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**Cyclic imine toxins survey in coastal European shellfish samples: Bioaccumulation and
Mode of Action of 28-O-palmitoyl ester of pinnatoxin-G.
First Report of portimine-A bioaccumulation.**

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Abstract

Cyclic imine toxins exhibit fast acting neurotoxicity and lethality by respiratory arrest in mice explained by their potent antagonistic activity against muscular nicotinic acetylcholine receptors. We performed a survey of gymnodimine-A, 13-desmethyl spirolide-C, 13,19-didesmethyl spirolide-C, 20-methyl spirolide-G, pinnatoxin-A, pinnatoxin-G, portimine-A and 28-*O*-palmitoyl ester of pinnatoxin-G in 36 shellfish samples collected in coastal areas of 8 European countries using a microplate receptor binding assay and UPLC-MS/MS for toxin identification and quantification. The major toxins found in these samples were pinnatoxin-G, 20-methyl spirolide-G, 13-desmethyl spirolide-C, gymnodimine-A and portimine-A. Traces of 13,19-didesmethyl spirolide-C, pinnatoxin-A and 28-*O*-palmitoyl ester of pinnatoxin-G were also detected. The rapid death of mice was correlated with higher pinnatoxin-G concentrations in mussel digestive gland extracts injected intraperitoneally. Our survey included non-toxic control samples that were found to contain moderate to trace amounts of several cyclic imine toxins. Shellfish may bioaccumulate not only cyclic imine toxins but also a large number of acyl derivatives as a product of metabolic transformation of these neurotoxins. This is the first report in which portimine-A and 28-*O*-palmitoyl ester of pinnatoxin-G were detected in shellfish extracts from digestive glands of mussels collected in Ingril lagoon. The bioaccumulation of portimine-A is particularly of concern because it is cytotoxic and is able to induce apoptosis. The mode of action of 28-*O*-palmitoyl ester of pinnatoxin-G was studied by receptor binding-assay and by two-electrode voltage clamp electrophysiology. The antagonistic behavior of the acylated pinnatoxin-G towards nicotinic acetylcholine receptor of muscle type is shown here for the first time. Since cyclic imine toxins are not regulated further monitoring of these emerging toxins is needed to improve evidence gathering of their occurrence in shellfish commercialized for human consumption in

Europe given their potent antagonism against muscle and neuronal nicotinic acetylcholine receptors.

Keywords: Microplate receptor-binding assay; cyclic imine toxins; nicotinic acetylcholine receptors; pinnatoxin-G; 28-*O*-palmitoyl ester of pinnatoxin-G; portimine-A; two-electrode voltage clamp electrophysiology.

1. Introduction

The emerging cyclic imines toxins (CiTXs) of global distribution across coastal environments and produced by dinoflagellates move fast up the human food chain through shellfish. The CiTX family includes 44 compounds: 7 gymnodimines (A-E, 12-methylgymnodimine and 16-desmethyl gymnodimine D) (Harju *et al.*, 2016; Miles *et al.*, 2003; Seki *et al.*, 1995; Van Wagoner *et al.*, 2011; Zurhelle *et al.*, 2018); 16 spirolides (A-D, G-I, 13-desmethyl spirolide-C, 27-Hydroxy-13-desmethyl spirolide-C, 13,19-didesmethyl spirolide-C, 20-hydroxy-13,19-didesmethyl spirolide-C, 27-hydroxy-13,19-spirolide-C, 27-Oxo-13,19-didesmethyl spirolide-C, 13-desmethyl spirolide-D, 20-hydroxy-13,19-didesmethyl spirolide-D and 20-methyl spirolide-G) (Gueret *et al.*, 2010; Harju *et al.*, 2016; Molgó *et al.*, 2013; Zurhelle *et al.*, 2018); 8 pinnatoxins (A-H) (Selwood *et al.*, 2010; Selwood *et al.*, 2014; Takada *et al.*, 2001a; Uemura *et al.*, 1995); 3 pteriatoxins (A-C) (Takada *et al.*, 2001b); 6 prorocentrolides (A-B, 4-hydroxyprorocentrolide, 9,51-dihydro prorocentrolide, 30-sulphate prorocentrolide and 14-O-acetyl-4-hydroxyprorocentrolide) (Amar *et al.*, 2018; Hu *et al.*, 1996; Lu *et al.*, 2001; Torigoe *et al.*, 1988); 1 spiroporocentrimine (Lu *et al.*, 2001) and 2 portimines (A and B) (Fribley *et al.*, 2019; Selwood *et al.*, 2013). The spiroimine moiety is the major feature characterizing cyclic imine toxins. The exception to this rule are prorocentrolides in which a hexahydroisoquinoline group replaces the spiroimine motif (Hu *et al.*, 1996; Torigoe *et al.*, 1988). Portimines display a 5-membered cyclic imine ring, gymnodimines, prorocentrolides and spiroporocentrimines a 6-membered imine ring, while spirolides, pinnatoxins and pteriatoxins display a 7-membered imine ring (Hu *et al.*, 1995; Seki *et al.*, 1995; Selwood *et al.*, 2013; Uemura *et al.*, 1995). Recently, a novel CiTX harboring a 6-membered imine ring was isolated from a *Vulcanodinium rugosum* strain collected in Kabira Bay, Okinawa, Japan, that was named kabirimine (Hermawan *et al.*, 2019).

The chemical diversity of CiTX may be even higher if considering metabolic transformations of these toxins by shellfish. A common strategy developed by shellfish to reduce the impact of dinoflagellate toxins on their physiology is the acylation of such compounds (Doucet *et al.*, 2007). Concerning CiTX, shellfish are able to generate a large number of acyl derivatives of gymnodimine-A (de la Iglesia *et al.*, 2013), 20-methyl spirolide-G (Aasen *et al.*, 2006) and pinnatoxin-G (McCarron *et al.*, 2012).

Prorocentrolide, the first ever-described CiTX, was discovered in 1988 from extracts of the marine dinoflagellate *Prorocentrum lima* (Torigoe *et al.*, 1988). In 1995 the physicochemical characterization of gymnodimines was reported following HABs outbreaks dominated by *Karenia selliformis* in New Zealand (Seki *et al.*, 1995). In the same year, several spirolides were characterized associated with blooms of *Alexandrium ostenfeldii* in Canada (Hu *et al.*, 1995), and pinnatoxin-A was characterized after intoxication cases following the consumption of *Pinna muricata* razor clams in China and Japan (Uemura *et al.*, 1995). In 2010, *V. rugosum*, the producer of pinnatoxins was identified in New Zealand and France (Nezan *et al.*, 2011; Rhodes *et al.*, 2010). Some of *V. rugosum* isolates were shown to produce portimine and kabirimine (Hermawan *et al.*, 2019; Selwood *et al.*, 2013).

Cyclic imine toxins exhibit fast-acting neurotoxicity causing rodent death within 3-5 minutes after intraperitoneal or oral administration at lethal doses characterized by transient mouse hyperactivity, followed by a decrease of the respiratory rate with prominent abdominal breathing leading to death by respiratory arrest (Gueret *et al.*, 2010; Takada *et al.*, 2001b; Uemura *et al.*, 1995). CiTXs display potent antagonism and broad selectivity towards skeletal muscle and neuronal nicotinic acetylcholine receptors (nAChRs) explaining rodent death by respiratory arrest (Aráoz *et al.*, 2011; Bourne *et al.*, 2010; Stivala *et al.*, 2015). Radio-ligand-binding assay experiments demonstrated the high affinity and selectivity profile of cyclic

imine toxins for muscle and neuronal nAChRs and showing their very low affinity to muscarinic acetylcholine receptors (M1-M5) (Aráoz *et al.*, 2015).

Although not regulated, these neurotoxic compounds are frequently found in European environments and shellfish samples (Garcia-Altares *et al.*, 2014; McCarthy *et al.*, 2015; Otero *et al.*, 2019; Rambla-Alegre *et al.*, 2018; Van De Waal *et al.*, 2015), constituting a latent threat for public health given their potent antagonistic activity against human muscular and neuronal nicotinic acetylcholine receptors (nAChR) (Aráoz *et al.*, 2011; Stivala *et al.*, 2015), and their capacity to cross the intestinal and brain-blood barriers (Alonso *et al.*, 2013; Munday *et al.*, 2012b). **Vulnerable populations suffering from neuromuscular disorders such as myasthenia gravis, an autoimmune disease characterized by the loss of muscle nAChRs or Lambert-Eaton myasthenic syndrome caused by the autoimmune destruction of voltage-gated calcium channels resulting in a reduced acetylcholine release at the neuromuscular junction (Vincent, 2002), might be particularly sensitive to shellfish contaminated with CiTXs.** In the present work we performed a survey of four families of CiTXs using a microplate receptor-binding assay (microplate-RBAssay) specific for the detection of agonists and antagonists of nicotinic acetylcholine receptors (Aráoz *et al.*, 2008; Aráoz *et al.*, 2012; Rubio *et al.*, 2014) and UPLC-MS/MS for identification and quantification of the referred neurotoxins in shellfish samples collected in the coastal areas of eight European countries. We have also studied the mode of action of **28-O-palmitoyl ester of pinnatoxin G** by using two-electrode voltage clamp (TEVC) electrophysiology.

