

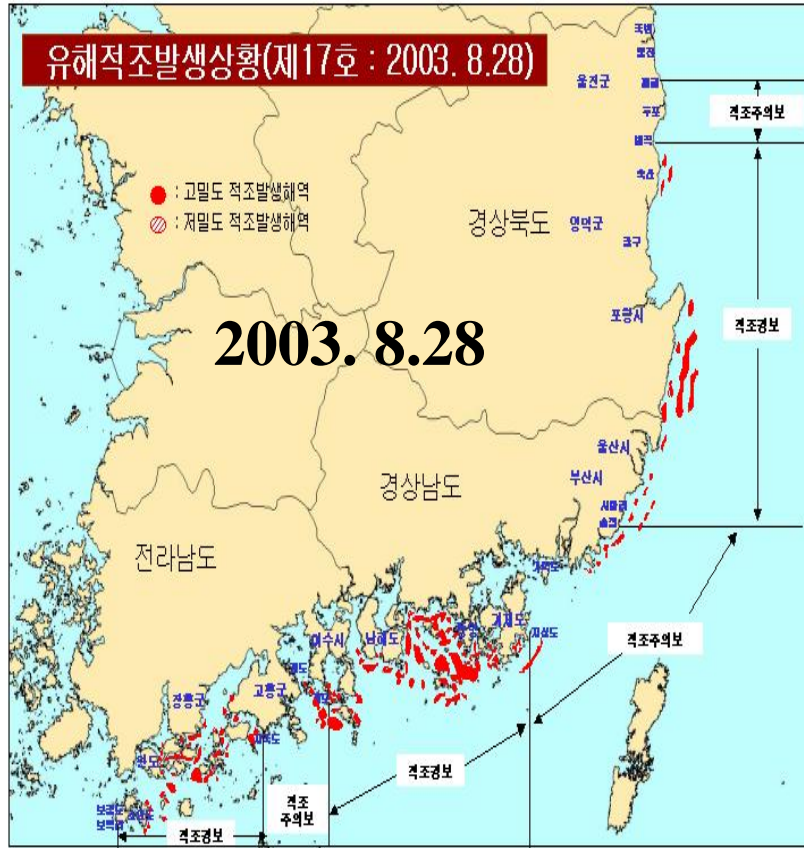
On the conditions for
Cochlodinium blooms:
A study using an individual-based model

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KORDI

Ansan, South Korea

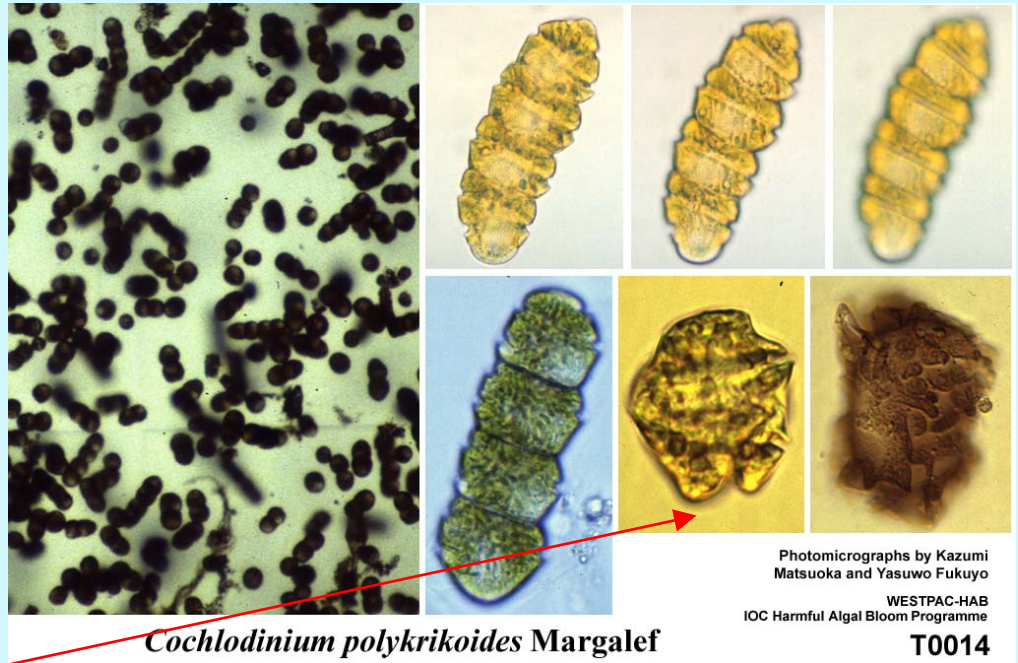
Fish-killing *Cochlodinium* HAB



From Redtide watch (NFRDI)

Cochlodinium polykrikoides

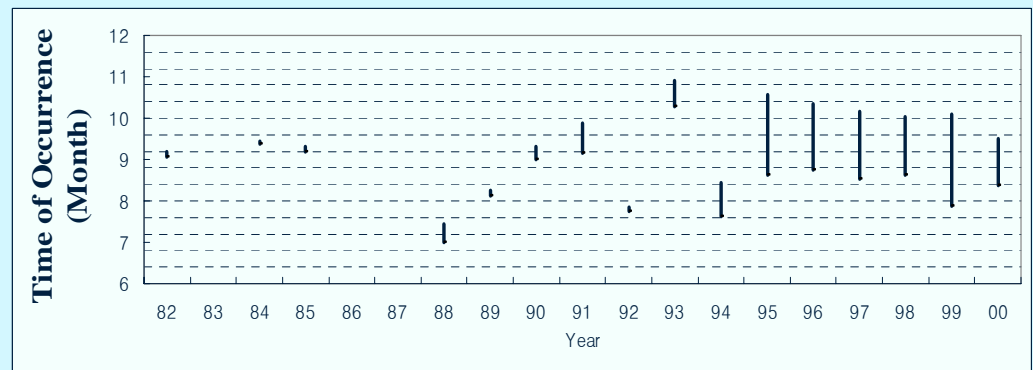
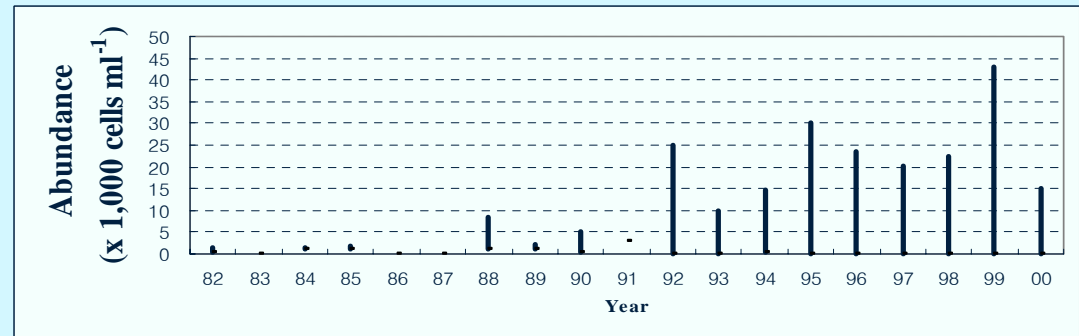
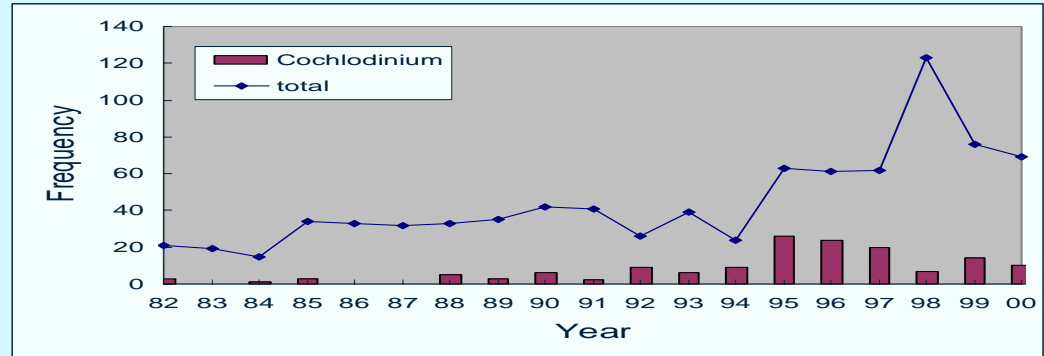
- An photosynthetic athecate planktonic dinoflagellate
- Cosmopolitan warm-water species
- Chain forming (usually ≤ 8 cells)
- cell size: 30~40 μm in length, 20~30 μm in width.
- binary fission (asexual reproduction) and **cyst** forming (probably as a result of sexual reproduction)
- Extensive fish kills in Korea and Japan (damage: US\$50,000,000 in Korea in 1995)



(by Matsuoka and Fukuyo)

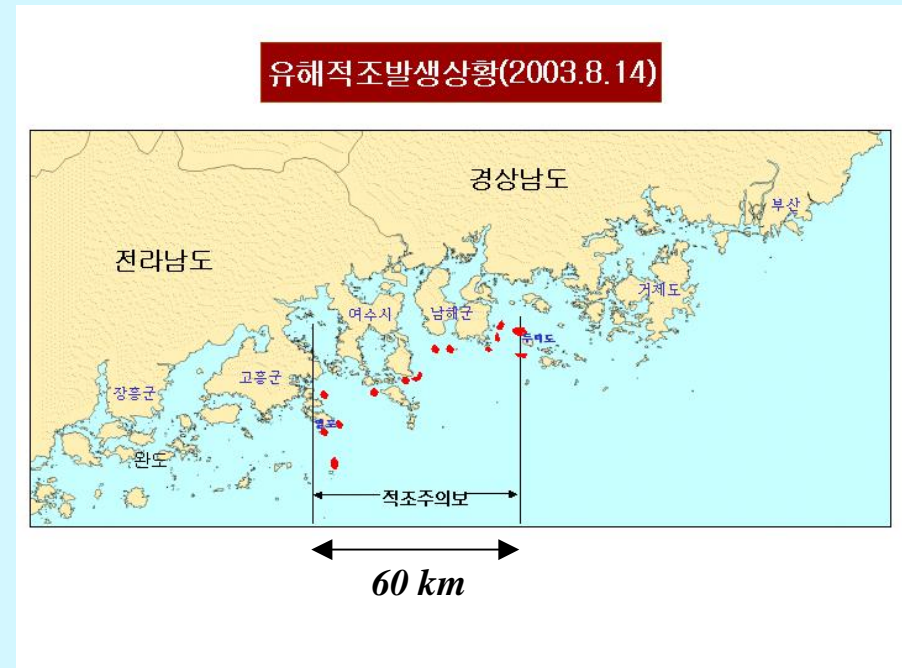
Trend in *Cochlodinium* blooms

- ❖ Since the first appearance in 1982, *Cochlodinium Polykrikoides* blooms increased in frequency, magnitude, duration and areal extent constituting a major threat to Korean mariculture industry (Kim *et al.*, 2001).

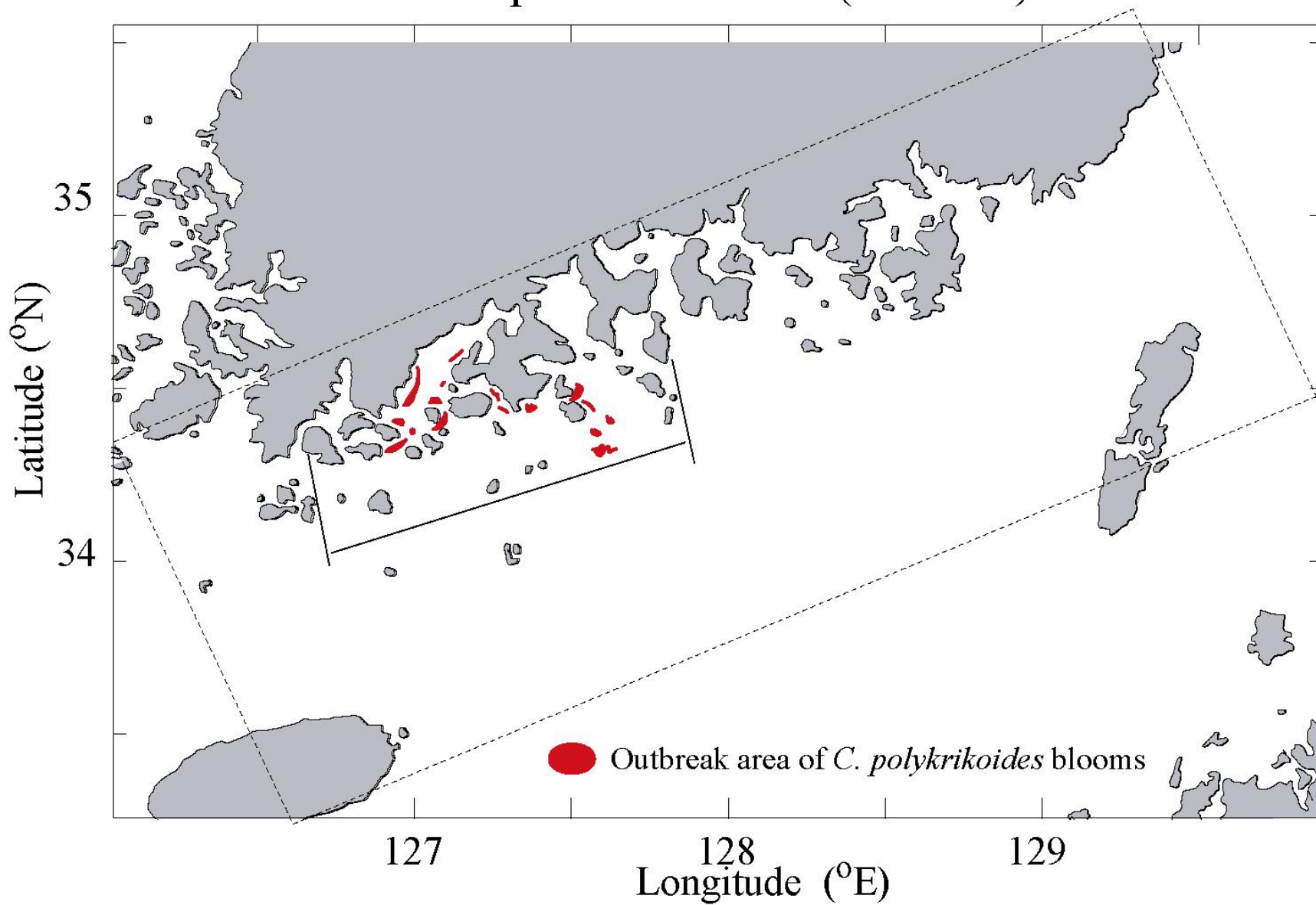


Features of *Cochlodinium* blooms in Korean waters

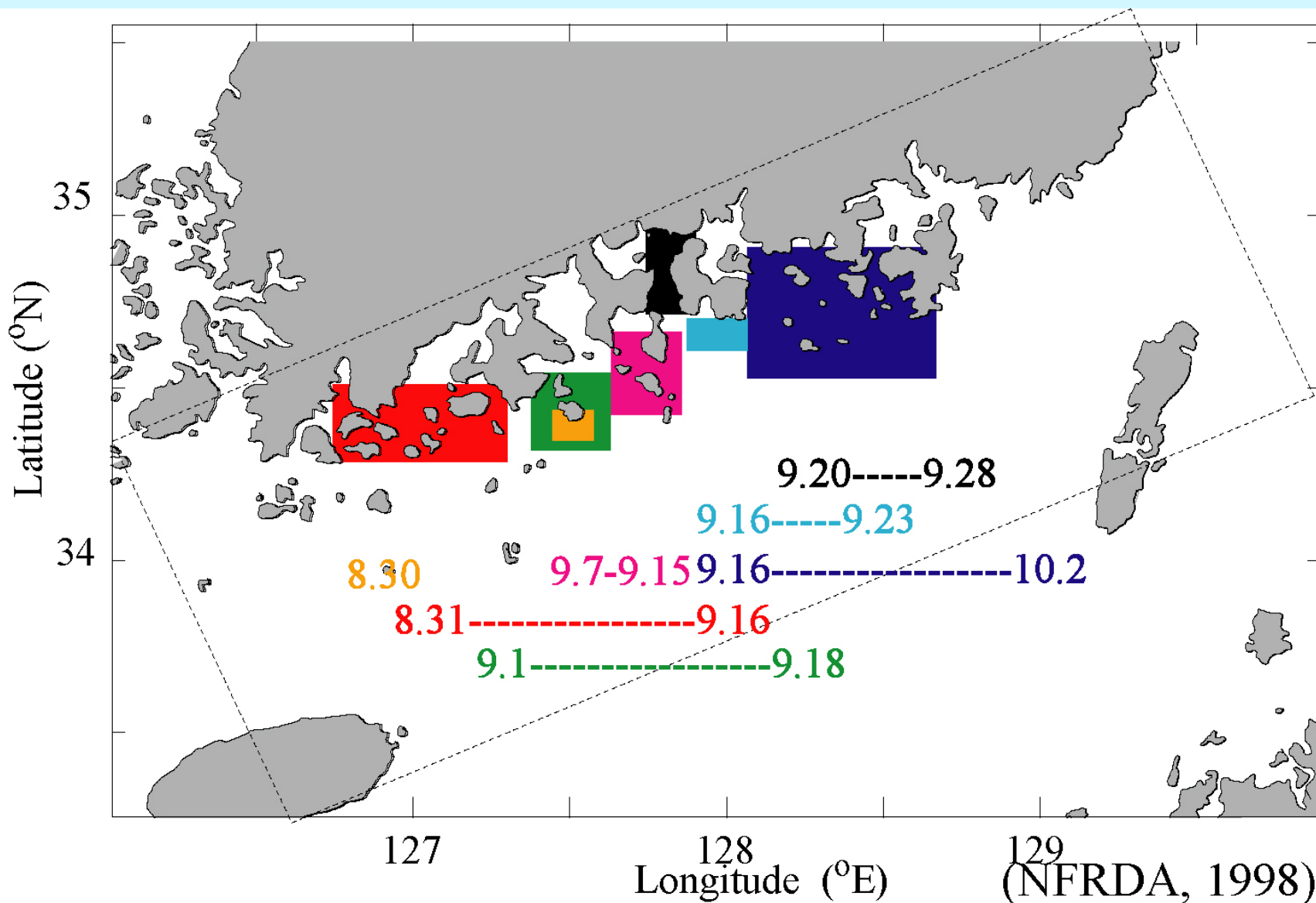
- ❖ Appear in Aug and continue until Sep/Oct (Although cysts have been identified, exact life cycle is not known).
- ❖ Usually begin in outskirts region
- ❖ Eastward progression in a season
- ❖ Westward expansion in the past three decades

