Potential risk from and prevention of phytoplankton outbreaks in blocking the cooling water system in a nuclear power plant on the Southeast China coast

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The invasion or aggregation of marine biota in coastal waters has become a common problem, threatening the safety of cooling water systems in the context of environmental and climate changes. However, studies on issues of coastal nuclear power plants from the view of marine ecology have mostly focused on the thermal effluent, entrainment effect on organisms, residual chlorine, radioactivity, etc. There is a lack of information on the safety of cooling water systems in nuclear power plants blocked by marine biota. Based on field and historical datasets, this study screened phytoplankton species that may potentially block the cooling water system of the Fuqing nuclear power plant on the Southeast China coast, identified the phytoplankton risk and a risk calendar, and established a catalog of risk phytoplankton species for the cooling water system. The screening criteria were simultaneously required as follows: (1) particle size of cells or group >2000 μm; (2) phytoplankton bloom or aggregation events occurred in surrounding areas; (3) and species abundance maintained at >0.5×10⁵ cells/L in the surrounding area. Nine high-risk species were screened out that can easily form colonies, clusters, and groups, including *Thalassiosira diporocyclus*, *Thalassiosira subtilis*, *Noctiluca scintillans*, *Phaeocystis globosa*, *Pseudo-nitzschia pungens*, *Pseudo-nitzschia delicatissima*, *Skeletonema costatum*, *Skeletonema dohrnii*, and *Paralia sulcata*. Due to small cells or the formation of short chains, some dominant phytoplankton species, even those with advantages in abundance and occurrence, had difficulty threatening the reliability of cooling water systems in nuclear power plants. The outbreak of different phytoplankton species varied daily, and the risk of blockage was highest from April to July. The biological characteristics, ecological mechanisms underlying outbreaks of high-risk phytoplankton species are reviewed and discussed, suggesting a high degree of temporal and spatial variability of blockage risk. Finally, main prevention and control measures used...
against high-risk phytoplankton species are provided to imply new ideas and methods for nuclear power plants. These strategies have very important practical and scientific significance.

KEYWORDS
phytoplankton outbreak, cooling water system, blockage risk, nuclear power plants, ecological mechanism, preventive measures

Highlights

1. Nuclear power plants face a high risk of blockage by marine biota outbreaks in cooling water systems.
2. Blockages risk from phytoplankton was concentrated from April to July.
3. Nine high-risk species of phytoplankton that can easily form colonies, clusters, and groups were screened out.
4. The biological characteristics, ecological mechanisms underlying outbreaks of high-risk phytoplankton species suggested a huge temporal and spatial variability of blockage.
5. Main prevention and control measures used against high-risk phytoplankton species are provided.

1 Introduction

With the depletion of traditional power sources, such as coal and oil, nuclear energy has been developed and utilized by more countries as a clean and efficient alternative (Dagnall, 2005; Yang et al., 2020). Nuclear power plants are mainly located on the coast, and water supply systems using seawater as the final cooling water source are widely used. The main cooling water system, also known as the circulating water system, is the cold end system for operating the power plant (Dagnall, 2005; Liu, 2019). The operational state of the cooling water system directly affects the safety of the power plant (Alsaffar and Zheng, 1999; Ruan, 2015).

The reliability of the cooling water system in nuclear power plants is crucial for their safe operation. The intrusion or aggregation of marine biota into cooling water systems seriously threatens nuclear safety in the context of environmental and climate changes (Alsaffar and Zheng, 1999; ERPI, 2015; Fu et al., 2020). The increasing amount of waste, marine biota, sediment, sea ice, etc. can block cooling water systems (Chen, 2009; Ruan, 2015; Arefev et al., 2015; Yan and Lu, 2016). If a large number of blockages are pumped into the system, clogging the water intake components, it will seriously threaten the normal operation and cause huge hidden dangers to the safe operation of nuclear power plants (Mark, 2012; Kumar et al., 2017; Fu et al., 2020). The World Association of Nuclear Operators (WANO) announced that 38% of the 60 blocking events from 2008 to 2015 were caused by marine biota; 38% were sand, silt, and debris; and 24% were other sources (sediment, plastic). Among the blocking events caused by marine biota since 1980, jellyfish account for 59%, phytoplankton for 11%, fish and shrimp for 10%, and microalgae and aquatic plants for 7% (ERPI, 2015; Han et al., 2018). Therefore, the aggregation and outbreak of marine biota is an important threat to cooling water systems of coastal nuclear power plants (Table 1). Table 1 indicates the most common issue occurs when jellyfish block the water intake filter screen and even cause the reactor to shut down (Matsumura et al., 2005; Mark, 2012; Florin et al., 2013). In addition to jellyfish, other marine biota have caused blockages, such as that caused by an outbreak of pelagic tunicates (Salpa fusiformis) and Euphausiids (Chae et al., 2008). In recent years, coastal nuclear power plants in China have also experienced incidents that threaten their cooling water systems, due to outbreaks of marine biota (Table 1). In July 2014, the water intake of the Hongyanhe nuclear power plant in northern China was blocked by A. aurita, resulting in a 10 day shutdown (Zhang et al., 2019). From December 2014 to February 2015, a bloom of Phaeocystis globosa erupted in the area of the Fangchenggang nuclear power plant, blocking some filtration equipment to varying degrees (He et al., 2019b; Kang et al., 2020). In January 2015, Acetes chinensis, Acetes erythraeus, and Acetes japonicus tripped the single circulating pump in Daya Bay nuclear power plant (Meng et al., 2018). In August 2015, the blocking of the Ningde nuclear power plant in Fujian by benthos, sea cucumbers Acaudina molpadioidea, was related to typhoons (Chen, 2018; Tang, 2018). Therefore, it can be seen that for phytoplankton or algae species, the blockage risk is relatively low, with relatively short time during blockage.