2. Material and Methods

2.1. Laboratory animal handling.

The transport and handling of *Torpedo marmorata* and *Xenopus laevis* were done according to the guidelines of the Ethical Committee for Animal Experimentation C2EA-59 of Paris

Center-South. Mouse bioassay has been performed as part of the Survey Program of REPHYTOX of the French General Direction for Food, Program 306. Ifremer data stored at <https://www.seanoe.org/data/00361/47251/>.

2.2. Collection and processing of shellfish extracts.

Digestive gland (DG) extracts from mussels collected at Ingril lagoon were provided by IFREMER. **Digestive gland was used as this was the only tissue available due to sampling procedures.** Digestive glands (30 g) were removed and homogenized with a blender. Methanol (9 ml) was added and the toxins were extracted using a Polytron homogenizer (15,000 rpm, 2 min). The homogenate was centrifuged (3,700g for 8 min at 4°C) and the supernatant was recovered (Hess *et al.*, 2013). The extraction was repeated twice and the supernatants were filtered through 2 µm membranes prior to **Ultra-Performance Liquid Chromatography-Tandem Mass Spectrometry (UPLC-MS/MS)** analysis or microplate-RBAssay.

Shellfish homogenates provided by Agri-food and Biosciences Institute (AFBI) were from the UK NRL archive **that were** retained positive materials from historical toxin events of known regulated toxins: **Amnesic Shellfish Poisoning (ASP)**, **Okadaic acid (OA)**, **Diarrhetic Shellfish Poisoning (DSP)**, **Azaspiracids (AZA)**, **Paralytic Shellfish Poisoning (PSP)** or **Domoic Acid (DA)**, and were processed as previously described (Aráoz *et al.*, 2012). Ten grams of shellfish homogenate were suspended in 40 ml acetone to precipitate the protein content. The sample was vortexed until obtaining a homogeneous mixture and roller-mixed for 2 h at room temperature. The homogenate was centrifuged (3,500g for 15 min at 4°C), the supernatant was recovered and the precipitate was re-extracted with acetone, as described. Both supernatants were pooled and evaporated at 40°C under a stream of N₂ (Organomation Associates Inc, Berlin, MA, USA). The dried matter was solubilized in 4 ml

methanol and filtered through a 10-kDa MWCO filter (Millex, Millipore, Billerica, MA, USA) prior to **microplate**-RBAssay or UPLC-MS/MS.

2.3. *Torpedo* electrocyte membrane purification

T. marmorata was purchased from the Biological Station of Roscoff, (University of Rennes, France). Upon arrival, the electric ray was conditioned for 2-4 days in the dark at 15°C in a 180 liters tank containing reconstituted marine water (Instant Ocean, Europrix, Lens, France). The water was oxygenated with compressed air; it was recycled and filtered prior to re-entry. The electric fish was **sacrificed** and the electric organ was dissected and immersed in 500 ml of *Torpedo*-physiological medium (280 mM NaCl, 3 mM KCl, 1.8 mM MgCl₂, 3.4 mM CaCl₂, 5 mM NaHCO₃, 5.5 mM glucose, 300 mM urea, 100 mM sucrose, 1.2 mM sodium phosphate buffer, pH 6.8). The solution was oxygenated and the final pH was 7 - 7.2. All further steps were performed at 4°C. Electrocyte membranes enriched in embryonic muscle-type nAChRs were purified according to (Hill *et al.*, 1991) with some modifications. The freshly dissected electric organ (500 g) was sliced into small pieces and immersed in 500 ml of *Torpedo*-membrane extraction buffer (TMEB) (50 mM Tris-HCl, 3 mM EDTA, 1 mM EGTA, protease inhibitors mix (Complete Roche Diagnostics GmbH, Mannheim, Germany), pH 7.5. The electric tissue was homogenized with a Waring blender (Appareils Scientifiques O.S.I., Paris, France) at maximal speed for 1 min (3 times with 1 min interval). The homogenate was centrifuged at 4,000g for 10 min at 4°C (Sorvall, Dupont Instruments, Newtown, USA; rotor GSA). The supernatant (S1) was collected and filtered through several layers of medical gauze. The remaining pellet was re-homogenized and centrifuged and processed as described. The supernatants S1 and S2 were centrifuged at 25,000g for 50 min at 4°C (Sorvall, rotor GSA). The pellets containing post-synaptic electrocyte membranes were homogenized with a Potter-Elvehjem (glass/Teflon) in 300 ml TMEB supplemented with 35% sucrose (w/w). A volume of 16 ml of this crude membrane suspension was carefully layered

onto 10 ml of 43% sucrose (w/w) TMEB. The samples were centrifuged at 106,000g for 3 hr at 4°C (Beckman, Palo Alto, Ca, USA; rotor 50Ti). ***Torpedo*-electrocyte membranes rich in nAChRs (*Torpedo*-nAChR)** were collected from the interface between 35% and the 43% sucrose layers. The electrocyte membranes were precipitated by centrifugation at 106,000g for 1 hr at 4°C (Beckman, rotor 50Ti). The precipitated membranes were homogenized in 150 ml TMEB using a Potter-Elvehjem and centrifuged at 106,000g for 1 h at 4°C (Beckman, rotor 50Ti). The membranes were homogenized in 100 ml of 5 mM glycine using a Potter-Elvehjem and centrifuged at 106,000g for 1 hr at 4°C (Beckman, rotor 50Ti). The latter washing step was repeated once. The purified electrocyte membranes were resuspended in 40 ml of 5 mM glycine. Protein concentration was adjusted to 2.5 - 3.5 mg/ml using Bradford method (Bio-Rad, Munich, Germany). The electrocyte *Torpedo*-membranes were aliquoted and stored at -80°C until use.

2.4. Microplate receptor-binding assay.

Maxisorp ELISA plates were purchased from NUNC (Kamstrupvej, Denmark). OPD tablets (***o*-phenylenediamine dihydrochloride**) were from DAKO (Glostrup, Denmark). Biotin- α -bungarotoxin was from Molecular Probes (Eugene, OR, USA). Streptavidin- **horseradish peroxidase** and α -bungarotoxin (α -BgTx) were from Sigma (St. Louis, MO, USA). Non-radioactive microplate-RBAssay for the detection of cyclic imine toxins directly on shellfish extracts was done as described (Ar  oz *et al.*, 2012). Briefly, 100 μ l shellfish extracts prepared in TBS-BSA buffer (50 mM Tris-HCl, 150 mM NaCl, 0.5% BSA pH 7.4) were incubated overnight with *Torpedo*-electrocyte membranes coated in a 96-wells plastic microplate at 4°C. The concentration of the methanolic shellfish extract in the incubation mix was kept lower than 5% to avoid methanol interference with the assay (Rubio *et al.*, 2014). Thereafter, 50 μ l of biotin- α -BgTx (2.4×10^{-7} M, Life Technologies, Eugene, OR, USA) was added to each well and further incubated for 30 min at room temperature and constant shaking. The

microplates were washed three times with 250 μ l washing buffer (TBS, 0.1% Tween 20, pH 7.4). A volume of 100 μ l of streptavidin- **horseradish peroxidase** (220 ng/ml protein) was added to each well. Following 30 min incubation at room temperature, the wells were washed three times as described, and 100 μ l of freshly prepared peroxidase substrate (OPD, Sigma, St. Louis, MO, USA) were added to each well. After 5 min, 100 μ l of 0.5 M H₂SO₄ were added to each well to stop the enzymatic reaction. The data were recorded using a **microplate** reader (CLARIOstar Plus, BMG Labtech, Ortenberg, Germany) at a wavelength of 492 nm. The data was transformed into inhibition percent by using Eq. 1:

$$\text{Inhibition \%} = 100 \times (100\% \text{ signal} - \text{signal sample}) \div (100\% \text{ signal} - 100\% \text{ inhibition})$$

where: **100% signal** is the absorption data from control wells in which *Torpedo*-electrocyte membranes were incubated in the absence of toxins/extracts; **signal sample** is the absorption data of test wells; **100% inhibition**, is the absorption data obtained from control wells in which *Torpedo*-electrocyte membranes were incubated with 1×10^{-5} M α -BgTx. The experiments were performed three times in replicates of six. The concentration of inhibitor at which biotin- α -BgTx binding to *Torpedo*-nAChRs is inhibited by 50% (*IC*₅₀) was determined by nonlinear regression (GraphPad Prism 5.0, San Diego, CA, USA).

2.5. Toxin standards

Gymnodimine-A and 13-desmethyl spirolide-C were purchased from NRC–CNRC (Institute for Marine Biosciences, NRC, Halifax, NS, Canada). 13,19-didesmethyl spirolide-C and 20-methyl spirolide-G were obtained from CIFGA (Lugo, Spain). Pinnatoxin-A, pinnatoxin-G and **28-O-palmitoyl ester of pinnatoxin-G** were synthesized in the laboratory of Professor Armen Zakarian (University of California, Santa Barbara California). Portimine was purified from an ethanolic extract of *V. rugosum*. Physicochemical characterization for its use as toxin standard was performed by NMR (Lamoise *et al.*, 2016). Briefly, the extracts were fractionated by **solid phase extraction** using 5g-C-18 cartridges (Sep-Pak, Waters).