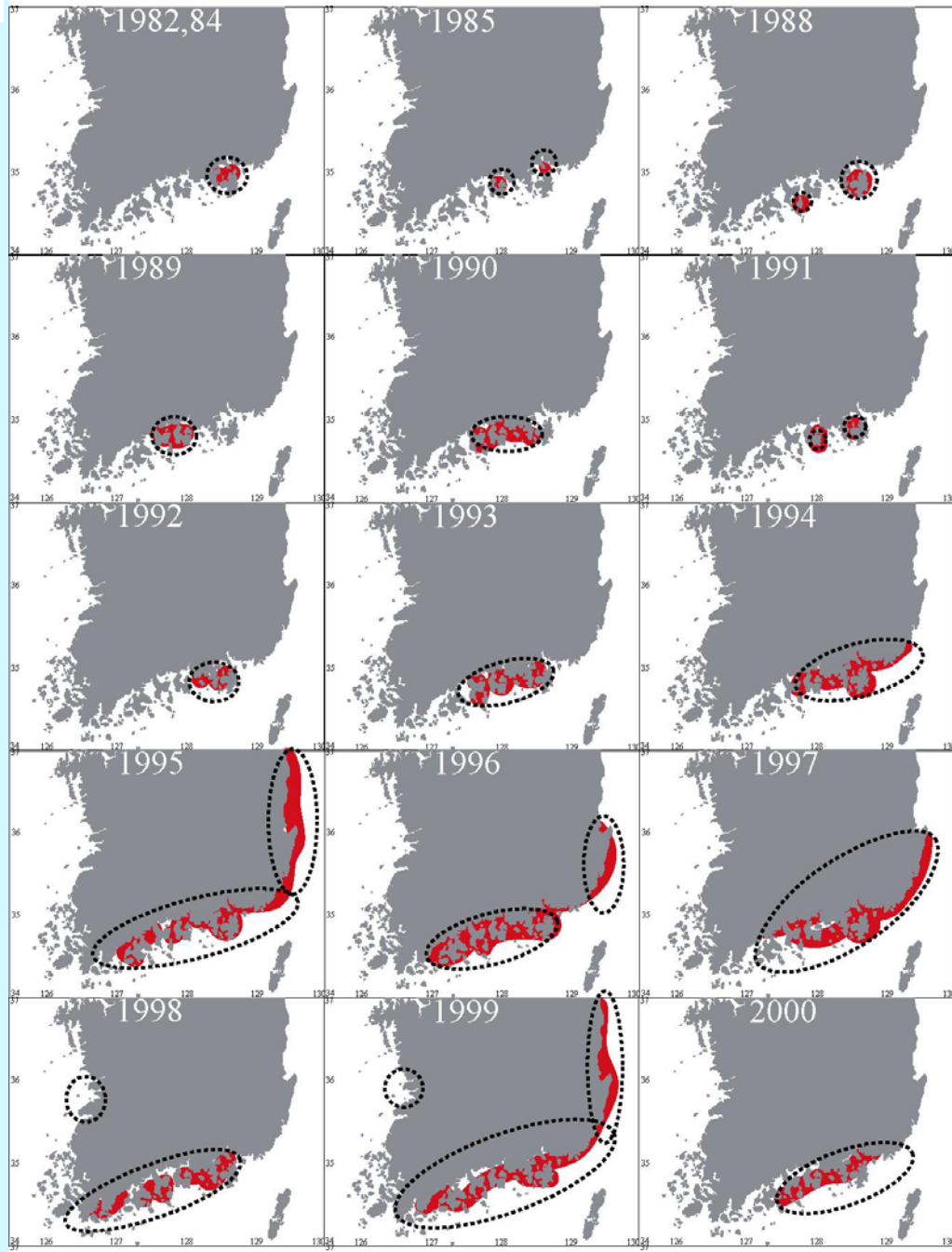


September 8. 1998 (NFRDA)




Eastward progression (1998)





Areal
expansion of
C. polykrikoides
blooms

 Outbreak area of *C. polykrikoides* blooms.

Objectives of this study

- ❖ Can we explain spatial features of *Cochlodinium* blooms in Korean waters?
 - Usually begin in outskirts region
 - Eastward progression in a season
 - Westward expansion in the past three decades

(Alternative hypotheses that have been proposed involve oil spills, nutrients, zooplankton, etc)

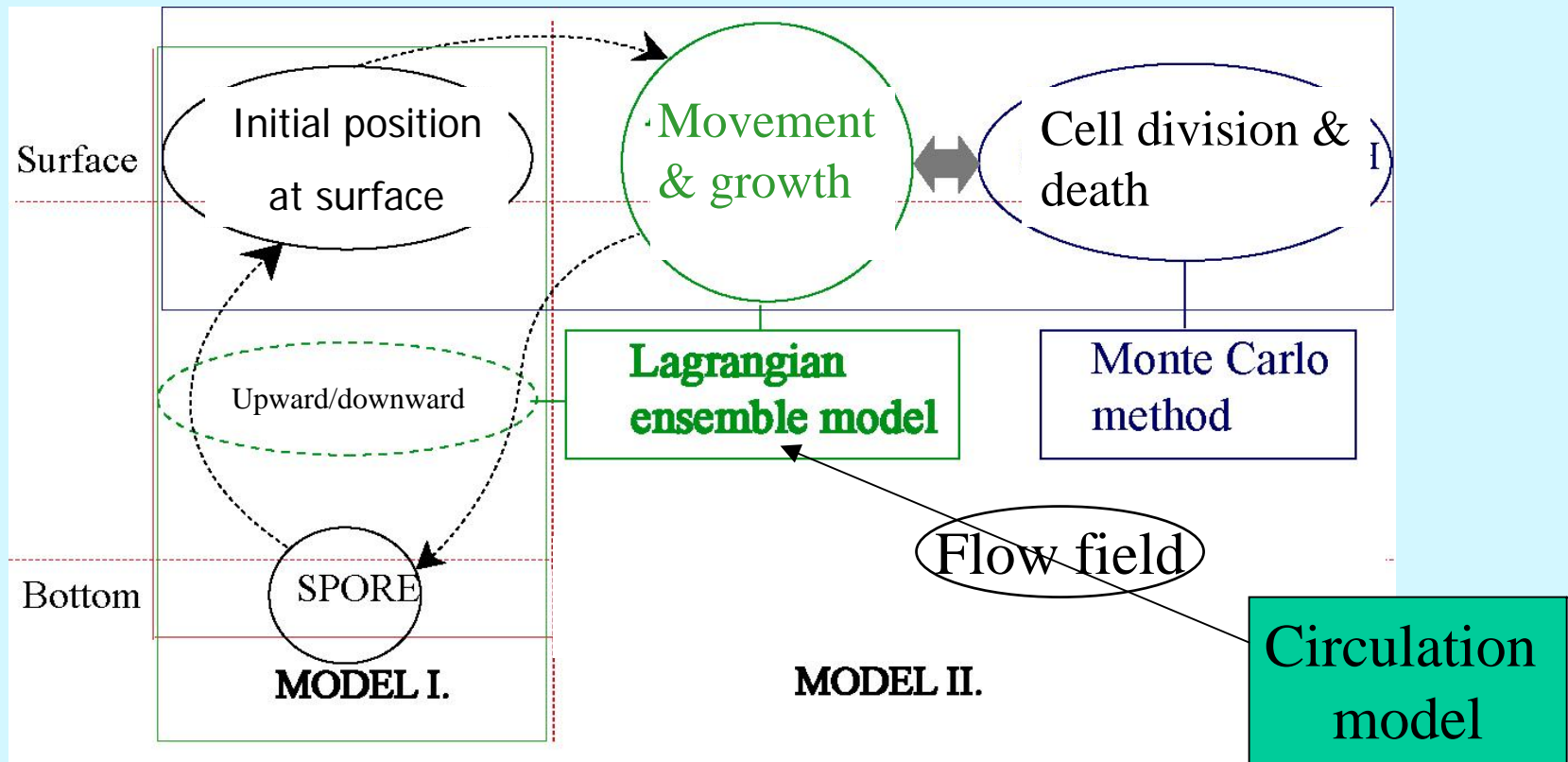
- ❖ Allochthonous seeding or autochthonous cycles?

IBMs (Individual-Based Models) could be more useful for HAB modeling

- ❖ Aggregated (“Chl”) models are not appropriate for HAB study.
 - Biomass increase is a regional, seasonal, and species-specific aspect. Therefore, biomass is not an appropriate indicator of HAB (Smayda, 1997).
 - Chlorophyll conc. is a community property representing mass-balance which cannot resolve the dynamics of each population.
 - Bloom condition itself does not necessarily mean harmfulness

Questions addressed by the Model

- How the current field affects settling of cysts and surfacing of the germinated cells?
- How the current field affects the spatial pattern of HAB growth?



Schematic diagram of modelled processes. The model consists of two part; Model I is for vertical movement of spores and Model II is for spatial development of HAB due to currents and reproduction of organisms.

Circulation Model

- . **Three-dimensional POM (Princeton Ocean Model)**
- . **Arakawa-C grid**
- . **Incompressible, hydrostatic, Boussinesq, mid-latitude β -plane approximations**
- . **σ -coordinate in the vertical, and cartesian coordinate in the horizontal**
- . **K_M, K_T : turbulence closure**
- . **A_M, A_T : Smagorinsky formula**
- . **Free surface**

- . **Model domain includes three open boundaries.**
- . **Spatial resolution: $3\text{km} \times 3\text{km} \times 8$ layers**
- . **Synoptic modeling to simulate circulation in September, 1998**

Circulation Model: Boundary conditions

❖ *Open boundary conditions*

- Specification of tides (M2, S2, O1, K1) based on tidal chart
- **Specification of steady inflow/outflow volume transport based on diurnally-averaged currents across the Korea ($V_{east}=1.72$ Sv) and Cheju Straits ($V_{west}=0.65$ Sv) observed in September, 1998;
 $V_{south}=V_{east}-V_{west}$**
- **Specification of climatological 30-year mean T, S distribution**

❖ *Surface boundary conditions*

- Spatially-uniform wind based on the observation at a buoy station at the end of August 1998
- **Wind stress is gradually increased towards the observed value over 7 days after the model reaches to a statistically equilibrium state, and is decreased to zero afterwards.**
- **Relaxation of surface T, S based on 30-year mean climatology**

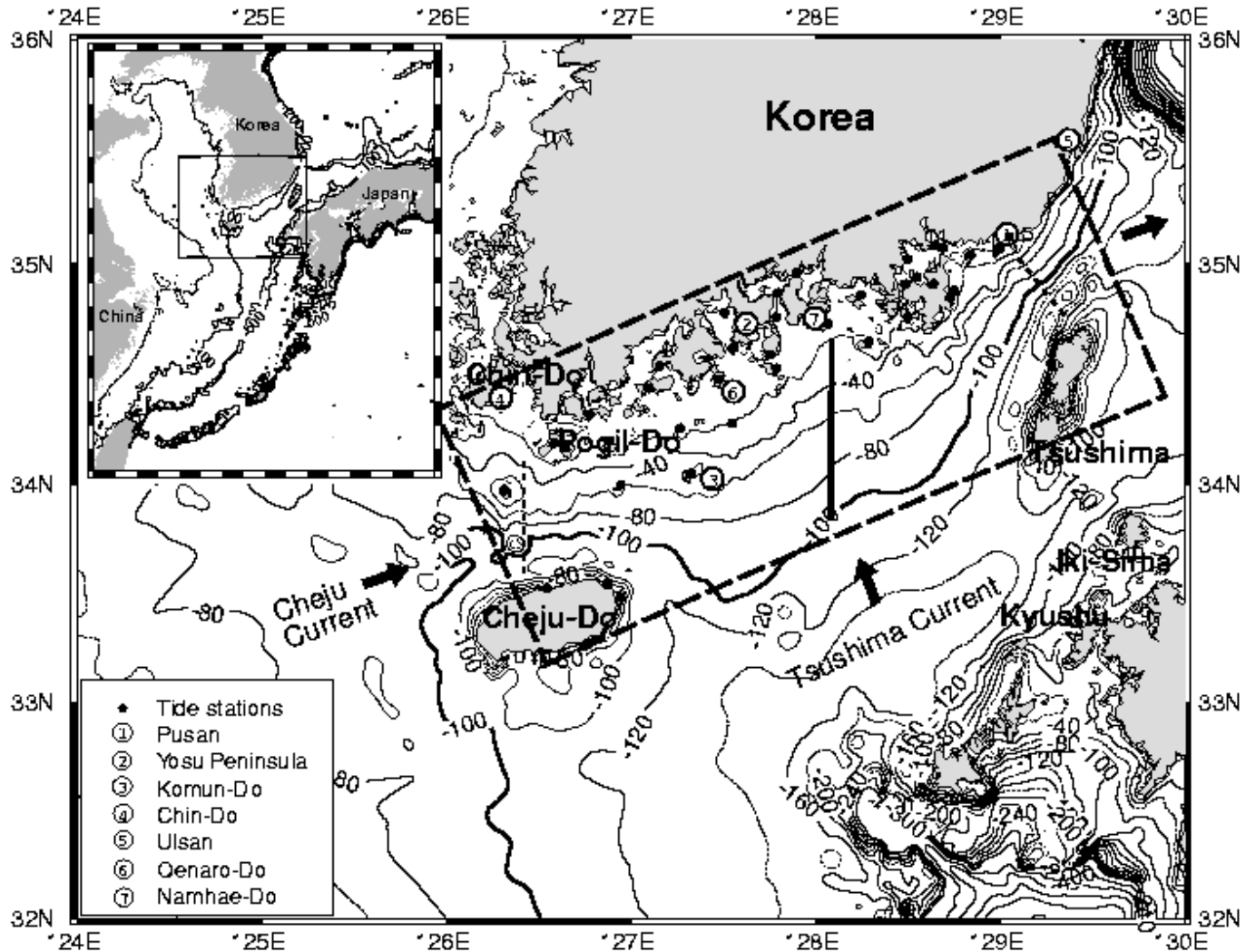
❖ *Bottom boundary conditions*

- **Quadratic bottom friction with $C_d=0.001$**

❖ *Initial conditions*

- Climatological mean T, S in September based on 30-year mean T, S in August and October
- **$U=V=0$**

Model domain



Biology model

❖ Growth terms

- Light: saturation assumed
- Nutrient: DIN - Michaelis-Menten eq.
- Intrinsic growth rate: from culture exp. (T, S)

❖ Loss terms

- Constants (0.02, 0.05, 0.1, 0.2)
 - Turbulence: effects are ambiguous
 - Predation: difficult to parameterize the spatial heterogeneity

Diel vertical migration of *Cochlodinium polykrikoides*

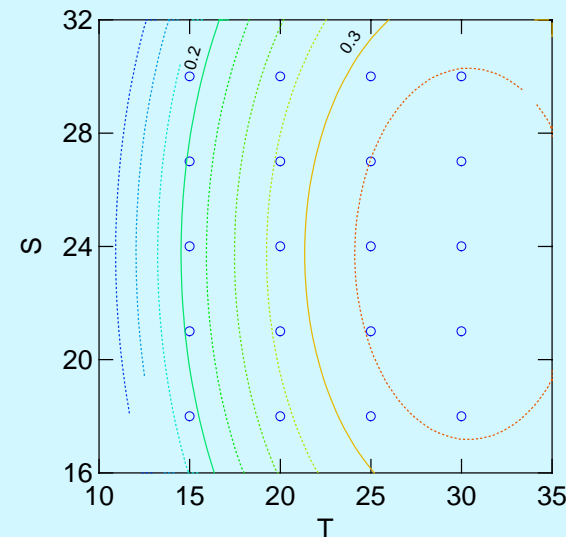
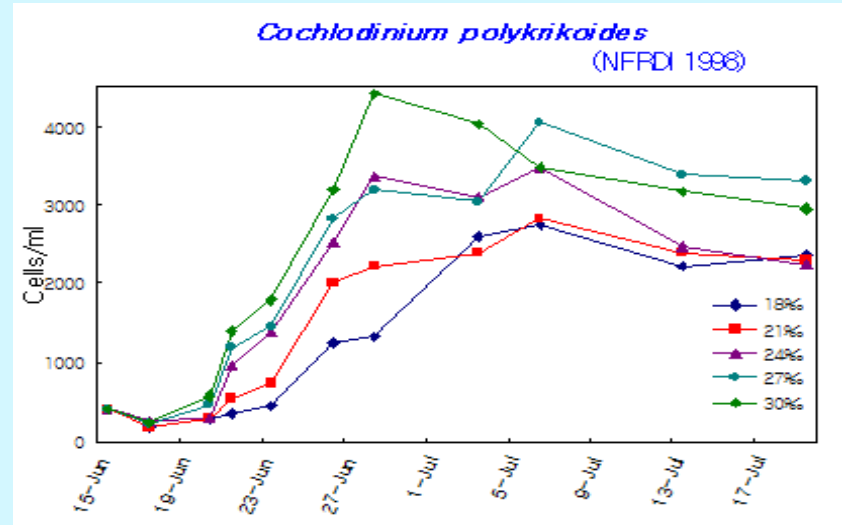
- ❖ Diel vertical migration (NFRDI, 2002)
 - 10:00~16:30: > 80% stayed in the upper 3m layer
 - 16:30~17:00: Downward movement begins
 - Swimming speed: mean = 1036 (± 84), max 1449 $\mu\text{m/s}$ (Jeong et al., 1999)
- ❖ Possibly for dark consumption of nutrients
- ❖ No light limitation was assumed.
 - I_m (saturation light intensity) = 100~200 $\text{OE m}^{-2} \text{s}^{-1}$ (NFRDI, 2002).