The monitoring and prevention of organisms that typically block coastal nuclear power plants aims to provide an analysis of contributing factors and prevention methods for outbreaks or aggregations (Chae et al., 2008; Moreau et al., 2013; Florin et al., 2013; Arefev et al., 2015; Fu et al., 2020). The ecological mechanisms of outbreaks of some marine biota in areas near nuclear power plants have been preliminarily analyzed. Chae et al. (2008) found...
that the warm surface current that occasionally extends to the nearshore area, makes *Salpa fusiformis* enter the cooling waters of Uljin nuclear power plant in South Korea. Prakash et al. (2012) used hydrological and seaweed migration models and found that the Salem power plant was invaded by seaweed from the intertidal swamp 3 miles away. Florin et al. (2013) traced the mussels, *Mytilus edulis*, that first appeared around the Bothnian Sea nuclear power plant in Sweden. Chen (2018) speculated that the block of the Ningde nuclear power plant may have been caused by the combined effect of ocean currents caused by typhoons, the disturbance of the bottom habitat, and the self-reproduction and growth of *A. molpadioidea*. Consequently, the biggest threat to the cooling water system of nuclear power plants is the uncertainty of outbreaks of marine biota in the context of environmental and climate changes (Deretz, 2007; King et al., 2010; Prakash et al., 2012; Kumar et al., 2017; Fu et al., 2020). In general, these outbreaks have no obvious signs or specific times, which causes problems for studying bio-blocking in cooling water systems. Changes in the marine environment and climate, the impact of thermal effluent, and species distribution are all potential factors for the aggregation or outbreak of marine biota (Kumar et al., 2017; Wu et al., 2017; Chen, 2018; Tang, 2018; Fu et al., 2020; Yu et al., 2021).

Harmful algae blooms (HABs), mainly composed of phytoplankton, are an ecological anomaly caused by the proliferation of tiny unicellular organisms under specific environmental conditions (Berdalet et al., 2017). Fujian in Southeast China has a long coastline with numerous bays. As a result of global change and human activities, marine environment pollution is increasing in this area, making Fujian prone to HABs, with the occurrence of toxic HABs increasing rapidly (Li, 2012). Specifically, in 2012, a toxic *Karenia mikimotoi* bloom in Fujian resulted in the death of numerous more than 70% of cultured abalones and a direct economic loss of 2.1 billion yuan, which had a serious impact on fishery production, coastal tourism, and human health. This was the most serious economic loss caused by HABs reported in China (Lin et al., 2016). Once a large number of HABs organisms gather, the rotating filter screen (usually 3 or 5 mm) of the circulating water filtration system can be blocked, tripping the seawater circulating pump of nuclear power plants (Deretz, 2007; Mark, 2012; Fu et al., 2019; He et al., 2019b; Zhang et al., 2019; Kang et al., 2020). Therefore, as a potential risk organism, phytoplankton may accumulate, with a high number of cells that then deposit on the filter screen surface. Based on field monitoring data of phytoplankton in the 15 km around Fuqing nuclear power plant, combined with historical HABs datasets from the Marine Environmental Quality Bulletin in Fujian that covered 16 years from 2006 to 2021, phytoplankton species were screened and evaluated that may potentially block the cooling water system, a three principle principle of screening risk phytoplankton specie was established, a catalog of risk phytoplankton species and monthly risk calendar were determined. Finally, the ecological mechanisms of high-risk phytoplankton outbreak were reviewed and discussed, and an

| Year | Nuclear power plant | Country    | Caused blockage                  | Operations                        |
|------|---------------------|------------|----------------------------------|-----------------------------------|
| Jan. 1997 | FitzPatrick | USA | Sticklebacks | reactor emergency shutdown |
| Jun. 2003 | Uljin | Korea | *Euphausia pacifica* and *Salpa fusiformis* | reactor emergency shutdown |
| Aug. 2005 | Stockholm’s Oskarshamn | Sweden | *Aurelia aurita* | reactor emergency shutdown |
| Aug. 2005 | Oyster Creek | USA | Sea weeds | reactor emergency shutdown |
| Sep. 2007 | FitzPatrick | USA | Callophora | multiple shutdown |
| Oct. 2008 | Diablo in California | USA | Jellyfish | canyon 2 reactor shutdown |
| Dec. 2009 | Cruas | France | Sea weeds | power reduction of 20% |
| 2011 | Shimane prefecture | Japan | Jellyfish | outlet generation reduced by 6% |
| Jun. 2011 | Torness nuclear power station | UK | Jellyfish | reactor emergency shutdown |
| Jul. 2011 | Hadera | Israel | Jellyfish | facing the risk of shutdown |
| Aug. 2011 | St. Lucie Nuclear Plant in Florida | USA | *Aurelia aurita* | reactor emergency shutdown |
| Jul. 2012 | San Luis Obispo in California | USA | *Salpa fusiformis* | reactor emergency shutdown |
| Sep. 2013 | Stockholm’s Oskarshamn | Sweden | *Aurelia aurita* | reactor emergency shutdown |
| Jul. 2007 | Tianwan in Guangdong | China | Wheat straw | power reduction |
| 2009 | Dayawan in Guangdong | China | Mussels | power reduction |
| Jun. 2011 | Qinshan in Zhejiang | China | Flotsam and jetsam | power reduction |
| Jul. 2014 | Homhuyane in Niaoning | China | Jellyfish | reactor automatic shutdown |
| Dec. 2014 | Fangchenggang in Guangxi | China | *Phaeocystis globosa* | power reduction |
| Jan. 2015 | Linao in Guangdong | China | *Acetes spp.* | power reduction |
| Jul. 2015 | Homhuyane in Niaoning | China | *Aurelia aurita* | power reduction |
| Aug. 2015 | Ningde in Fujian | China | *Acudes molpadioidea* | reactor emergency shutdown |
| Jan. 2016 | Linao in Guangdong | China | *Acetes spp.* | reactor emergency shutdown |
An effective solution for prevention was provided. These strategies have very important practical and scientific significance.

2 Material and methods

2.1 Data recourse

Field monitoring data of phytoplankton species were obtained by water sampling quantitative method that followed the UNESCO Manuals and Guides (Karlson et al., 2010). Twelve stations (Figure 1) were set within 15 km around Fuqing nuclear power plant area according to technical guidelines for sea area utilization demonstration in China (General Administration of Quality Supervision, Inspection and Quarantine of the People’s Republic of China 2010). Water samples (500 mL) from the surface (0.5 m) and bottom (0.5 m above the seabed) were taken in December 2020, February 2021, April 2021, May 2021, June 2021, August 2021, and November 2021. The samples were fixed with 1% neutral formalin and brought back to the laboratory for analysis. Data on phytoplankton species included species composition, abundance, and dominant species. Dominant species were defined as the dominance ≥ 0.02 (Xu, 2006). HABs organisms and outbreak events were screened out from historical datasets of the Marine Environmental Quality Bulletin in Fujian on Southeast China coast between 2006 and 2021. Based on the field monitoring and historical datasets, the potential phytoplankton species of risk in the target area were screened, and the risk of blockage was assessed.

