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# Characterization of H<sub>2</sub>S positive and negative strains of Shewanella baltica isolated from spoiled whiting (Merlangius merlangus)

## Alexandre DEHAUT<sup>1</sup>, Graziella MIDELET-BOURDIN<sup>1</sup>, Jean-Sébastien PY<sup>2</sup>, Benoît GASSILLOUD<sup>2</sup>, Anne BRISABOIS<sup>1</sup> & Guillaume DUFLOS<sup>1</sup>

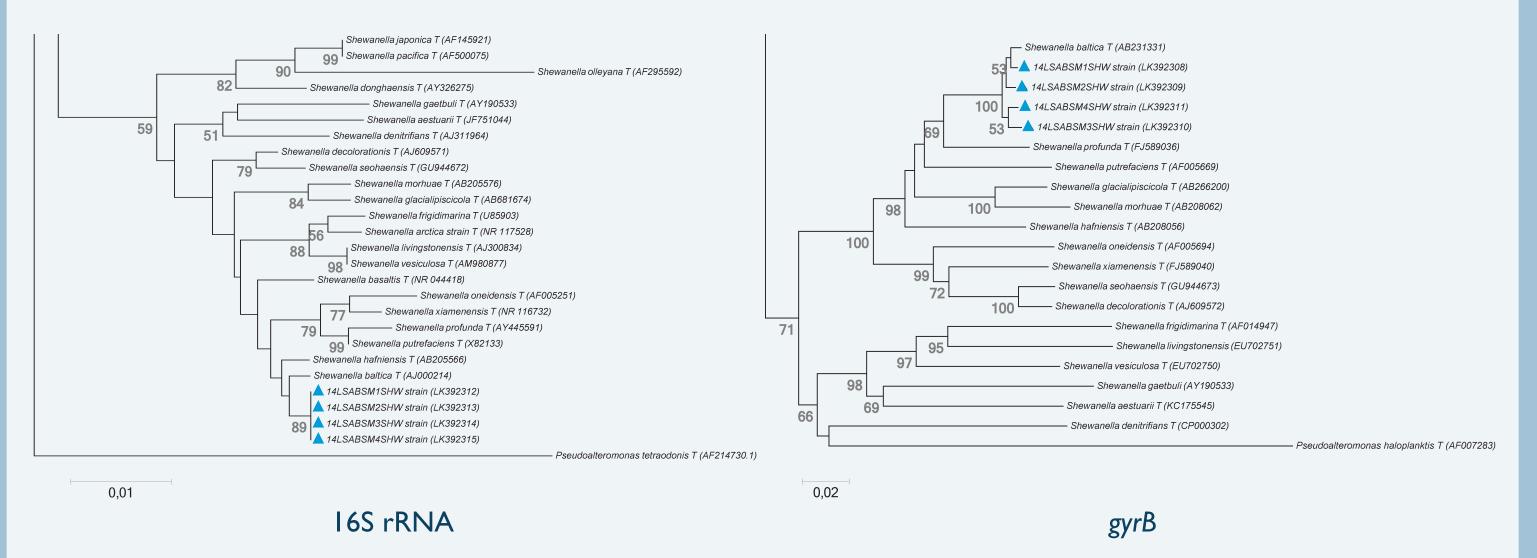
I. Anses, Laboratoire de Sécurité des Aliments - Département des Produits de la Pêche et de l'Aquaculture, Boulevard du Bassin Napoléon, 62200 Boulogne-sur-Mer, France

2. Anses, Laboratoire d'Hydrologie de Nancy, 40 rue Lionnois, 54000 Nancy, France

# Introduction

Among the numerous Shewanella species described in marine animals and sea water, Shewanella putrefaciens has been described as the spoiling agent for fresh fish stored on melting ice (Gram & Dalgaard, 2002).

## Genotypic approaches



Nonetheless S. putrefaciens is more likely a bacterial group composed of four sub-groups than a single species.