Growth rate of *C. polykrikoides* as a function of T and S

1. Exponential growth rates were calculated from 20 (4T x 5S) culture experiments (NFRDI, 1998).

2. Then a quadratic function was fitted:

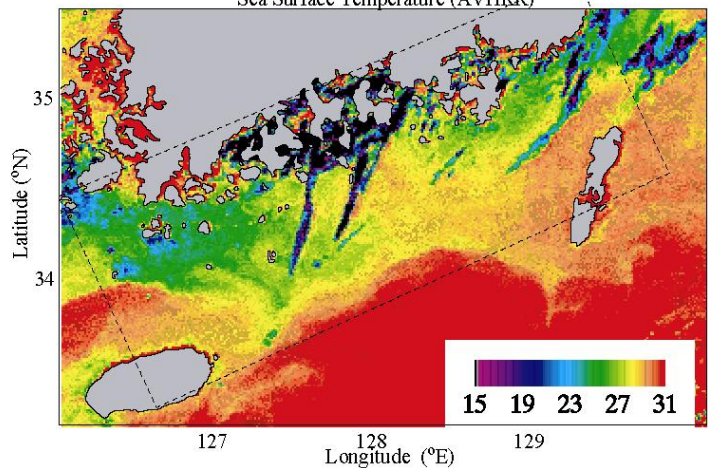
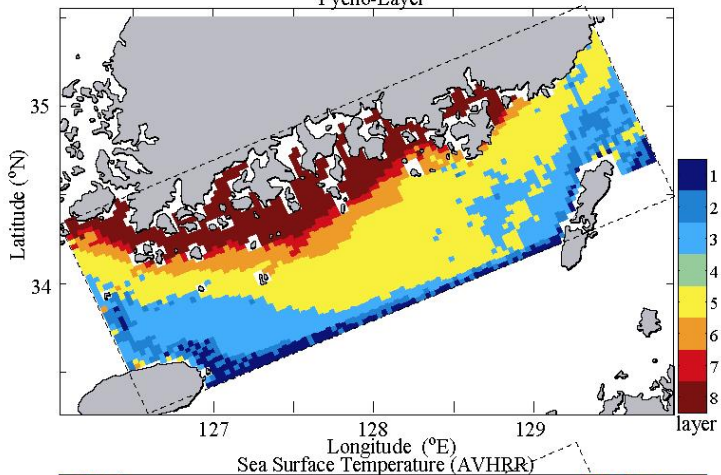
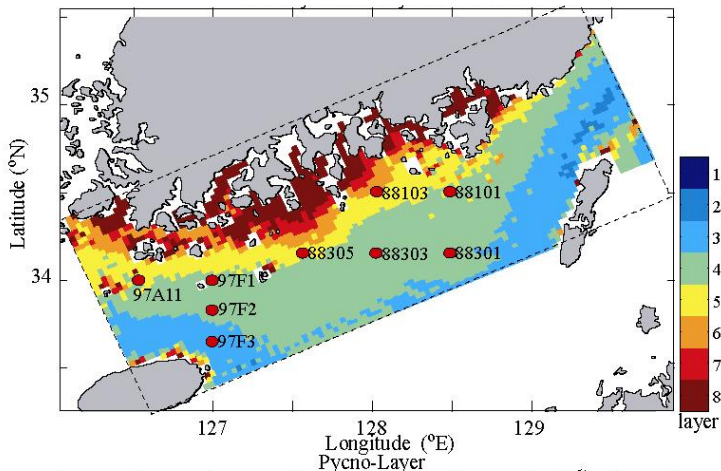
$$\begin{aligned} \mu = & -0.499 + 0.036T \\ & -0.001T^2 + 0.026S \\ & -0.001S^2 \\ & (R^2 = 0.488) \end{aligned}$$



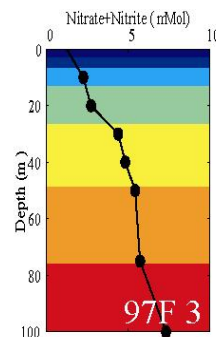
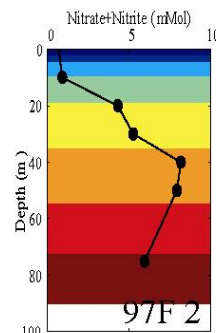
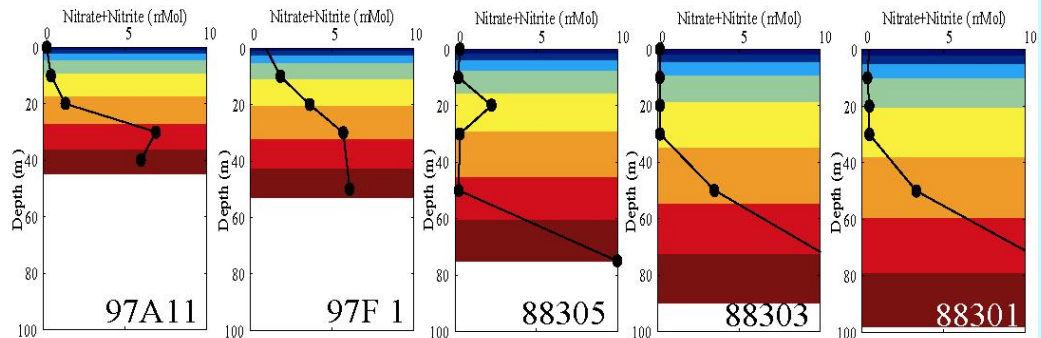
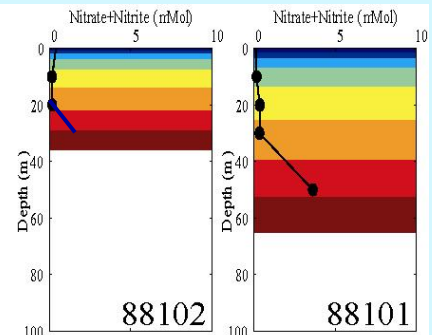
Computation of Nutrient limitation

- ❖ Dark consumption of nutrients is assumed.
- ❖ Vertical movement of the cells for six hours are calculated from swimming speed and vertical current velocity.
- ❖ Representative summer values are taken for upper and lower values of nutricline (only nitrogen is considered).
- ❖ $K_s = 2.1 \mu\text{M}$ (nitrate), $1.03 \mu\text{M}$ (ammonia), $0.57 \mu\text{M}$ (phosphates) (NFRDI, 2002)

Layer reached within 6 hours of vertical migration



DIN in summer



Cell division and death

- ❖ Synchronous cell divisions are assumed (Lipp, 1993).
- ❖ Divisions and deaths are counted every 24 hours using Monte Carlo method.

$$\frac{dN}{dt} = rN = (g - m)N$$

$$N_{t+1} = N_t(1 + p_g)(1 - p_m)$$

$$p_g = e^g - 1.0$$

$$p_m = 1.0 - e^{-m}$$

g : growth rate per day

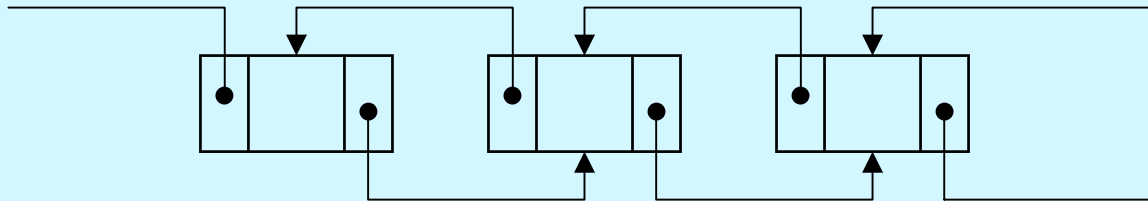
m : loss rate

p_g : proportion of the cells that divide

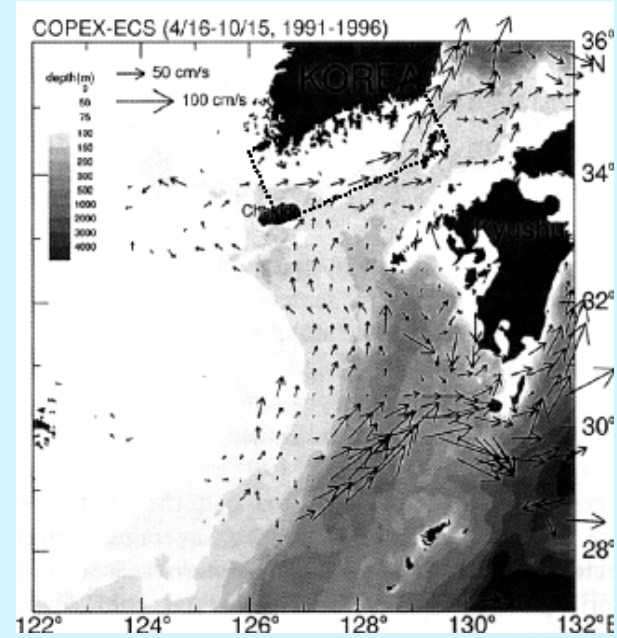
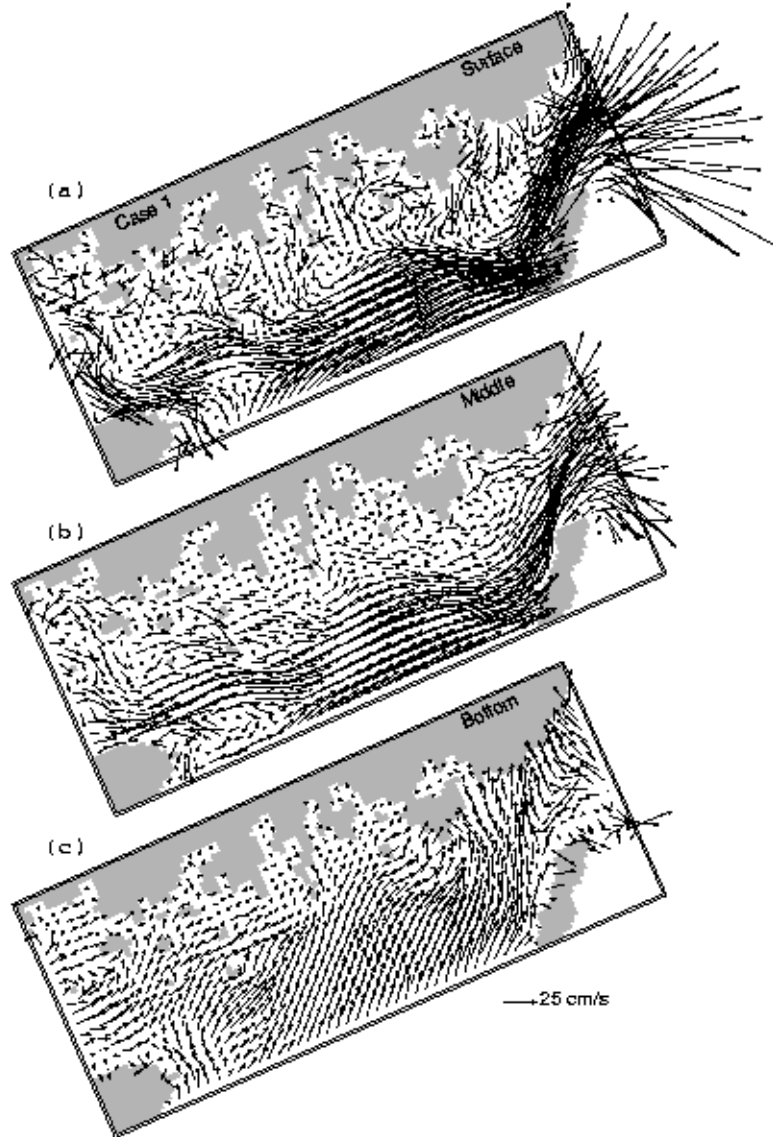
p_m : proportion of the cells that are lost

Computation

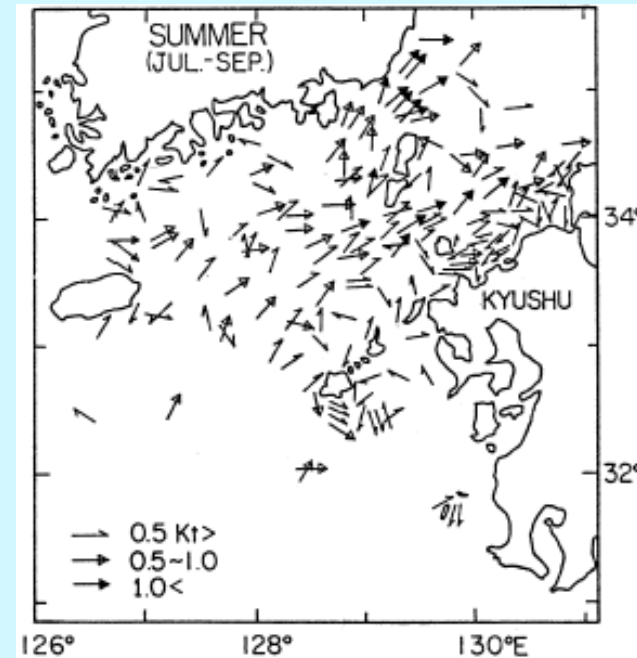
- ❖ Lagrangian movement of cysts and cells are calculated every 30 min using the current field from the circulation model.
- ❖ Linked-list technique was used for efficient memory management: $\sim 10^7$ cells are created but only $\sim 10^4$ cells are alive after 30 days.



Modelled flow field 15 days average

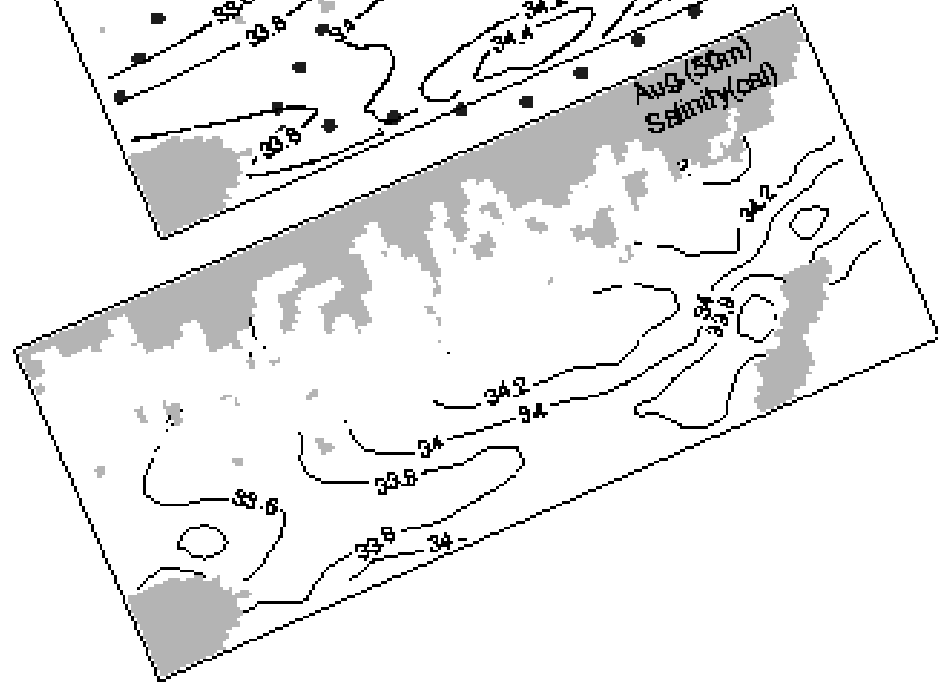
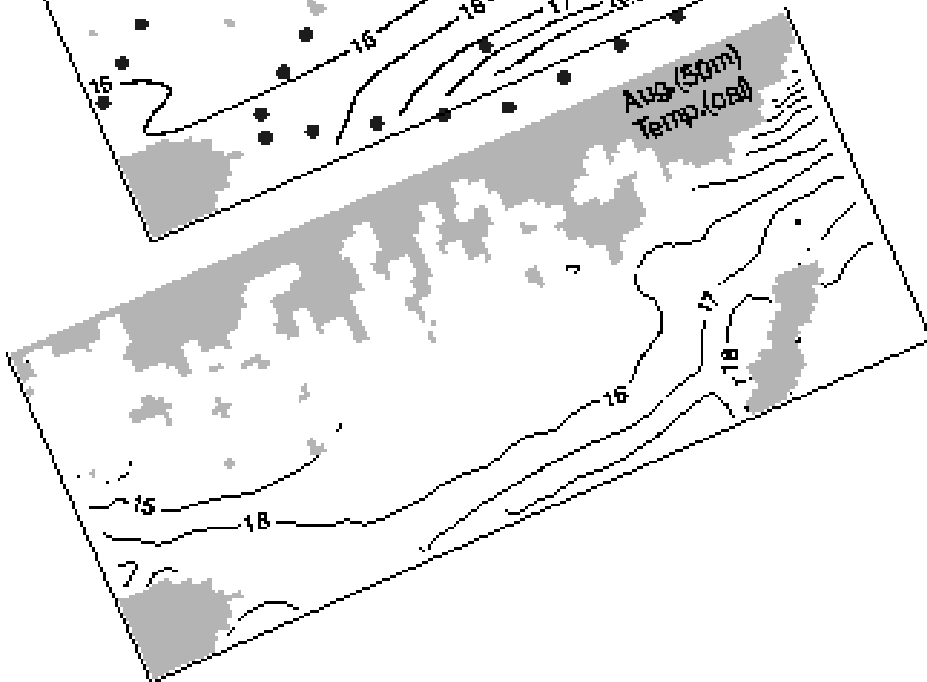
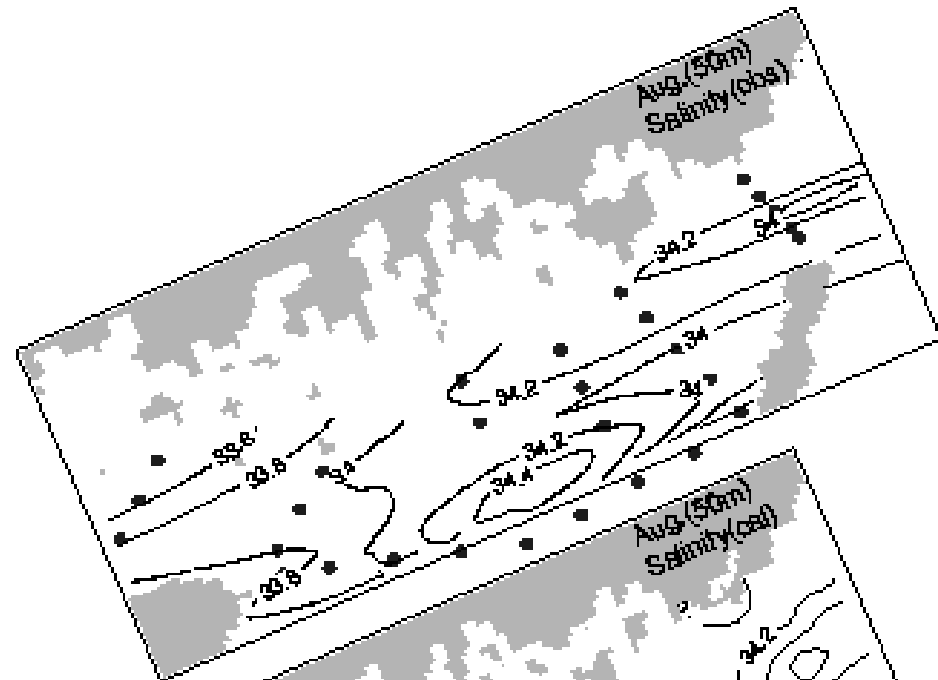
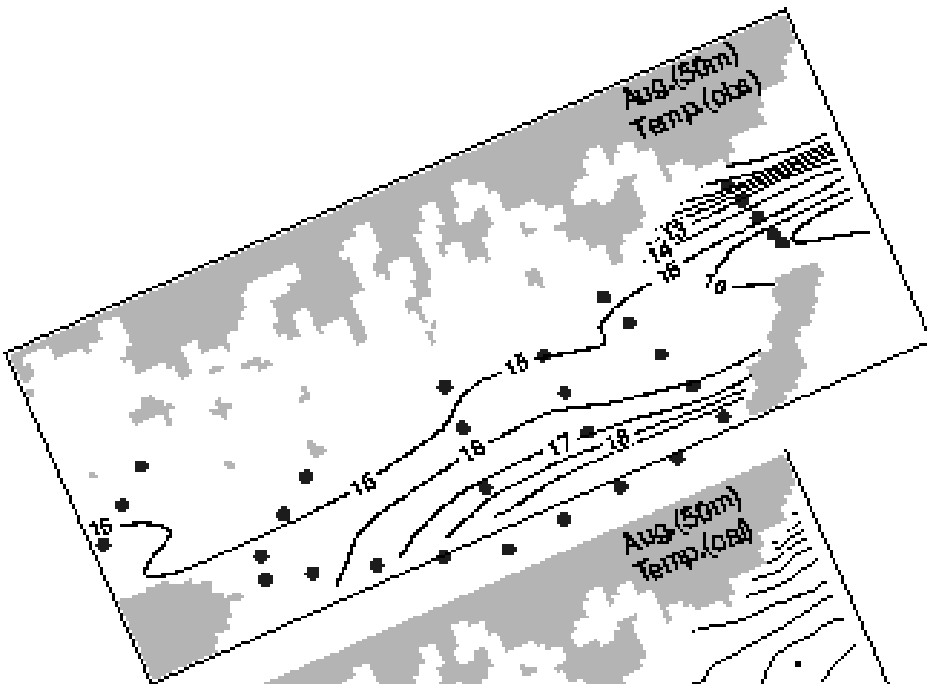