Analytical chromatography of portimine-A was performed using an X-Bridge BEH 300 (**ethylene-bridged hybrid particles**) C18 column (5 μm 4.6 \times 250 mm). Purification of portimine-A was achieved using a preparative X-Bridge Prep C18 column (5 μm OBD - **Optimum Bed Density**- 19 \times 150 mm) **using as solvent A: H₂O, 0.1% formic acid, and as solvent B: CH₃CN, 0.1% formic acid**, at a flow rate of 25 ml/min. NMR was performed at 20°C on a Bruker DRX700 spectrometer equipped with a triple resonance cryoprobe. NMR data were processed with Topspin 1.3 software (Bruker Biospin, Germany) and analyzed with CcpNmr Analysis software. ¹H and ¹³C resonance of portimine were previously assigned by (Selwood *et al.*, 2013). ¹H NMR quantification of portimine was performed as described in (Aráoz *et al.*, 2011). Briefly, 7.0 μl of CHCl₃ and 7.0 μl of C₆H₆ were added to 7.0 ml of **deuterated methanol**. Portimine was dissolved in 0.5 ml of the previous solution in a 2 ml glass flask. The entire sample was added to a NMR tube for quantification.

2.6. UPLC - Triple Quadrupole Detector (UPLC-TQD)

Simultaneous detection and quantification of 7 cyclic imine toxins and 28-*O*-palmitoyl ester of pinnatoxin-G was performed on a UPLC-TQD (Waters, Milford, MA, USA). A calibration curve for each CiTX was obtained by injecting onto a BEH C18 (Waters, 2.1 \times 100 mm; 1.8 μm particle size) 5 μl of a toxin standard mixture containing gymnodimine-A, 13,19-didesmethyl spirolide-C, pinnatoxin-A, 13-desmethyl spirolide-C, 20-methyl spirolide-G, pinnatoxin-G, 28-*O*-palmitoyl ester of pinnatoxin-G and portimine-A, in the concentration range of 1 pM to 1 μM . The samples were chromatographed at a flow rate of 0.6 ml/min using as solvent A: H₂O, 0.1% formic acid, and as solvent B: acetonitrile, 0.1% formic acid. The gradient profile was 0-1 min: isocratic 15% B, 1-9 min: linear gradient 15-36.5% B, 9-9.2 min: linear gradient 36.5-100% B, 9.2-11 min: isocratic 100% B, 11-11.2 min: linear gradient 100-15% B and 11.2-18 min: isocratic 15% B. All the standard mixes were run in triplicate and two blank samples (5 μl methanol) were intercalated between different samples to avoid

carryover **since repetitive injection of standard toxins and shellfish extracts impinge the chromatographic column and the mass detector**. TQD mass spectrometer was operated in the **electrospray ionization** positive mode. The capillary potential was set at 3.5 kV; desolvation temperature at 450°C; source temperature at 150°C; desolvation gas flow at 800 l/hr and cone gas flow at 50 l/hr. Data acquisition was performed at a multiple reaction-monitoring mode (MRM; **Table 1**). Methanolic shellfish extracts (5 µl) were analyzed in triplicate and two blank samples (5 µl methanol) were intercalated between different samples to avoid carryover. Toxin quantification was performed by integration of the area under each toxin peak of three independent MRM chromatograms using MassLynx 4.1 software (Waters).

2.6. Two-electrodes voltage clamp (TEVC) electrophysiology

TEVC electrophysiology was performed on *Xenopus laevis* oocytes having incorporated *T. marmorata* electrocyte membranes rich in nAChR in their plasmatic membrane. Adult *X. laevis* females were from the Biological Resource Center (University of Rennes, France). Laparotomy of *X. laevis* mature female frogs for oocyte extraction was performed according to the protocol APAFIS#5310-2016042016067330 v3 approved by the Ethical Committee for Animal Experimentation C2EA-59 of Paris Center-South. A female donor tagged with an electronic chip was anesthetized by immersion in a 1 g/l ethyl-3-amino benzoate methane sulfonate solution (Sigma-Aldrich, Saint Quentin, France) for 20 min. Following a small incision below 5 mm of the termination of the sternum, the ovarian lobes were excised and immediately immersed in OR2 solution containing 88 mM NaCl, 1 mM MgCl₂, 5 mM HEPES 5; pH 7.6. The muscle and dermic planes were sutured and the amphibian was placed in a small aquarium for recovery and observation for 24 h. After three washes with OR2, the ovarian lobes were incubated in Barth's solution containing 88 mM NaCl, 1 mM KCl, 0.41 mM CaCl₂, 0.82 mM MgSO₄, 2.5 mM NaHCO₃, 0.33 mM Ca(NO₃)₂ and 7.5 mM Hepes (pH

7.6) at 18°C. Fifty nanoliters of *Torpedo*-electrocyte membranes (3.5 mg/ml total protein) were microinjected into stage V–VI defollicularized oocytes using a Nanoliter 2000 Micro 4 Controller (WPI, Sarasota, FL, USA). Injected oocytes were incubated at 18°C.

TEVC electrophysiology recordings were performed 1 to 4 days after microinjection using an OC-725B TEVC amplifier (Warner Instrument, Hamden, CT, USA). Voltage and current microelectrodes were filled with 3 M KCl (4-10 MΩ electrode resistance). ACh-evoked currents were acquired using a pCLAMP-9/ Digidata-1322A system (Molecular Devices, San Jose, CA, USA). The *Xenopus* oocyte was constantly clamped at -60 mV and was perfused with Ringer-Barium containing 100 mM NaCl, 2.8 mM KCl, 0.3 mM BaCl₂ and 5 mM HEPES (pH 7.4), at a flow rate of 8 ml/min.

Dose-response curves for agonist activation were fitted to equation 2:

$$I = I_{max}[L]^{nH}/(EC_{50} + [L])^{nH}$$

where I is the measured agonist-evoked current, $[L]$ is the agonist concentration, EC_{50} is the agonist concentration that evoked half the maximal current (I_{max}), and nH is the Hill coefficient. For antagonist inhibition, current (I) values were normalized to the I_{max} ACh value recorded from the same oocyte to yield fractional (%) response data. IC_{50} values were determined from dose-response curves by fitting to the equation 3:

$$F = 1/[1 + ([X]/IC_{50})^{nH}]$$

where F is the fractional response obtained in the presence of the inhibitor at concentration $[X]$ and IC_{50} is the inhibitor concentration that reduced the ACh-evoked amplitude by half. Data was analyzed with Clampfit 10.2 (Molecular Devices) and GraphPad Prism 5.0.

3. Results

3.1. Pinnatoxin-G occurrence in mussels (DG) and mice survival time

Primary screening for CiTXs was performed using the microplate-RBAssay, a high throughput test specific for the detection of competitive agonists/ antagonists of nAChRs. **Thus, the microplate RBAssay can be easily adapted to pipetting platforms for automated liquid handling intercalating incubation and washing steps, signal development and automatic data handling (WO2012101378A1).** When present in the extracts, CiTXs compete with the toxin tracer for binding the ACh-site of *Torpedo*-nAChRs in a concentration dependent manner. Most of the digestive gland extracts (DG) from mussels collected at Ingril lagoon were highly active on *Torpedo*-nAChR with inhibition values higher than 55% (Fig. 1A, B, Table 2 and 4). Only sample DG-15 showed an inhibition binding of ~25%. **Calibration curves were obtained for CiTXs quantification by UPLC-MS/MS with R^2 values ranging from 0.9797 to 0.9998 (Fig. 1D, E), with limits of detection (LOD) in the sub-nanomolar range and with limits of quantification (LOQ) in the nanomolar range (Table 1).** UPLC-MS/MS analysis showed high concentrations of pinnatoxin-G in 87% of the samples. Traces of 13-desmethyl spirolide-C, gymnodimine-A, pinnatoxin-A and 20-methyl spirolide-G were also present in these extracts (Table 2). The acylated 28-*O*-palmitoyl ester of pinnatoxin-G and portimine-A were not detected in these samples.

The digestive gland extracts from mussels collected in Ingril were highly lethal to mice. The survival time of mice following intraperitoneal injection of DG extracts ($n = 3$) ranged between 4 to 80 min for ~70% of the tested animals (Table 2). **The survival time was monitored for 24 h (1440 min), after which, surviving mice were sacrificed.** The Pearson's correlation coefficient between mice survival time *versus* pinnatoxin-G concentrations in DG extracts was significant ($r = -0.4433$; one-tailed) suggesting that the rapid death of mice was associated with higher concentrations in pinnatoxin-G.

