Publication of a paper describing the possible effect of a MELK inhibitor,

OTS167, on small cell lung cancer

Important biological roles of a protein kinase MELK (Maternal Embryonic

Leucine-zipper Kinase), on which our small molecule inhibitor OTS167 targets, were

reported by a collaboration between Prof. Yusuke Nakamura's group in The University

of Chicago and our research development group in OncoTharapy Science (OTS).

MELK is known to be highly expressed in various types of cancer and play critical roles

in maintenance of cancer stem cells. OTS is currently conducting clinical trials of

OTS167 in the United States and Australia. In this study, the collaborative group

demonstrated that MELK was overexpressed in the majority of small cell lung cancer

(SCLC) cell lines and primary tumors, and that either knockdown of MELK or

treatment with OTS167 exhibited strong growth inhibitory effect on those SCLC cell

lines.

SCLC, a biologically different type of lung cancer from non-small cell lung cancer

(NSCLC), comprises approximately 15% of all lung cancers. SCLC usually exhibits

aggressive behavior, rapid growth, and early spread to distant sites. Most of the SCLC

cases are diagnosed at an advanced stage for which surgical resection is not applicable.

The five-year survival rate for SCLC is as low as 10% and little improvement was

observed in the last three decades.

Based on the results in this manuscript, we plan further clinical development of

OTS167 in SCLC.

The paper has been published online in the journal *Oncotarget*.

h%5B%5D=7297)