2.2 Analytical method

The treatment and analysis of phytoplankton species of risk were carried out in accordance with the relevant provisions of the Survey of Marine Biology (General Administration of Quality Supervision, Inspection and Quarantine of the People’s Republic of China, 2008), Technical Guidelines for Treatment with Red Tide Disaster (General Administration of Quality Supervision, Inspection and Quarantine of the People’s Republic of China, 2014). Phytoplankton species were identified and counted at 200× to 400× magnification under an inverted optical microscope (Zeiss Z1, Germany). Phytoplankton cell or group or chain length was measured using AxioVision Rel. 4.8 link software connected with microscopy. Phytoplankton abundance was calculated as cells/L. HABs events in Fujian coastal waters over 16 years, including phytoplankton causing species, occurrence, and duration days, were classified. According to the ecological status of phytoplankton around Fuqing nuclear power plant and the HABs events previously recorded, combined with the quantitative assignment risk level assessment method suggested by Tang (2018); Zhang et al. (2019) and Yu et al. (2021), the screening criteria for blockage organisms, catalog of potential phytoplankton species of risk, and the monthly risk calendar were obtained. Furthermore, ecological mechanisms of high-risk species outbreaks and preventive measures are discussed in the context of a comprehensive literature review. This study provides a scientific basis to carry out purposeful monitoring to control marine biota of risk for cooling water system, and also provides a reference for further research.

3 Results

3.1 Ecological status of phytoplankton based on field monitoring around Fuqing nuclear power plant in Southeast China

A total of 147 phytoplankton species were recorded in the waters 15 km around Fuqing nuclear power plant, including 117 species of diatoms that dominated the community, 25 species of dinoflagellates, two species of Cyanobacteria, and two species of Chrysophyta (Figure 2A). In terms of monthly variation, the maximum species richness was 74 in June 2021, which was the
FIGURE 2
Monthly variation of phytoplankton composition (A), species richness (B) and abundance (C) around Fuqing nuclear power plant on southeast China coast.
same as in October 2021. The minimum species richness was 43 in February 2021. The monthly variation of diatom species richness was basically the same as that of the total phytoplankton. However, dinoflagellates had the maximum species richness in May 2021 (Figure 2B). The abundance of phytoplankton was the highest in June 2021, followed by May 2021, and the lowest in October 2021 (Figure 2C).

The monthly variation of dominant species was evident (Table 2). The dominant species taking a greater advantage in months were chain-forming, mainly including *Paralia sulcata*, *Skeletonema costatum*, *Skeletonema dohrnii*, *Pseudo-nitzschia pungens*, *Pseudo-nitzschia delicatissima*, *Eucampia zodiacus*, *Chaetoceros curvisetus*, *Leptocylindrus danicus*, and *Lauderia annulata*. The major dominant species were diatoms. From December 2020 to April 2021, the major dominant species remained as chain-forming *P. sulcata* and *S. costatum*. From May 2021 to July 2021, chain-forming diatoms maintained high abundance and occurrence in the surrounding waters. These species are an important component of potential blocking organisms.

The spatial distribution of major dominant species varied significantly (Figure 3). The most highly abundant species found in waters close to the nuclear power plant were *S. dohrnii* in December, *L. annulata*, *E. zodiacus*, *C. curvisetus* in June, and *P. delicatissima* and *Bacillaria paradoxa* in July. While the most highly abundant species found offshore from the nuclear power plant was *Thalassiosira diporocyclus* in May, *P. pungens* in June, *P. sulcata*, and *S. costatum* in February (Figure 3). Specifically, the maximum abundance of *S. dohrnii* in December was 1.10×10^5 cells/L with an average of 0.57×10^5 cells/L; the maximum abundance of *L. annulata*, *E. zodiacus*, *C. curvisetus* in June were 0.54×10^5 cells/L, 0.90×10^5 cells/L, 1.10×10^5 cells/L, respectively; and *P. delicatissima* and *B. paradoxa* in July had the maximum abundance of 1.07×10^5 cells/L and 0.97×10^5 cells/L, respectively. Subsequently, *T. diporocyclus* in May had the maximum abundance of 1.13×10^5 cells/L with an average of 3.01×10^4 cells/L, while *P. pungens* in June had 1.11×10^5 cells/L in maximum. The maximum abundance of *P. sulcata*, and *S. costatum* in February were 1.03×10^5 cells/L and 1.11×10^5 cells/L, respectively.

