

The 19th JFCR-ISCC

New Antitumor Agents under Development in the US, Europe and Japan



2014

December 10, (Wed)
13:00 - 17:40

December 11, (Thu)
9:00 - 17:35

Miraikan

National Museum of
Emerging Science and Innovation
2-3-6, Aomi, Koto-ku, Tokyo, Japan

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<http://iscc.umin.jp/>

Dec. 10

**New Drugs and
Clinical Design for
Rare Types of
Lung Cancers**

13:10-

**DNA Repair Pathways
as Targets for
Cancer Therapy**

15:10 -

Poster Session

16:40 -

Dec.11

**Cancer Immuno-
therapy Using
Checkpoint Inhibitor
Antibodies and
CAR-modified T Cells**

9:00 -

Poster Discussion Session 11:15-

Luncheon Seminar

12:25-

**Meet-the-Expert
Seminar** 13:15-

**Targeting Epigenetics
and Micro-
environment for
Cancer Therapy**

14:25-

**New Drugs and
Combination Therapy
under Clinical Trials**

15:25-

Jeffrey A. Engelman *MGH Cancer Center*
Resistance mechanisms to kinase inhibitors

Ryohei Katayama *Cancer Chemotherapy Center, JFCR*
New mechanisms of acquired resistance to ALK-inhibitors

Koichi Goto *National Cancer Center Hospital East*
Development of a nationwide genomic screening network
and clinical trial for lung cancer with rare driver mutations

Yves G. Pommier *NCI, NIH*
Novel topoisomerase I inhibitors on the horizon and
topoisomerase repair

Johann de Bono *Institute of Cancer Research*
Targeting PARP: What have we learnt?

Koji Tamada *Yamaguchi Univ.*
Novel strategies of chimeric antigen receptor T cells for
cancer immunotherapy

Jean Viallet *Bristol-Myers Squibb*
Emerging principles of combination immunotherapy for
cancer: Focus on anti-CS1/SLAMF7 Elotuzumab and
anti-CD137 Urelumab

Gregg Fine *Genentech inc.*
MPDL3280A: an engineered anti-PDL1 antibody specifically
recognize tumor immune system and destroy tumor cells

Masami Suzuki *Chugai Pharmaceutical Co., Ltd.*
Establishment of human colon cancer stem cell lines and
drug discovery targeting cancer stem cells

Bruce A. Chabner *MGH Cancer Center*
Rapid approval of targeted cancer drugs: Lessons from
the first decade

Kenichi Nomoto *Eisai Inc.*
Selective inhibition of EZH2 by EPZ-6438 (E7438) as
personalized cancer therapy

Morihiro Watanabe *Merck Serono Ltd.*
TH-302, a hypoxia-activated prodrug with potential
antineoplastic activity

Keith T. Flaherty *MGH Cancer Center*
Learning from selective pressure of MAP kinase pathway
inhibition in BRAF and RAS mutant cancer to develop
rational combinations

Sylvie Assadourian *Sanofi S.A.*
c-Met inhibitor SAR125844

Nobuya Ishii *Chugai Pharmaceutical Co., Ltd.*
Debio1347/CH5183284: a novel selective FGFR inhibitor