Hello,
I am Wendy,
a designer/anthropologist,
an interdisciplinary
researcher in the artscience and an educator.

こんにちは。 私はウェンディです。 デザイナー兼人類学者、 芸術科学の学際的研究者、 そして教育者です。



Making Sense of Beautiful Toxicities

美しい毒性 き を理解する

Anthropocene Curriculum

Research

CONVERSATION CARBON CRITICAL MATERIALS DEEPTIME DEGRADATION KNOWLEDGE TRANSFORMATION RADIOACTIVITY

Bernadette Bensaude-Vincent Andy Cundy Irka Hajdas Susan Schuppli Colin Waters

FINGERPRINTS OF THE NUCLEAR AGE

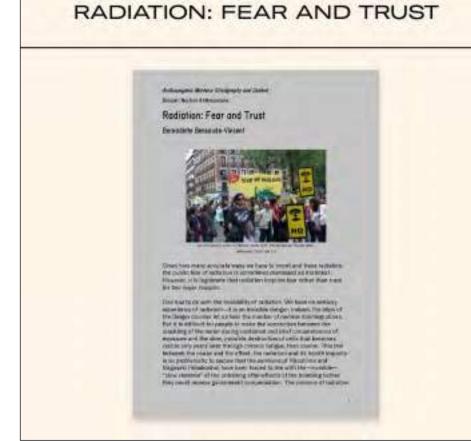
A seminar held during Unearthing the Present

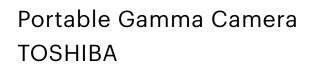
AC-CS #49097

The spike in artificial radiation from nuclear bomb tests in the 1950s might mark the beginning of the Anthropocene. These radioactive products have left their residues in soil, sediment, ice, and even our bodies. How is it possible to measure, let alone visualize, these invisible, minuscule amounts of radiation that escape all sensory data? The materials gathered below are from the ninar Fingerprints of the Nuclear Age, which was held during *Unearthing the Present* in May 2022. The seminar explored how the (in)visibility of radiation is linked to the legacy of the nuclear age, histories of public fear, political secrecy, and counter-expertise.

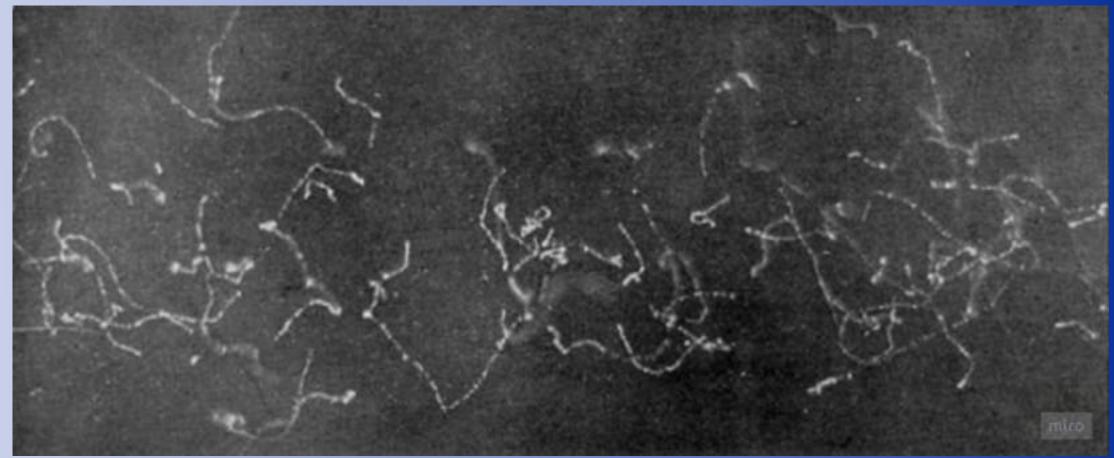


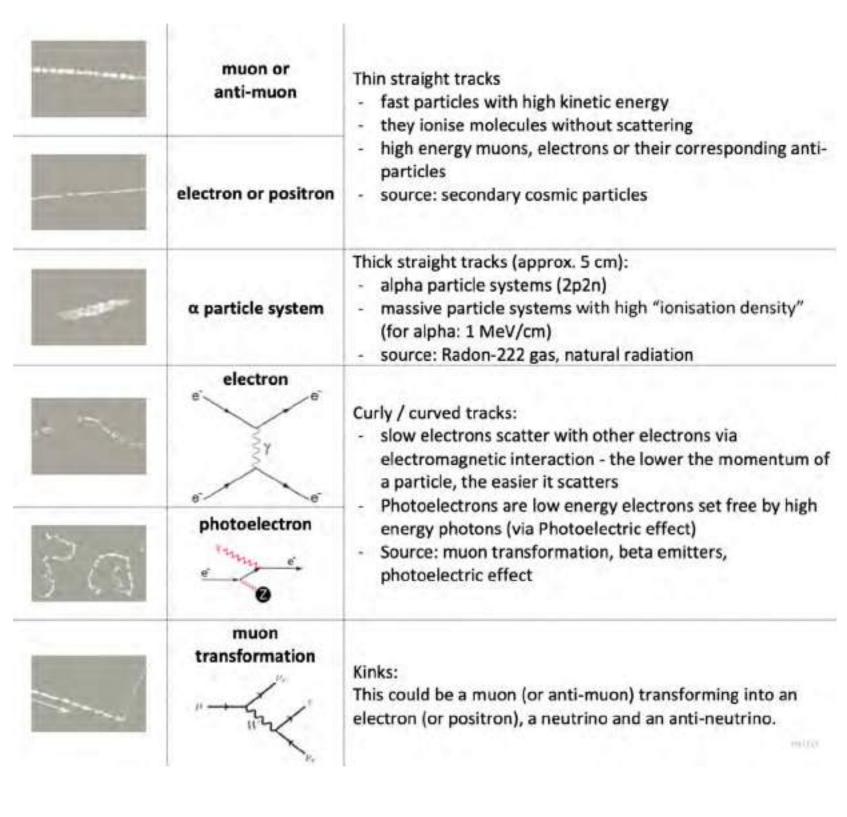
Diffusion Cloud Chamber

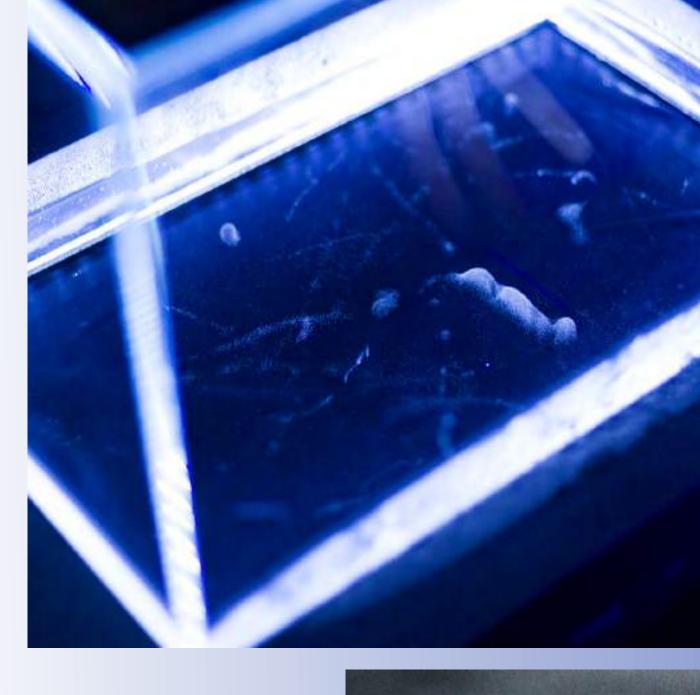


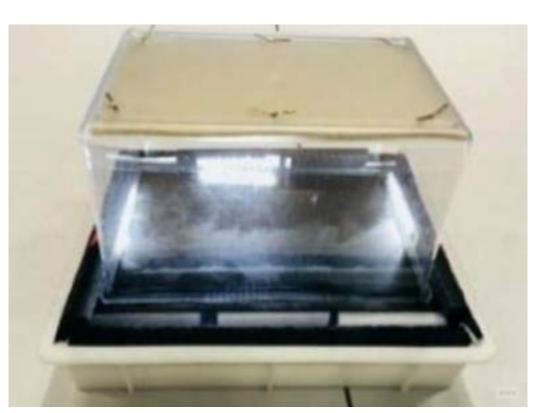












Cloud Traces
Humboldt Labor, Humboldt Forum Berlin, 2022
In collaboration with Andres Gatto,
Belen Palacios, Julia Mieles



