France

Identification of *Pseudo-nitzschia australis* and *P. multiseries* in the Bay of Seine. Was there a relation to presence of domoic acid in king scallops in autumn 2004?

In late 2004, the French phytoplankton monitoring network (REPHY of Ifremer) detected domoic acid (DA) above the EU-regulatory limit of 20 μg DA/g tissue in king scallops (Pecten maximus) from the Bay of Seine in the Channel (Fig. 1). Shellfish harvesting sites were closed for several months due to slow depuration of the scallops.

Accumulation of DA in French shellfish was previously correlated (as in June 1999) with blooms of the pennate diatom *Pseudo-nitzschia*, dominated most often by species of the *«pseudodelicatissima/cuspidata* complex» [1] or exceptionally by species of the *«seriata* complex». Scanning electron microscopy (SEM) analyses were performed and *P.* (*Cont'd on p. 2*)

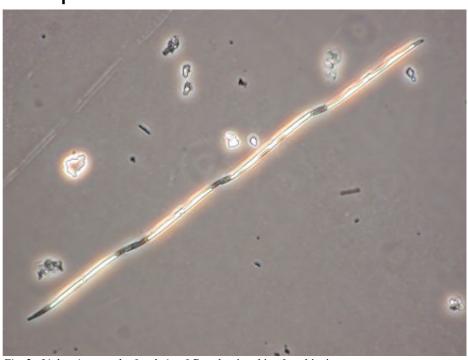


Fig. 2. Light micrograph of a chain of Pseudo-nitzschia cf. multistriata.

Mexico

Bloom of Chattonella subsalsa in an impacted coastal lagoon in the Gulf of California

As in many countries, Mexico, with high population growth and increasing agriculture, mariculture, and tourism, has contributed to coastal eutrophication, providing conditions that favour development of harmful algae blooms (HABs). HABs are expected in areas with high nutrient input, and an increase in their occurrence in the last few years is often cited [1]. This has also been the case for Mexico [2]. An apparent increase of raphidophycean species (*Fibrocapsa*

japonica and *Chattonella* spp.) along Asian and European coasts has been reported [3]. Their presence is of concern because of their toxins. Massive poisoning of economically important fishes has been reported to be caused by the small *F. japonica*; toxicity of *Chattonella* spp. has also been related to fish kills [4].

The presence of *Chattonella* species in a limited number of Mexican coastal areas [5, 6] is noted in two previous reports of blooms in the Gulf of Cali-

fornia [7, 8], one of them associated with loss of marine resources [7]. However, the toxicity of these blooms has not been demonstrated in Mexican waters.

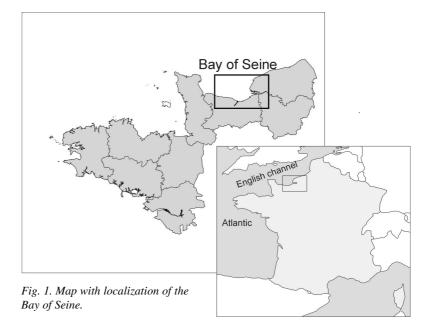
A bloom of *Chattonella subsalsa* was observed in Laguna Navachiste, Sinaloa on the eastern coast of the Gulf of California (Fig. 1). Laguna Navachiste is a shallow, semi-enclosed coastal lagoon characterized by harsh environmental conditions and severe (*Cont'd on p. 4*)

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(Cont'd from p. 1)

pseudodelicatissima and P. multiseries were identified respectively [2]. More recently, the toxic species P. calliantha (which belongs to the «pseudodelicatissima/cuspidata complex») was shown to be also present in French coastal waters (Billard, unpublished data).

At the end of October 2004, just before the opening of the harvesting period, and when DA was detected in king scallops from the Bay of Seine, field samples were examined to look for the diatoms suspected to be responsible for this contamination. A first investigation by light microscopy (LM) of bottom seawater did not detectthe DA-producing benthic diatoms Amphora coffaeformis or *Nitzschia navis-varingica*. On the other hand, cells and empty frustules of at least four species of Pseudo-nitzschia were present in low numbers. The most abundant featured narrow cells with sigmoid ends (Fig. 2) as in P. multistriata or wide symmetrical valves as in P. fraudulenta, and the least abundant showed wide asymmetrical (type A) or narrowly fusiform valves with coarse fibulae and interstriae (type B). The usually toxic



«pseudodelicatissima/cuspidata complex» was absent.

During November 2004 a specific study was undertaken in several Bay of Seine sites and samples examined included: contents of scallop gastrointestinal tracts, sediment, bottom seawater and surface plankton (20 µm mesh size nets). Thorough LM analyses showed that only rare frustule fragments were present in scallops whereas rare sigmoid cells and a few empty frustules of the four species of *Pseudo-nitzschia*

were recorded in the sediment samples; in bottom waters and surface nets, neither of the two morphotypes A and B was recorded. Three Bay sites sampled on November 29-30th for isolation of living cells and EM identification showed scarcity of Pseudo-nitzschia: two types only were recognized by LM, thin sigmoid cells and larger cells with symmetrical valves. The latter species, more abundant at the most coastal of the three sites, was isolated into unialgal culture and unambiguously identified by SEM as P. fraudulenta (Fig. 3).

Since *P. fraudulenta* is generally considered

non-toxic and other putative toxic *Pseudo-nitzschia* species were scarce in autumn of 2004, and in order to understand what had been responsible for the DA contamination, phytoplankton samples collected routinely in the Bay of Seine by the REPHY network during the previous summer were examined. A moderate *Pseudo-nitzschia* bloom was detected from mid-July to September, with a peak on September 14th, 2004 at 56,000 cells 1⁻¹.