Drifter
(Lie&Cho,
1997)

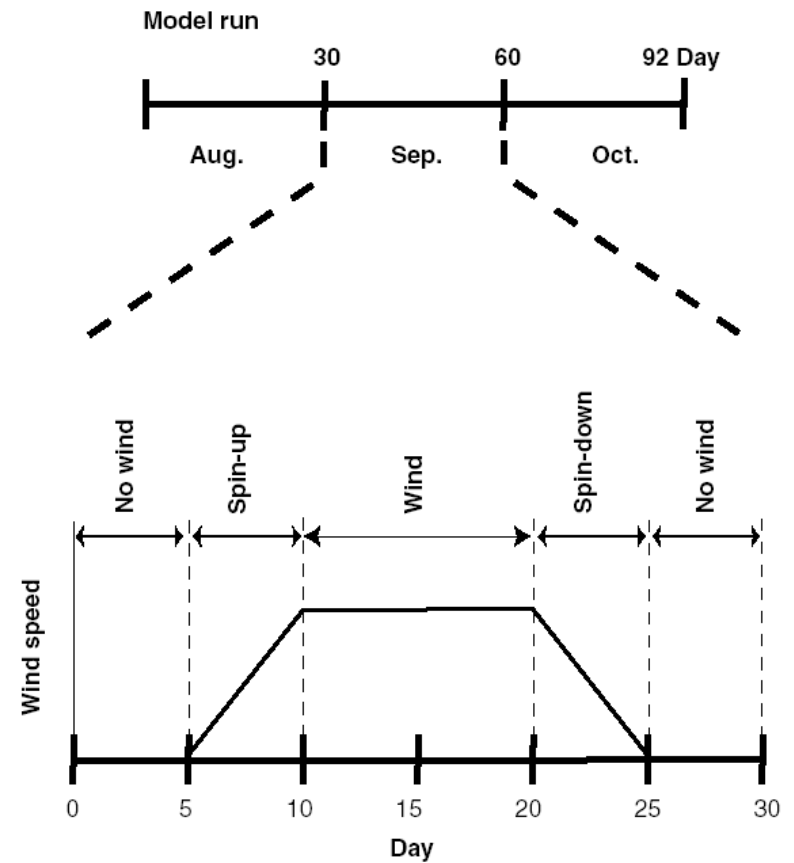
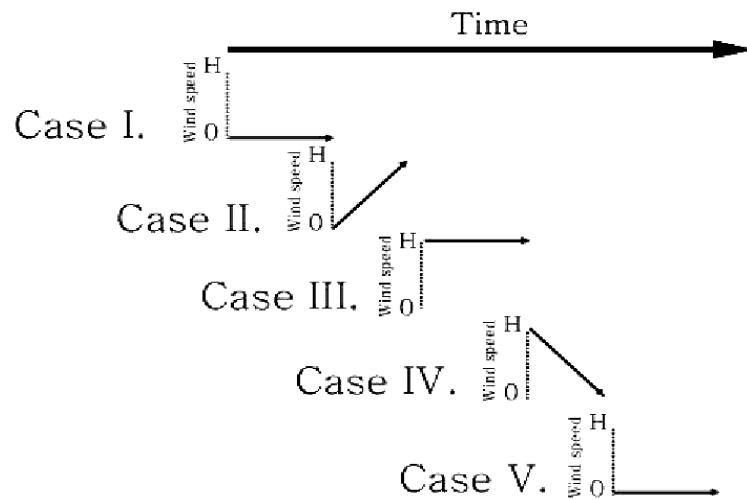


Current meter
(Mitta &
Ogawa, 1984)

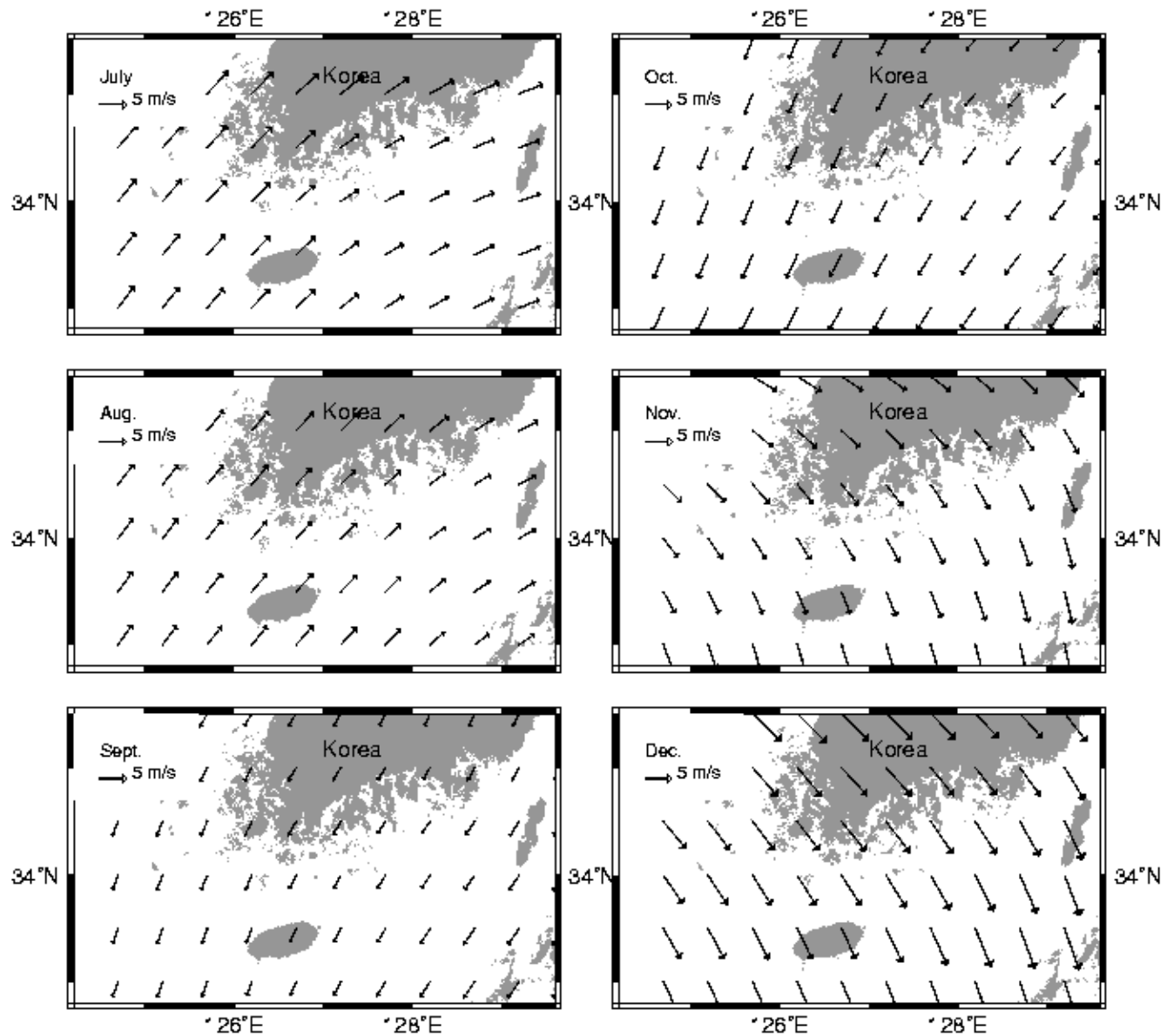
Comparison of model output and actual (T, S)



Five Scenarios of wind forcing

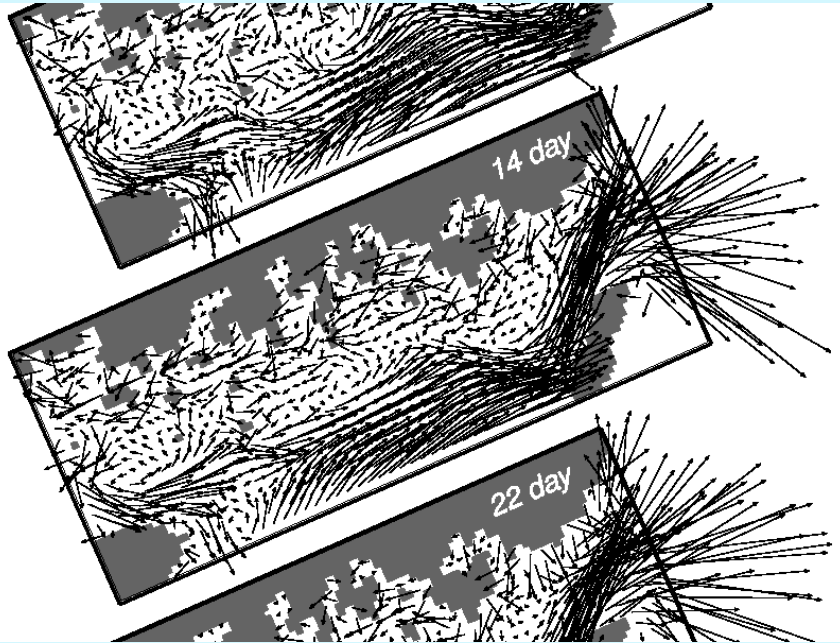


Monthly mean sea surface wind (Na and Seo, 1998)

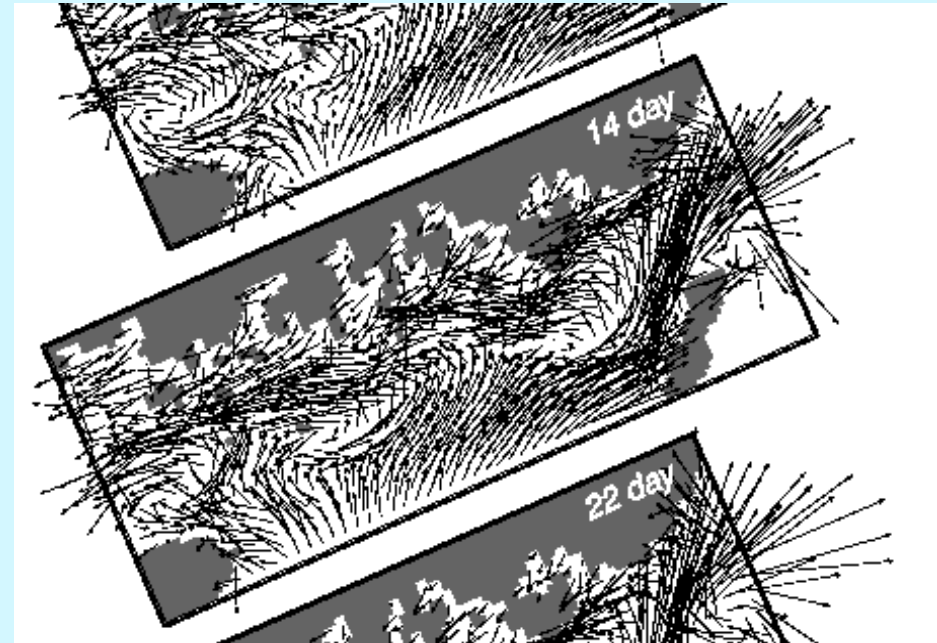


Wind and surface current

— 25 cm s⁻¹

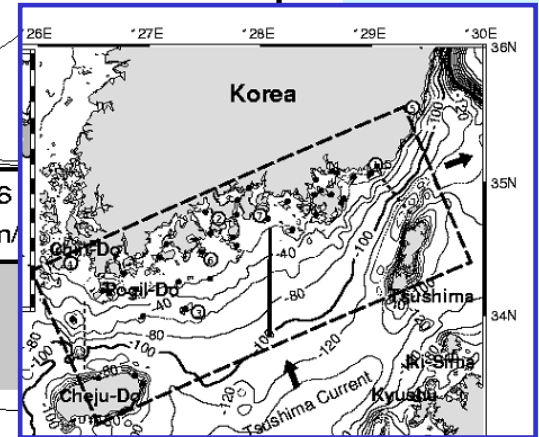
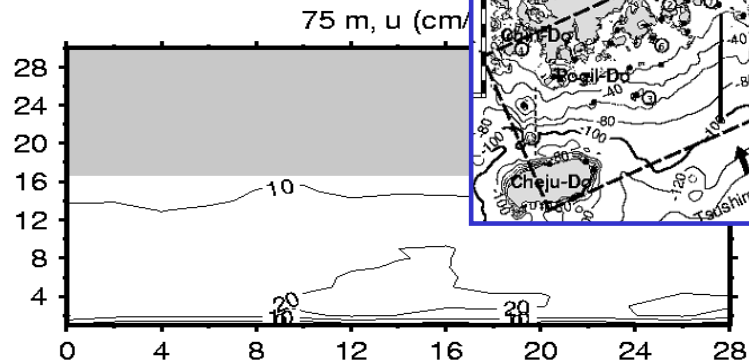
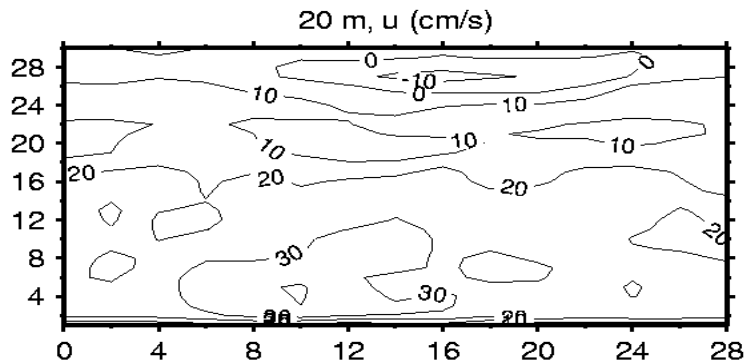
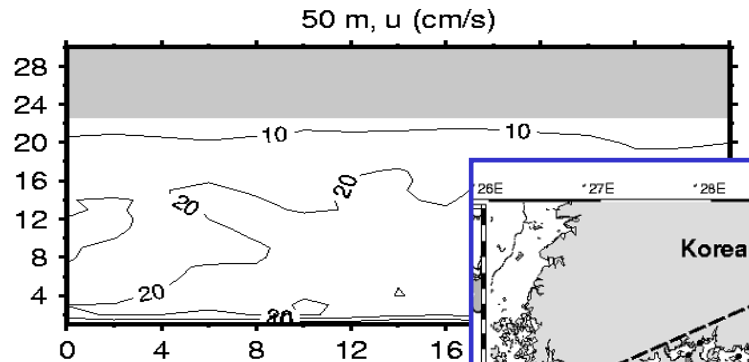
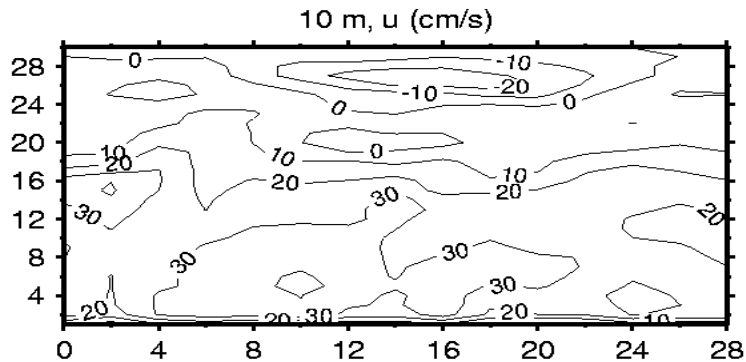
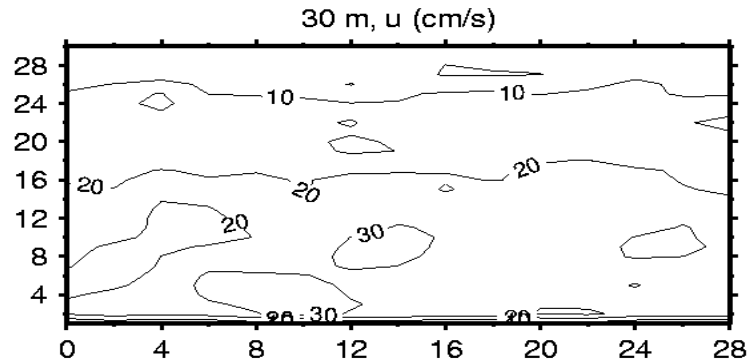
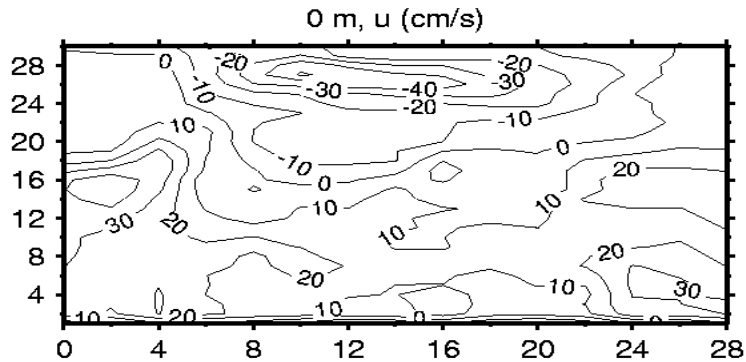


