

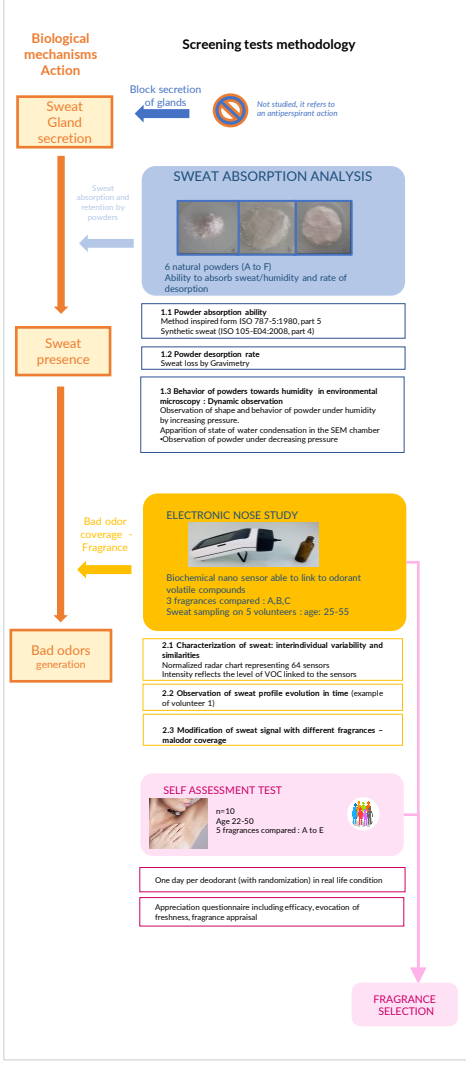
How a combination of innovative methods can help for the development of new deodorant Products

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Introduction

Sweating is an essential process to maintain the human body at an optimal temperature. Nevertheless, it may induce two major inconveniences: wet sensation and malodor. This is why we developed a screening test strategy for the selection of deodorant ingredients. This methodology is oriented through the evaluation of humidity sorption by powders and malodor covering with fragrances. Humidity sorption and desorption studies were conducted on several powders by gravimetry, quantifying the maximal amount of synthetic sweat absorbed by powder.
We characterized with Environmental Scanning Electron Microscopy (ESEM) the behavior of selected powders under humidity by increasing and decreasing pressure. Sweat sampling from volunteers was analyzed using an electronic nose technology. Comparison of sweat signals treated or not with fragrances allowed us to discriminate their efficacy by their ability to mask odor. Self-assessment of a deodorant with different fragrances was proposed to volunteers for appreciation.

Materials & Methods:



Results & Discussion:

1. Powder ability to Sweat/humidity absorption and retention

1.1 Sweat-humidity absorption

Graph 1: Classification of the level of sweat absorption, in µg of powder/µg sweat

Powder C absorbed progressively a high level of sweat, until a state of saturation. Powder F is rapidly saturated and doesn't seem to swell.

1.2 Powder desorption rate

Graph 2: Rate of desorption, % of weight loss

Gravimetric measurement shows the higher rate of desorption for powder F, with a 50% loss of sweat absorbed at 105 min. Powders C, D, F present the lowest desorption rate. After 7 hours, they lose less than 50% of their sweat.

1.3 Dynamic humidity absorption observation via ESEM

Graph 3: Rate of desorption, % of weight loss

Powder C appears composed of fibers with various size length. Fibers do not exceed 300µm. During the rise in pressure (hydration) the fibers get bigger, this swelling kinetics is observed from 5.40 to 5.65 Torr. Water droplets appear at 5.65 Torr. During drying, water disappears and the volume of the fibers decreases. The shape of the fibers is very similar to its initial dry state (not shown). At the same magnification Powder F size appears smaller and has a spherical shape. During the hydration sequence, water spreads quickly (5.55 Torr) before covering the field of observation. After drying, we observed aggregation of particles more compact than initially.

2. Bad odor coverage using electronic nose technology

2.1 Variability of olfactive signal between volunteers

Graph 3: Variability of olfactive signal between sample of panel 1 to 5, 01, at 37°C

2.2 Evolution of sweat signal in time

Graph 4: Olfactive profile of panel 1 sweat (%). Kinetic evolution of sweat signal

After 6 hours at 37°C, we observed for each volunteer a specific profile showing a variability inter individuals in term of sweat composition (extinct sensor). Some peaks were present in all the profiles (10,18,47).

To follow the evolution of sweat in time, we focused on volunteer 4 by comparing data at T0, T6h, T24h. Volatiles compounds are linked particularly to sensors 10, 15, 17, 31, 46, 47, 53, 61 at T0. Kinetics measurements showed a growing intensity appearing at T6h and more intensely at T24h on sensors 16,17,31 and 47.

2.3 Sweat odor coverage with fragrances

Graph 5: Olfactive signal of sweat after 6h at 37°C, with and without deodorant, with fragrance A,B,C, volunteer 4

Graph 6: PCA for volunteer 1. Spatial representation of untreated and treated sweat with fragrances A,B,C

Focusing on volunteer 1, radar of sweat shows the most important signal measured on sensors 6, 17, 46. A new profile is generated when sweat is mixed with different fragrances. New sensors are solicited. To evaluate the fragrance with the most efficiency in its capacity of masking odor, we represented on Principal Component Analysis (PCA) the untreated sweat and sweat treated with fragrance.

Data were analyzed individually for each panel. Observation was made to detect which fragrance appears more different than sweat signal (meaning far on PCA). We obtain most volunteers with higher distance between untreated and treated sweat for product with fragrance B (3/5) followed by fragrance C (2/5).

3. Self assessment test

Focusing on A,B,C fragrances, product C with better global notes than A and B presents a pleasant fragrance and freshness evocation for 80% of volunteers. Improvement should be the persistence of perfume in time. The results seems aligned with electronic nose study where the B fragrance appears more efficient. However, we noticed that the preferred formula was A, which reveals the importance of individual hedonic appreciation, despite the perceived efficacy.

Conclusions:

With this step-by-step approach we developed a screening strategy to help the selection of ingredients for efficient anti humidity and anti-odor deodorant. The properties of powders under humidity can be assessed with a simple gravimetric protocol. Observation at a microstructural level, thanks to real time monitoring ESEM allows a better understanding of interaction of these powders with water.
The second step was to compare fragrance efficacy in terms of odor covering. We chose to use a particular and still not very documented electronic nose technology. We clearly observed a variability between subjects, and a modification of the sweat signal in time, testifying of the presence of more volatile compounds, probably responsible for malodor. Finally, we were able to appreciate fragrance proposals in terms of covering effect by their ability to transform the signal of odorous sweat. This latter approach is very promising and needs more data to deepen the phenomenon of malodor generation and covering. The self-perception evaluation is essential to give us confirmation of the ingredients selected and warns us to consider the consumer preference.

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