

**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**

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# **The negative effect of High Fat Diet on the olfactory system in *Drosophila melanogaster***

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High-fat diet (HFD) often causes obesity, heart disease, cancer and it has detrimental effects on the sensory system. In particular, sensory-mediated responses are crucial for maintaining energy balance, as they are involved in a metabolic regulation; however, little is known about the impact of HFD-induced stress on sensory system. We have used a *Drosophila melanogaster* model to gain insight on how HFD-induced stress affects physiological, behavioral, and transcriptional changes related with olfactory and nutrient-related signaling. We demonstrated that lifespan and locomotor activity in HFD-treated flies decreased and that olfactory sensitivity and behavioral responses to odorants were changed. Olfactory sensitivity to eight of ten odorants after 14 days on HFD treatment were reduced, while behavioral attraction was increased to benzaldehyde and 1-hexanol in flies that were treated with HFD for 7 days. This behavioral and physiological modification by HFD treatment was accompanied by a significant decrease in DmOrco gene expression in a peripheral olfactory organ, suggesting that it could be involved in the action of metabolic and sensory signal. Gene expression profiles of antennae showed significant differences on the olfactory receptors, odorant-binding proteins, and insulin signaling. Our results suggested that olfactory sensitivity and behavioral responses to HFD-induced stress are mediated through olfactory and nutrient-related signaling pathways.