Wind speed: 3 cm s⁻¹

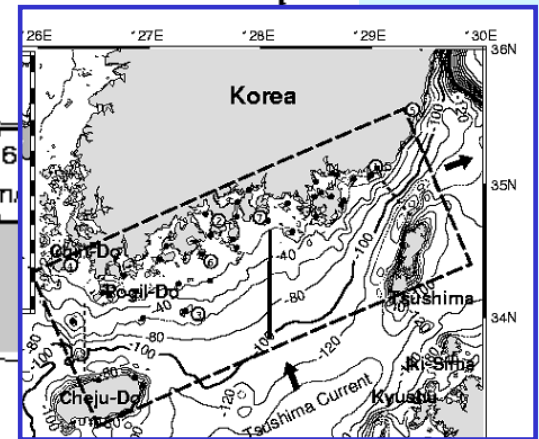
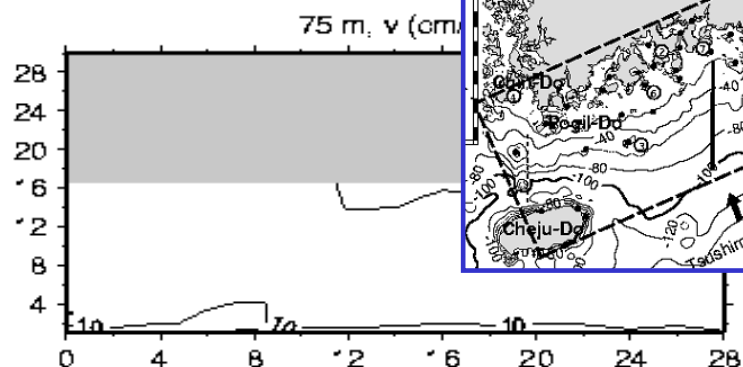
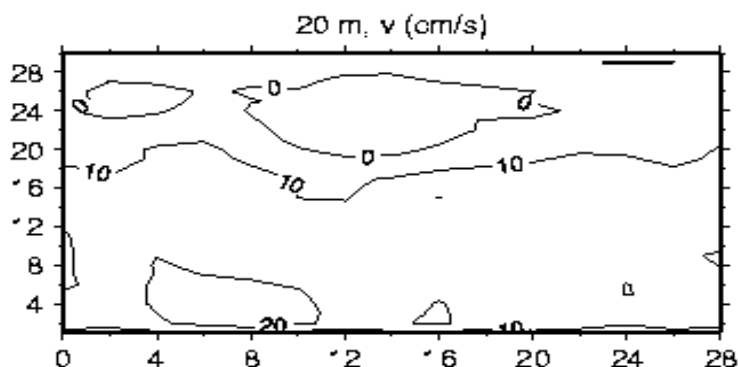
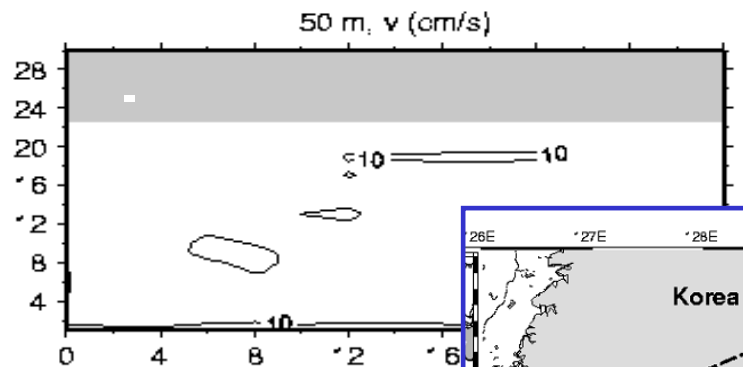
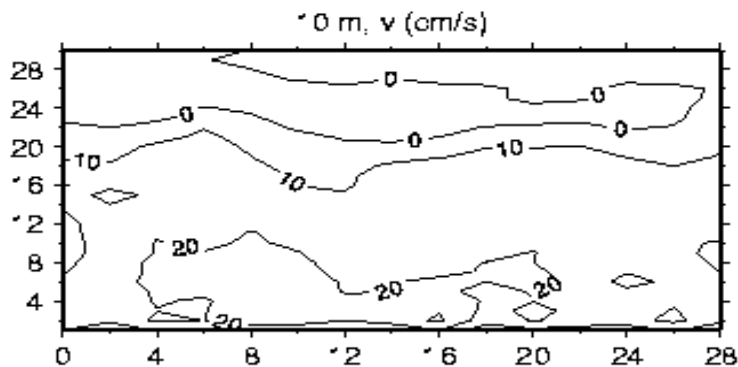
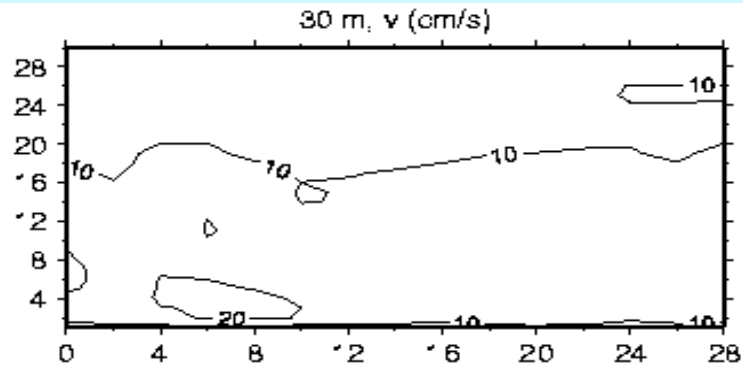
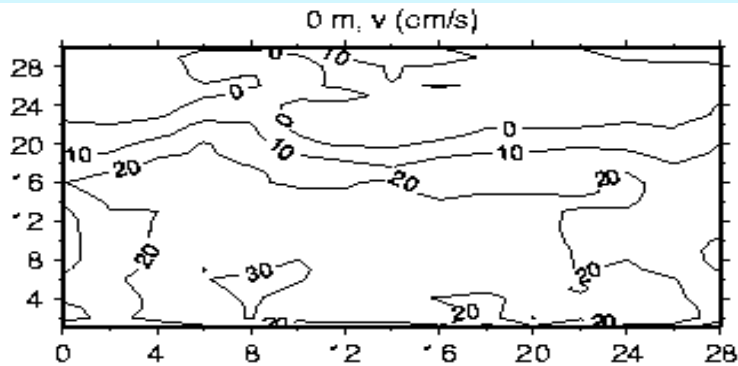


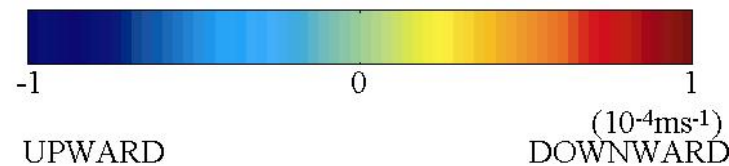
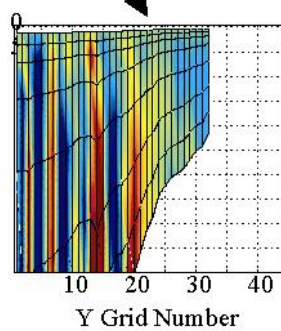
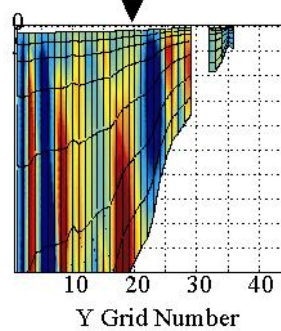
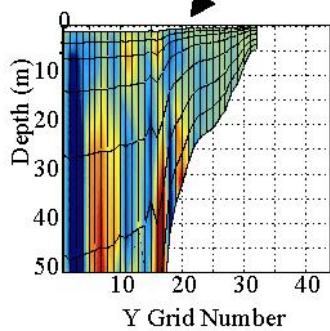
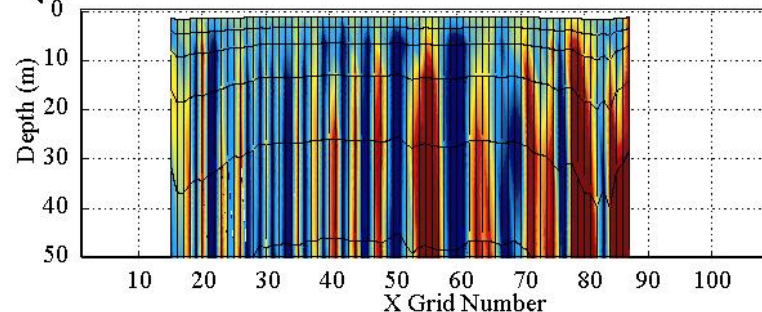
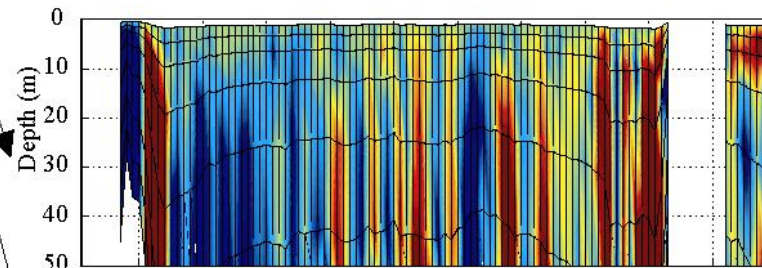
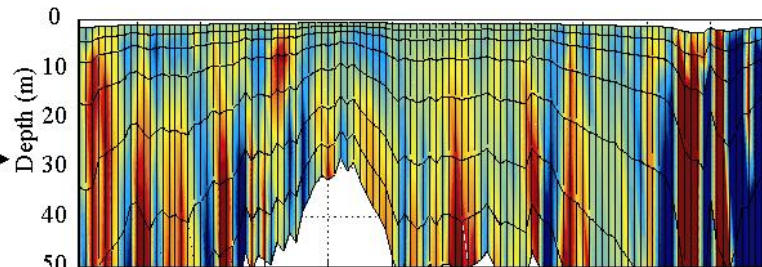
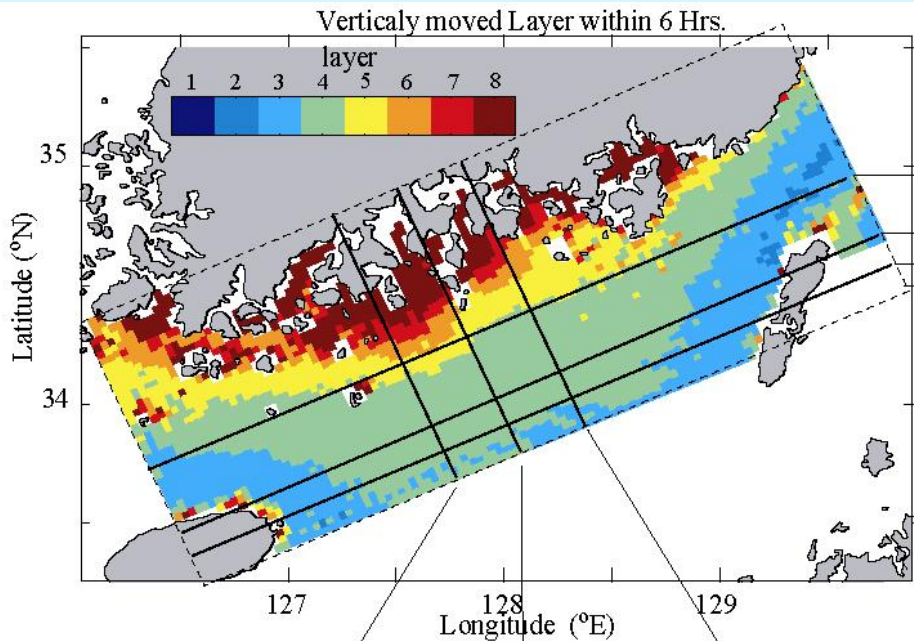
Wind speed: 10 cm s⁻¹

Time-space plot for the diurnally-averaged east-west component of currents along a meridional section with wind speed 10 m s^{-1}

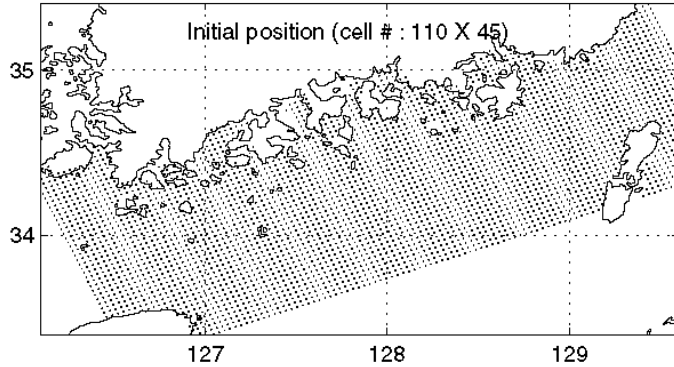


Time-space plot for the diurnally-averaged north-south component of currents along a meridional section with wind speed 10 m s^{-1}

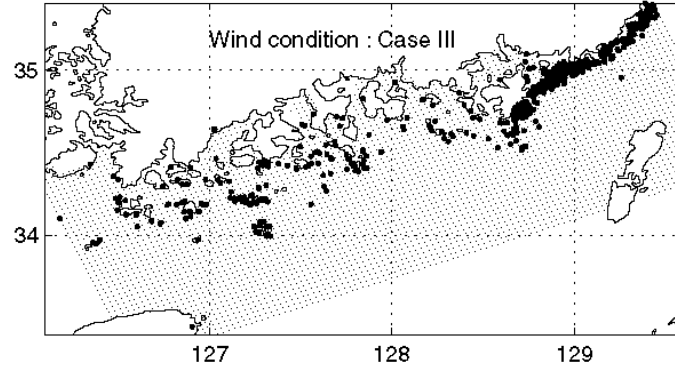




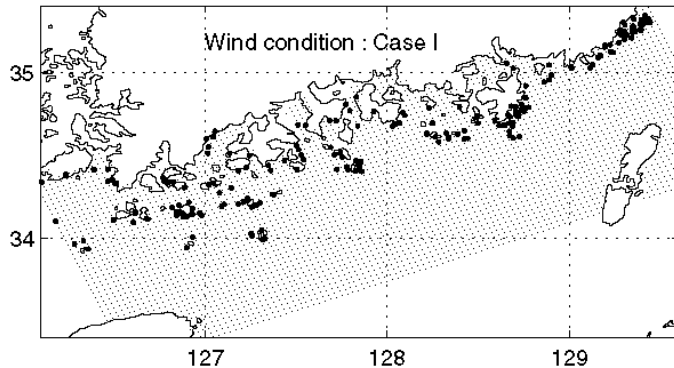
Settlement of cysts



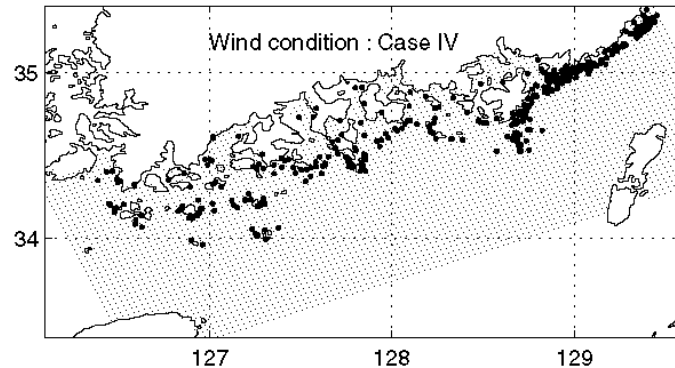
b)