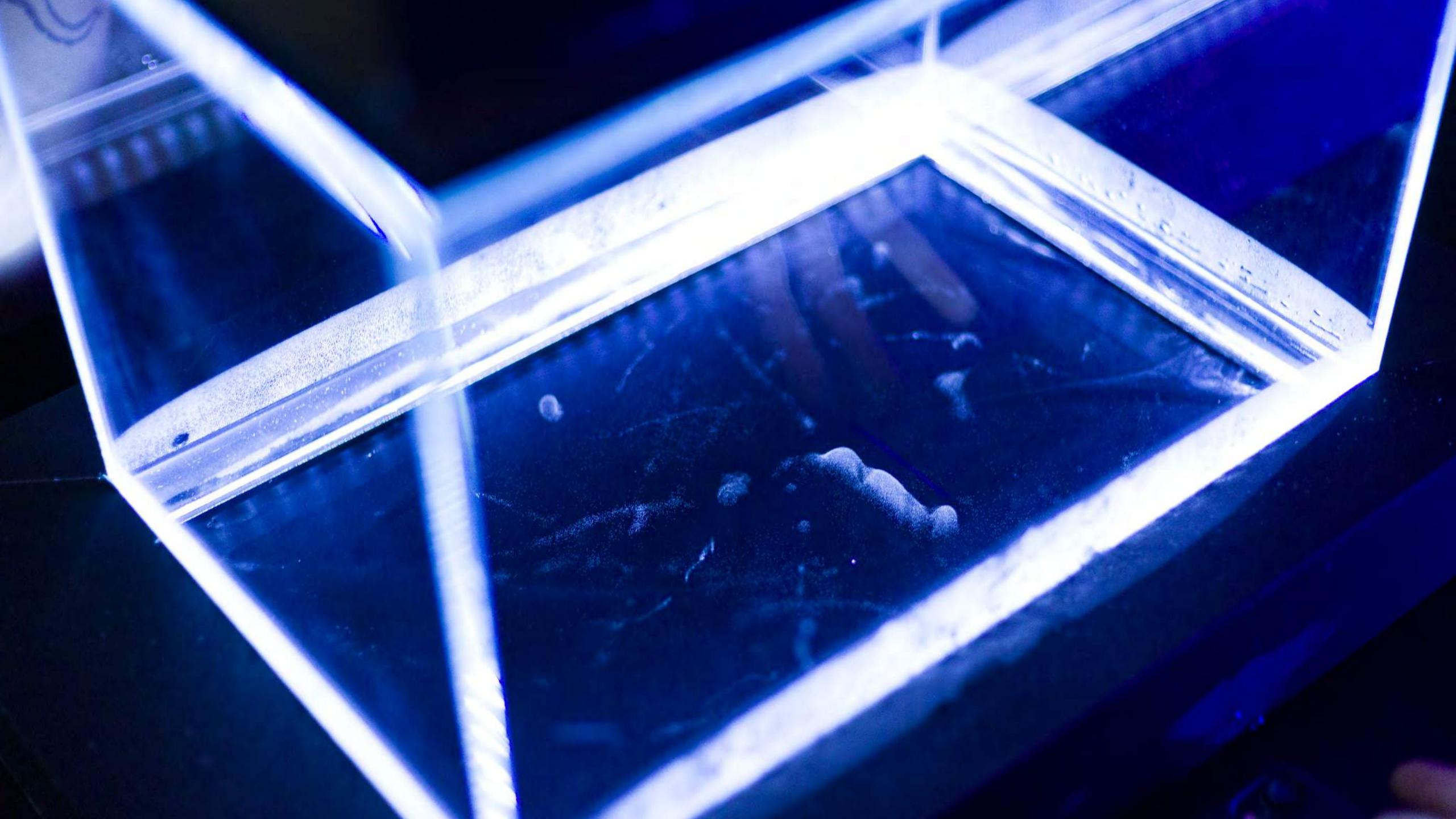






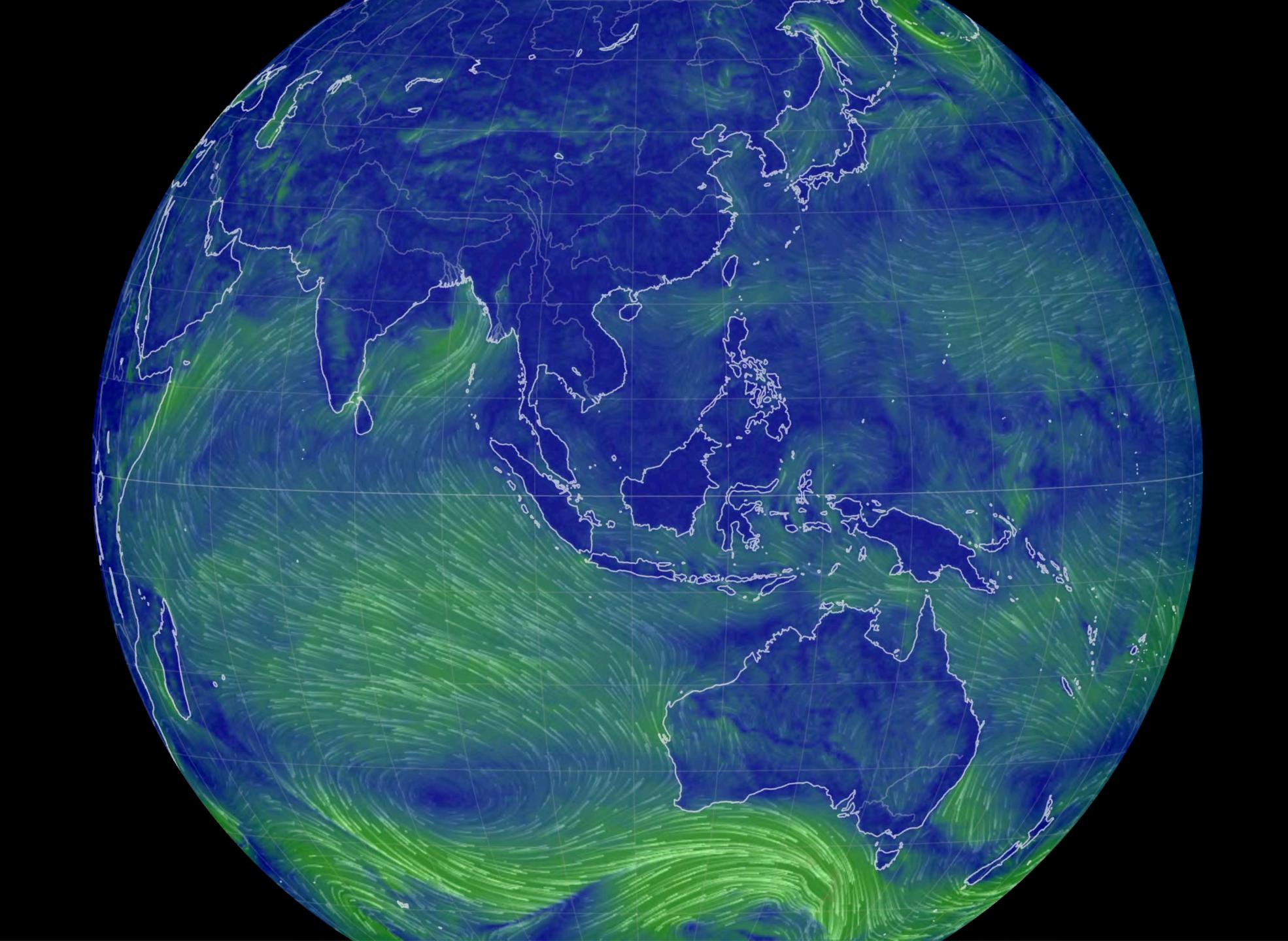




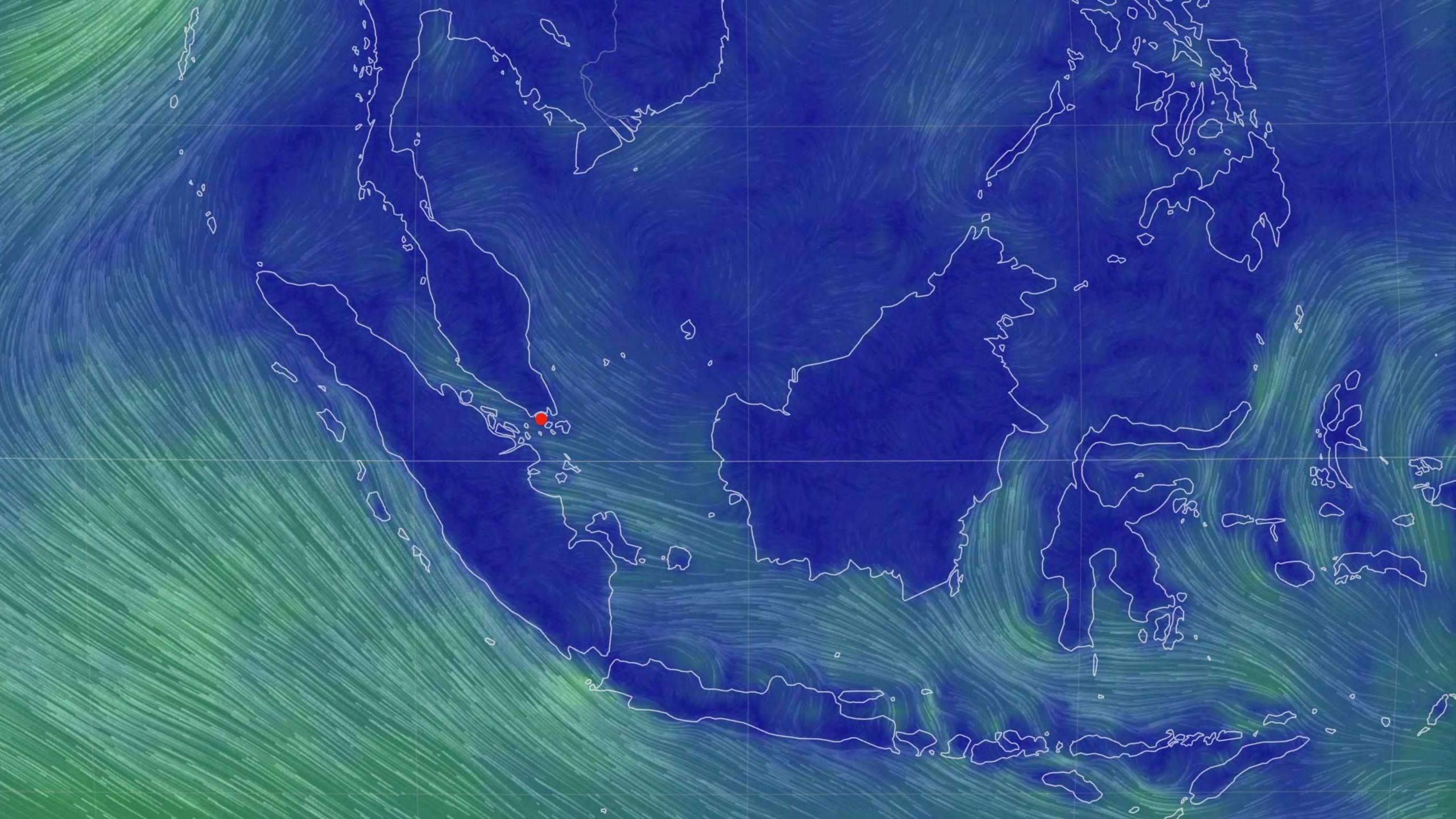


Tieranatomisches Theater TA T, BERLIN

7 May – 28 June 2025



earth ≡









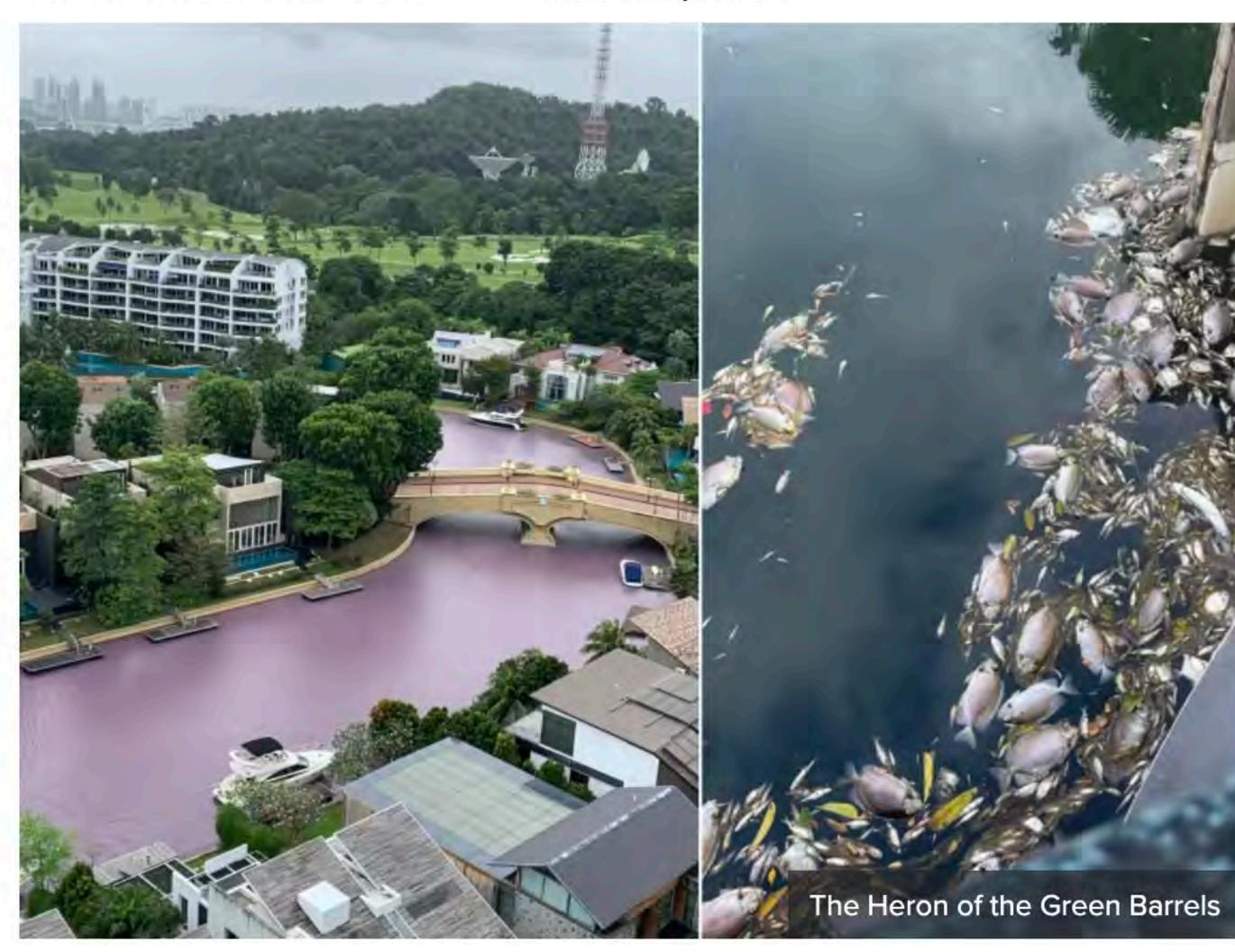
Fish deaths: Plankton bloom causing fish deaths 'likely to recur'