Light microscopy observations showed that the Pseudo-nitzschia summer assemblages included only three morphotypes: *P.* cf. *fraudulenta*, but also types A and B. Either type A or B was dominant during the period of mid-July to September as shown by estimation of cell densities of individual types. Type A was not only characterized by asymmetrical valves but also by 7.5-8 µm wide cells, with important overlapping ends (25-28% of cell length) as in P. australis or P. seriata; type B with narrowly fusiform (4.5-6 µm wide) and coarsely silicifed valves showed overlap of cell ends about 27-30% of length, as in *P. pungens* or P. multiseries.

Species identification by transmission electron microscopy (TEM) was carried out on acid-cleaned frustules following the method of Lundholm & Moestrup [3]. Examination of a subsample of the September 14th 2004 surface water collection allowed us to formally

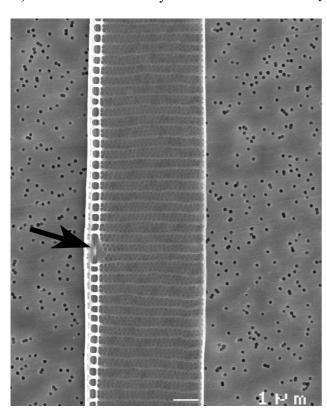


Fig. 3. *Pseudo-nitzschia fraudulenta*, SEM micrograph of valve showing central interspace, fibulae, interstriae and the typical poroid arrangement.

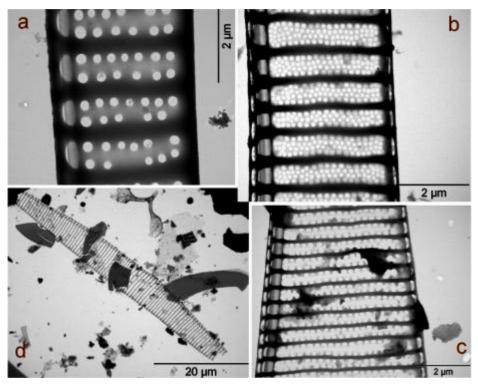


Fig. 4. TEM micrographs of valve views of Pseudo-nitzschia spp. with arrangement of poroids. a) P. pungens. b) P. multiseries. c-d) P. australis, with d) showing asymmetrical outline of the valve.

identify three species of Pseudo-nitzschia: P. pungens, P. multiseries and P. australis (Fig. 4). The first two, which are roughly similar in shape, are easily distinguished by their ornamentation, P. pungens with 2 rows of poroids between thick interstriae and P. multiseries with striae showing 3-4 rows of tightly packed poroids. P. australis, with large asymmetrical valves, features thin interstriae (18 interstriae in 10 µm and approximately the same number of fibulae) and 2 rows of poroids per valve

stria (4-5 pores in 1 μ m). Following EM analyses, type A cells may therefore be assimilated to *P. australis*, while type B cells were either *P. pungens* or *P. multiseries*.

Individual chains of type A cells were isolated from a Lugol-fixed water sample collected in mid-August 2004 and processed for molecular analysis. The ITS1, 5.8S and ITS2 region and part of the 28S of the nuclear ribosomal DNA were sequenced following PCR amplifications. The resulting sequences

Fig. 5. Pair-wise distance calculations using MEGA 3.1 (Neighbor joining, Kimura 2-parameters), accession numbers of the aligned sequences, and one of the two similar obtained matrices.

	[1]	[2]	[3]	[4]	[5]	[6]	[7]	[8]
[1]								
[2]	0.000							
[3]	0.000	0.000						
[4]	0.000	0.000	0.000					
[5]	0.000	0.000	0.000	0.000				
[6]	0.000	0.000	0.000	0.000	0.000			
[7]	0.000	0.000	0.000	0.000	0.000	0.000		
[8]	0.000	0.000	0.000	0.000	0.000	0.000	0.000	

[1] P. australis Bay of Seine-28S partial, [2] AY452530-P.australis-28Spartial, [3] AY452529-P.australis-28Spartial, [4] AF440768-P.australis-28Spartial, [5] AF417651-P.australis-28Spartial, [6] U92260-P.australis-28Spartial, [7] U41393-P.australis-D1-D3, [8] U40850-P.australis-D1-D3.

[1] P. australis Bay of Seine-ITS1-2, [2] AY452528-P.australis-ITS1-2, [3] AY452527-P.australisITS1-2, [4] AY559850-P.cf.australis-18Spart-28Spart, [5] AY257842-P.australis-18Spart-28Spart

were compared with known sequences of Pseudo-nitzschia species from other geographic regions available in databases (Fig. 5) and shown to be identical (100 % identity) with two sequences of Scottish strains of P. australis [4] and with other strains of P. australis from U.S.A [5, 6], New Zealand [U92260, Haywood, A.J., et al., direct submission] and Denmark [AF417651, Lundholm, N. et al., direct submission]. Results of the molecular analyses therefore confirm once more that type A cells were of P. australis and, furthermore that this species was present in mid-August in the Bay of Seine.

Pseudo-nitzschia multiseries had never been recorded before in the Bay of Seine, while this is the first report for the occurrence of *P. australis* in France. Both these species, known to be highly toxic, co-occurred during the summer bloom of 2004 in the Bay. They are suspected to be the main source of DA in the king scallops, while *P. fraudulenta* and the *P. multistriata*-like cells could be potential secondary sources of DA.

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