| Dominant species and dominance | Time     |
|--------------------------------|----------|
|                                | Dec. 2020| Feb. 2021| Apr. 2021| May 2021| Jun. 2021| Jul. 2021| Oct. 2021|
| Paralia sulcata                | 0.21     | 0.39     | 0.20     | ND      | < 0.02   | 0.13     | 0.25     |
| Skeletonema costatum           | < 0.01   | 0.37     | 0.17     | ND      | < 0.02   | < 0.01   | 0.20     |
| Skeletonema dohrnii            | 0.31     | < 0.01   | ND       | ND      | ND       | ND       | ND       |
| Coscinodiscus bipartitus       | 0.06     | 0.04     | 0.05     | ND      | < 0.01   | < 0.01   | < 0.01   |
| Coscinodiscus jonesianus       | ND       | ND       | ND       | ND      | < 0.01   | < 0.01   | < 0.01   |
| Nitzschia longissima v. reversa| < 0.01   | < 0.01   | < 0.01   | < 0.01  | < 0.01   | < 0.01   | < 0.01   |
| Nitzschia closterum            | < 0.01   | 0.04     | 0.07     | < 0.01  | < 0.01   | < 0.01   | < 0.01   |
| Nitzschia sigma                | 0.04     | < 0.01   | < 0.01   | 0.04    | < 0.01   | 0.03     | < 0.01   |
| Pleurosigma affine             | 0.02     | 0.02     | < 0.01   | ND      | < 0.01   | < 0.01   | < 0.01   |
| Pleurosigma elongatum          | 0.04     | < 0.01   | < 0.01   | ND      | < 0.01   | < 0.01   | < 0.01   |
| Ditylum brightwellii           | 0.02     | ND       | ND       | ND      | ND       | 0.06     | < 0.01   |
| Cyclotella caspia              | < 0.01   | 0.02     | ND       | ND      | ND       | ND       | ND       |
| Leptocylindrus danicus         | ND       | ND       | ND       | ND      | < 0.01   | < 0.01   | < 0.01   |
| Leptocylindrus minus           | ND       | ND       | ND       | ND      | < 0.02   | 0.13     | < 0.01   |
| Pseudo-nitzschia delicatissima| < 0.01   | < 0.01   | 0.02     | < 0.01  | 0.02     | 0.06     | < 0.02   |
| Pseudo-nitzschia pungens       | < 0.01   | < 0.01   | 0.02     | 0.02    | 0.07     | 0.02     | < 0.02   |
| Chaetoceros curvisetus         | ND       | < 0.01   | < 0.01   | ND      | 0.14     | < 0.01   | < 0.01   |
| Eucampia zodiacus              | ND       | < 0.01   | < 0.01   | ND      | 0.18     | < 0.01   | < 0.01   |
| Eucampia cornuta               | ND       | ND       | ND       | ND      | 0.05     | < 0.01   | ND       |
| Guinardia striata              | ND       | ND       | ND       | ND      | 0.02     | < 0.01   | < 0.01   |
| Lauderia annulata              | ND       | ND       | ND       | ND      | 0.21     | < 0.01   | < 0.01   |
| Bacillaria paradoxa            | ND       | ND       | ND       | ND      | 0.13     | < 0.01   | < 0.01   |
| Asterionella japonica          | ND       | ND       | ND       | ND      | 0.06     | < 0.01   | < 0.01   |
| Thalassionema nitzschiosides   | < 0.01   | < 0.01   | < 0.01   | ND      | < 0.01   | < 0.01   | 0.02     |
| Thalassionema nitzschiosides v. parva| ND | < 0.01   | < 0.01   | ND      | < 0.01   | < 0.01   | 0.02     |
| Thalassiosira diporocyclus     | ND       | ND       | 0.68     | ND      | ND       | ND       | ND       |

ND indicates non-detected.
FIGURE 3
Horizontal distribution of 10 major phytoplankton dominant species around Fuqing nuclear power plant on southeast China coast.
3.2 Phytoplankton bloom events and annual changes based on historical datasets in the Fujian coast, Southeast China

Over 16 years, HABs outbreaks along the coast of Fujian in Southeast China were seasonal (Figure 4B), with 90.5% occurring from April to July. The highest incidence period of HABs was in May and June, with dinoflagellates as the majority of HABs causing species. The occurrence of HABs in May and June accounted for 60.0% of the total number, while the duration days accounted for 46.4% during the whole year. HABs showed a downward trend from 2014 to 2021, and there was a small increase by 2021 (Figure 4A). There were 13 HABs causing species from 2006 to 2021 in common, including S. costatum, Chaetoceros sp., Thalassiosira sp., E. zodiacus, P. pungens, Prorocentrum micans, Karenia mikimotoi, Scrippsiella trochoidea, Noctiluca scintillans, Gymnodinium catenatum, Chattonella marina, Akashiwo sanguine, and Phaeocystis globosa. The phytoplankton species that repeatedly cause HABs were P. micans, S. costatum, Thalassiosira sp., N. scintillans, and A. sanguine. Specifically, P. micans caused outbreaks from April to July each year along the coast of Fujian in Southeast China (Figure 4B).

3.3 Blockages caused by phytoplankton species: Screening criteria, catalog, and monthly calendar of potential risk species

