinic, Danville, PA.
- A24 Functional annotation of the Hippo somatic mutations in human cancer.** Han Han, Department of Development and Cell Biology, University of California, Irvine, Irvine, CA.
- A25, PR06 Mechanistic Insights for TEAD/YAP Activation.** Jeffrey Holden, Genentech, South San Francisco, CA.

- A26 Classification of glioblastoma tumorsphere depending on the regulatory mechanisms of the Hippo pathway.** Seok-Gu Kang, Departments of Neurosurgery, Brain Tumor Center, Severance Hospital, Yonsei University College of Medicine, Seoul, Seoul, Korea.
- A27 Increasing proximity triggers Mst2 autophosphorylation.** Jennifer Kavran, Johns Hopkins School of Public Health, Baltimore, MD.
- A28 DEAD-box RNA helicase DP103 enhances YAP sumoylation for YAP-TEAD dependence and statin sensitivity in triple negative breast cancer.** Xianning Lai, Cancer Science Institute of Singapore, National University of Singapore, Singapore, Singapore.
- A29 Genome-wide CRISPR/Cas9 screens for the identification of novel YAP1/TAZ modulators.** Martin Lange, Bayer AG, Research & Development, Pharmaceuticals Division, Berlin, Germany.
- A30 Super-enhancer-associated Long Noncoding RNA UCA1 Interacts Directly with AMOT to Inhibit Hippo Signaling Pathway in Epithelial Ovarian Cancer.** Xianzhi Lin, Women's Cancer Program at the Samuel Oschin Comprehensive Cancer Institute, Cedars-Sinai Medical Center, Los Angeles, CA.
- A31 Transcriptional addiction to YAP1 -a major driving force of oral cancer carcinogenesis and evolution ?** Muneyuki Masuda, Department of Head and Neck Surgery, National Kyushu Cancer Center, Fukuoka, Fukuoka, Japan.
- A32 The small GTPase Rac1 controls the stability of Yes-Associated Protein (YAP) independently of the LATS1/2 kinases.** Chitra Palanivel, University of Nebraska Medical Center, Omaha, NE.
- A33, PR01 Regulation of TEAD by p38 MAPK-induced cytoplasmic translocation.** Hyun Woo Park, Yonsei University, Seoul, South Korea.
- A34 Identification of a MAP kinase that regulates YAP abundance.** Sanghyun Park, KAIST, Daejeon, Republic of Korea.
- A35 Title: Regulation of the Hippo signaling pathway through ubiquitin-mediated degradation of TEAD transcription factors.** Trang Pham, Genentech, South San Francisco, CA.
- A36 Paracrine orchestration of intestinal tumorigenesis at the mesenchymal-epithelial interface.** Manolis Roulis, Yale School of Medicine, New Haven, CT.
- A37 Implication of targeting YAP1 in KRAS-mutant lung cancer cells.** Iwao Shimomura, National Cancer Center Research Institute, Tokyo, Japan.
- A38 A 4-gene YAP-related pathway expression signature informs about dependence of tumors on Hippo pathway signaling.** Dirk Wienke, Merck KGaA, Biopharma, R&D, Darmstadt, Germany.

A39, PR09 **The Hippo pathway integrates PI3K-Akt signals with mechanical cues to control tissue growth.** Barry Thompson, Francis Crick Institute, London, England, United Kingdom.

A40 **Hippo signaling in cancer development.** Wenqi Wang, University of California, Irvine, Irvine, CA.

A41 **Inhibition of aberrant YAP and TAZ activity to prevent metastasis formation and growth.**
Janine Warren, Albany Medical College, Albany, NY.

A43 **Identification of novel YAP/TAZ regulators in metastatic cancer.** Yuxuan Xiao, Albany Medical College, Albany, NY.

A44 **PR55 α regulatory subunit of PP2A inhibits the MOB1/LATS cascade and activates YAP in pancreatic cancer cells.** Ying Yan, University of Nebraska Medical Center, Omaha, NE.