# The 17th International Symposium on Molecular and Neural Mechanisms of Taste and Olfactory Perception (ISMNTOP/YRUF/AISCRIB 2018)

in conjunction with

Special Events celebrating the 50<sup>th</sup> Anniversary of

Monell Chemical Senses Center

#### 第17回国際シンポジウム"味覚嗅覚の分子神経機構"

(うま味若手フォーラム/アジア国際シンポジウム"化学受容と摂食行動" 2018)/米国モネル化学感覚センター50周年記念イベント

#### Organizer: Yuzo Ninomiva

Research and Development Center for Five-Sense Devices
Kyushu University

#### PROGRAM & ABSTRACTS

November 30-December 2, 2018

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Research (Maidashi campus), Fukuoka, Japan

#### Hosted by

Kyushu University, Research and Development Center for Five-Sense Devices

九州大学・五感応用デバイス研究開発センター

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# Malfuction through single point mutation in an odorant receptor of the major disease vector, *Aedes aegypti*

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The Aedes aegypti (Ae. aegypti) is a primary vector for dengue virus, the most widespread vector-borne viral disease. Ae. aegypti also acts as a vector of yellow fever, chikungunya and zika virus. Although odorant receptors of Ae. aegypti (AaOrs) has been identified 131 including co-receptor (AaOrco) by bioinformatics analysis, there is a lack of studies about function and structure. We have previously reported that AaOr8 and AaOr49 in the mouth part of mosquito play a major role in mosquito blood-feeding behavior by as detecting the volatile compounds in blood. In this study, we found the ligand binding site of AaOr8 and AaOr49 using SYBYL-X 2.1 packages and homology protein model by the computer program. For demonstrated ligand binding site, we also performed calcium imaging using Sf9 cells with site-directed point mutation AaOrs. AaOr8-S167A in which serine residue in position 167 amino acid was substituted to non-polar and hydrophobic amino acid, alanine, represented a loss of binding affinity to 1-octen-3-ol and cyclohexanol, while AaOr49-V168M where valine in the position 168 amino acid residue was replaced with methionine and AaOr49-Y190A where tyrosine of the position 190 amino acid was substituted to alanine showed complete loss of binding affinity to 2-ethyl-1-hexanol with intact responses to cyclohexanone and benzyl alcohol. Taken togethr, We identified major active residues of AaOr8 and AaOr49 using both in silico and in vitro approach, which should be useful in future research about odorant receptors and control of disease vector insects.