# A whole-body approach to understanding chemosensory cells

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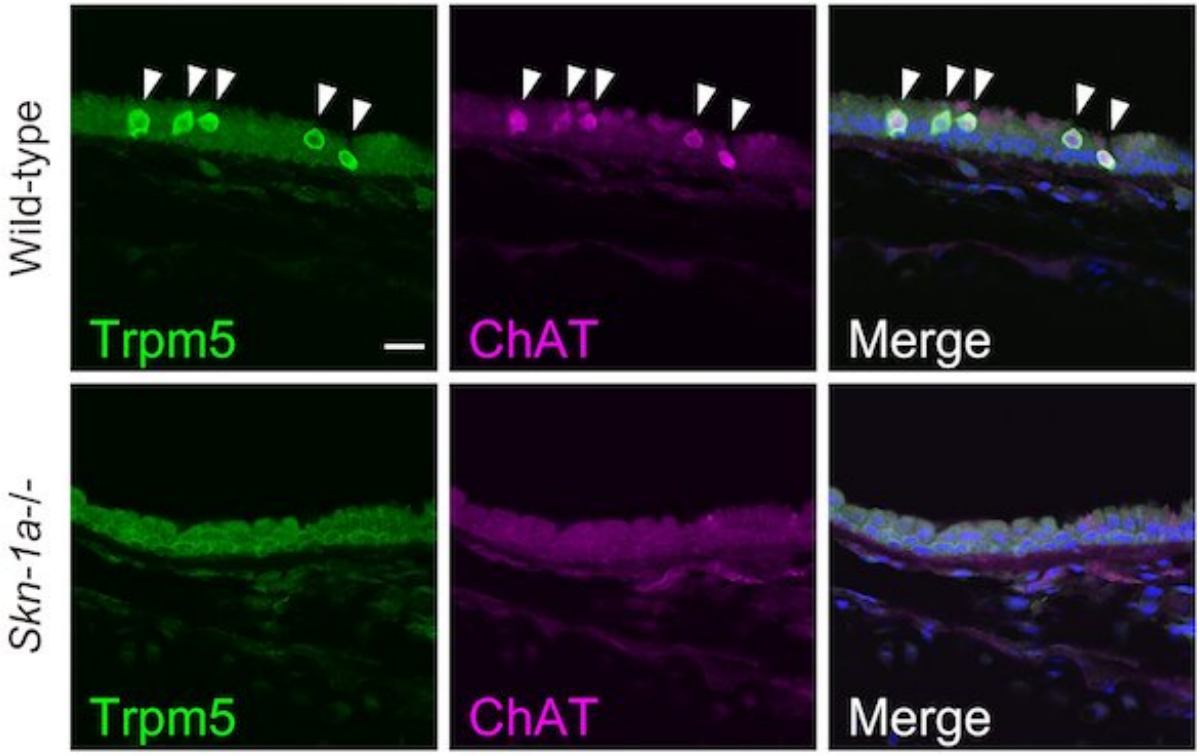


Immunostaining of Trpm5 and choline acetyltransferase (ChAT) on coronal sections of the trachea of wild-type and Skn-1a-deficient mice. The key point is that compared to the wild-type, no signals for Trpm5 and ChAT were observed in the Skn-1a-deficient mice. Thus, Skn-1a is essential for the functional differentiation of Trpm5-positive tracheal brush cells. Credit: Junji Hirota

Growing evidence shows that sensory cells which enable us to taste sweetness, bitterness and savoriness (umami) are not limited to the tongue. These so-called Trpm5-expressing chemosensory cells are also found in the respiratory system, digestive tract and other parts of the body.

Although their precise function in areas other than the mouth are not fully known, these sensory cells are thought to play an important "gatekeeper" role, protecting the body against bacteria and potentially harmful substances.

Now, researchers have found that a protein called Skn-1a behaves as a master regulator for the generation of these cells across multiple tissues and organs.

"Based on our previous studies, we knew that Skn-1a plays an essential role in generating these cells, for example, in the nose," says Junji Hirota, associate professor at the Center for Biological Resources and Informatics, Tokyo Institute of Technology (Tokyo Tech).

In the new study published in *PLOS ONE*, the researchers comprehensively analyzed multiple tissues using knockout mice and bio-imaging techniques. "One by one, we found that without Skn-1a, the sensory cells were not generated," says Hirota. "All of our results indicated that Skn-1a is a master regulator for the generation of these cells throughout the body."

The study arose from a collaboration between two teams—one led by Hirota, a specialist in olfactory systems, and the other by Ichiro Matsumoto, an expert on taste receptors at the Monell Chemical Senses Center in Philadelphia.

Hirota says: "Our collaboration is very fruitful—by working together, we

can extend our knowledge beyond the nose and tongue to the whole body." Following Matsumoto's original discovery of Skn-1a, published in *Nature Neuroscience* in 2011, the two teams found that Skn-1a is vital for generating chemosensory cells in the nasal respiratory epithelium (in 2013) and the main olfactory epithelium (in 2014).

The latest study goes further by revealing that Skn-1a controls the generation of chemosensory cells in the trachea (see Figure 1), auditory tube, urethra, thymus, pancreatic duct, stomach, and large intestine.

Many questions remain about why these cells are found in such a wide range of organs.

"For example, in the trachea, we think there may be at least two or three types of chemosensory cells," Hirota says. "We're interested in their characterization—this would contribute to fundamental knowledge of biological systems."

The thymus is particularly intriguing, says Hirota, as it is different to the respiratory and digestive systems, and could lead to new research directions in immunology.

In the urethra, chemosensory cells may help protect the body against infections, for example by sending signals to release more urine, thus ridding the body of potentially dangerous bacteria or toxins.

"If we can identify the receptor types expressed by these chemosensory cells, we can enhance our understanding of how they detect hazardous compounds," Hirota says. "Then, by studying which ligands or substrates bind to these receptors, it may be possible to identify new candidate drugs in future."

**More information:** Junpei Yamashita et al, Skn-1a/Pou2f3 functions

as a master regulator to generate Trpm5-expressing chemosensory cells in mice, *PLOS ONE* (2017). DOI: 10.1371/journal.pone.0189340

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