PUBLISHED 9 MAR 2015, 5:53 AM SGT

AVA and farmers must discuss best way to tackle challenge: Vivian

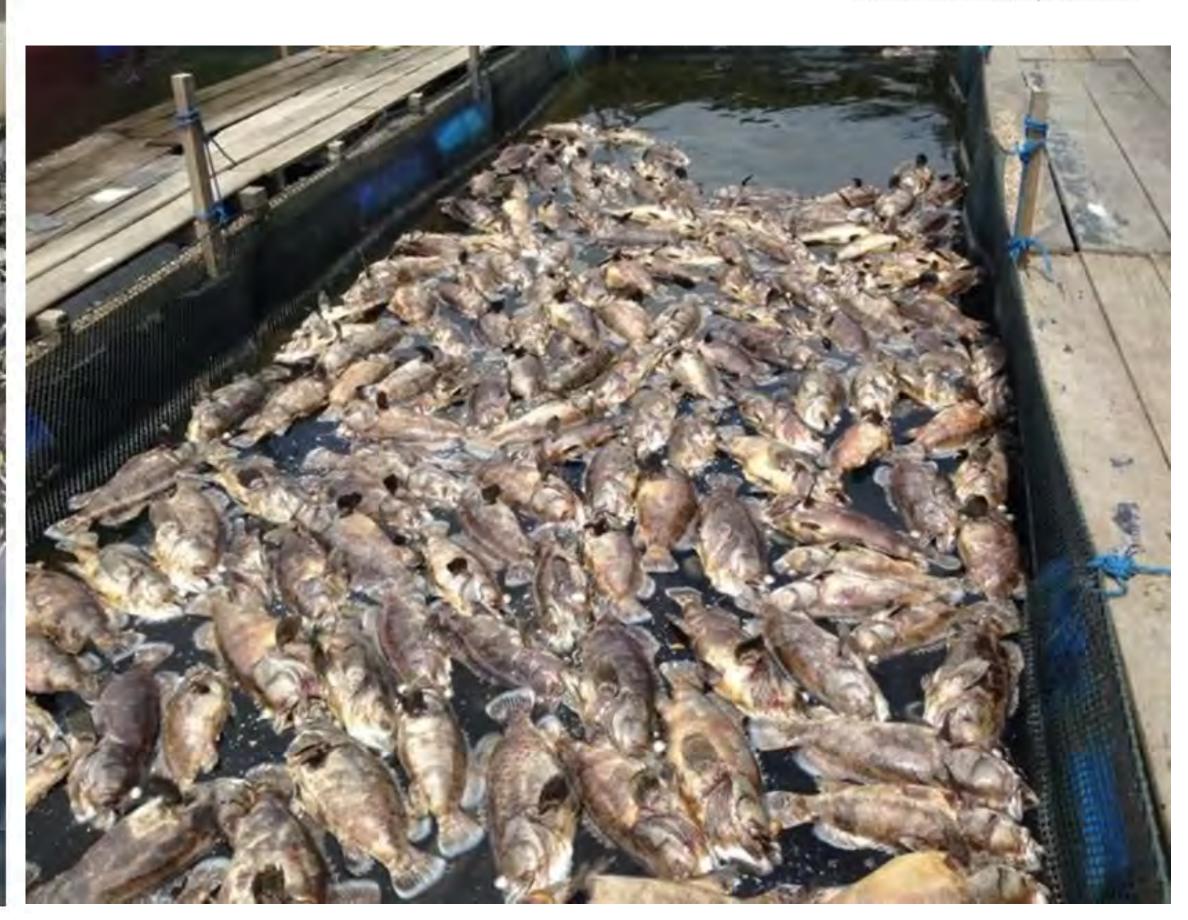
Sentosa Cove waters turn pink, smell rotten; one resident calls it an 'environmental disaster'