3.2. CiTX occurrence in shellfish samples collected in Europe

AFBI shellfish samples were from six European countries with an Atlantic and Mediterranean coastline that included Scotland, Northern Ireland, Ireland, Italy, Norway and Portugal. In addition, as control samples, four commercial shellfish were purchased from a Fish market in the Paris Region: *Glycymeris glycymeris* (clams), *Venerupis philippinarum* (Japanese cockle) and *Ostrea edulis* (oysters) that were produced in the Brittany Region of France, and *Mytilus galloprovincialis* (mussels) that was imported from Netherlands (Table 3). The inhibition binding potency of the whole group of samples varied from ~25 to 100%, as tested by microplate-RBAssay, suggesting that all samples were contaminated with CiTXs (Fig. 1C, Table 3). UPLC-MS/MS analysis confirmed the presence of pinnatoxin-G, 20-methyl spirolide-G, 13-desmethyl spirolide-C, pinnatoxin-A, 13,19-didesmethyl spirolide-C and gymnodimine-A in these extracts. The acylated compound 28-*O*-palmitoyl ester of pinnatoxin-G and portimine-A were not detected. The most abundant CiTXs found in these European shellfish extracts were, pinnatoxin-G, 20-methyl spirolide-G and 13-desmethyl spirolide-C. **It is to be noted that the samples AFBI 01-08 contained high levels of OA, DTX, YTX, AZA (Supplementary Table 1), while AFBI 13-14 mussel samples contained PSP and DA, respectively. Samples AFBI-09 (C) – AFBI-12 (C) were non-toxic control entries. The marine phyocotoxins OA, DTX, YTX and AZA were tested by microplate-RBAssay at a concentration of 10 μ M without significant effects on the performance of the bioassay (Supplementary Fig. 1). The presence of other toxins not acting on nAChRs did not interfere with inhibition binding by microplate-RBAssay as it was previously shown for saxitoxin and domoic acid (Aráoz *et al.*, 2012).**

Ligand-binding assay showed that the control samples **AFBI-09 (C) – AFBI-12 (C) and the commercial samples FR-01 (C) – FR-04 (C)** were contaminated with CiTXs (inhibition binding values: 29 – 81%). CiTXs in these extracts were quantified by UPLC-

MS/MS (Table 3). Since cyclic imine toxins are not regulated in Europe, it is a common issue to retrieve them in commercial shellfish. The presence of these neurotoxins in shellfish destined for human consumption is of concern given their high affinity for human nAChRs and their capacity to cross the intestinal and brain-blood barriers.

3.3. Detection of 28-*O*-palmitoyl ester of pinnatoxin-G -A and portimine-A

We previously detected 28-*O*-palmitoyl ester of pinnatoxin-G in the mussel digestive gland samples 13/414-2 and 13/599-2 collected in Ingril lagoon (Mondeguer et al., 2015). The availability of synthetic 28-*O*-palmitoyl ester of pinnatoxin-G allowed us to determine its retention time and MRM conditions for toxin quantification. Portimine was characterized and quantified by NMR (Fig. 2A). The collision energy of the triple quadrupole detector of the UPLC/MS-MS was optimized to obtain a robust MRM transition pair for the quantification of 28-*O*-palmitoyl ester of pinnatoxin-G and portimine-A (Fig. 1E, 2B, 3B). While portimine eluted from the column at 2.56 min (18% B), the acylated 28-*O*-palmitoyl ester of pinnatoxin-G eluted at 11.09 min (100% B). Since the acylated compound was strongly retained by the BEH C18 column, the chromatographic run was longer in order to clean up the column (see Material and Methods). It is worth mentioning that 28-*O*-palmitoyl ester of pinnatoxin-G emitted strong MS signals even at low concentrations under the experimental conditions here used (Fig. 1D). Both toxins were detected in the digestive glands of mussel samples 13/414-2 and 13/599-2 collected from Ingril lagoon (Fig. 2C, Table 4). The inhibition binding potency of these extracts were 98% and 72%, respectively, as measured by microplate-RBAssay. UPLC-MS/MS quantification showed that DG-sample 13/414-2 contained 0.67 µg/kg of 28-*O*-palmitoyl ester of pinnatoxin-G and 69.31 µg/kg of portimine-A, while sample 13/599-2 contained 0.07 µg/kg of 28-*O*-palmitoyl ester of pinnatoxin-G and 32.08 µg/kg of portimine-A (Fig. 2C, Table 4). Both DG extracts were also contaminated with high levels of

gymnodimine-A and pinnatoxin-G. Low levels of 13-desmethyl spirolide-C, pinnatoxin-A were also detected (Fig. 2C, Table 4).

3.4. Mode of action of 28-*O*-palmitoyl ester of pinnatoxin-G

The affinity of 28-*O*-palmitoyl ester of pinnatoxin-G for *Torpedo*-nAChR of embryonic muscle type was verified by microplate-RBAssay (Mondeguer et al., 2015; Fig. 3C). The acylation of pinnatoxin-G at C-28 provoked a ~700 decrease of the inhibition binding of 28-*O*-palmitoyl ester of pinnatoxin-G with an $IC_{50} = 13.1 \mu\text{M}$ (95% interval confidence: 6.20 – 27.37 μM) when compared to its parent compound pinnatoxin-G whose IC_{50} was 18.65 nM (95% interval confidence: 8.8 – 39.6 nM). TEVC electrophysiological recordings showed the mode of action of 28-*O*-palmitoyl ester of pinnatoxin-G: it acts as an antagonist of *Torpedo*-nAChR of embryonic muscle type ($\alpha 1$)₂ $\beta 1\gamma\delta$. Thus, the perfusion of **28-*O*-palmitoyl ester of pinnatoxin-G** did not elicit inward currents on *X. laevis* oocytes carrying *Torpedo* ($\alpha 1$)₂ $\beta 1\gamma\delta$ nAChR suggesting no agonist activity (Fig. 3D). However, it competitively antagonized ACh-evoked currents in a dose dependent manner ($IC_{50} = 969.3 \text{ nM}$; 95% interval confidence: 752.6 – 2129 nM, Fig. 3E). The interaction between 28-*O*-palmitoyl ester of pinnatoxin-G and the *Torpedo* ($\alpha 1$)₂ $\beta 1\gamma\delta$ nAChR was reversible: following perfusion of 2.5 μM **28-*O*-palmitoyl ester of pinnatoxin-G**, the ACh-evoked current was 30% of the initial value. After 300 s washing with Ringer-Ba, the ACh-evoked current recovered to 76% of the control current (Fig. 3D). The esterification of pinnatoxin-G with palmitic acid (16:0) anhydride at C-28 provoked a ~300 decrease of the antagonistic potency of 28-*O*-palmitoyl ester of pinnatoxin-G relative to pinnatoxin-G ($IC_{50} = 3.15 \text{ nM}$; 95% interval confidence: 2.9 – 4.9 nM, Fig 3E).

4. Discussion

This is the first report describing the mode of action of 28-*O*-palmitoyl ester of pinnatoxin-G and the **bioaccumulation** of portimine-A **by** shellfish extracts thanks to the use of synthetic and purified toxins as standards. The monitoring of CiTXs occurrence was performed on 36 shellfish samples using a target-directed functional method and UPLC-MS/MS for the identification and quantification of 8 CiTX members in a single run. The most frequently detected CiTXs were pinnatoxin-G, 20-methyl spirolide-G and 13-desmethyl spirolide-C, which showed a similar distribution across the European coastal regions tested. The latter may reflect the European-wide distribution of *Alexandrium ostenfeldii* and *A. peruvianum* (Aasen *et al.*, 2005; Brown *et al.*, 2010; Krock *et al.*, 2008; McCarthy *et al.*, 2015; Suikkanen *et al.*, 2013), producers of spirolides, and of *V. rugosum*, producer of pinnatoxins (Garcia-Altares *et al.*, 2014; Hess *et al.*, 2013; McCarthy *et al.*, 2015; Rundberget *et al.*, 2011).

Although highly responsive by microplate-RBAssay, the quantification of the monitored CiTXs by UPLC-MS/MS gave, in some cases, lower toxin concentrations (*i.e.* DG-05, DG-09, DG-11; Table 2), suggesting that the referred samples may contain other CiTXs not monitored in this work. **An alternative for detecting untargeted CiTXs may be the use of high resolution mass spectrometry however; no quantitative data could be obtained by this approach.** Along the same line, the low correlation between pinnatoxin-G content of Ingril samples and mice survival time (Table 2) may indicate the presence of other CiTXs or acylated derivatives in these extracts. A delayed mouse mortality provoked by mussel extracts collected several weeks after an algal bloom dominated by *A. ostenfeldii* in Skjer location (Norway) led to the discovery of 21 fatty acid ester metabolites of 20-methyl spirolide-G (Aasen *et al.*, 2006). Initially, the mouse survival time after intraperitoneal administration of shellfish extract samples dominated by 20-methyl spirolide-G was very short (3-5 min). However, the survival time of mouse became longer as the mussels gradually detoxified

(Aasen *et al.*, 2006). Acylated derivatives of CiTXs may constitute an inhibition background noise for the microplate-RBAssay if present. Taking all together, the results presented here show that the microplate-RBAssay is highly sensitive, and provides an alternative high-throughput method for monitoring neurotoxins targeting nAChRs on environmental samples that could facilitate LC/MS analysis (Rubio *et al.*, 2014).