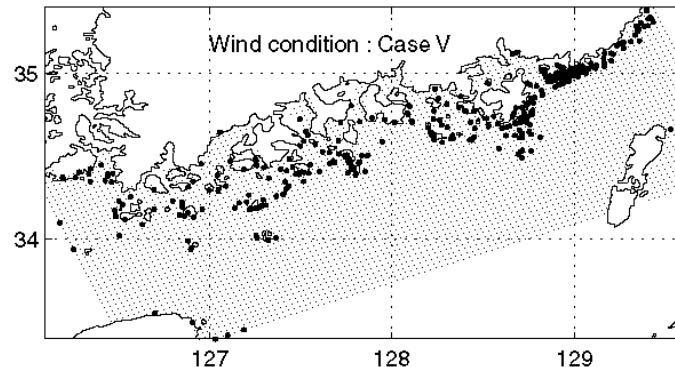
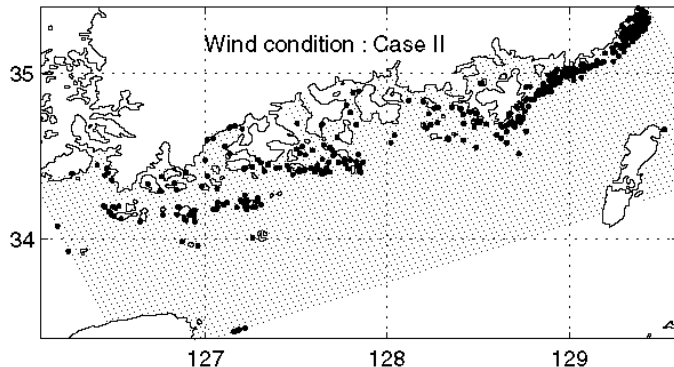
e)



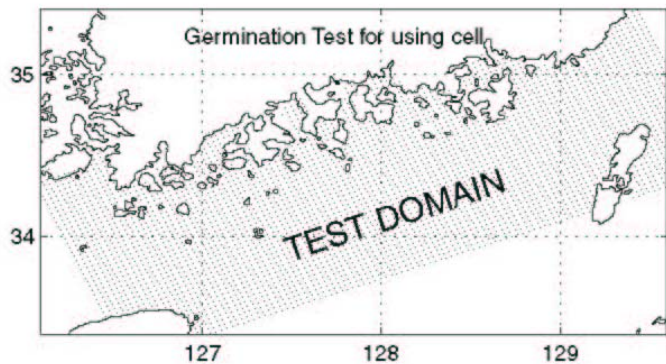
c)



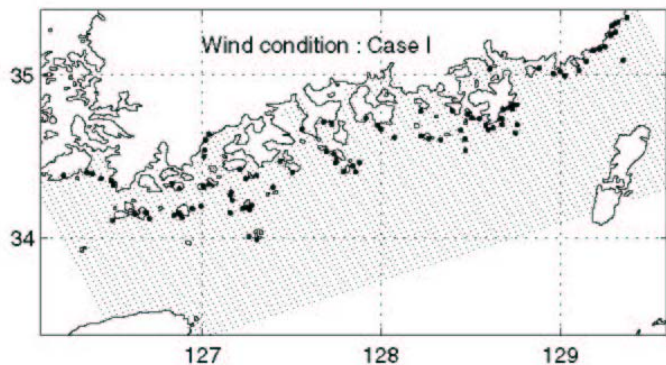
f)



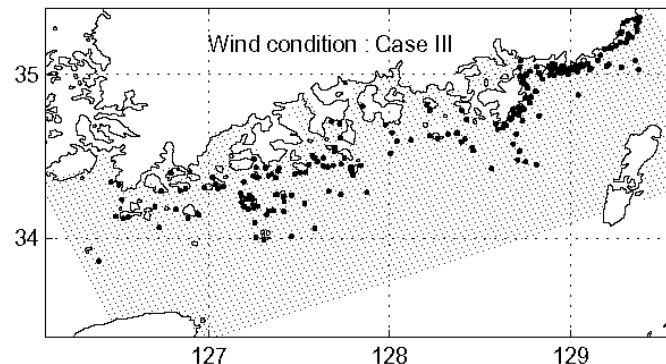
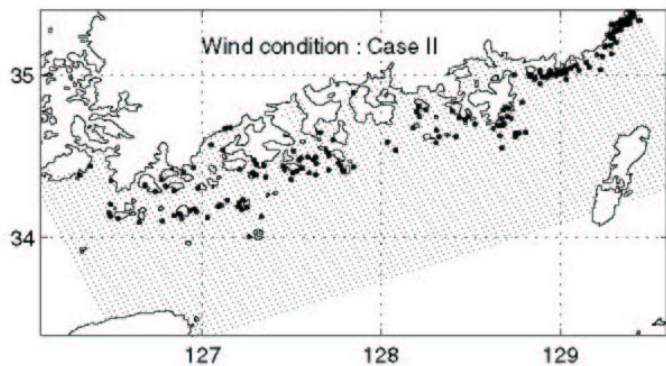
Ascending of germinated cells



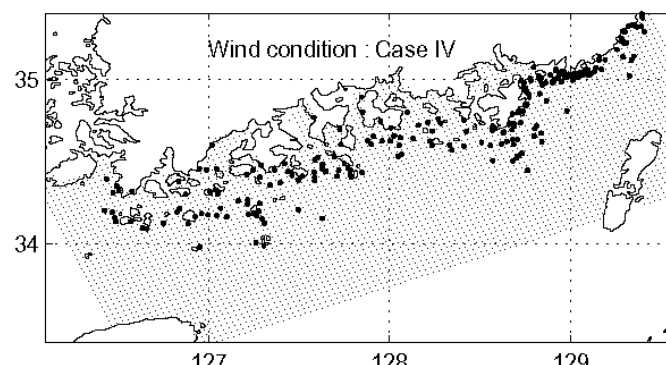
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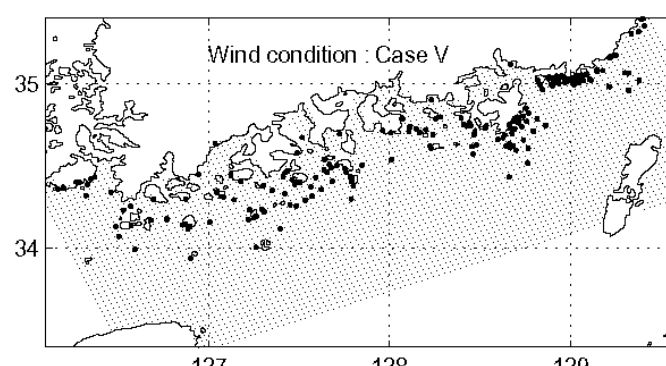
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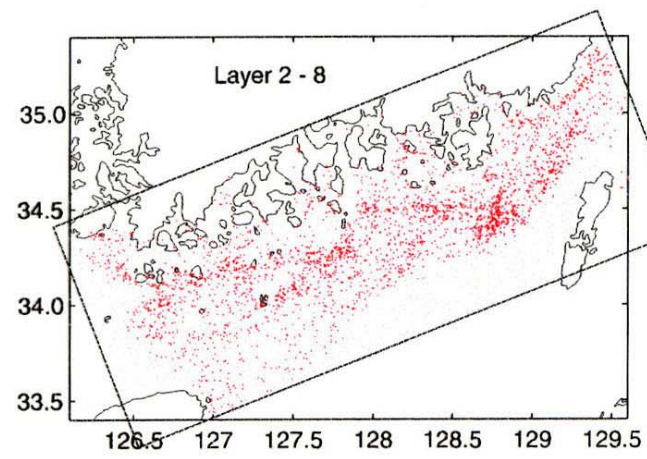
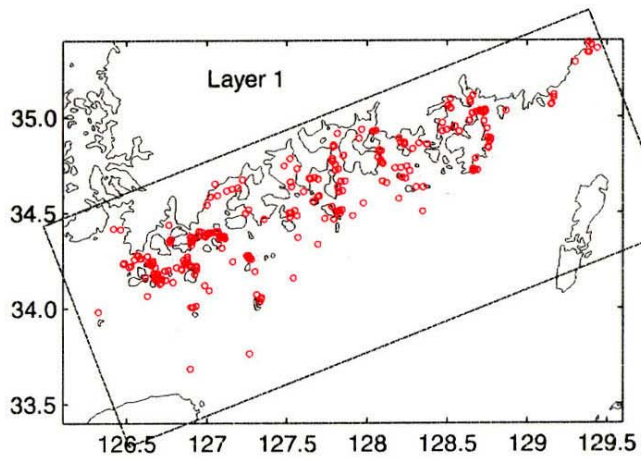
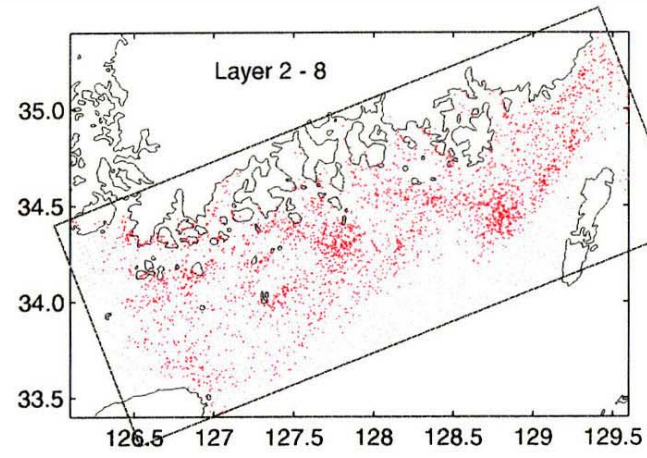
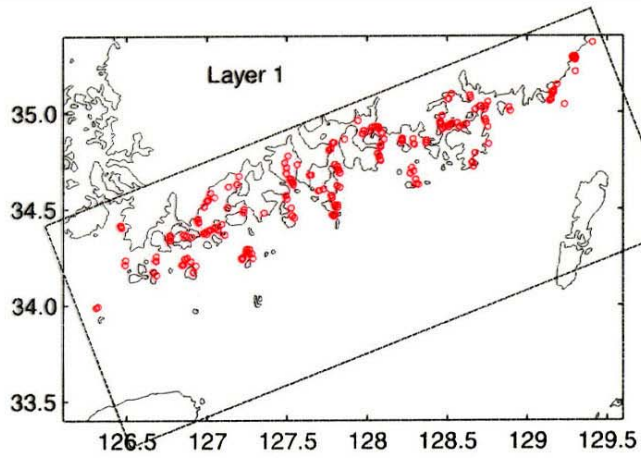


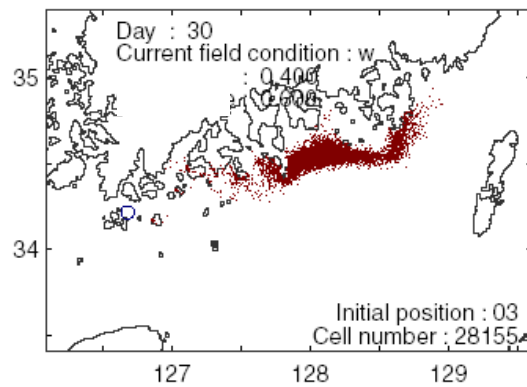
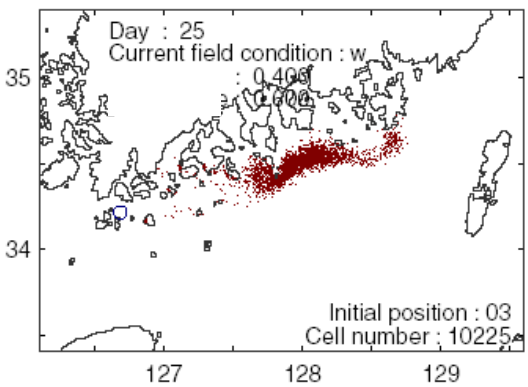
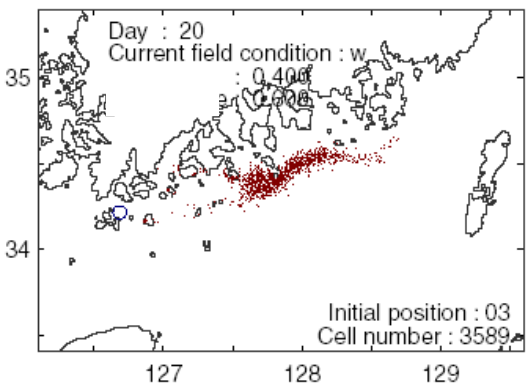
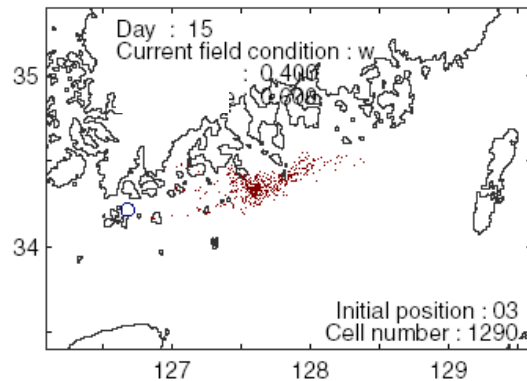
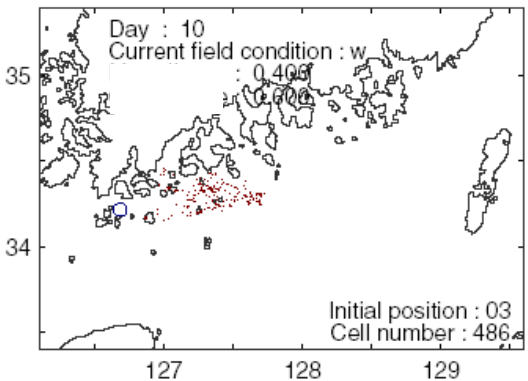
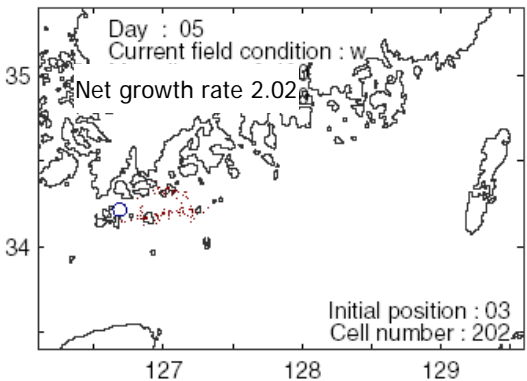
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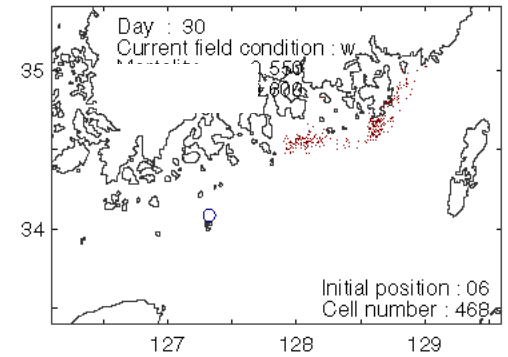
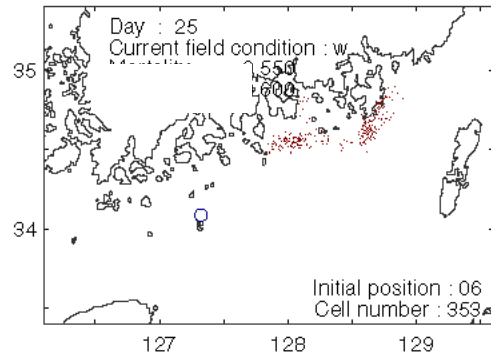
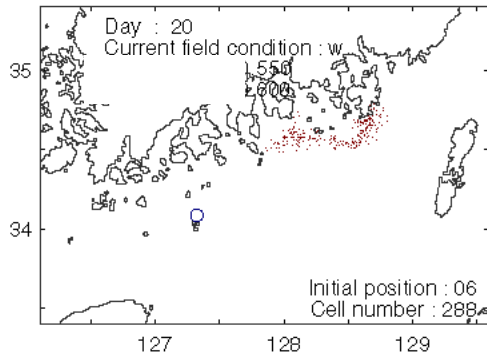
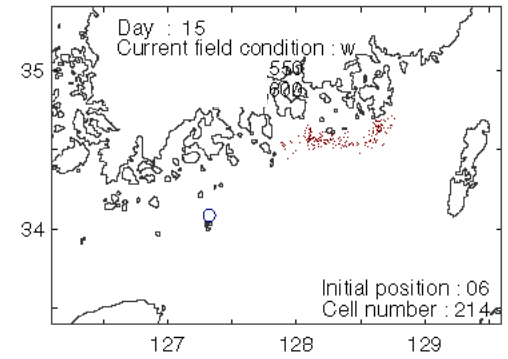
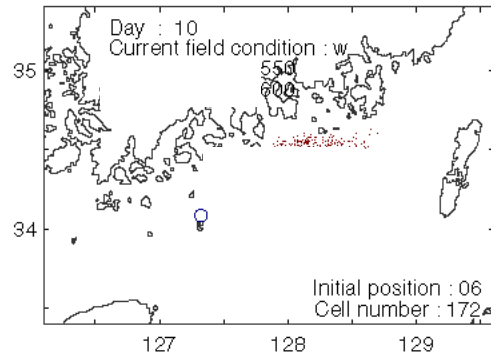
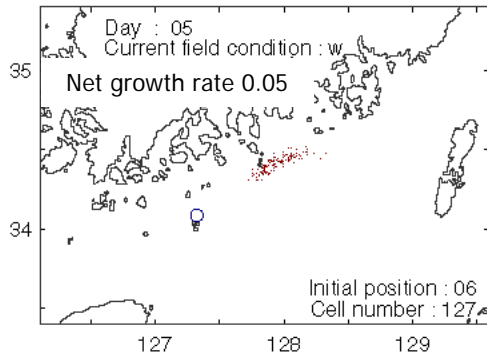