Published January 13, 2021

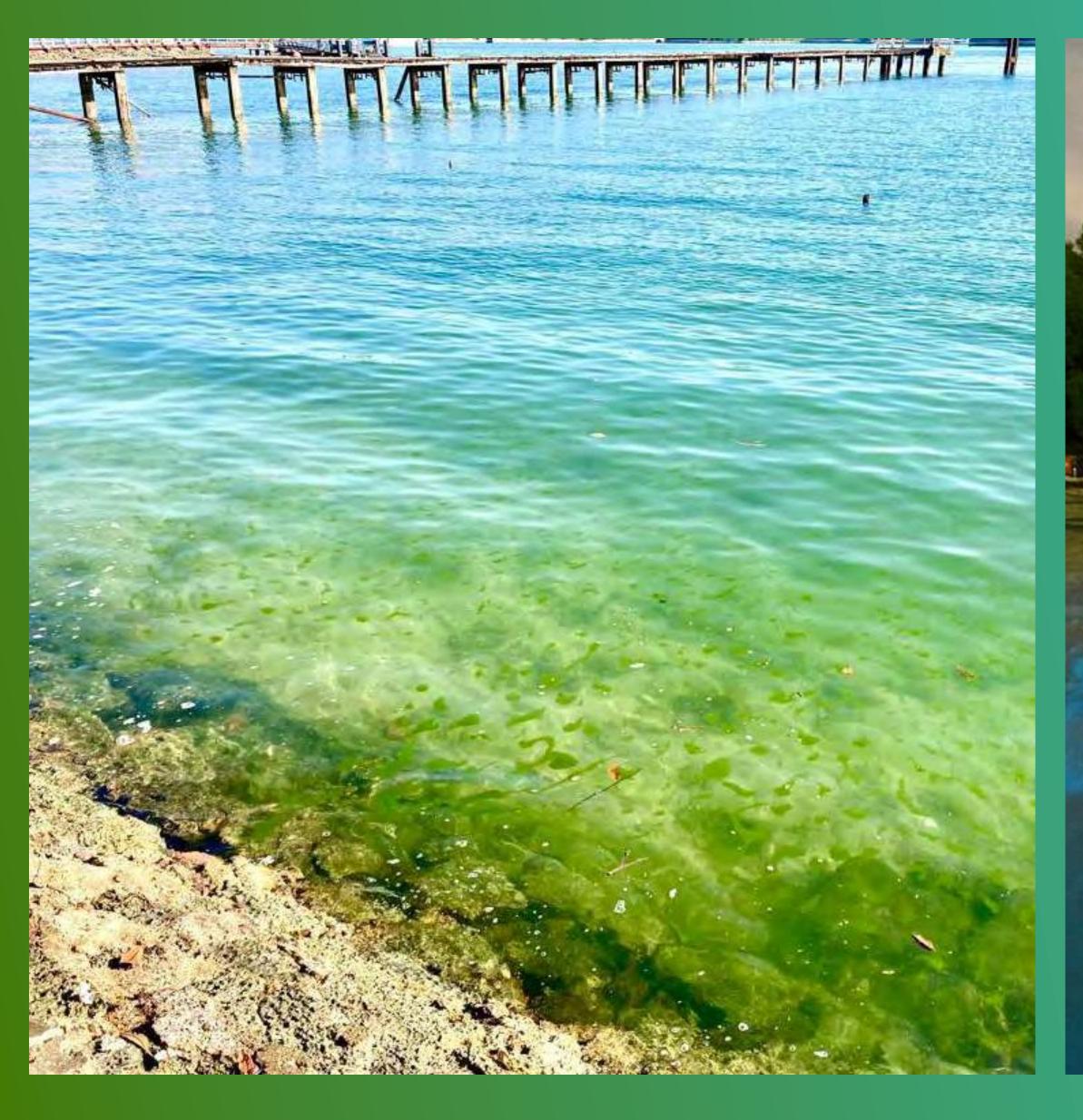


Pasir Ris mass fish deaths may lead over \$1 million loss for farmers

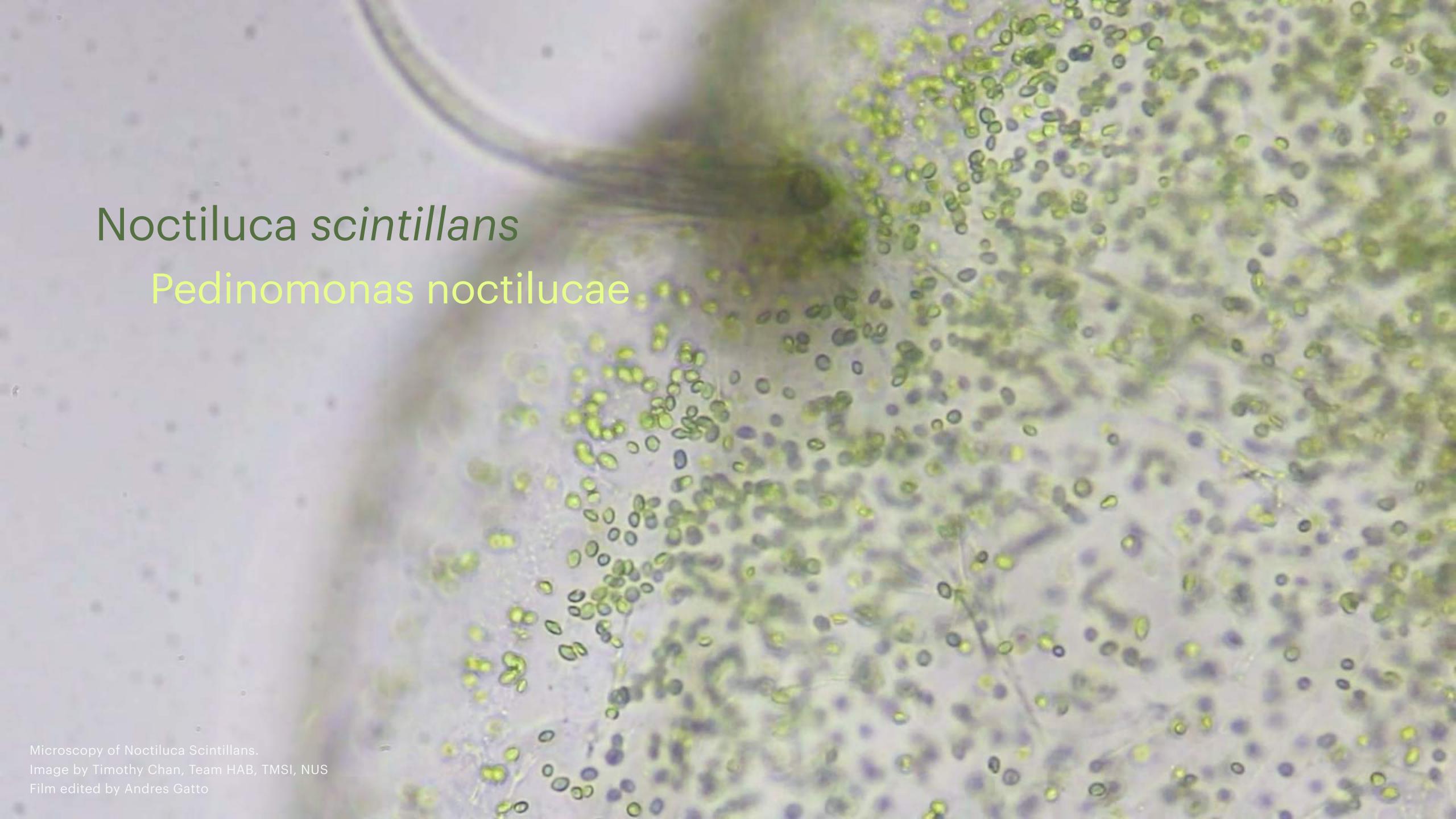
PUBLISHED 11 FEB 2014, 8:31 AM SGT



Beauty — Toxicities 美しさ — 毒性

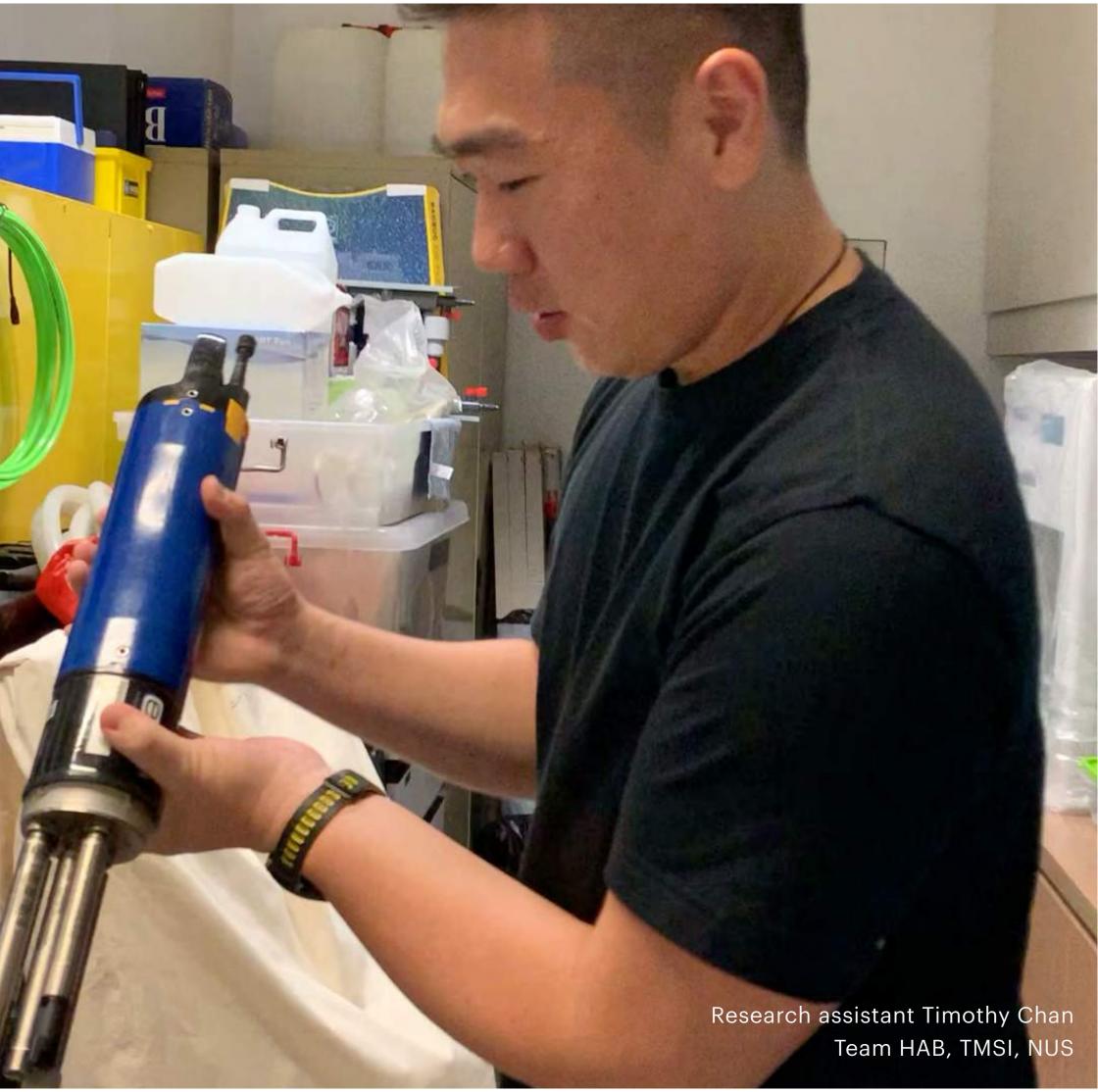






Measuring Blooms 藻類ブルームの測定





Fieldwork with Harmful Algal Bloom scientists in the Johor Strait

Credit: Team HABs, Tropical Marine Science Institute, National University of Singapore



藻類ブルームの測定

ジョホール海峡における有害藻類ブル ームの研究者とのフィールドワーク

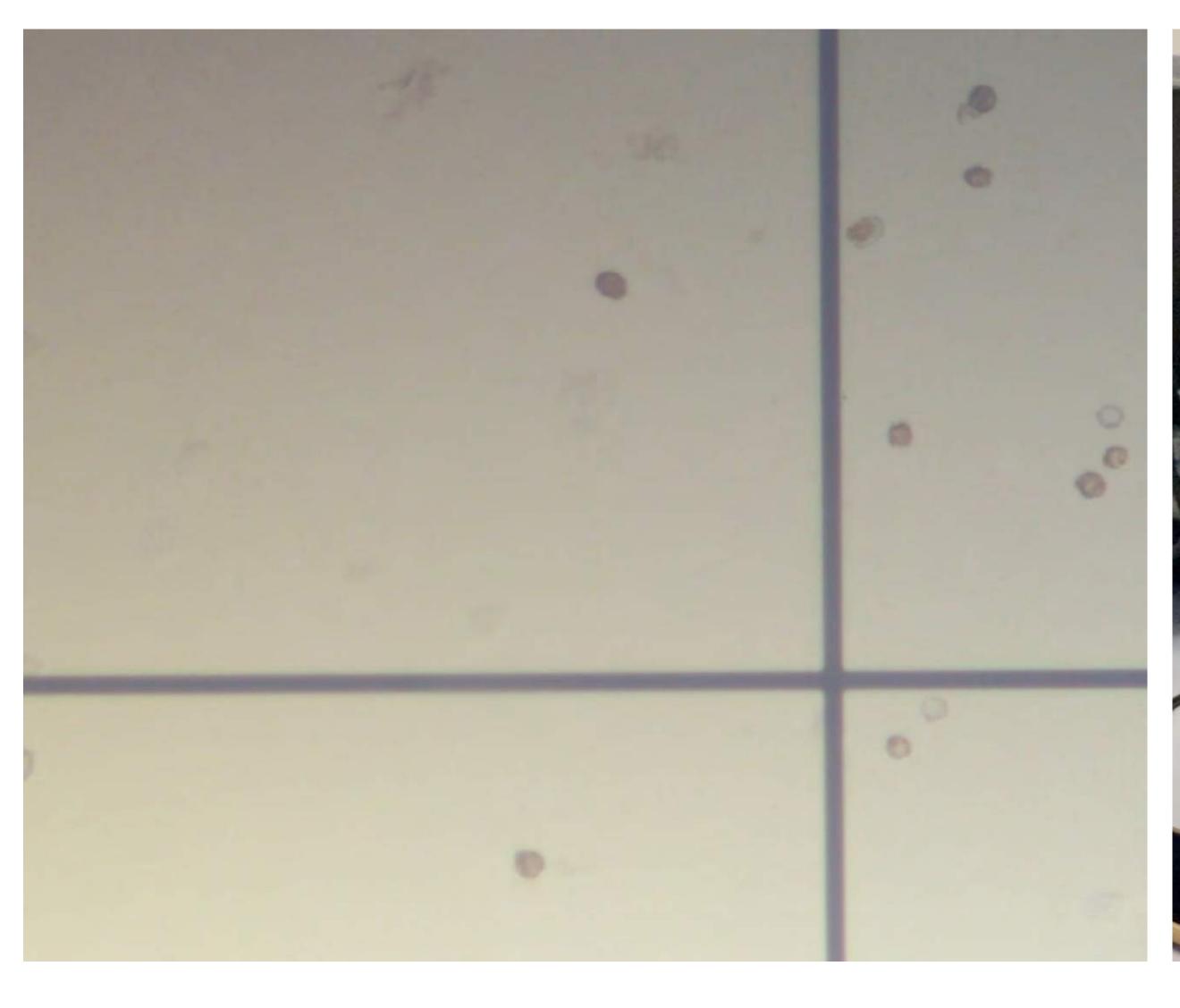




藻類ブルームの測定

Saturation Thresholds

飽和閾値



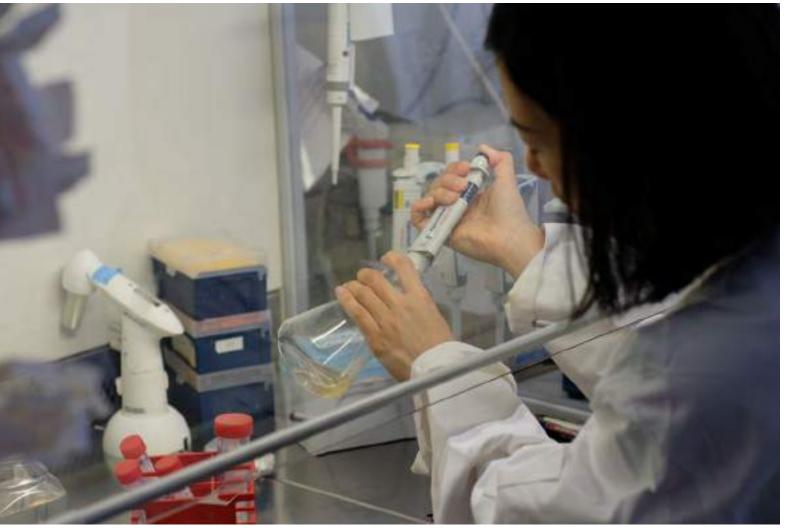


Temperature Thresholds

藻類ブルームの測定

温度閾値













Temperature Thresholds of Algae in Marine Heatwaves

藻類ブルームの測定

海洋熱波における藻類 の温度閾値

Hypothesis

How would the toxic algal species Gonyaulax spinifera survive temperature spikes in marine heat waves?

Culture Strains:

Gonyaulax Spinifera 10 ml preculture tubes x 4

Culture Medium

F2 filtered seawater (nutrients)

Cool white, or cool/warm white mix, fluorescent tubes or bulbs with full spectrum of 400 -700 nm

30-40 µmol m-2s-1 (CCAP culturing conditions)

100 µmol m-2s-1 to induce growth for incubation ex

Avoid stressing or inhibit growth by keeping to a maxAcclimatization Avoid going beyond 2500 µmol m-2s-1 which is equivale May suggin 2023 (16 days)

Light Cycle

12h light : 12h dark (for faster growth try 16h:8h)

15 - 20 °C (CCAP)

Subculture Ratio 1:10 or 1:5

Culture Vessel

Conical flasks containing approx. 500ml culture

Culture Chamber

Each chamber measures 385 x 310 x 380 mm Light bulb wired in chamber, set to timer.

ACDP Hazard Group 1

Non pathogenic / non hazardous. Unlikely to cause h Found to be a carrier of Yessotoxins in 20061 in New Zerala Provide air flow with foil cover

Algae 5, no. 2 (2006): 148.