The growing CiTX group includes seven families of toxins totaling by now 44 CiTX analogs (Lamoise *et al.*, 2016), and very few of them are available as commercial standards (Rambla-Alegre *et al.*, 2018). CiTXs were considered as false negatives for saxitoxin monitoring by mouse bioassay (Vilariño *et al.*, 2009). Thus, the mouse lethality of 13-desmethyl spirolide-C, 13,19 didesmethyl spirolide-C, 20-methyl spirolide-G or pinnatoxin-F, to mention some of them (Munday *et al.*, 2012a; Munday *et al.*, 2012b), compares the mouse lethality of saxitoxin by gavage or intraperitoneal administration (Llewellyn, 2006). Still, since no human fatalities are associated with CiTXs worldwide, they are not internationally regulated (EFSA Panel on Contaminants in the Food Chain, 2010). Consequently, it is common to detect CiTXs in commercial shellfish in Europe (Aráoz *et al.*, 2012; Otero *et al.*, 2019; Rambla-Alegre *et al.*, 2018). Here it is shown that shellfish control samples provided as part of the study were contaminated with 13-desmethyl spirolide-C, 20-methyl spirolide-G, pinnatoxin-G and with traces of pinnatoxin-A, and that shellfish commercial samples purchased in Paris Region were contaminated with 13-desmethyl spirolide-C and with traces of pinnatoxin-G, gymnodimine-A and 20-methyl spirolide-G (Table 3).

Of particular interest for food safety is that shellfish bioaccumulate not only cyclic imine toxins but also a large number of acyl derivatives as product of shellfish metabolism. 28-O-palmitoyl ester of pinnatoxin-G is one of the 26 derivatives described for pinnatoxin-G (McCarron *et al.*, 2012). Binding experiments showed that 28-O-palmitoyl ester of pinnatoxin-G inhibited tracer binding to *Torpedo*-nAChR in the

micromolar range and with a lower affinity compared to pinnatoxin-G (Aráoz *et al.*, 2012; Hess *et al.*, 2013). Further, TEVC electrophysiological recordings showed that 28-*O*-palmitoyl ester of pinnatoxin-G is a competitive antagonist of *Torpedo*-nAChR. The chemically controlled esterification of pinnatoxin-G with palmitic acid occurred at the secondary OH at C28 close to the spirocyclic group (Fig. 3A). It was predicted that the palmitate acyl chain at position C28 in the bridged 5,6-bicycloketal (EF) substructure of pinnatoxin-G may produce steric clashes with amino acid residues in loop F of the orthosteric site of the nAChR reducing significantly the potency of 28-*O*-palmitoyl ester of pinnatoxin-G compared to pinnatoxin-G (Bourne *et al.*, 2015). The results presented here support the previous hypothesis; *i.e.*: the IC_{50} of 969.3 nM determined for 28-*O*-palmitoyl ester of pinnatoxin-G was 300-fold higher than the IC_{50} determined for pinnatoxin-G (3.15 nM). Overall, even slight structural changes impact the affinity and the antagonistic potency of CiTXs towards nAChR subtypes as it is shown for pinnatoxin-G and pinnatoxin-A (Fig. 3A, 3C, 3E, Bourne *et al.*, 2015). Further work is needed to determine the passage of acylated CiTX through the intestinal barrier and their impact on human muscle and neuronal nAChRs.

V. rugosum may now be considered an endemic dinoflagellate of Ingril lagoon (Mediterranean, France), since early observation of this strain go back to 1990 (E. Nézan, *personal communication*). The highest abundances for *V. rugosum* were recorded in summer time (June – September). This species is thermophilic (Abadie *et al.*, 2018), nonetheless, it can adapt to colder lagoons such as Lough Hyne, (Ireland) where pinnatoxin-G was reported (McCarthy *et al.*, 2015). Indeed, a *V. rugosum* bloom episode was recently observed (December 2019) at Ingril lagoon prompting the **prohibition** of shellfish harvesting (thau-infos.fr/index/php/commune/frontignan/69083-etang-d-ingril-peche-aux-coquillages-interdite). Actually, Ingril is not intensively used for shellfish farming. Given that shellfish at

Ingril are frequently exposed to toxic *V. rugosum*, higher levels of pinnatoxin-G (1,200 µg/kg) were recorded in mussels growing in this lagoon (Hess *et al.*, 2013). The high levels of gymnodimine-A and 13-desmethyl spirolide-C found in these samples (Table 4) suggest that Ingril lagoon may also be a hot spot for other CiTXs-producing dinoflagellates. The bioaccumulation of portimine by mussels –described here for the first time– is also of concern, since this toxin not only acts on nAChRs (Lamoise *et al.*, 2016), but it is a potent apoptotic inducer molecule (Cuddihy *et al.*, 2016; Fribley *et al.*, 2019; Selwood *et al.*, 2013).

5. Conclusion

The microplate-RBAssay is a robust technology providing an alternative high-throughput test for CiTXs in environmental samples with very low matrix effect. A UPLC-MS/MS protocol was designed for the simultaneous detection of seven CiTXs and 28-*O*-palmitoyl ester of pinnatoxin-G in a single run taking advantage of commercial, synthetic and purified toxin standards. Pinnatoxin-G, 20-methyl spirolide-G, 13-desmethyl spirolide-C, gymnodimine-A and portimine-A were the major neurotoxins found in this survey. The rapid death of mice was associated with higher pinnatoxin-G concentrations in mussel digestive gland extracts injected intraperitoneally. 28-*O*-palmitoyl ester of pinnatoxin-G was detected in two mussel extracts collected in Ingril lagoon. The mode of action of 28-*O*-palmitoyl ester of pinnatoxin-G is here shown for the first time: 28-*O*-palmitoyl ester of pinnatoxin-G is an antagonist of muscle-type nAChR. The lipid acylation at position C-28 of pinnatoxin-G provoked a serious decrease of the antagonistic potency of 28-*O*-palmitoyl ester of pinnatoxin-G against *Torpedo*-nAChRs when compared to its precursor neurotoxin. This is also the first report in which the bioaccumulation of portimine-A in mussels was demonstrated. Portimine-A may represent a different toxicological risk since it is a potent apoptotic molecule and a nAChR antagonist. The presence of CiTXs in commercial shellfish is of

concern given the high affinity of these neurotoxins for human nAChR of muscle and neuronal subtypes, and their capacity to cross the intestinal and brain-blood barriers in particular for vulnerable populations suffering from neuromuscular junction autoimmune disorders such as myasthenia gravis or Lambert-Eaton myasthenic syndrome. Since CiTXs are emergent toxins in Europe a systematic monitoring of this family of toxins is needed calling for general guidelines regulating CiTXs concentrations in shellfish.