S. putrefaciens group II, reclassified as Shewanella baltica, is a very heterogeneous group recently characterized by MultiLocus Sequence Typing (MLST) in nine clades (Ziemke et al., 1998; Deng et al., 2014).

S. baltica has been described as a potential spoilage organism and the most  $H_2S$  producing bacteria during storage of marine fish (Vogel et al. 2005).

→ The present study describes the phenotypic, genotypic and proteomic characterization of four S. baltica strains isolated from spoiled whiting (Merlangius merlangus).



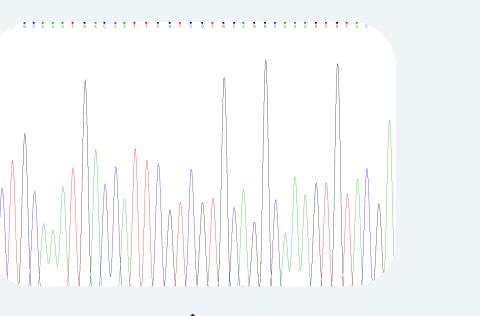
# Materials & Methods

**Phenotypic and Biochemical** Bacterial cultures and biochemical tests



Genotypic

### 16S rRNA and gyrB genes sequencing



**Proteomic** MALDI-TOF profiles



All the strains were identified as S. baltica by both 16S rRNA and gyrB partial sequencing. The second gene, gyrB, gave more informations on the diversity of the isolated isolates.

#### **Proteomic approaches**

|    |      |       |      |       |       |       |                                       |                  | Strains                      |
|----|------|-------|------|-------|-------|-------|---------------------------------------|------------------|------------------------------|
|    |      |       |      |       |       |       |                                       |                  | 14LSABSMISH                  |
|    |      |       |      |       |       |       |                                       |                  | I4LSABSM2SH                  |
|    |      | uluhh | held | Milul |       |       | · · · · · · · · · · · · · · · · · · · |                  | I 4LSABSM3SH<br>I 4LSABSM4SH |
| 0  | 4000 | 6000  | 8000 | 10000 | 12000 | 14000 | 16000                                 | 14LSABSM4SHW     | Överall, isola               |
|    |      |       |      | 444   | i i   |       |                                       | -8               | profiles                     |
|    |      |       |      |       | 1     |       |                                       | ■14LSABSM3SHW    | 3                            |
|    |      |       |      |       |       |       |                                       | -5<br>-5         | , MALDI-TOF                  |
|    |      |       |      |       |       |       |                                       | ■ 14LSABSM2SHW   | 2.3) and low                 |
|    |      |       |      |       |       |       |                                       | -3               | strain.                      |
|    |      |       |      |       |       |       |                                       | ■ 14LSABSM1SHW _ | Using the Br                 |
|    |      |       |      |       |       |       |                                       |                  | closer to S. b               |
| 00 | 4000 | 6000  | 8000 | 10000 | 12000 | 14000 | 16000                                 | 18000 Da         |                              |

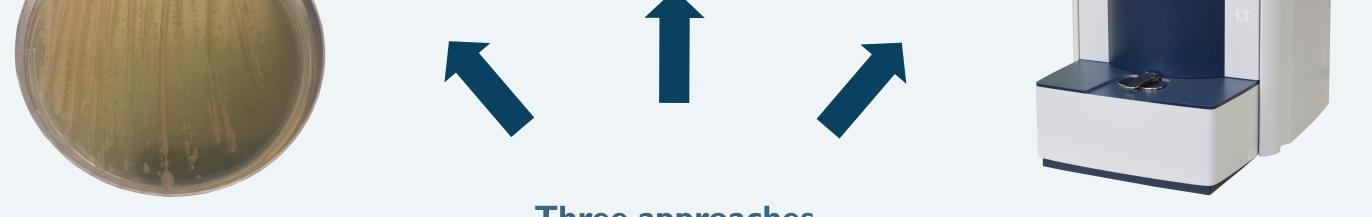
More deeply, some differences have been highlighted between the

| Strains      | Mean  | %RSD | Identification on   |
|--------------|-------|------|---------------------|
| 14LSABSM1SHW | 2.402 | 2.8% | S. baltica OSSI 55  |
| I4LSABSM2SHW | 2.355 | 2.7% | S. baltica OSS I 55 |
| I4LSABSM3SHW | 2.330 | 2.0% | S. baltica OSS I 55 |
| I4LSABSM4SHW | 2.294 | 3.1% | S. baltica OSSI 55  |

Overall, isolates display homogeneous MALDI-TOF profiles.