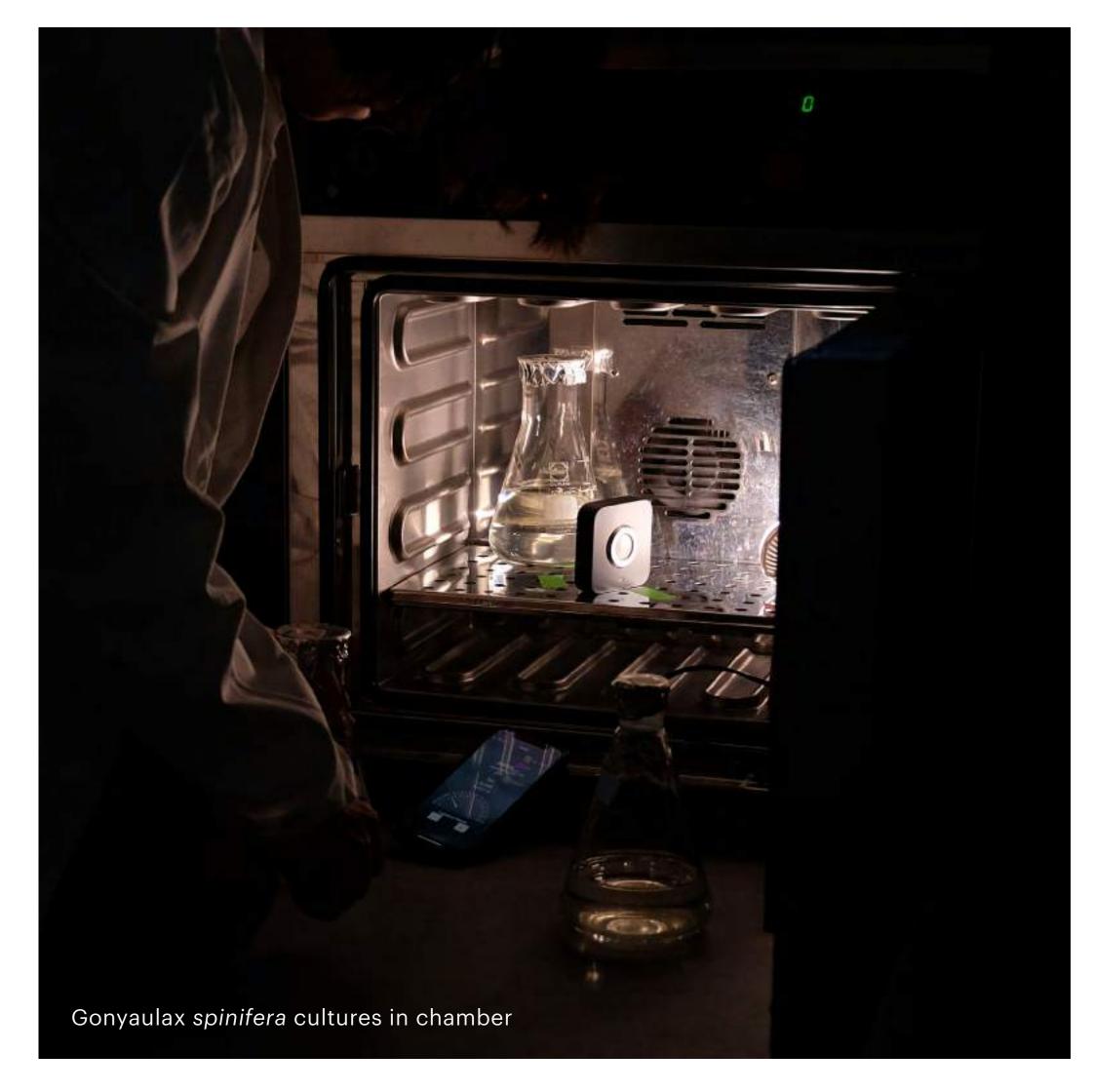
Aim: Keep culture alive and multiply, do not rush to grow cultures. Dino grow slower. Wait undisturbed 3-5 days.

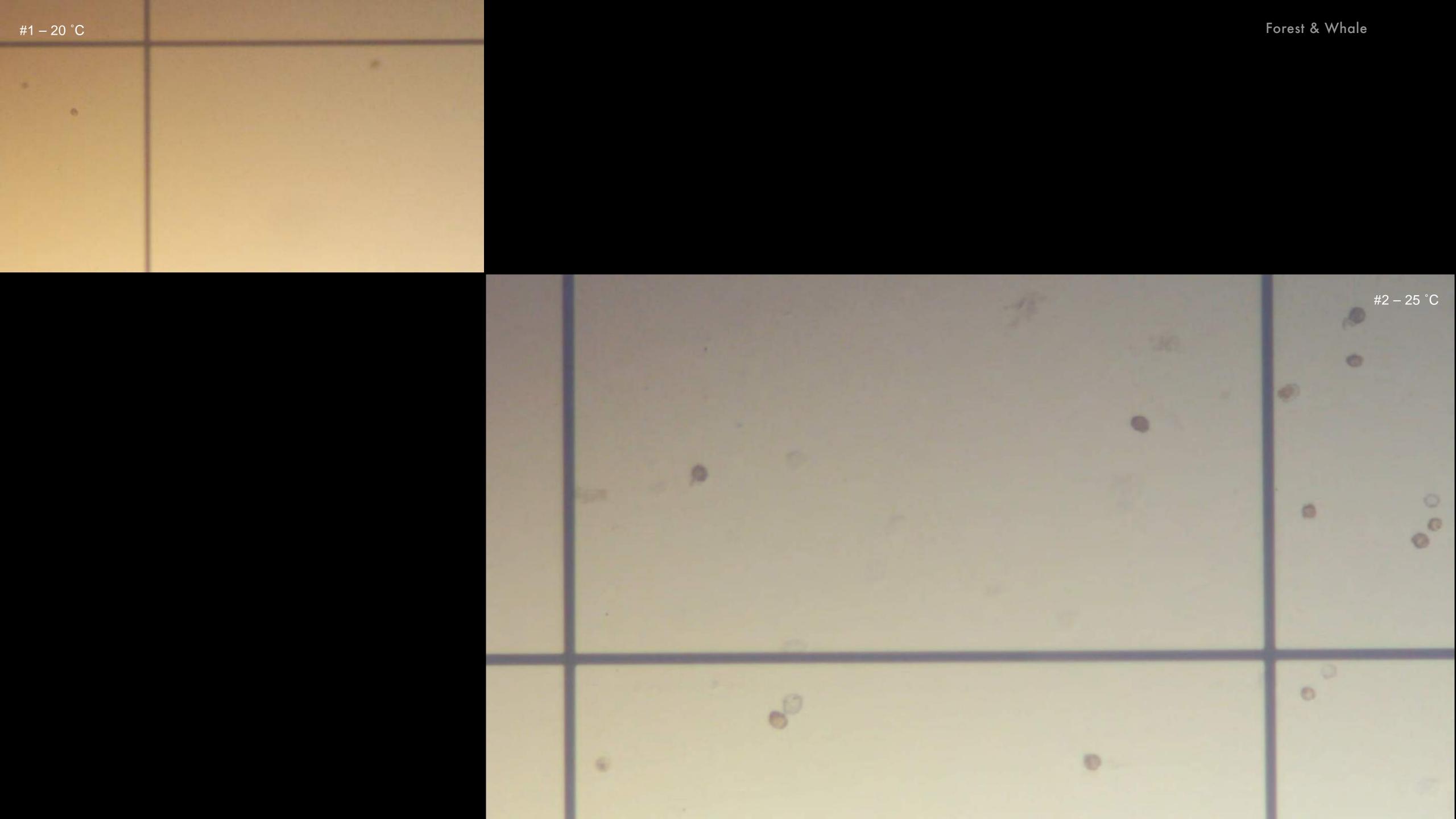
- 1. Upon arrival of cultures (4 tubes of 10ml), open and close caps.
- 2. Pour cultures (1-2ml) onto slides to check health.
- 3. Inoculate two tubes into fresh sterile medium in the ratio of 1:10 e.g. 5mls culture into no more than 50mls medium. If the culture is not in optimal condition or bacteria are obvious 5. Counting then 1:5 may be necessary. Particularly dense cultures can be added
- to slightly larger volumes of medium. 4. Place 2 culture tubes (2 x 10ml) and 2 inoculated flasks in the chamber at 20 °C (see table)
- 5. Light cycle of 12h:12h hours (10am 10pm)
- 6. Adjust lighting to irradiance at 100 μmol m⁻²s⁻¹
- 8. After 3 days, pipette 20-50 μL to observe. Leave it untouched until the cultures recovered (hundreds moving in 1 ml).
- 1 Lesley Rhodes et al., "Yessotoxin Production by Gon aulf all titibes are the afthy, combine into one stock culture of 1000 ml. 10. If cultures have recovered, subculture every 3 days to grow cultures to a₅. Flush with tap water, wipe with a soft tissue
 - cell density of 100 cells per ml.

Use strict aseptic techniques throughout all subculturing within a laminar flow cabinet. Subculture cultures in fresh sterile medium after it arrives.

Experiment Procedure:

- 1. Once experiment begins, check cultures every 2 days on Monday, Wednesday, Friday at the same time (930am) each day.
- 2. Prepare four tube flasks mark them 1-4 to correspond to chamber #.
- 3. Take out flasks from chambers to pour approx. 10 ml into respective tubes
- 4. Protocol for extracting cultures without dilution
 - 1. Take out flask from chamber, swirl in gentle big circles.
 - 2. Pour 5 ml into tube and return flask to chamber
 - 3. Shake tube gently in big circles, in both directions (10 seconds) to remove aggregates
 - 4. Pipette 1ml/ 1000 μL from culture tube onto gridded rafter by moving in zig zag manner, maintain same action
 - 5. Slide slide cover to close rafter
 - 6. Place rafter in the fridge for 2-3 mins
 - 7. Remove rafter and place under microscope
 - 8. Observe under x5 to count live cells, brown cytoplasm. Do not count dead, transparent cells or green cysts
 - 9. Count cells based on counting protocol below
 - 10. Plot cells/ml every 2 days on Monday, Wednesday, Friday 11. Plot growth curve
 - Protocol: Counting cells by moving grid to grid
 - Zig zag up and down, counting 10 columns of 200 squares. Count 3 different sets of columns to take an average.
 - if the whole rafter < 400 cells, then count the entire rafter. Density is too low to take an average count.
- 6. Return flask back to the chamber 7. Wash gridded rafter for next count
 - Discard used glass slide cover Pour it into a waste container
 - Rinse with water Repeat 2 times
 - 6. Use clean rafter for next count and new glass slide cover
- Repeat steps till Day 10 or 14





Temperature Thresholds of Algae in Marine Heatwaves

Results 20 °C

Constant growth

25 °C

Initial dip then pre-bloom trends as growth spikes under optimum conditions

28 °C

Shock to algal communities, slow cell death

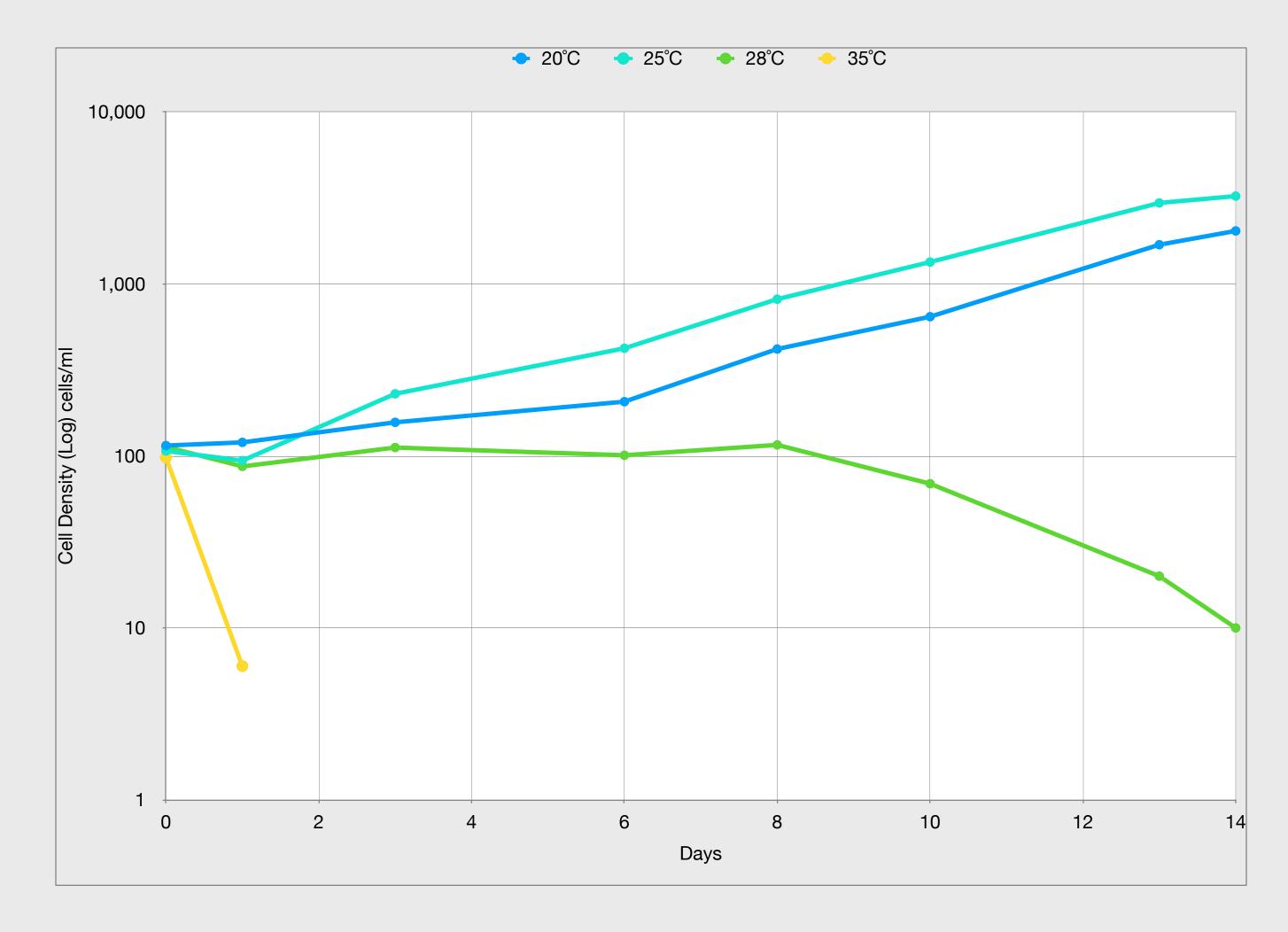
35 °C

Shock and catastrophic death

藻類ブルームの測定

海洋熱波における藻類





As invisible wild fires rage unseen, we ask: What are we not hearing?