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FIGURE 1

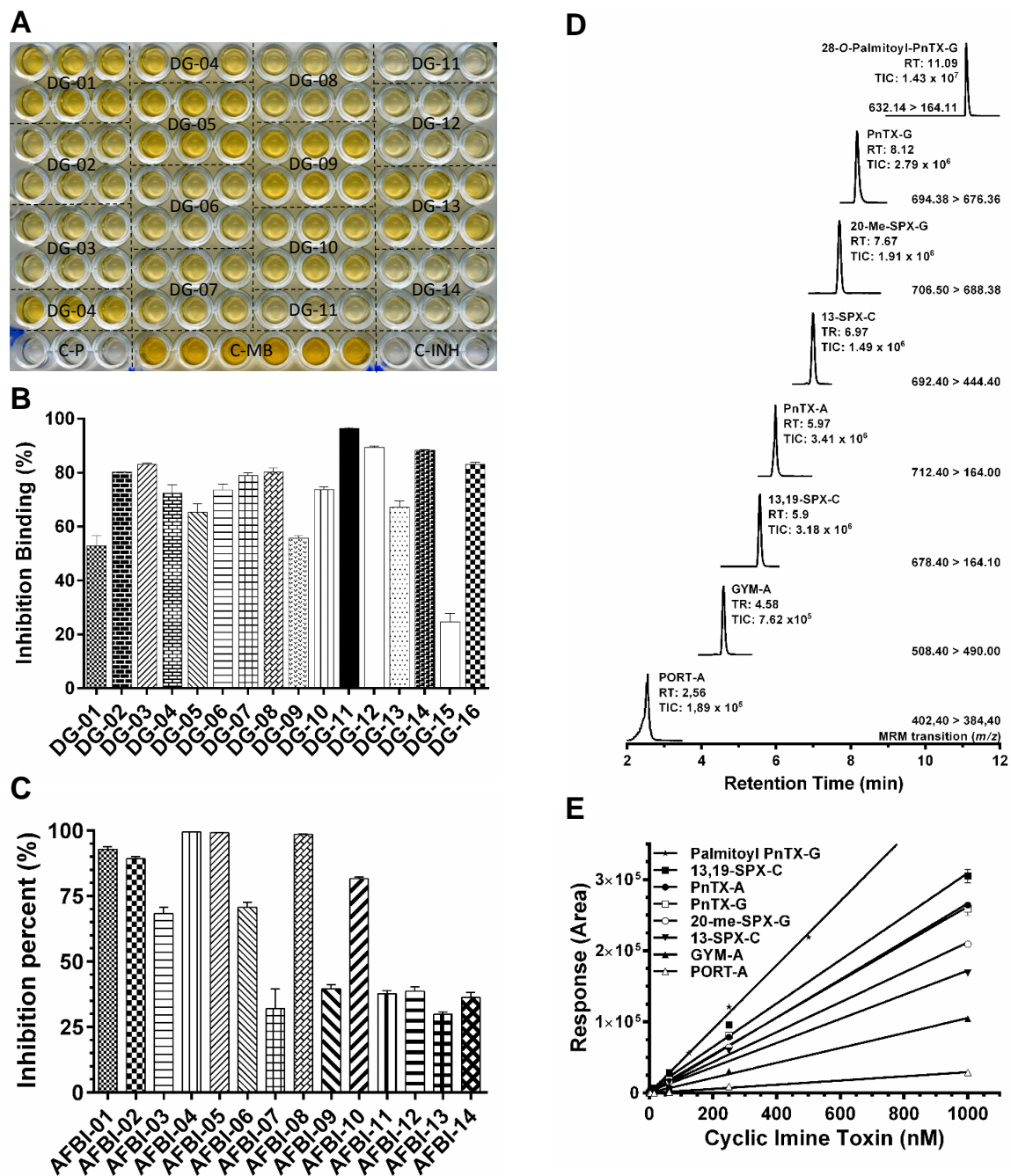


Figure 1

1A. Microplate receptor-binding assay showing the inhibition of biotin- α -bungarotoxin by digestive-gland extracts from mussels collected in Ingril lagoon. DG-samples were analyzed by sextuplicate. **Plate signal (C-P):** control wells processed without membrane coating. **100% signal (C-MB):** control wells in which *Torpedo*-nAChRs were incubated in the absence of toxin or extract samples. **100% inhibition (C-INH):** control wells in which *Torpedo*-nAChRs were incubated with 1×10^{-6} M α -BgTx. **1B.** Inhibition binding potency of digestive-gland extracts from mussels collected in Ingril lagoon. OD₄₉₂ data were expressed in inhibition percent using Equation 1 (Material and methods). Data are mean values \pm SEM of sextuplicate assays of at least two independent experiments. **1C.** Inhibition binding potency of shellfish extracts provided by Agri-food and Biosciences Institute (AFBI). OD₄₉₂ data were expressed in inhibition percent using Equation 1 (Material and methods). Data are mean values \pm SEM of sextuplicate assays of at least two independent experiments. **1D.** UPLC-MS/MS chromatogram showing the simultaneous analysis of seven cyclic imine toxins and 28-*O*-palmitoyl ester of pinnatoxin-G. The MRM conditions, retention time and MS-signal are indicated in the figure. **1E.** Calibration curves for seven cyclic imine toxins and 28-*O*-palmitoyl ester of pinnatoxin-G. 5 μ l of a toxin standard mix in the concentration range of 1 pM to 1 μ M was loaded to the BEH C18 column (see Material and Methods). Toxin quantification was done by integration of the MRM peak area of each analyte. **Abbreviations:** DG: digestive glands C-P: Control plate signal; C-MB: Control membrane signals. C-INH: Control inhibition signals; AFBI: Shellfish samples provided by Agri-food and Biosciences Institute; PORT-A: Portimine-A; GYM-A: gymnodimine-A; 13,19-SPX-C: 13,19-didesmethyl-spirolide-C; PnTX-A: pinnatoxin-A; 13-SPX-C: 13-desmethyl spirolide-C; 20-Me-SPX-G: 20-methyl spirolide-G; PnTX-G: Pinnatoxin-G; Palmitoyl-PnTX-G: 28-*O*-palmitoyl pinnatoxin-G.

FIGURE 2

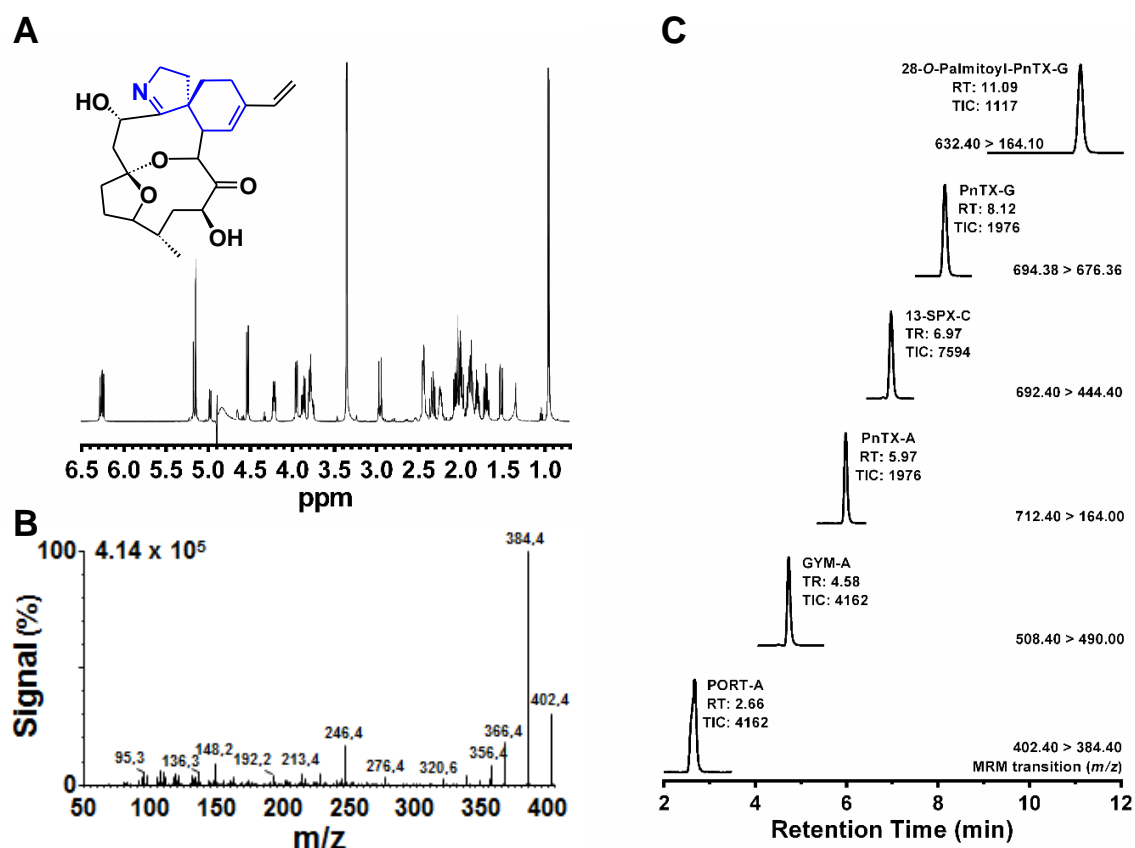
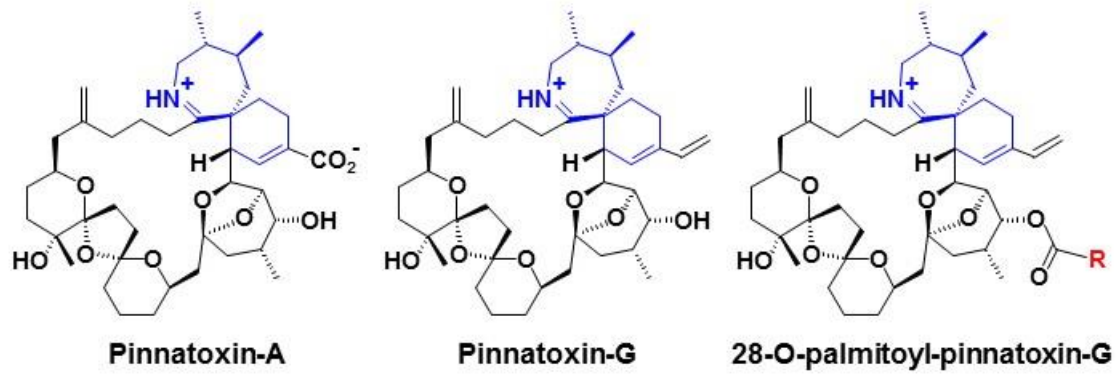


Figure 2

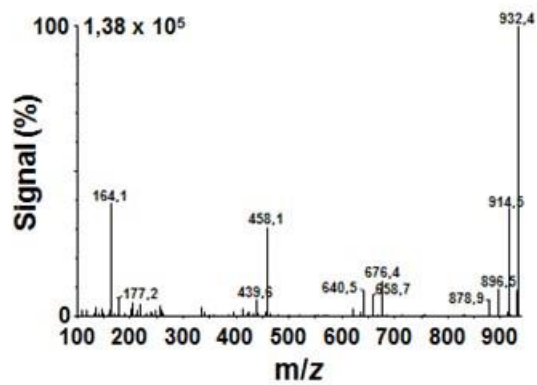
2A. NMR spectrum of portimine-A (inset). NMR was performed at 20°C on a Bruker DRX700 spectrometer equipped with a triple resonance cryoprobe. **2B.** MS/MS spectrum of portimine-A showing its fragmentation pattern at the collision-induced dissociation condition for MRM. **2C.** UPLC-MS/MS chromatogram showing the simultaneous detection of portimine-A (PORT-A), ; gymnodimine-A (GYM-A), pinnatoxin-A (PnTX-A), 13-desmethyl spirolide-C (13-SPX-C) pinnatoxin-G (PnTX-G) and 28-*O*-palmitoyl pinnatoxin-G (Palmitoyl-PnTX-G) in the digestive gland extract sample N°13414-2 collected in Ingril Lagoon (South France). The MRM conditions, retention time and MS-signal are indicated in the figure. *Abbreviations:* m/z : mass-to-charge ratio.