According to the HABs events previously recorded within 16 years along the coast of Fujian, 87 phytoplankton species belonging to 39 genera of four phyla were selected as HABs organisms around Fuqing nuclear power plant (Figure 5A). Of these, 68 species of 27 genera belonging to diatoms accounted for 78% of the total species; 15 species of 10 genera from dinoflagellate accounted for 17%; two species of two genera from Chrysophyta accounted for 2%; and two species of one genus from cyanobacteria accounted for 3% (Figure 5A). Based on the diameter of the drum net of the circulating water system (usually 3 or 5 mm) in nuclear power plants,
combined with blockage events in cooling water system documented by marine biota, the screening criteria for potential risk phytoplankton species were as follows: (1) particle size of cells or groups >2000 μm; (2) phytoplankton bloom or aggregation events recorded in surrounding areas; and (3) high species abundance of >0.5×10^5 cells/L in the surrounding sea area. Notably, the entire three principles should be simultaneously required instead of optional selection. Consequently, 38 potential risk species among the above 87 HABs organisms and a monthly risk calendar were established for Fuqing nuclear power plant (Table 3). This includes species with high dominance over multiple months, such as *S. costaum*, *S. dohrnii*, *P. sulcata*, *P. pungens*, and *L. danicus*, as well as other non-dominant species with high abundance in the adjacent waters close to the nuclear power plant. Based on field monitoring and historical datasets, the phytoplankton of risk in the sea area around Fuqing nuclear power plant could be roughly determined (Table 3). Furthermore, the HABs events recorded in the adjacent waters or offshore from the nuclear power plant were referred to (Figure 5B). The potential risk from phytoplankton species may exist in a certain area in real time. Blockages in cooling water systems may occur when phytoplankton species abundance reaches a high level. Phytoplankton species that can easily form colonies, clusters, filaments, and groups were selected as high-risk organisms (Table 3).
| Species                        | Dec. 2020 | Feb. 2021 | Apr. 2021 | May 2021 | Jun. 2021 | Jul. 2021 | Oct. 2021 | Morphology     | Particle size (cm) | Abundance (10^5 cells/L) | Comprehensive risk level |
|-------------------------------|-----------|-----------|-----------|-----------|-----------|-----------|-----------|----------------|----------------------|--------------------------|--------------------------|
| Asterionella japonica         |           |           |           |           |           |           |           | catenoid colony | < 0.03               | 0.03–0.77                | ★                       |
| Bacillaria paradoxa           |           |           |           |           |           |           |           | catenoid colony | 0.01–0.03           | 0.03–0.97                | ★                       |
| Chaetoceros affinis           |           |           |           |           |           |           |           | long chains    | 0.01–0.03           | 0.01–1.10                | ★                       |
| Chaetoceros curvisetus        |           |           |           |           |           |           |           | long chains    | > 0.1               | 0.03–1.10                | ★★                      |
| Chaetoceros tortissimus       |           |           |           |           |           |           |           | long chains    | 0.01–0.03           | 0.01–1.00                | ★                       |
| Chaetoceros densus            |           |           |           |           |           |           |           | long chains    | > 0.1               | 0.01–1.00                | ★★                      |
| Eucampia zodiacus             |           |           |           |           |           |           |           | catenoid colony | 0.01–0.03           | 0.07–0.90                | ★                       |
| Guinardia striata             |           |           |           |           |           |           |           | catenoid colony | > 0.5               | 0.02–0.90                | ★★                      |
| Guinardia delicatula          |           |           |           |           |           |           |           | catenoid colony | 0.01–0.03           | 0.02–0.91                | ★                       |
| Lauderia annulata             |           |           |           |           |           |           |           | long chains    | 0.01–0.03           | 0.02–0.66                | ★                       |
| Leptocylindrus danicus        |           |           |           |           |           |           |           | long chains    | > 0.1               | 0.02–0.93                | ★★                      |
| Leptocylindrus minus          |           |           |           |           |           |           |           | long chains    | > 0.1               | 0.02–0.55                | ★★                      |
| Odontella mobilensis          |           |           |           |           |           |           |           | large cell     | 0.0–0.03             | 0.01–0.65                | ★                       |
| Odontella regia               |           |           |           |           |           |           |           | large cell     | > 0.1               | 0.02–0.55                | ★                       |
| Odontella sinensis            |           |           |           |           |           |           |           | large cell     | > 0.1               | 0.2–0.75                 | ★                       |
| Paralia sulpata               |           |           |           |           |           |           |           | long chains    | > 0.1               | 0.7–1.03                 | ★★                      |
| Pseudo-nitzschia pungens      |           |           |           |           |           |           |           | long chains    | > 0.1               | 0.04–1.1                 | ★★                      |
| Pseudo-nitzschia delicatissima|           |           |           |           |           |           |           | long chains    | > 0.1               | 0.04–1.07                | ★★                      |
| Rhizosolenia setigera         |           |           |           |           |           |           |           | long cell      | > 0.5               | 0.02–0.61                | ★                       |
| Rhizosolenia alatae           |           |           |           |           |           |           |           | long cell      | > 0.5               | 0.02–0.61                | ★                       |
| Skeletonema costatum          |           |           |           |           |           |           |           | long chains    | > 0.1               | 0.7–1.11                 | ★★                      |
| Skeletonema dohrnii           |           |           |           |           |           |           |           | long chains    | > 0.1               | 0.2–1.10                 | ★★                      |
| Stephanopyxis palmeriana      |           |           |           |           |           |           |           | long chains    | 0.01–0.03           | 0.02–1.03                | ★                       |
| Thalassiosira diporocyclus    |           |           |           |           |           |           |           | colony         | 1–2                 | 0.01–1.13                | ★★                      |
| Thalassiosira rotula          |           |           |           |           |           |           |           | long chains    | > 0.1               | 0.02–0.83                | ★★                      |
| Thalassiosira subtilis        |           |           |           |           |           |           |           | colony         | 1–2                 | 0.02–0.71                | ★★                      |
| Thalassiothrix frauenfeldii   |           |           |           |           |           |           |           | catenoid colony | 0.01–0.03           | 0.02–0.72                | ★                       |
| Thalassionema mitzschneideri   |           |           |           |           |           |           |           | catenoid colony | 0.01–0.03           | 0.07–0.62                | ★★                      |
| Akashiwo sanguinea            |           |           |           |           |           |           |           | large cell     | < 0.003             | 0.03–0.62                | ★                       |
| Ceratium furca                |           |           |           |           |           |           |           | large cell     | 0.01–0.03           | 0.03–0.62                | ★                       |

(Continued)
4 Discussion

4.1 Assessment on potential blocking risk from phytoplankton species for a nuclear power plant on the Southeast China coast

In terms of the relationship between nuclear power plants and the marine environment, the impact of thermal effluent on marine biology is the most well studied (Kaplan et al., 2016; Dvorniko et al., 2017; Wei et al., 2018). However, with changes in the marine environment and climate, the safety of cooling water systems has become a new research area. The safe operation of cooling water systems requires establishing a catalog of risk organisms in the surrounding area, and developing monitoring systems, early warning technologies, and preventive countermeasures (Moreau et al., 2013; Tang et al., 2017; Zhang et al., 2018; Kirsti et al., 2018; Lu et al., 2018; Yu et al., 2021). It is a complex problem, and factors such as population composition, distribution, abundance of marine biota, trends and intensity into water intake components are important issues for evaluating the threat or risk. Tang (2018) used the Analytic Hierarchy Process, which initially proposed the principle of Ningde nuclear blockage causing organisms screening and determined the possibility blockage in accordance with the criteria: 0.000 to 0.249 for low risk, 0.250 to 0.499 for medium risk, 0.500 to 0.749 for high risk, 0.750 to 1.000 for very high risk. Some studies have established a catalog of risk organisms and a biological screening index system for jellyfish. For example, Zhang et al. (2019) according to relative biomass, i.e. the number of jellyfish caught per net per hour, suggested the risk level was “High” when relative density of jellyfish was ≥ 500 ind./(net.h), called the red level; level “Medium” was when the relative density was > 300 but < 500 ind./(net.h), called the yellow level; level “Low” was > 100 and < 300 ind./(net.h), called the blue level. Subsequently, over the past 5 years, several monitoring plans, early warning technologies, and decision support systems have been developed for marine biota that block cooling water systems (Tang et al., 2017; Zhang et al., 2018; Kirsti et al., 2018; Lu et al., 2018; Meng et al., 2018).