目に見えない山火事が猛威を振るう中、私たちは問いかけます。私たちは何を聞いていないのでしょうか?

'Unheard of' marine heatwave off UK and Irish coasts poses serious threat

Sustained high temperatures over summer could trigger mass mortality of fish and oysters, say scientists



What you need to know about the 'extreme' heatwave hitting our oceans - video explains

Helena Horton Environment reporter

Mon 19 Jun 2023 12.21 CEST



A humpback whale, nicknamed Festus, who died near Glacier Bay in June 2016 during a marine heatwave in the north-east Pacific. Starvation was given as the primary cause of death. Photograph: Craig Murdoch, taken under authority of NOAA Marine Mammal Health and Stranding

The age of extinction

Did a marine heatwave cause 7,000 humpback whales to starve to death?

Populations were recovering, but a new study reveals that numbers dropped by 20% coinciding with a period of record temperatures in the North Pacific

The age of extinction is supported by



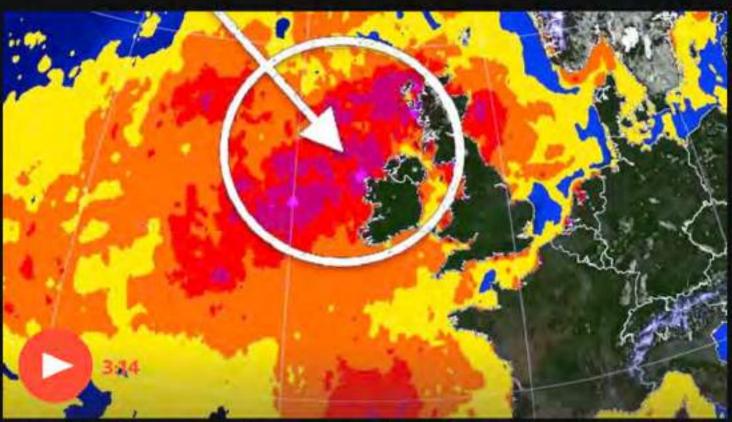
About this content

Brianna Randall

Wed 28 Feb 2024 08.01 CET

What you need to know about the 'extreme' heatwave hitting our oceans - video explainer

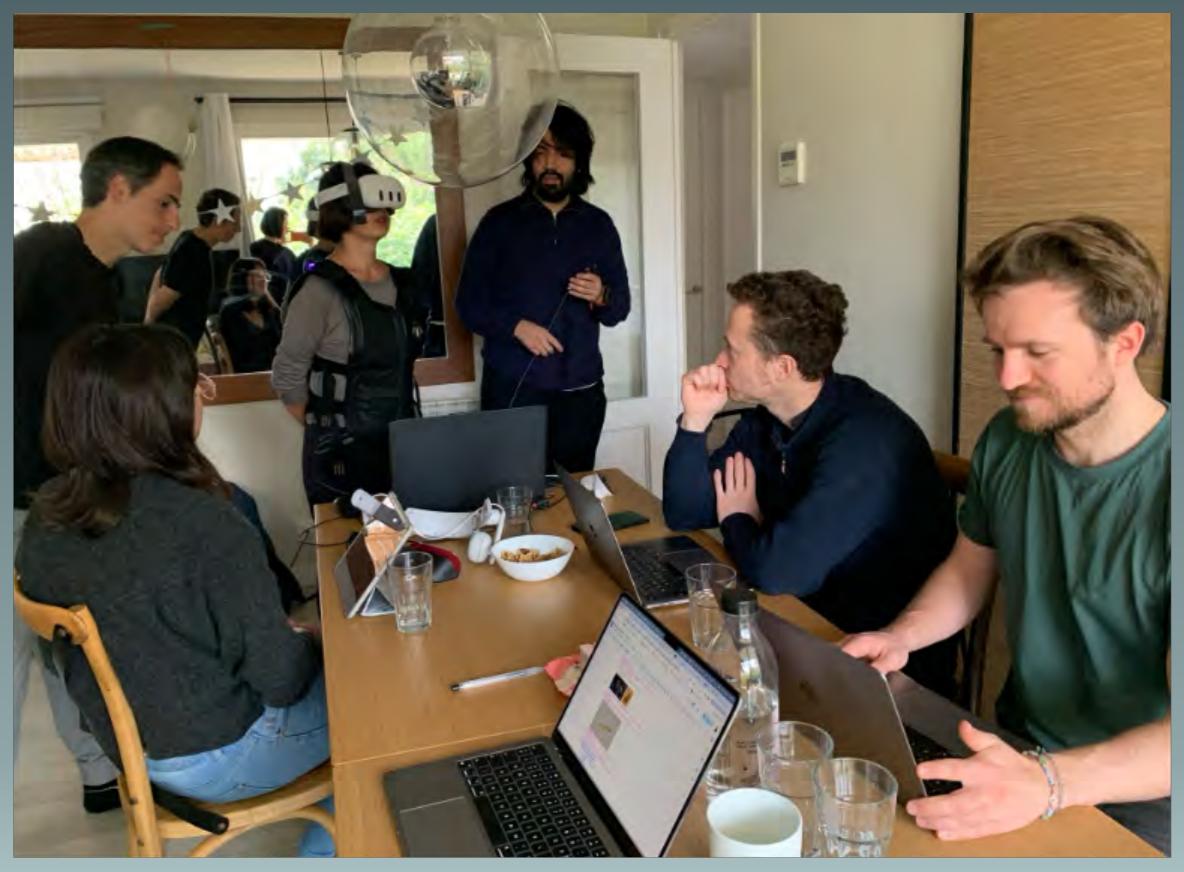
Wendy Chua | forest & whale



Scientists have warned that a marine heatwave off the coasts of the UK and Ireland poses a serious threat to species. Sea temperatures, particularly off the north-east coast of England and the west of Ireland, are several degrees above normal, breaking records for late spring and early summer. The Met Office has said that according to records dating to 1850, global sea surface temperatures in April and May reached an all-time high, and June is likely to follow suit. A professor of Earth sciences, Daniela Schmidt, said 'the extreme and unprecedented temperatures show the power of the combination of human-induced warming and natural climate variability like El Niño'. Experts said marine heatwaves have a similar impact on the environment as wildfires on land, destroying organisms that store carbon such as kelp. The damage caused is also harmful to humanity, which relies on oceans for oxygen, storm protection and food

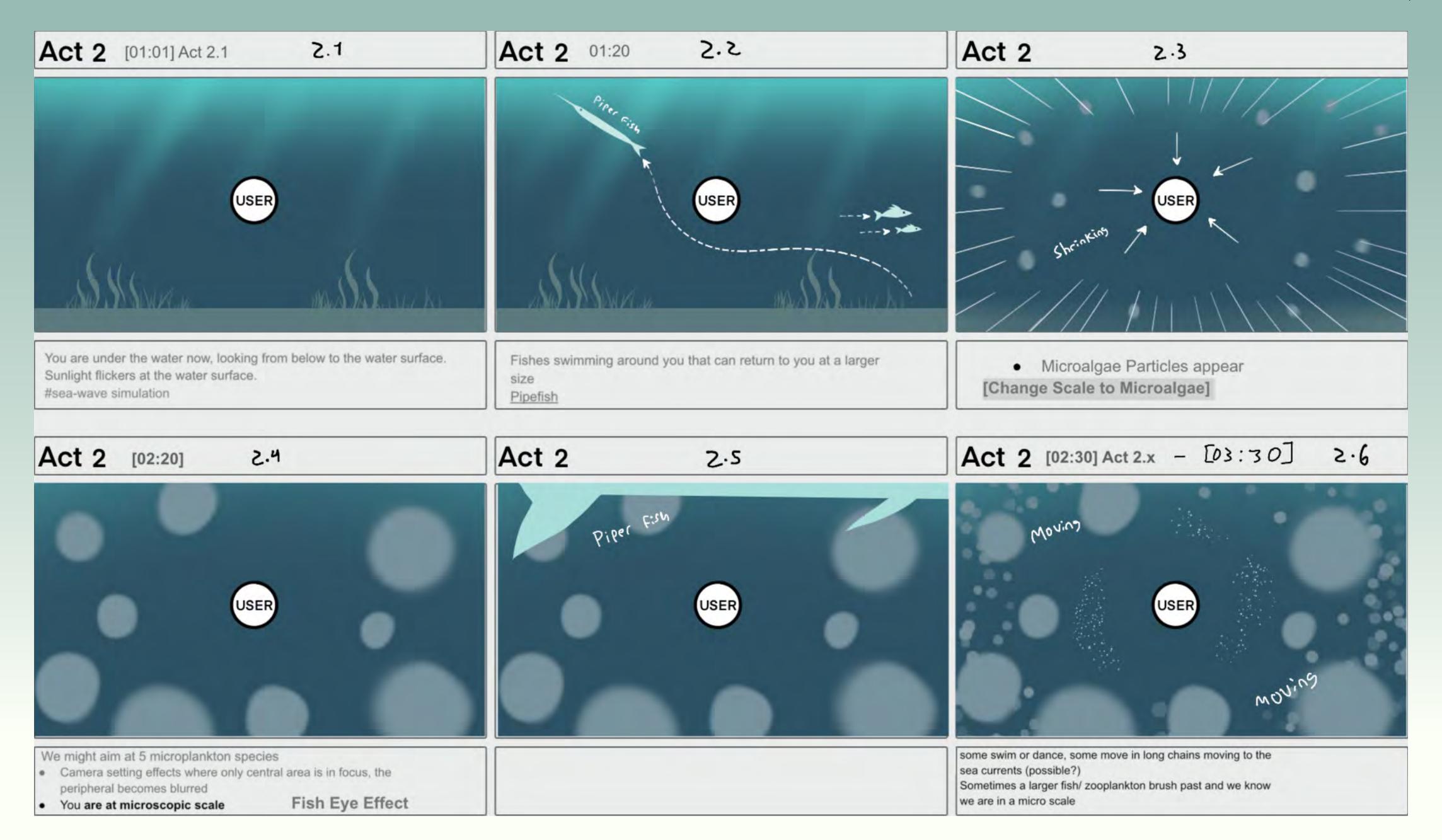
 'Unheard of' marine heatwave off UK and Irish coasts poses serious threat

Helena Horton Bryony Moore Monika Čvorak, Source: The Guardian / As credited; Thumbnail image: Scott Duncan
Thu 22 Jun 2023 16.22 CEST