FIGURE 3

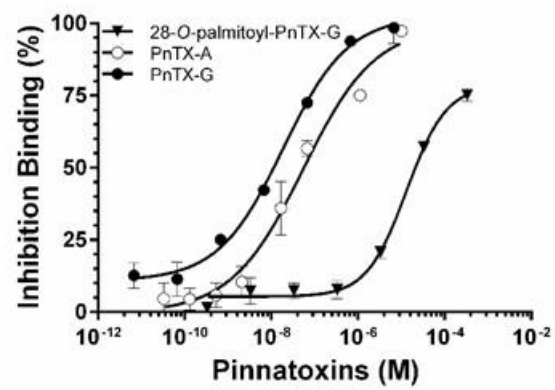
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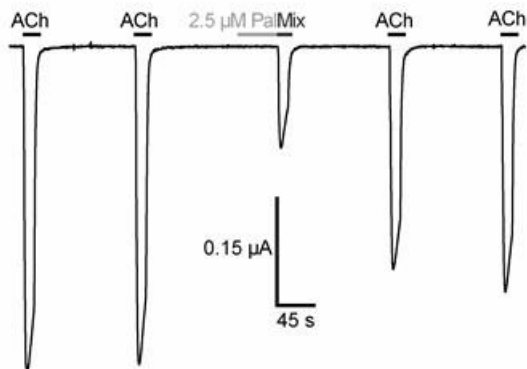
B



C



D



E

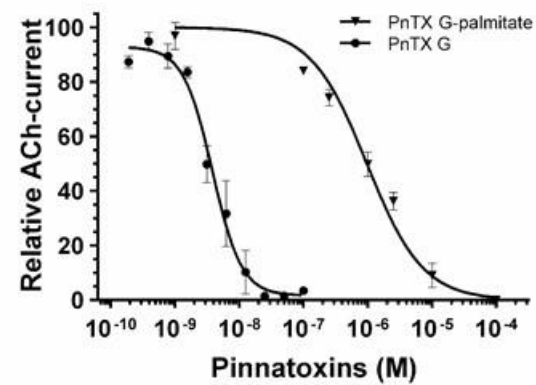


Figure 3

3A. Chemical structure of pinnatoxin-A, pinnatoxin-G and 28-*O*-palmitoyl pinnatoxin-G; R = ester 16:0. **3B** Mass spectrum of 28-*O*-palmitoyl ester of pinnatoxin-G showing its fragmentation pattern at the collision-induced dissociation condition for MRM. **3C.** Inhibition binding potency of 28-*O*-palmitoyl pinnatoxin-G compared to its precursor molecule pinnatoxin-G and to pinnatoxin-A. Each point represents the mean value \pm SEM of three independent experiments using the Microplate-RBAssay. **D.** Antagonistic effect of 2.5 μ M 28-*O*-palmitoyl ester of pinnatoxin-G on *Torpedo*-nAChR of muscle embryonic type recorded using two-electrode voltage clamp electrophysiology. The perfusion protocol was as follows: A clamped oocyte at a holding potential of -60 mV was perfused with 25 μ M ACh (ACh) for 15 s at 150 s intervals. Thereafter, the oocyte was perfused with 2.5 μ M 28-*O*-palmitoyl ester of pinnatoxin-G (Pal) for 45 s and immediately after it was exposed to a mixture of 25 μ M ACh and 2.5 μ M 28-*O*-palmitoyl ester of pinnatoxin-G (Mix). The oocyte was washed with Ringer-Ba and pulses of 25 μ M ACh were recorded twice at 150s intervals. **E.** Concentration-dependent antagonistic effect of 28-*O*-palmitoyl ester of pinnatoxin-G (filled triangles) and pinnatoxin-G (filled circles) on *Xenopus laevis* oocytes having incorporated *Torpedo* muscle-type ($\alpha 1$)₂ $\beta\gamma\delta$ nAChR into their plasma membranes. The peak amplitudes of the ACh-evoked current values (mean \pm SEM; 5 oocytes per concentration) were normalized to control ACh-peak amplitudes recorded from the same oocyte to yield fractional response. The data were fitted to the equation 3 to obtain the IC_{50} values for 28-*O*-palmitoyl ester of pinnatoxin-G (IC_{50} = 969.3 nM; 95% interval confidence: 752.6 – 2129 nM) and pinnatoxin-G (IC_{50} = 3.15 nM; 95% interval confidence: 2.9 – 4.9 nM, Fig 3E). *Abbreviations:* m/z: Mass-to-charge ratio; PnTX-A: Pinnatoxin-A; PnTX-G: Pinnatoxin-G; PnTX-G-palmitate: 28-*O*-palmitoyl pinnatoxin-G.

Table 1.

UPLC-MS/MS Conditions for the Detection/ Quantification of CiTXs.

	PORT-A	GYM-A	13,19-SPX-C	PnTX-A	13-SPX-C	20me-SPX-G	PnTX-G	Palmitoyl-PnTX-G
MRM transition (m/z)	402.4 > 384.4	508.4 > 490.4	678.4 > 164.1	712.4 > 164.0	692.4 > 444.4	706.5 > 688.4	694.4 > 676.4	932.4 > 164.1
Cone voltage (V)	27	50	50	65	60	65	75	90
Collision energy (eV)	24	32	40	48	35	30	30	50
Retention time (min)	2.56	4.58	5.90	5.97	6.97	7.67	8.12	11.09
<i>R</i>²	0.9848	0.9959	0.9965	0.9797	0.9969	0.9970	0.9956	0.9998
LOD (nM)	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.6
LOD (pg) in xolumn	1.0	1.2	1.7	1.8	1.7	1.8	1.7	2.8
LOQ (nM)	2.5	1.0	1.0	1	1	1.2	1.2	1.5
LOQ (pg) in column	5.2	2.5	3.4	3.6	3.5	4.2	4.2	6.9

Abbreviations: PORT-A: portimine-A; GYM-A: gymnodimine-A; 13,19-SPX-C: 13, 19 didesmethyl spirolide-C; PnTX-A: pinnatoxin-A; 13SPX-C: 13-methyl spirolide-C; 20me-SPX-G: 20-methyl spirolide-G; PnTX-G: pinnatoxin-G; Palmitoyl-PnTX-G: 28-*O*-palmitoyl ester of pinnatoxin-G ; MRM: Multiple Reaction-Monitoring; LOD, limit of detection (obtained from low-level standards with a signal-to-noise ratio equal to 3); LOQ, limit of quantification (obtained from low-level standards with a signal-to-noise ratio equal to 10).

Table 2.

Lethality, CiTX content and inhibition binding potency of digestive-gland extracts from mussels collected in Ingril lagoon **that were contaminated with high levels of pinnatoxin-G.**

Extr	Surv. time (min) mean \pm SEM; <i>n</i>	GYM-A (μ g/kg)	13SPX-C (μ g/kg)	20meSPX-G (μ g/kg)	PnTX-A (μ g/kg)	PnTX-G (μ g/kg)	Inh. binding (%) mean \pm SEM
DG-01	67.0 \pm 0.7 ; 3					11.6	52.8 \pm 3.8
DG-02	34.0 \pm 0.7 ; 3					77.4	80.3 \pm 0.1
DG-03	79.3 \pm 0.4 ; 3					52.1	83.1 \pm 0.6
DG-04	< 1440 ; 3	1.4				18.8	72.3 \pm 3.2
DG-05	70.5 \pm 0.5 ; 2					2.1	65.3 \pm 3.2
DG-06	12.0 \pm 2.0 ; 3					40.3	73.6 \pm 2.3
DG-07	4.0 \pm 0.7 ; 3		1.8		1.3	37.2	78.9 \pm 1.0
DG-08	22.7 \pm 2.9 ; 3			1.8	2.1	12.1	80.3 \pm 1.4
DG-09	< 1440 ; 3						55.8 \pm 0.9
DG-10	27.7 \pm 2.2 ; 3					12.0	73.8 \pm 0.9
DG-11	< 1440 ; 3	2.8	3.1				96.4 \pm 0.4
DG-12	32.7 \pm 3.6 ; 3		4.8			2.0	89.3 \pm 0.6
DG-13	< 1440 ; 3					3.5	67.2 \pm 2.4
DG-14	17.0 \pm 0.7 ; 3					60.5	88.4 \pm 0.2
DG-15	< 1440 ; 3		10.6			2.2	24.7 \pm 3.2
DG-16	73.7 \pm 0.4 ; 3					65.5	83.1 \pm 0.7

The survival time is the mean value obtained after intraperitoneal injection of shellfish extracts on three mice (*n*). Abbreviations: GYM-A: gymnodimine-A; 13SPX-C: 13-methyl spirolide-C; 13,19SPXC: 20meSPX-G: 20-methyl spirolide-G; PnTX-A: pinnatoxin-A; PnTX-G: pinnatoxin-G. Toxin quantification was performed by UPLC-MS/MS in the MRM mode. Inhibition binding was performed by microplate-RBAssay. Inhibition binding percent are expressed as mean values of sextuplicate assays of at least two independent experiments.