MALDI-TOF study led to good identification scores (≥ 2.3) and low %RSD among the 10 replicates per strain.

Using the Brüker database, isolates were identified as closer to S. baltica OSS 155 MALDI-TOF profile.



**Three approaches** 

Four strains of Shewanellaceae (14LSABSM1SHW, 14LSABSM2SHW, 14LSABSM3SHW & 14LSABSM4SHW)

Isolated from a spoiled whiting (Merlangius merlangus) and selection on tryptone soya agar yeast extract with 1.5 % (w/v) NaCl (TSA-Ye-NaCl).

# Results

## Phenotypic and Biochemical approaches

#### Strains growth characteristics:

|                      |       | 14LSABSM1SHW | I4LSABSM2SHW | I4LSABSM3SHW | I4LSABSM4SHW |
|----------------------|-------|--------------|--------------|--------------|--------------|
| Aerobic growth       | (n=2) |              |              |              |              |
| (TSA-Ye medium)      | 4°C   | +            | +            | +            | +            |
|                      | 8°C   | +            | +            | +            | +            |
|                      | I5°C  | +            | +            | +            | +            |
|                      | 25°C  | +            | +            | +            | +            |
|                      | 30°C  | +            | +            | +            | +            |
|                      | 37°C  | -            | -            | -            | -            |
| Anaerobic growth     | (n=2) |              |              |              |              |
| (TMAO-TSA-Ye medium) | 25°C  | +            | +            | +            | +            |

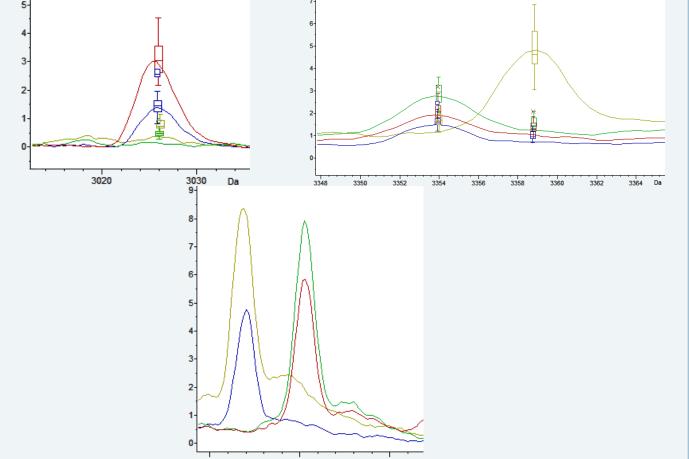
Aerobic growth was achieved on TSA-Ye-NaCI plates for 72H (4°C), 48H (8°C) and 24H (15°C, 25°C, 30°C and 37°C). Anaerobic growth was carried out on TSA-Ye-

four isolates :

.lon 3025 m/z differentiate 14LSABSM1SHW & 14LSABSM3SHW from 14LSABSM2SHW & 14LSABSM4SHW.

.lons 3359 m/z, 3767 m/z, & 3780 m/z differentiate I4LSABSM2SHW from I4LSABSM4SHW.

.lons 3767 m/z, 3780 m/z, 7536 m/z & 7562 m/z differentiate I4LSABSMISHW from I4LSABSM3SHW.



Confirmation of genotypic identification by the MALDI-TOF study. This technology allowed to visualize some slight differences between the proteomic profiles of the strains.