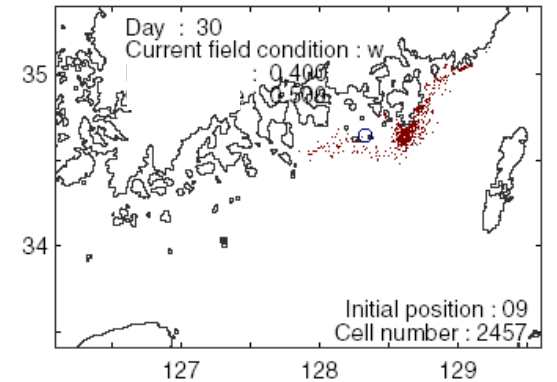
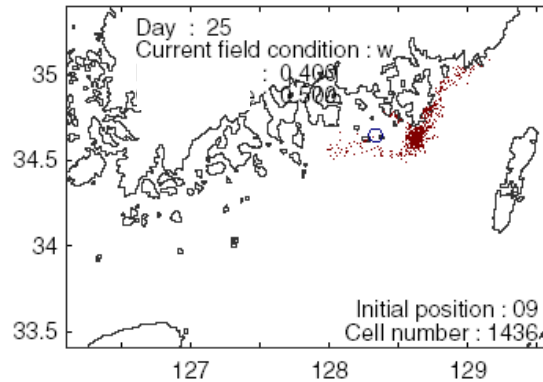
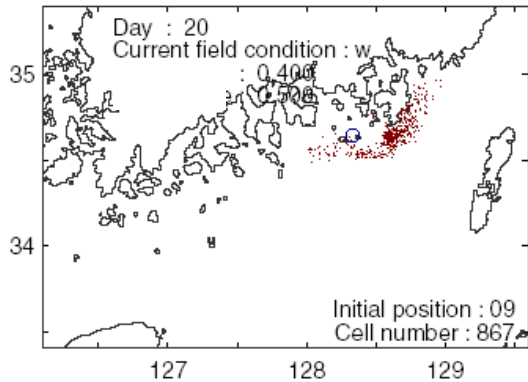
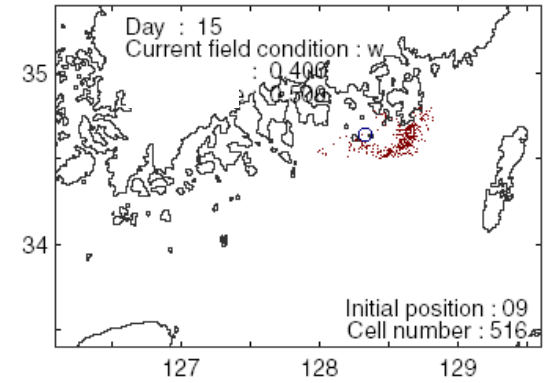
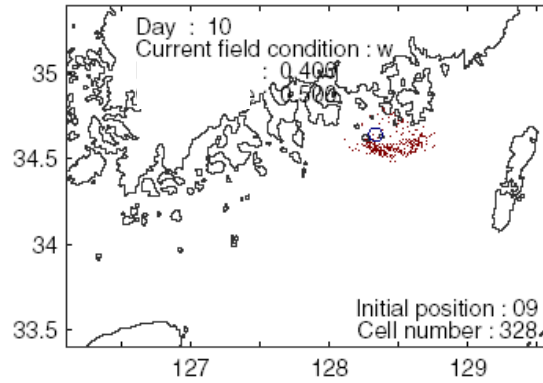
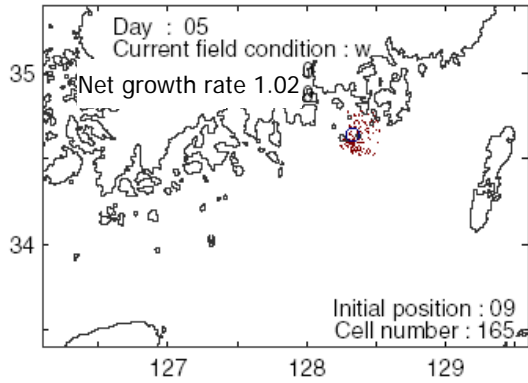
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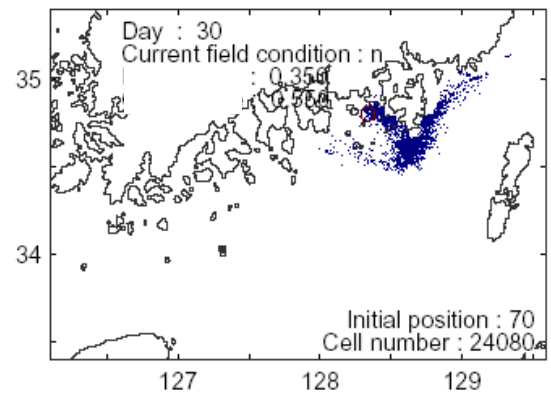
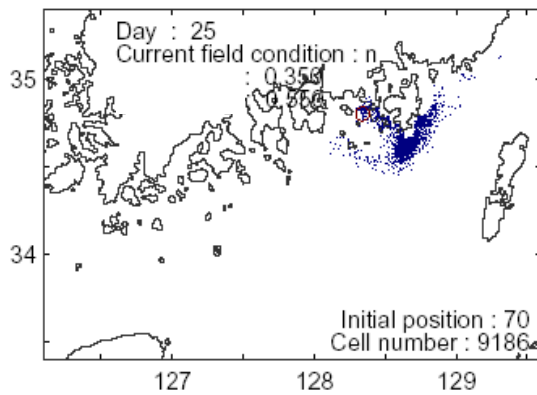
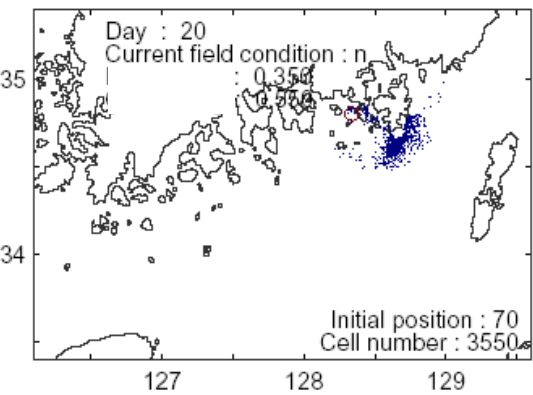
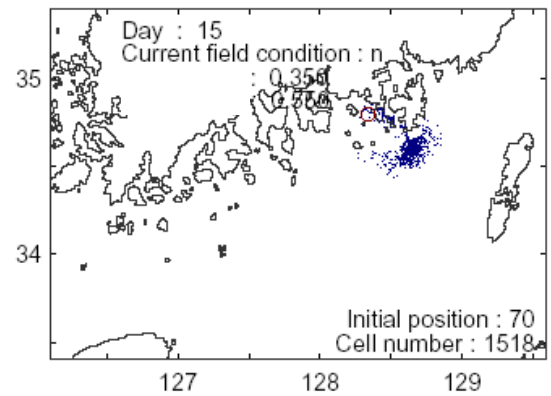
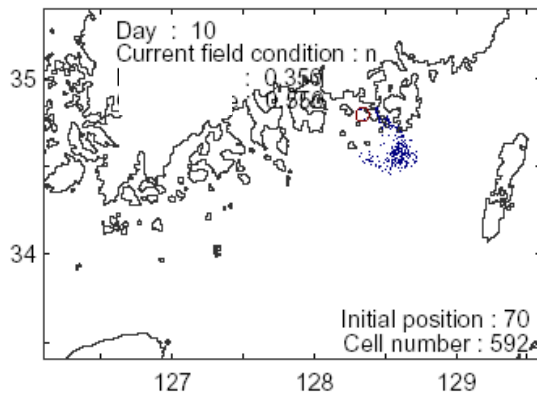
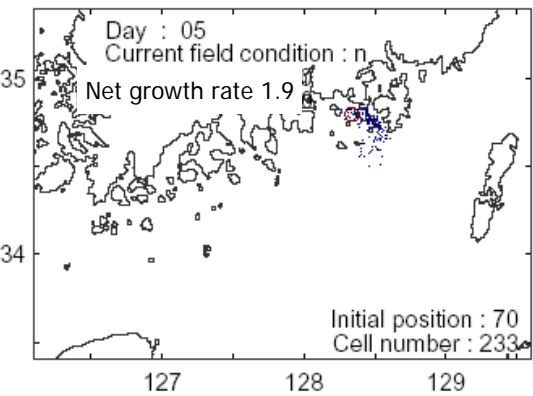




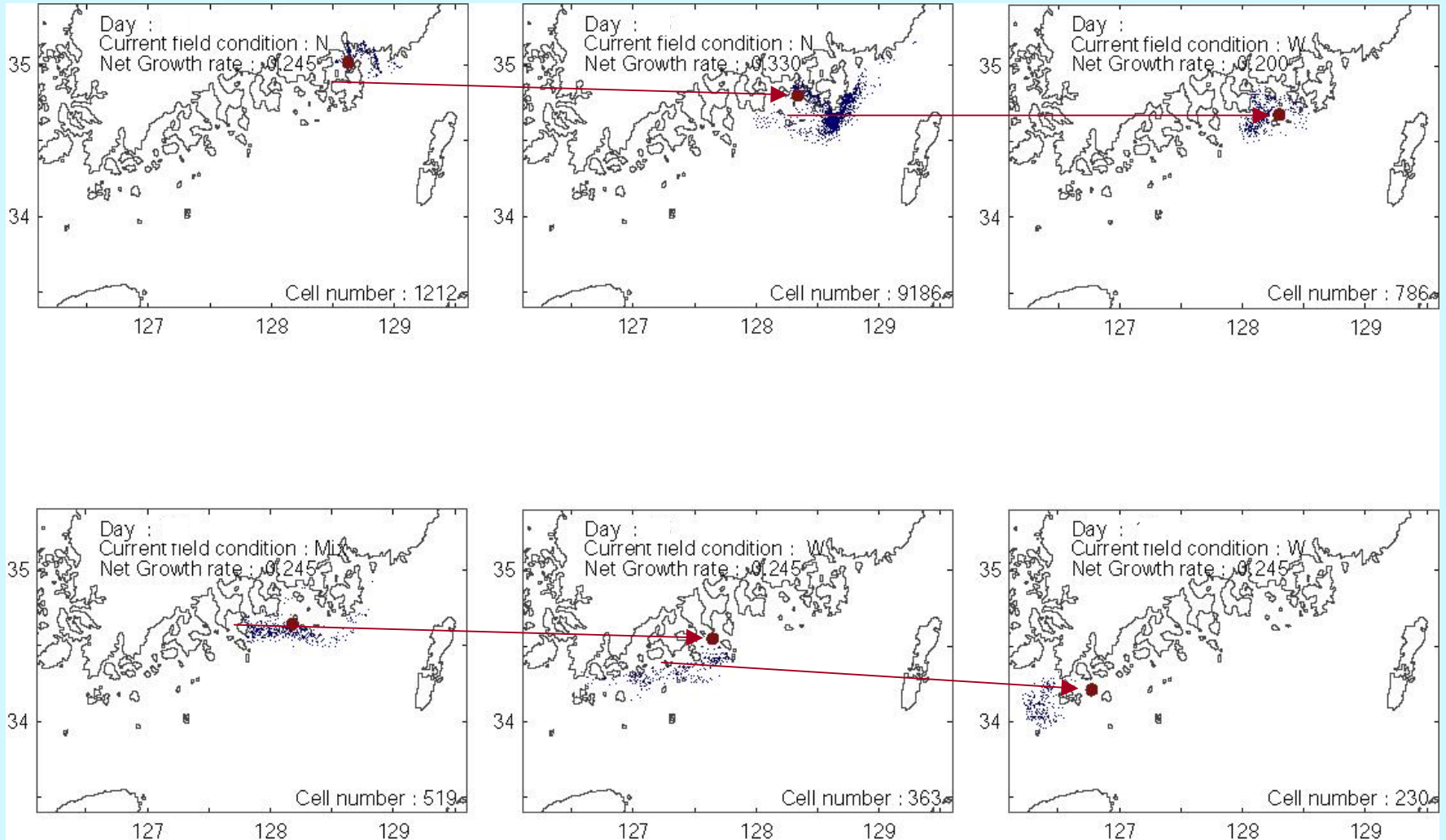








A possible scenario for Year-after-year westward procession



Conclusions (1)

- ❖ The spatial patterns of *C. polykrikoides* blooms can be explained by the characteristics of the circulation system in the southern coastal area.
 - Cysts and cells are not retained in the Tsushima Current region (fast eastward current: 25-85 cm s⁻¹)
 - In the outskirts region, northward and southward current converge when seasonal northeasterly wind blows. Cysts and cells are accumulated in the outskirts regions.
 - Cells cannot reach beneath the nutricline in the Tsushima Current region and could be nutrient-limited.
- ❖ Alternative hypotheses: oil spills, nutrients, etc

Conclusions (2)

- ❖ Net movement of cells in the surface is eastward due to eastward current.
- ❖ Strong north-easterly wind increases westward surface current and keep the bloom patches from moving eastward (as the case of 1998).
- ❖ Long-term westward expansion of the *C. polykrikoides* blooms can be explained by gradual spreading of cysts.
- ❖ Given accurate current field prediction, spatial propagation could be projected using IBM models.

Future directions

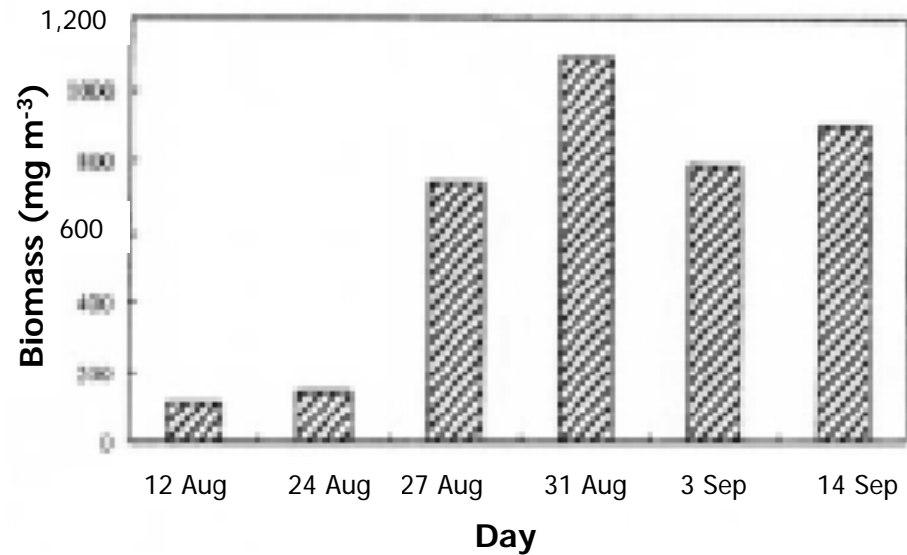
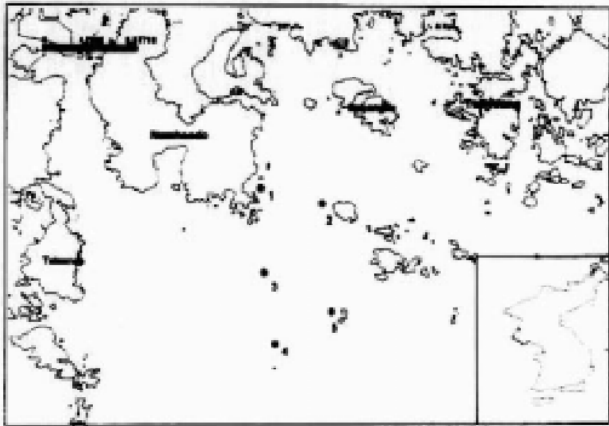
- ❖ Quantitative cyst maps could be used to test the model results.
- ❖ Grazers and grazing pressure on *C. polykrikoides*.
- ❖ Turbulence effects on the growth of *C. polykrikoides*.
- ❖ Circulation models with finer resolution could be used for forecasting the propagation of blooms.

Thank you for listening!