Table 3.

CiTX content and inhibition binding potency of toxic and control shellfish extracts collected in coastal Regions of 8 European countries

Sample	Species	CiTX Group (µg/kg)					Inh. binding (%)	
		GYM-A	13SPX-C	13,19SPX-C	20meSPX-G	PnTX-A	PnTX-G	mean ± SEM
AFBI-01	Mussel		1.2		20.5	0.9	15.7	93.0 ± 1.8
AFBI-02	Mussel		3.9		16.3	1.5	19.5	89.2 ± 1.7
AFBI-03	Mussel		9.9		11.2		11.1	68.3 ± 5.0
AFBI-04	Mussel		1.7		13.8	2.6	45.4	99.5 ± 0.1
AFBI-05	Mussel		12.9		13.2	1.4	11.0	99.1 ± 0.2
AFBI-06	Mussel		3.1	0.5	6.6	1.4	14.5	70.5 ± 4.0
AFBI-07	Mussel	6.2		5.3				24.5 ± 2.9
AFBI-08	Mussel		10.2	0.5	6.6	1.4	14.5	98.5 ± 0.1
AFBI-09 (C)	Mussel		3.8		2.9		5.8	39.0 ± 2.6
AFBI-10 (C)	Mussel		20.6		5.3			81.7 ± 1.3
AFBI-11 (C)	Mussel		5.6		2.3	1.5	3.8	37.6 ± 2.2
AFBI-12 (C)	Mussel		3.8		3.3	1.0	5.6	38.5 ± 3.7
AFBI-13	Clam		7.0		1.5		2.0	29.8 ± 1.9
AFBI-14	Scallop		4.1		11.6		4.2	35.9 ± 3.2
FR-01 (C)	Clam		0.4		1.4			28.5 ± 2.2
FR-02 (C)	Cockle		5.1					44.9 ± 1.1
FR-03 (C)	Oyster	0.8	8.8				0.1	47.7 ± 3.8
FR-04 (C)	Mussel		5.1				0.1	41.9 ± 1.2

Abbreviations: AFBI: Shellfish samples provided by the Agri-food and Biosciences Institute. AFBI-01 to AFBI-09, AFBI-13 and AFBI-14: Shellfish samples contaminated with regulated marine toxins. AFBI-09 (C) to AFBI-12 (C): control samples non-contaminated with regulated marine toxins. FR-01 (C) to FR-04 (C): Control samples purchased from a shellfish market in Paris Region. GYM-A: gymnodimine-A; 13SPXC: 13-methyl spirolide-C; 13,19SPX-C: 13,19 didesmethyl spirolide-C; 20meSPX-G: 20-methyl spirolide-G; PnTX-A: pinnatoxin-A; PnTX-G: pinnatoxin-G. Toxin quantification was performed by UPLC-MS/MS for cyclic imine

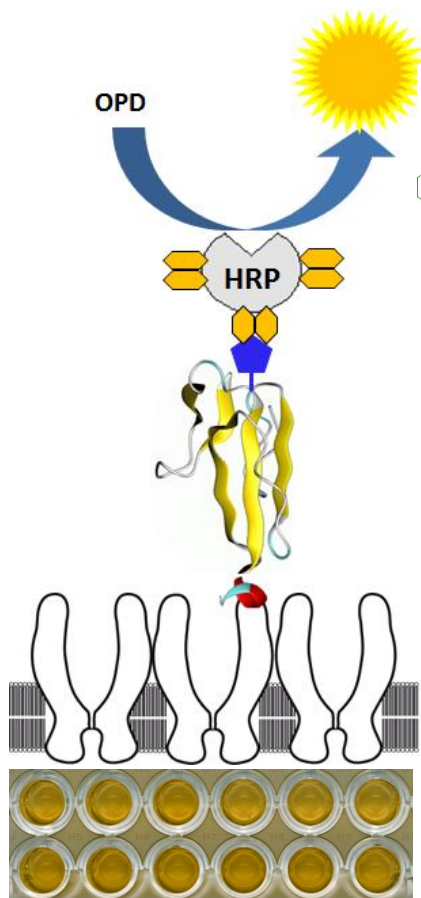
toxins. Inhibition binding was performed by microplate RBA assay. Data are mean values \pm SEM of sextuplicate assays of at least two independent experiments.

Table 4.

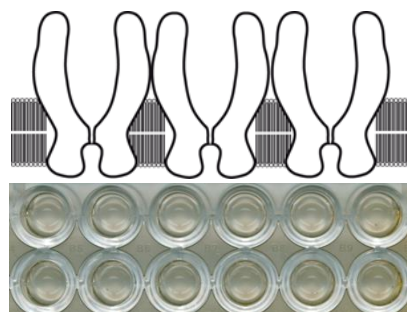
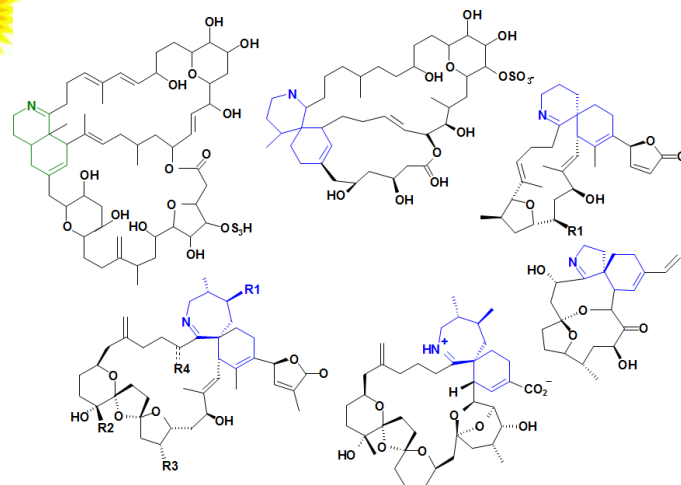
Detection of 28-*O*-palmitoyl ester of pinnatoxin-G and portimine-A in digestive gland extracts of mussels collected at Ingril lagoon.

Extract	PORT-A ($\mu\text{g/kg}$)	GYM-A ($\mu\text{g/kg}$)	13SPX-C ($\mu\text{g/kg}$)	PnTX-A ($\mu\text{g/kg}$)	PnTX-G ($\mu\text{g/kg}$)	Palmitoyl- PnTX-G ($\mu\text{g/kg}$)	Inh. binding (%) mean \pm SEM
13414-2	69.3 \pm 1.9	617.3 \pm 14.3	25.9 \pm 1.5	6.6 \pm 0.2	273.1 \pm 3.8	0.7 \pm 0.1	98.5 \pm 1.2
13/599-2	32.1 \pm 1.1	51.5 \pm 2.8	2.0 \pm 0.1	2.5 \pm 0.2	11.3 \pm 0.1	0.1 \pm 0.0	72.0 \pm 3.0

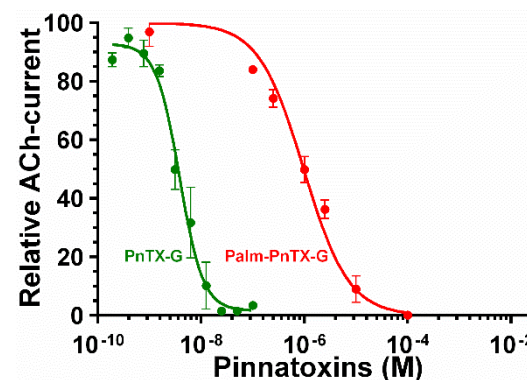
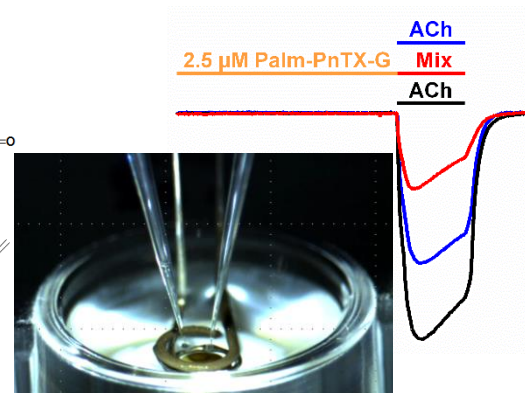
Abbreviations: PORT-A: portimine-A; GYM-A: gymnodimine-A; 13SPX-C: 13-methyl spirolide-C; PnTX-A: pinnatoxin-A; PnTX-G: pinnatoxin-G; Palmitoyl-PnTX-G: 28-*O*-palmitoyl ester of pinnatoxin-G. Toxin quantification was performed by UPLC-MS. Inhibition binding was performed by microplate receptor-binding assay. Inhibition binding percent are expressed as mean values \pm SEM of sextuplicate assays of at least two independent experiments.



NO INHIBITION



INHIBITION



MOA