In present study, utilizing the quantitative risk level evaluation method suggested by Tang (2018); Zhang et al. (2019) and Yu et al. (2021), the risk level of 38 potential risk phytoplankton species preliminarily screened were assigned to quantitative values of 1, 2, 3, and 4, with corresponding levels of low, medium, high, and extremely high. Consequently, a comprehensive quantitative analysis showed that the organisms of potential risk to Fuqing nuclear power plant mainly appeared between April and July, with occasional high abundance aggregation events occurring in May (Figure 6). At the species level, there were nine major high-risk species, including T. diporocyclus, T. subtilis, N. scintillans, Phaeocystis globosa, P. pungens, P. delicatissima, S. costatum, S. dohrnii, and P. sulcata (Figure 6A). There were seven species of medium risk, 22 species of low risk, and no species of extremely high risk. Temporally, the high-risk species were most abundant in June and July, and the risk of blockage was highest. In the other months, there was a low to medium risk of blockage (Figure 6B).

| Species                     | Dec. 2020 | Feb. 2021 | Apr. 2021 | May 2021 | Jun. 2021 | Jul. 2021 | Oct. 2021 | Morphology | Particle size (cm) | Abundance (10⁵ cells/L) | Comprehensive risk level |
|-----------------------------|-----------|-----------|-----------|----------|-----------|-----------|-----------|-------------|----------------------|--------------------------|--------------------------|
| Ceratium fusus              |           |           |           |          |           |           |           | large cell   | 0.01–0.03           | 0.03–0.61               | ★                       |
| Noctiluca scintillans       |           |           |           |          |           |           |           | large cell   | 1–2                 | 0.02–0.68               | ★★                      |
| Gymnodinium catenatum       |           |           |           |          |           |           |           | long chains  | < 0.4             | 0.03–0.66               | ★                       |
| Karenia mikimotoi           |           |           |           |          |           |           |           | large cell   | < 0.003            | 0.03–0.66               | ★                       |
| P. subtilis                 |           |           |           |          |           |           |           | large cell   | < 0.03             | 0.03–0.63               | ★                       |
| Scrippsiella trochoidea     |           |           |           |          |           |           |           | large cell   | < 0.003            | 0.03–0.62               | ★                       |
| Trichodesmium erythraeum    |           |           |           |          |           |           |           | long cell    | < 0.4             | 0.03–0.63               | ★                       |
| Phaeocystis globosa          |           |           |           |          |           |           |           | granular colony | < 3               | 0.01–0.38               | ★★                      |

The symbol ★ suggests the risk level of blockage from marine phytoplankton species, the more number of ★ showing, the high the blockage risk being.
Notably, the nine high-risk species in this study were only treated as a risk for the cooling water systems in nuclear power plants. For the entire ecosystem, these species had corresponding niches, and there was no distinction between mentions of high, medium, and low risk. The investigation and evaluation of marine biota should be guided by the data requirements of the calendar of marine organism outbreak risk (Mark, 2012; Florin et al., 2013; Tang, 2018; Zhang et al., 2019; He et al., 2019b; Yu et al., 2021).

4.2 Characteristics and ecological mechanism of outbreaks or aggregations of potential risk phytoplankton species for a nuclear power plant on the Southeast China coast

The increasing severity of HABs in the coastal waters of China has a close relationship with the intensified eutrophication, especially in the estuaries and bays (GEOHAB, 2010; Yu and
Chen, 2019). Besides, the alteration of natural habitat by mariculture industry and other anthropogenic activities also contributes to the occurrence of HABs events (Figure 7), such as the green tides in the southern Yellow Sea. Other factors like climate change also influence the distribution, dynamics, and impacts of HABs in the coastal waters of China directly or indirectly, through the effects on water temperature, stratification, and currents (Figure 7) (Anderson et al., 2012; Glibert et al., 2018; Yu and Chen, 2019).

The presence of phytoplankton species was the main cause of HABs. Appropriate external conditions, including physical, chemical, biological, hydrological, meteorological, and other factors were the environmental cause of HABs. The formation of HABs requires certain species and specific environmental factors (Figure 7). However, predicting and understanding the ecological mechanism of HABs formations remains difficult because it has a high degree of temporal and spatial variability (GEOHAB, 2010; Anderson et al., 2012; Glibert et al., 2018; Yu and Chen, 2019). In this study, the biological characteristics and ecological mechanism of outbreaks for major high-risk phytoplankton species are discussed below.

4.2.1 Biological properties and mechanisms underlying outbreak of *Phaeocystis globosa*

*Phaeocystis globosa* belongs to the genus *Phaeocystis* of Haptophyta. It is a major causative agent of HABs in oceans globally, featuring a complex polymorphic life cycle, alternating between free-living solitary cells (3–8 μm) and colonial cells (maximum 3 cm) (Shen and Qi, 2021). A colony is the dominant morphotype during *P. globosa* blooms (Figures 8A-C). The sticky and elastic carbohydrate skins of colonies can block the cooling water system of nuclear power plants (He et al., 2019b; Wang et al., 2021). Colony skin reportedly blocked the entrance of the cooling water filtration system of the Fangchenggang Nuclear Power Plant in China (Kang et al., 2020). According to the historical datasets and field investigation of this study, *Phaeocystis* blooms were not found in Fuqing nuclear power plant but were detected in July and October in lower abundance (0.07×10⁵ cells/L, 0.11×10⁵ cells/L), so may become a germplasm source of *Phaeocystis* blooms (Table 3, Figure 6). In general, *P. globosa* is prone to bloom in coastal waters with abundant or eutrophic nutrients. The restriction of nitrogen and phosphate will inhibit the growth and life cycle transformation of *P. globosa*. *P. globosa* colonies are more susceptible to phosphate restriction than free single cells, so colonies cannot form at low phosphate concentrations (Wang, 2006). When nitrate is the nitrogen source, colonies can easily develop, while other forms of nitrogen lead to the dissipation of colonies (Tungaraza et al., 2003). The colonies can be formed in a wide range of salinities, but temperature plays a key role in the colony size (He et al., 2019a). Higher external temperatures will accelerate colony
rupture, and make single cells rapidly form large colonies. Specifically, Wang and Tang (2010) found that the maximum size of the colonies can be obtained at a salinity of 28 and a temperature of 25°C or at a salinity of 25 and temperature of 20°C. In addition, the formation of *P. globosa* colonies also requires light intensity of more than 40 μmol/(m²s) (Deng, 2013). Water disturbance can promote the proliferation of *P. globosa* cells and prolong the growth time (Li et al., 2010; Cai et al., 2011). Therefore, water disturbance caused by monsoons may be one of the main factors of *P. globosa* blooms (Hai et al., 2010). Diatoms are often associated with the occurrence and dissipation of *P. globosa* blooms, especially chain diatoms, including *Chaetoceros* sp., *Skeletonema costatum*, and *Pseudonitzschia* sp. (Peperzak et al., 1998; Hamm and Rousseau, 2003). The formation and increase of *P. globosa* colonies causes a discrepancy in particle size between the food and the feeders, inhibits the feeding efficiency, and protects the colonies (Nejstgaard et al., 2007; Wang et al., 2011). Moreover, *P. globosa* can improve the inner space of colony formation when facing the feeding of mesozooplankton. In summary, the frequent occurrence of *P. globosa* blooms may be attributed to the competitive advantage of its life cycle transformation and colony formation.