## Conclusions

Phenotypic and biochemical approachs failed to accuretly identify the four strains of Shewanellaceae.

Genotypic approach, by sequencing the two genes 16s rRNA and gyrB, allowed to identify the whole isolates as members of S. baltica species.

Proteomic approach, using the MALDI-TOF, confirmed the conclusion of the genotypic study and highlits slight differences between strains.

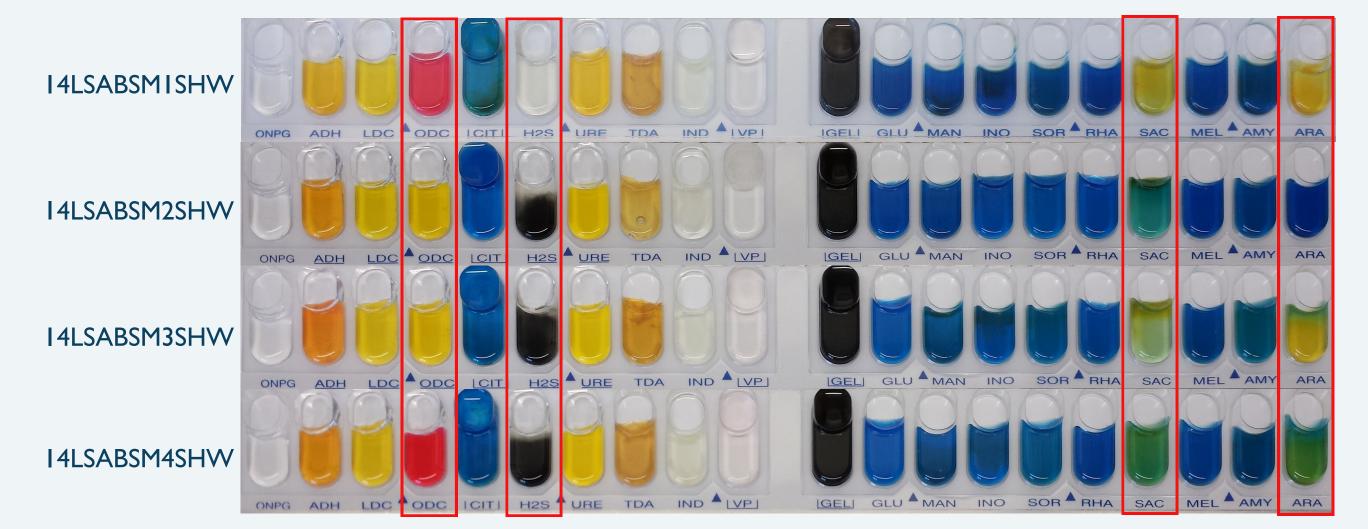
Isolate 14LSABSMISHW, unable to produce  $H_2S$ , displayed an atypic phenotype that is why it was filed in the collection of the Institut Pasteur with the identifier CIP 110804.

H<sub>2</sub>S production might not be the most appropriate screening parameter for isolation of Shewanella isolates.

NaCl plates supplemented with 40 mMTMAO for 24H at 25°C, anaerobic conditions having been generated thanks to Genbag system.

Strains grew on a large range of temperature (4-30°C) and are able to grow anaerobically, in presence of TMAO, with the development of the specific fish off-odor.

#### Strains biochemical patterns:



Heterogeneous biochemicals patterns observed. One strain is surprinsigly  $H_2S$  negative, two strains synthetize an ornithine decorboxilase and all the strains, except one, are able to use sucrose and arabinose.

#### The spoiling potential of I4LSABSMISHW has to be characterized.

## In-depth reading...

Dehaut, A., Midelet-Bourdin, G., Brisabois, A., & Duflos, G. (2014). Phenotypic and genotypic characterization of  $H_2S$ -positive and  $H_2S$ -negative strains of Shewanella baltica isolated from spoiled whiting (Merlangius merlangus). Letters in applied microbiology, 59(5), 542-548.



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