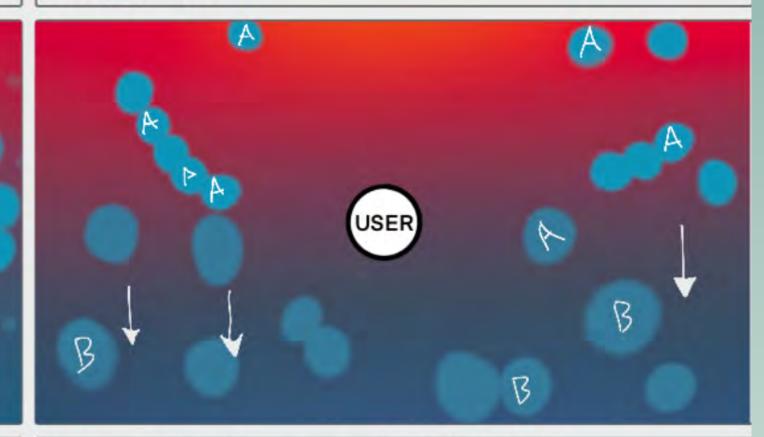
Gustavo Maggio, Joyce Koh, Christian Stein, Valentin Hanau, Marco, Romina, Andres Gatto



Act 3 [03:30] Act 3.1 MHW Heat Transition USER

- Act 3 [04:00] Act 3.2 Blooming begins Main Act: Competing Blooms
- Act 3 3.3

· Cells come closer to the camera



- [VR] Red gradient glow from surface and fade downwards
- [VR] heat haze effect
- [Global env] infra red light shining heat to feel it

All the dinos and zooplankton freeze

- We see crowding of species into the frame pairs of cells in

 * Space for participant feels slowly enclosing cell-division process, multiplying
- . We dont need to see the actual cell division unfolding but cells in the divided states appearing around you
- . Density of cells is increasing. = Reducing the visible space in

B can swim down (towards the cold). A stays in place

Act 3 3.4

Act 3 3.5



Act 3 3.6

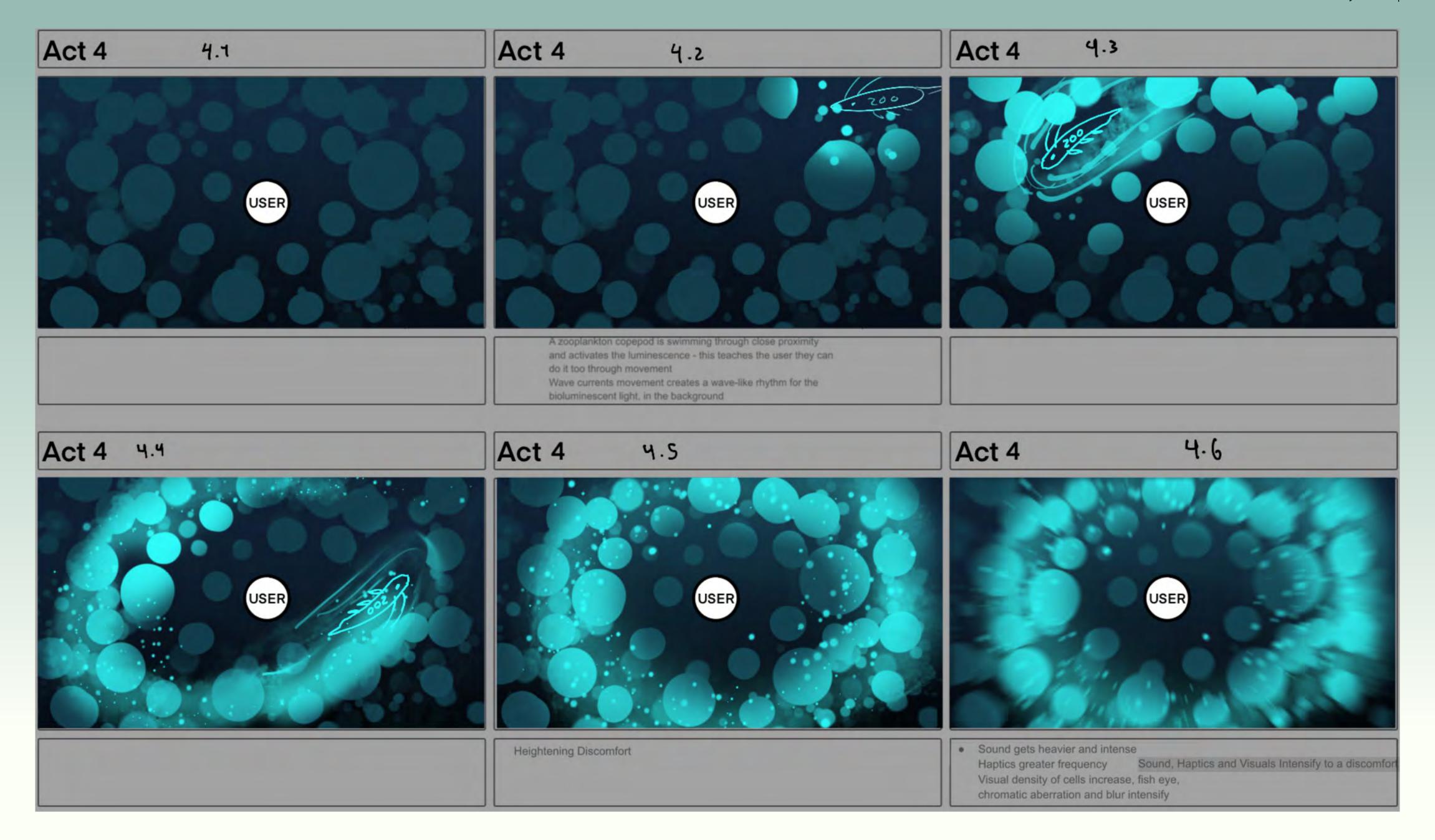


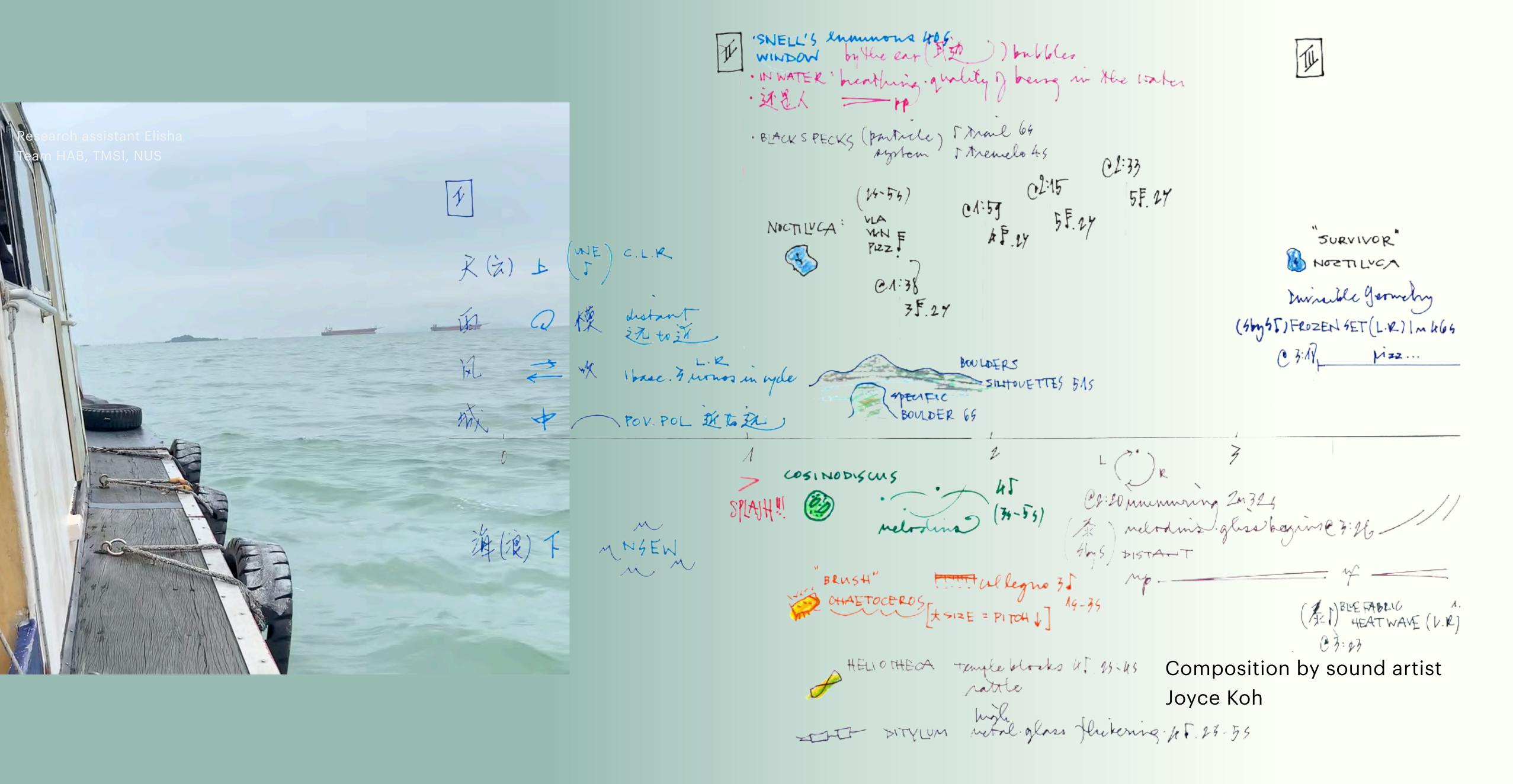
A dies and B survives.

Bloom #1 slowly die out, long chains start to detach and die off The diatom turn from green to grayscale crystalline structures They slowly break into smaller chunks with X's The crystalline cells fall gently as marine snow

- Bloom #2 is starting to take over, and fill the screen
 - They cannot form chains but their density increase
 - . They swim and proliferates and enter the field with the drift through the opacity change.

It gets darker and darker, it's getting dark





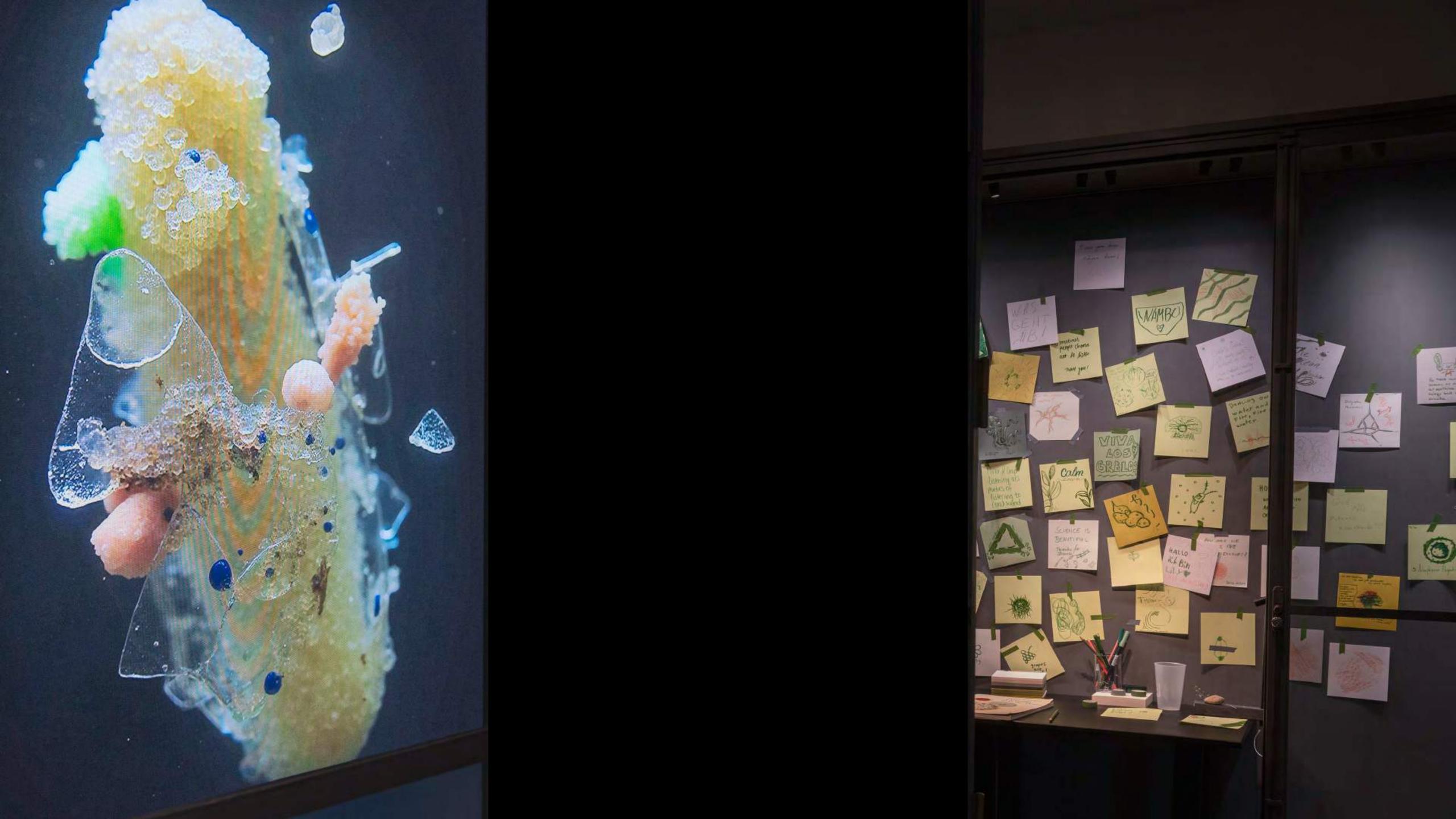
Tieranatomisches Theater TA T, BERLIN 7 May - 28 June 2025