### 4.2.2 Biological properties and mechanisms underlying outbreak of *Noctiluca scintillans*

*N. scintillans*, belonging to the genus *Noctiluca* of Pyrrophyta, is a unicellular species with a nearly spherical body and a cell diameter of 150–2000 μm, visible to the naked eye, especially giving off blue light at night (Figures 8D-F). It has both plant
attributes and some animal functions; therefore, its floristic classification includes red heterotrophic species and green amphotrophic species (Mikaelyan et al., 2014; Song et al., 2016). *N. scintillans* blooms frequently occur on the China coast, and its occurrence can reach about 50% of the total HABs (Tang et al., 2006). During the blooms, *N. scintillans* are concentrated on the surface or near the surface (Turkoğlu, 2013), distributed in strips or blocks of dark pink and orange, with bloom abundance of >10^6 cells/L (Qian et al., 1994). The *N. scintillans* cell is large and has colloidal substances on its surface and so it may block the cooling water system and cause potential safety hazards during outbreaks. This species has frequently caused HABs in Fujian since 2007, and near the Yangtze River Estuary in 2021, which can easily block Fuqing nuclear power plant on Southeast China coast (Table 3, Figure 6).

The population dynamics of *N. scintillans* are affected by many factors, such as the environment, biological characteristics, and population structure. Water temperature is the main factor affecting the seasonal variation of *N. scintillans* (Harrison et al., 2011). A study in Jangmok Bay, South Korea, showed that *N. scintillans* abundance decreased sharply when the water temperature was >28°C (Jang et al., 2010). Le Fèvre and Grall (1970) reported that, as *N. scintillans* is a cold-water species, high temperatures would inhibit population growth (e.g. Huang et al., 1997; Tada et al., 2004). *N. scintillans* is a heterotrophic dinoflagellate, and its population dynamics will also be affected by food availability. Harrison et al. (2011) reported that the abundance of zooplankton also controlled *N. scintillans* abundance due to competition. Miyaguchi et al. (2006) found that the large number of sexual-multiplication cells is one of the necessary conditions for *N. scintillans* blooms. Excessive nutrients or eutrophication are important prerequisites and foundations for *N. scintillans* outbreaks (Turkoğlu, 2013; Balıarsingh et al., 2016). Eutrophication provides a rich nutritional basis for the phytoplankton, which is beneficial for the proliferation of *N. scintillans* (Xu, 2009). Furthermore, Miyaguchi et al. (2006) documented that wind speed and precipitation can significantly affect the abundance of *N. scintillans*. Several recent studies have demonstrated the potential of satellite remote sensing in assessing *N. scintillans* blooms (Van Mol et al., 2007; Qi et al., 2019). *N. scintillans* cells have strong absorption in the blue–green wavelengths and strong scattering in all wavelengths (Astórcena et al., 2005), leading to distinguishable spectral reflectance shapes, for example, a sharp reflectance increases from 520 to 600 nm (Van Mol et al., 2007), where the enhanced reflectance continues to near-infrared wavelengths.

4.2.3 Biological properties and mechanisms underlying outbreak of *Thalassiosira diporocyclus*

*Thalassiosira diporocyclus* is a diatom with a rounding, convex, or flat, rarely concave, cell shell, and a circle of small or long thorns arranged radially around the shell surface (Hasle and Vertsen, 1997). In the center of the shell surface, there is a hole for secreting colloid, from which thick or thin colloidal filaments are generated, connecting the cells into a straight or slightly curved chain group (Figures 8G-I). Many *T. diporocyclus* cells live together in a colloidal colony (1–2 cm), while few single cells live. The colloidal colony of *T. diporocyclus* is solid spherical or ellipsoidal in shape and tawny or milky white. The tawny group is a normal healthy colony, while the milky white group is a colony in decline. Each colloidal colony is generally composed of *T. diporocyclus* cells distributed in the outer layer and pennatae diatoms embedded in the colloid (Figure 8G). This species must be identified by electron microscope, so previous historical datasets are often missed or misjudged. In this study, *T. diporocyclus* was listed as a high-risk species for blockages (Table 3, Figure 6). *T. diporocyclus* often developed into HABs along the coast of Fujian in recent years (Figure 3). Therefore, it was classified as a high-risk organism for Fuqing nuclear power plant on Southeast China coast.

In January 2002, a *T. diporocyclus* bloom occurred in the waters around Nanao Island on the southeast coast of China, which was the first record in China of an outbreak of *T. diporocyclus* (Chen et al., 2004). *T. diporocyclus* has a low bloom occurrence, originating from the open sea. The ecological mechanism of *T. diporocyclus* outbreaks may be related to the abnormal increase of water temperature caused by El Niño (Nagai et al., 1995; Miyahara et al., 1996; Chen et al., 2004). In normal years, the wind in autumn and winter seasons along the coast of China is generally northwesterly, forcing strong coastal currents from north to south. If the weather is abnormal and the northwest wind weakens while the southeast wind prevails, the southward coastal current may lessen accordingly, providing an opportunity for *T. diporocyclus* from the open sea to invade the coast of China (Yin et al., 1999). The occurrence of *T. diporocyclus* blooms is closely related to low salinity (23–28). Consequently, the sudden decrease in salinity during El Niño may also be one of the factors in the *T. diporocyclus* outbreak (Nagai et al., 1995; Miyahara et al., 1996; Chen et al., 2004).

4.2.4 Biological properties and mechanisms underlying outbreak of *Pseudo-nitzschia pungens*

*Pseudo-nitzschia pungens* is a typical species of the genus *Pseudo-nitzschia*, a chain-forming cosmopolitan marine pennate diatom (Casteleyn et al., 2008; Kimon et al., 2012). Cells of *P. pungens* are 74–142 µm long and 2.9–4.5 µm wide, with 9–15 striae in 10 µm and 9–15 fibulae in 10 µm, mainly lanceolate in both valve and girdle views, forming a long stepped colony (Hasle and Vertsen, 1997; Li, 2005) (Figures 8J, K).Blooms have been reported in Dalian, Qingdao, the Yangtze River estuary, Xiamen Port, and many harbors in the northern South China Sea (Lu and Qi, 1992; Zou et al., 1993; Qi et al., 1994; Yang and Hodgkiss, 2002). In addition, *P. pungens* is also one of the important toxigenic species, and reports from New Zealand (Rhodes et al., 1996), Washington (Trainer et al., 1998), Montreal Bay, and...
California (Bates et al., 1998) found that it produces toxicants and is poisonous to aquatic organisms and humans. Chinese scholars have also tested the toxins of *P. pungens* in the East China Sea, but no domoic acid has been detected (Li, 2005).

Bates et al. (1998) identified that *P. pungens* blooms usually occur in summer, in the form of a chain-like multicellular population, while Dorich et al. (1997) reported that *P. pungens* is euryhaline, with a tolerance range of 0–36, which confers an advantage in salinities of 15–20. *P. pungens* has a strong ability to absorb inorganic nitrogen, which may make it competitive over other species for nitrogen to form blooms. However, its ability to absorb phosphate was weak, which may be a factor limiting its proliferation in the natural environment (Zhang and Zou, 1997). Experiments suggested that the nitrogen and phosphate demand for *P. pungens* growth did not show a proportional increase, but a certain range was required, with a relatively rapid growth in the range of 10–32 for c(N)/c(P). When c(N)/c(P) = 5 or c(N)/c(P) = 100, *P. pungens* growth was significantly inhibited (Lu et al., 2006). A long-term field observation on the south China coast showed that *P. pungens* blooms were positively correlated with nitrite and total nitrogen (Chen et al., 2002). Overall in this study, *P. pungens* was a high-risk, potential blocking organism in the sea area around the Fujing nuclear power plant on Southeast China coast (Table 3, Figure 6).

### 4.3 Preventive measures against high-risk phytoplankton species for a nuclear power plant on the Southeast China coast

According to the Important Operation Event Report of WANO and relevant datasets, nuclear power plants usually adopt the “prevention first and control combined” method for blockages of marine biota in the cooling water system (Zhan and Zhao, 2005; ERPI, 2015; Tang et al., 2017; He et al., 2019b). The specific methods are mainly divided into physical, biological, and chemical prevention, according to the relevant mechanism (Mark, 2012; Wu et al., 2017; Tang, 2018; Fu et al., 2020; Yu et al., 2021). In view of the characteristics of phytoplankton outbreaks, in terms of early warning technology, focus should be on investigating phytoplankton distribution and dynamic monitoring, long-term water temperature monitoring in the thermal effluent mixed area, and satellite remote sensing monitoring (Kaplan et al., 2016; Dvorniko et al., 2017; Tang et al., 2017; Wei et al., 2018; Lu et al., 2018; Yu et al., 2021).

In terms of algae disinfection and sterilization, measures and methods with mature technology application, reasonable economic cost control, and low damage to biological resources and the marine environment are preferred (Cao et al., 2017; Yu et al., 2017; He et al., 2019b; Kang et al., 2020). Specifically, the modified clay method is a HABs emergency treatment technology that can be applied on a large scale in the field (GEOHAB, 2010; Yu et al., 2017; Getchis and Shumway, 2017). There have been many successful applications of modified clay, such as for the treatment of *P. globosa* in Fangchenggang nuclear power plant, in which it has been very effective in removing microalgae cells, with a removal rate of >90%. *P. globosa* sank into the seabed with the modified clay, where it decomposed and was inactivated (Yu et al., 2017; Cao et al., 2017). Based on the results of long-term dynamic monitoring and analyses of historical datasets, modified clay removal research should be carried out in the laboratory for potential phytoplankton outbreaks, and the modified clay formula and sterilization strategy should be optimized to provide technical support for the safety of cooling water systems in nuclear power plants (He et al., 2019b; Kang et al., 2020).

Chemical reagents are also a common method for algae bloom sterilization. These reagents include thiazolidinedione derivative TD49, suitable for semi enclosed bays, which has the highest inhibition rate of HABs at 2.0 μM (Baek et al., 2014); Sophorolipid, which significantly inhibits the activity and growth of algal cells between 20 and 5 mg/L, with no recurrence at higher concentrations (Lee et al., 2008); Hydrogen peroxide (Burson et al., 2014); Chitosan (Divakaran and Pillai, 2002), and NaOCl (Jeong et al., 2008). However, attention should be paid to the toxic effects of these chemical reagents on another marine biota (Cao et al., 2017; Yu et al., 2017).

### 5 Conclusions

A number of marine biota cause outbreaks, blocking cooling water systems, leading to the shutdown of nuclear power plants, while the invasion of marine biota seriously affects their safe operation. Based on the field monitoring data of phytoplankton in the sea area within 15 km of the Fujing nuclear power plant, in combination with HABs datasets in Fujian on Southeast China coast from 2006 to 2021, this study determined the risk from phytoplankton species that may potentially block the cooling water system. Three principles of screening blocking organisms were determined, and a catalog of risk organisms and a monthly risk calendar were established. Nine high-risk species of phytoplankton that can easily form colonies, clusters, and groups in case of outbreaks were screened out in this study. The outbreak of various phytoplankton species varied between days, and the risk of blockages from phytoplankton was concentrated from April to July. In addition, a preliminary discussion on the biological characteristics and ecological mechanism of four high-risk species is given, including *Phaeocystis globosa*, *Noctiluca scintillans*, *Thalassiosira diporocyclus*, and *Pseudo-nitzschia pungens*, suggesting that the ecological mechanism of HABs had a high degree of temporal and spatial variability. As for preventive measures against high-risk phytoplankton species, the modified clay method was an effective technology, while use of chemical reagents should be aware of their toxic effects on marine biota. The results of this study show that nuclear power
plants face a high risk of blockage by marine biota in cooling water systems. It is recommended that the administration of nuclear power plants establish a regular monitoring mechanism for cooling water intake safety.

Data availability statement

The data that support the findings of this study are available from the corresponding author or the first author upon reasonable request.

Author contributions

YW: conceptualization, data curation, investigation, methodology, writing—original draft, and funding acquisition. XC and YL: conceptualization, funding acquisition, and validation. SZ, LC, and XT: investigation, formal analysis, and data analysis. PX and HL: project administration and supervision. All authors contributed to the article and approved the submitted version.

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Conflict of interest

YL was employed by Fujian Fuqing Nuclear Power Company Limited.

The remaining authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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