Field Sampling Feldprobenahme

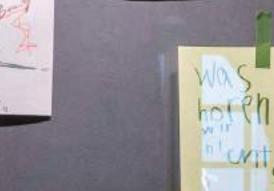
Site Name Sitename	Depth	Temperature Temperatur	Salinity Salzgehalt	Chlorophyll ug/L	Optical Dissolved	Turbidity Trubung NTI
	m	100	psu	-8/-	Oxygen	ET TOTAL
			10000000		optisch	
					gelöster	
					Sauerstoff	
					mg/L	
Sa5	0.003	29,609	29.34	0.38	3.64	2.08
Sa5	0.003	29.609	29.34	0.34	3.64	2.07
Sa5	0.004	29.607	29,34	0.36	3.64	2.07
Sa5	0.008	29.599	29.34	0.38	3.63	2.05
Sa5	0.013	29.587	29.35	0.34	3.63	2.05
Sa5	0.019	29.578	29.35	0.36	3.63	2.06
Sa5	0.037	29.571	29.35	0.34	3.63	2.06
Sa5	0.062	29.562	29.36	0.35	3.63	2.09
Sa5	0.086	29.554	29.36	0.34	3.62	2.09
Sa5	0.11	29.543	29.36	0.36	3.62	2.1
Sa5	0.14	29.533	29.37	0.41	3.62	2.12
Sa5	0.176	29.522	29.37	0.48	3.62	2.17
Sa5	0.232	29.511	29.38	0.45	3.61	2.19
Sa5	0.284	29.498	29.39	0.42	3.61	2.21
Sa5	0.339	29.485	29.4	0.42	3.61	2.24
Sa5	0.391	29.471	29.4	0.46	3.61	2.24
Sa5	0.433	29.461	29,41	0.49	3.6	2.24
Sa5	0.507	29.445	29.42	0.45	3.6	2.23
Sa5	0.556	29.434	29.43	0.51	3.6	2.24
Sa5	0.605	29.423	29.44	0.52	3.6	2.23
Sa5	0.651	29,412	29.45	0.52	3.6	2.23
Sa5	0.699	29.4	29.46	0.49	3.6	2.21
Sa5	0.735	29.386	29.46	0.53	3.6	2.23
Sa5	0.769	29.37	29.48	0.59	3.61	2.27
Sa5	0.802	29,348	29.49	0.61	3.61	2.31
Sa5	0.835	29.325	29.51	0.57	3.62	2.35
Sa5	0.876	29.295	29.52	0.55	3.63	2.39
Sa5	0.911	29.269	29.54	0.55	3.64	2.45
Sa5	0.951	29.225	29.57	0.51	3.65	2.52
Sa5	0.989	29.183	29.6	0.54	3.67	2.6
Sa5	1.029	29.117	29.64	0.52	3.69	2.65
Sa5	1.078	29.086	29.67	0.53	3.71	2.71
Sa5	112	29.026	29.7	0.53	3.73	2.74

















Base Milano 9 – 22 October 2025











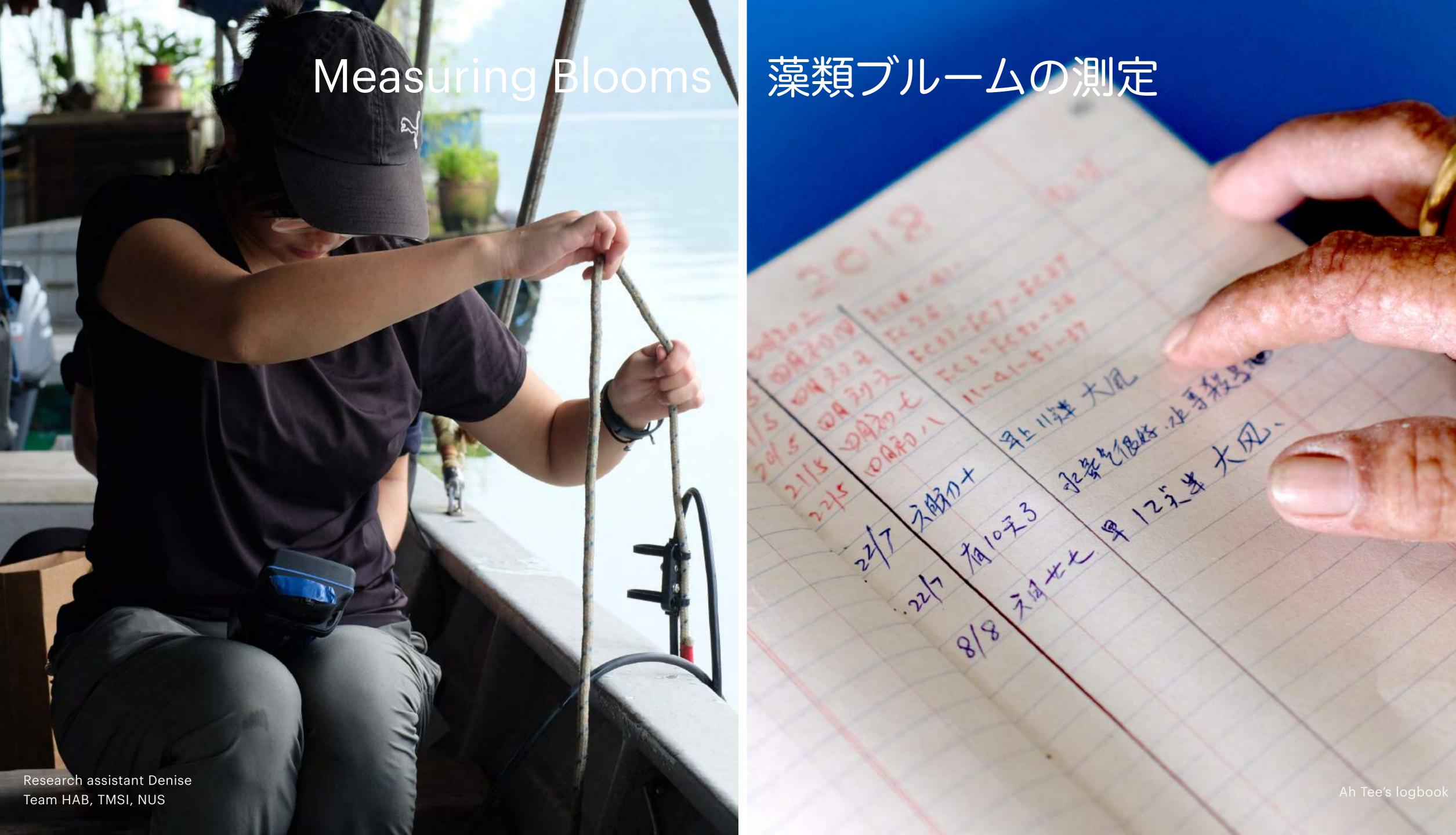






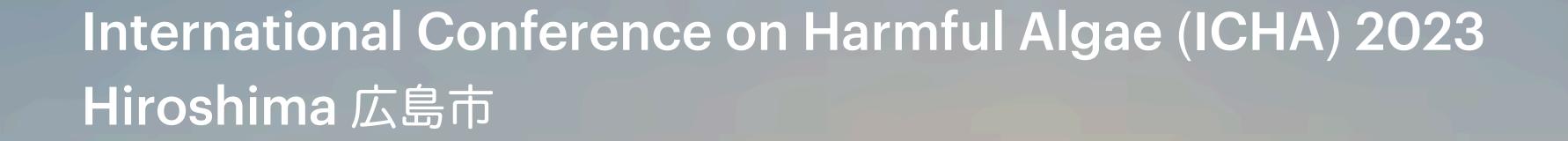














My longtime question to myself is: There must be no microalga that wishes to harm to others.

Maybe the micro algae have specific abilities to survive in the struggle for life among plankton world. The abilities are fast uptake of organic substances, fast growth rate ... it means the algae behave naughty in finding good condition for themselves then they can bloom.

Yasuwo Fukuyo
Professor Emeritus,
The University of Tokyo.

長年、私はこう自問しています。 他の生物に害を及ぼそうとする微細藻類は 存在しないはずだ。

もしかしたら、微細藻類はプランクトン界での 生存競争を生き抜くために、特別な能力を持っ ているのかもしれません。その能力とは、有機 物の吸収速度の速さ、成長速度の速さなどで す。つまり、藻類は自分にとって良い環境を見 つけるために、いたずら好きな行動をとること で、繁殖できるのです。

福代 康夫東京大学名誉教授

To people, the bloom shows unusual scenery, but it is not harm, unless people utilise coastal area and coastal seafoods. It means people change the (lovely) naughty algae to the one that has harm for others, becoming harmful algae.

Microalgae are friends struggling in the sea, in the same way as I do on land.

Yasuwo Fukuyo
Professor Emeritus,
The University of Tokyo.

人間にとって、藻類のブルームは異様な光景ですが、沿岸地域や沿岸の魚介類を利用しない限り、害にはなりません。つまり、人間が(愛らしい)いたずら好きな藻を、他者に害を及ぼす有害藻に変えてしまうのです。

微細藻類は、私が陸上で闘うのと同じよう に、海の中で闘う仲間なのです。

福代 康夫東京大学名誉教授



DANCE OF THE DIATOMS

Sound Composition by sound artist Joyce Koh Microscopy of Noctiluca Scintillans by Team HAB, TMSI, NUS Film edited by Andres Gatto

BLOOMS TEAM

Wendy Chua Research and art direction
Joyce Koh Sound artist
Gustavo Maggio Product/ Space designer
Andres Gatto Media artist

Christian Stein VR creative director
Valentin Hanau VR project manager
Julian Dietz VR developer
Arthur Melzow VR developer
Marco Garcia VR developer
Romina Valdivia Sanz 3D artist

Dr Sandric Leong Marine Scientist
Audrey Lee Marine Science Researcher
Timothy Chan Marine Science Researcher

Manuel Cirauqui EU S+T+ARTS ReSilence curator Berta Gutierrez EU S+T+ARTS ReSilence curator

wendy@forestandwhale.com