Turbulence and dinoflagellates

- ❖ Recently studies have challenged the conventional view that turbulence inhibit the growth of dinoflagellates.
- ❖ For example, using 10 autotrophic dinoflagellate species, Sullivan and Swift (2003) experimentally showed that the effects of turbulence on growth could be negative, neutral, or positive depending on species.
- ❖ *Gymnodinium catenatum*, a phylogenetically related species to *Cochlodinium*, exhibits a significant increase in growth rate in the high turbulence treatment

Zooplankton biomass in the bloom area in 1998



Calculation of Lagrangian movement

$$X_{n+1} = X_n + U \nabla t + \gamma \sqrt{2 \nabla t D}$$

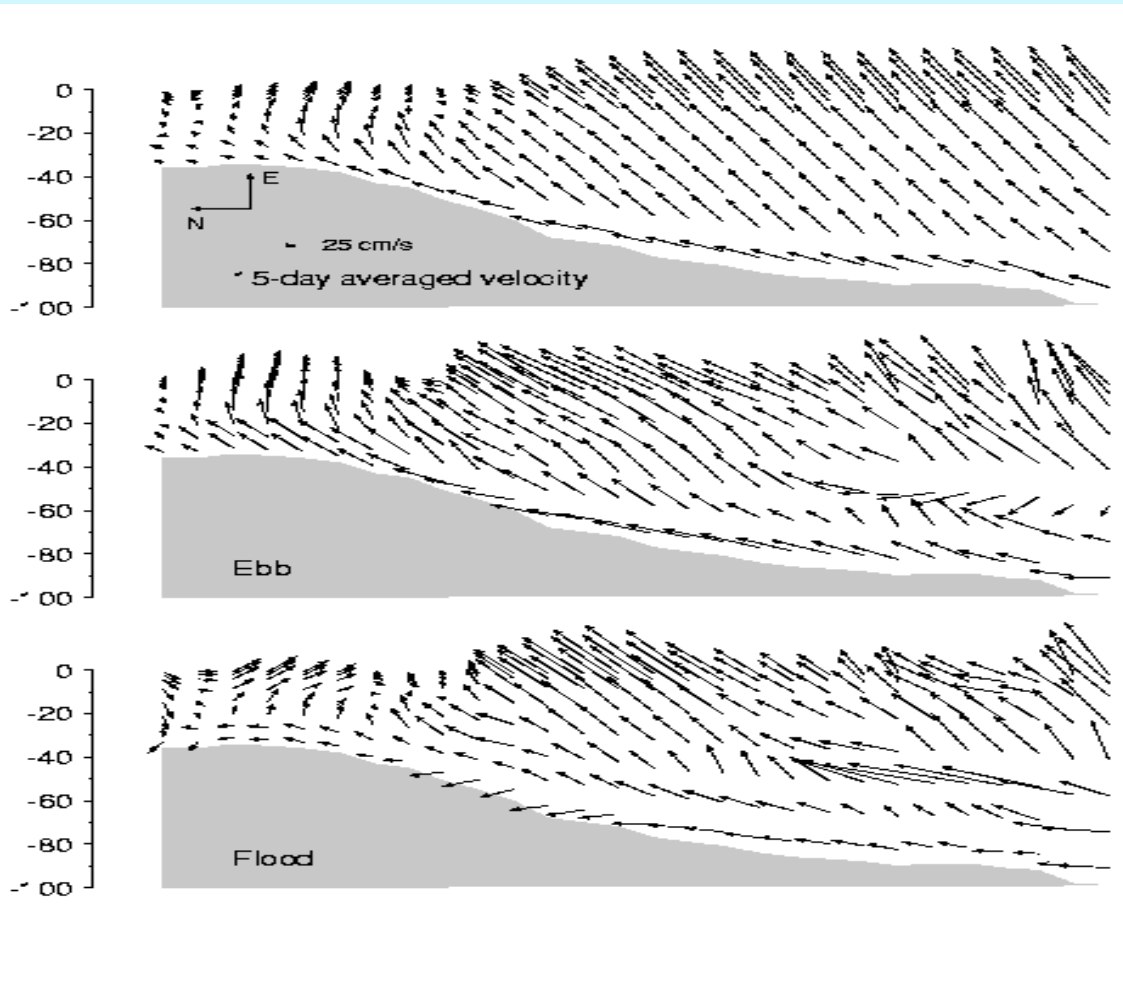
X_n : Position of a cell (x_n, y_n)

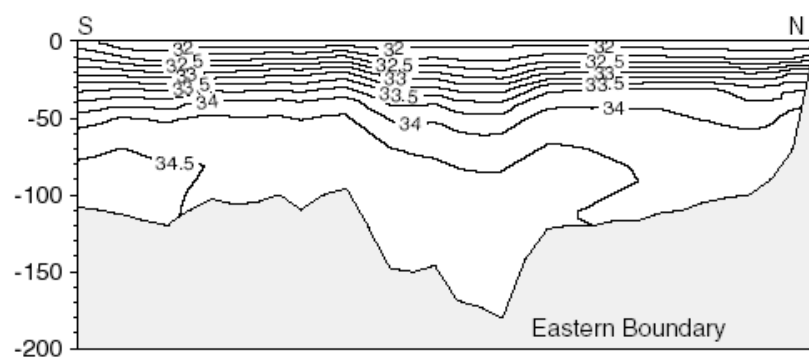
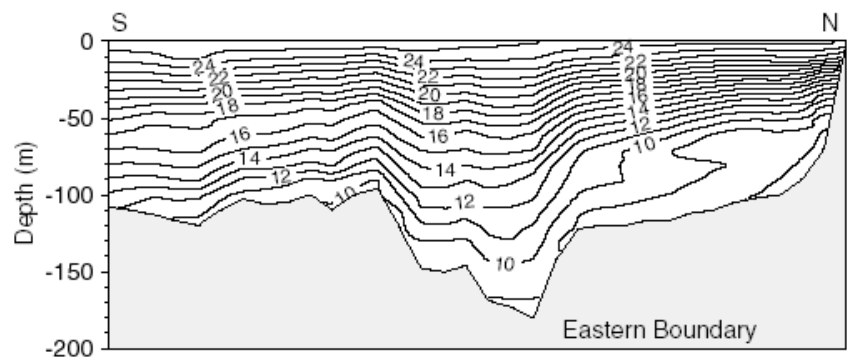
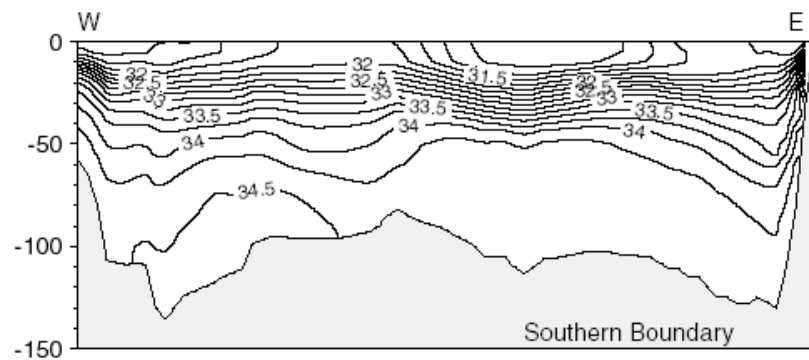
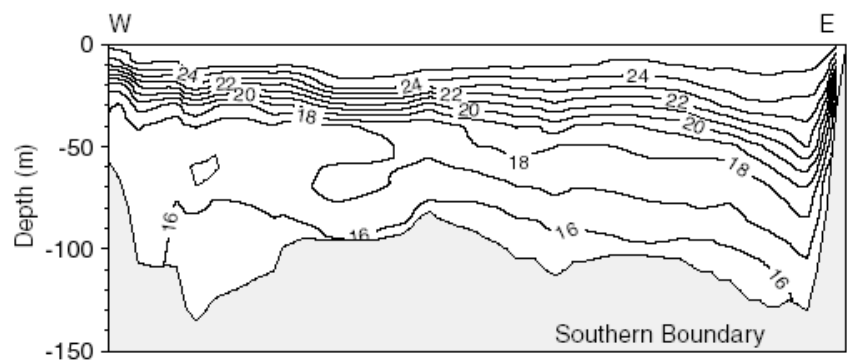
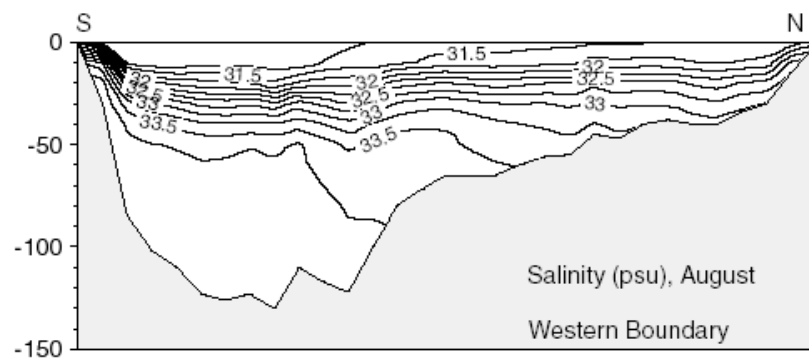
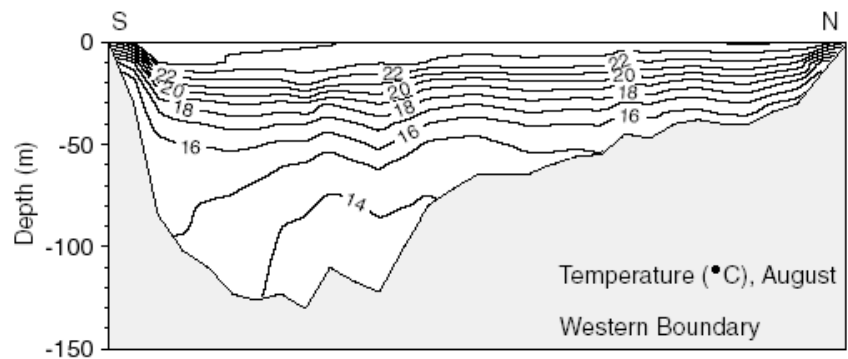
U_n : Current field (u_n, v_n)

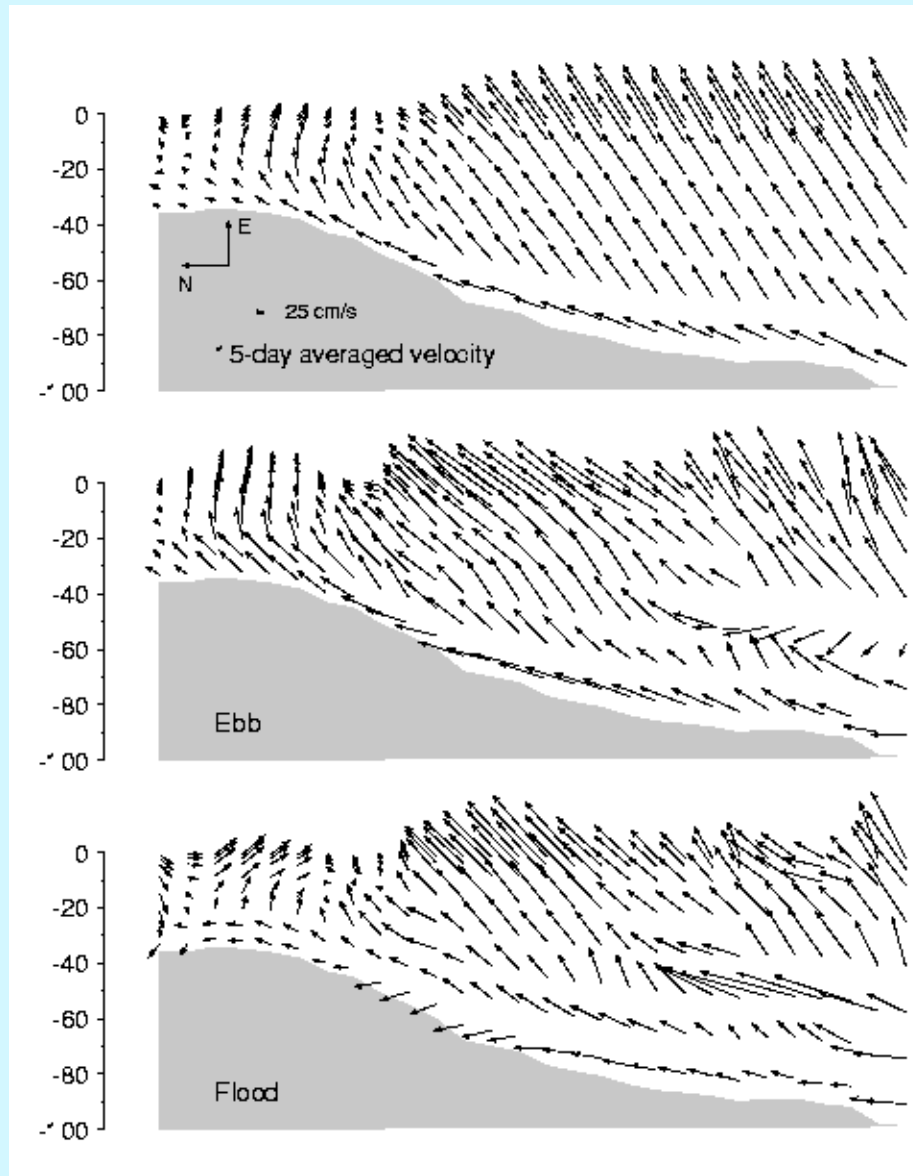
γ : random normal deviate

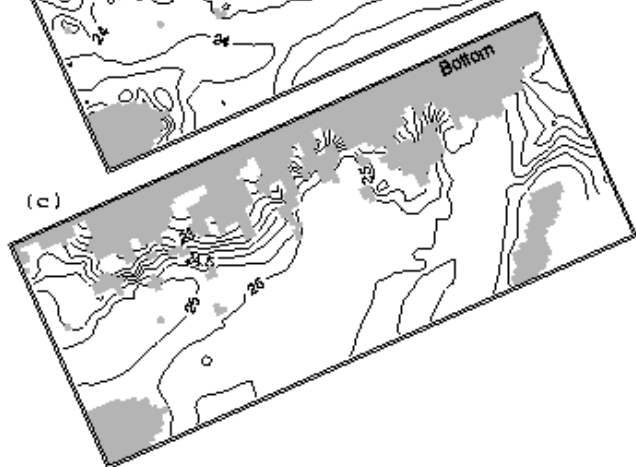
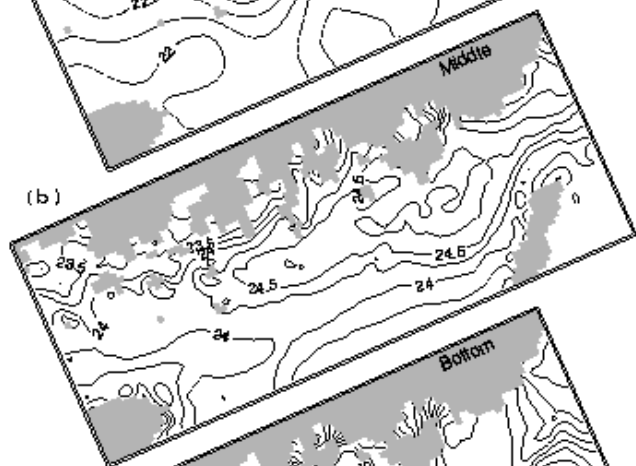
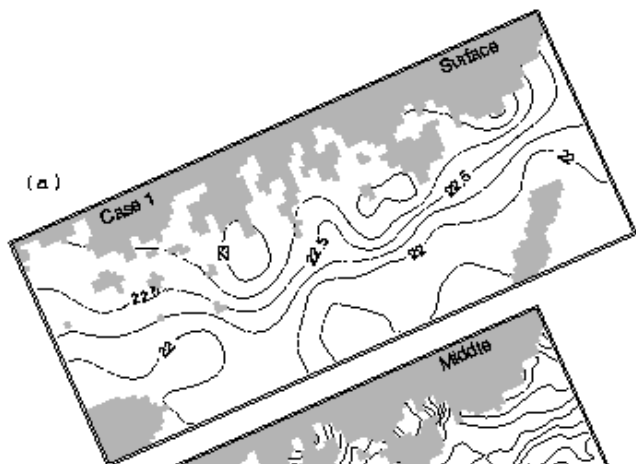
D : diffusivity

Vertical velocity

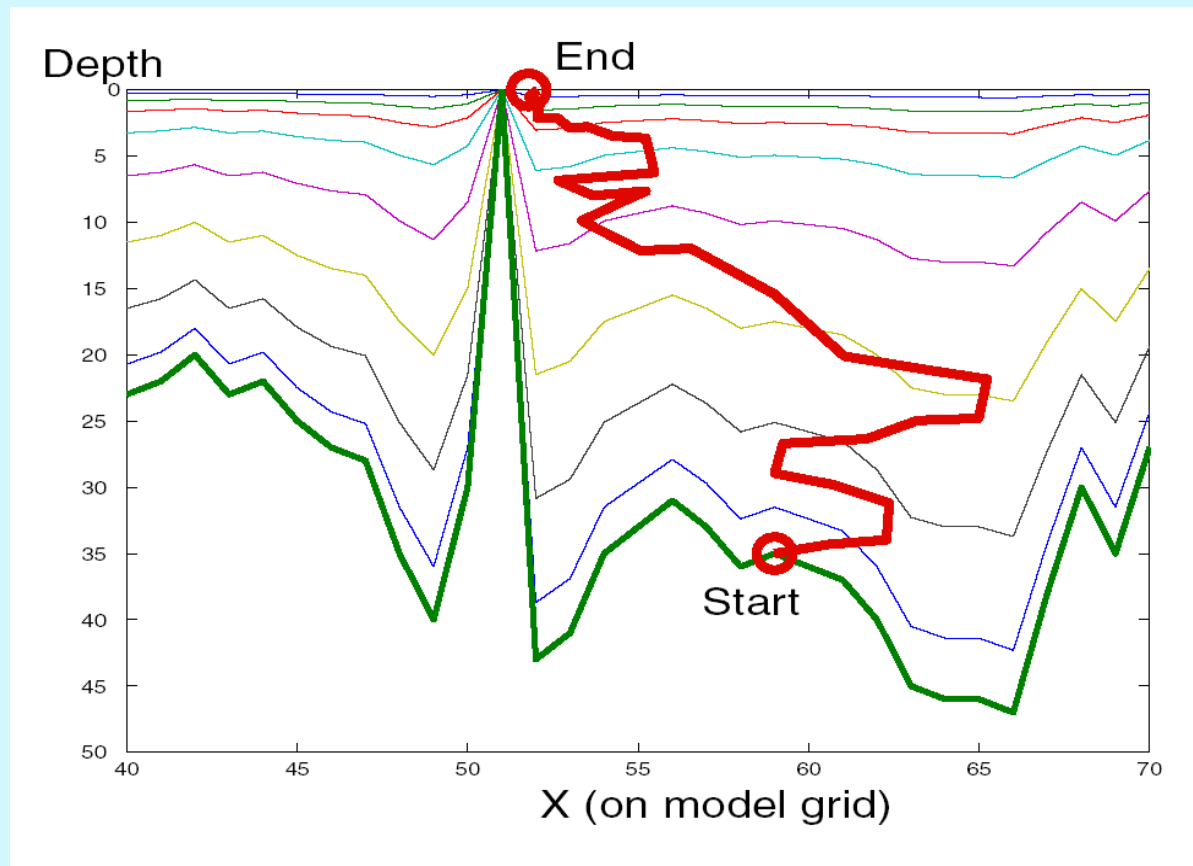


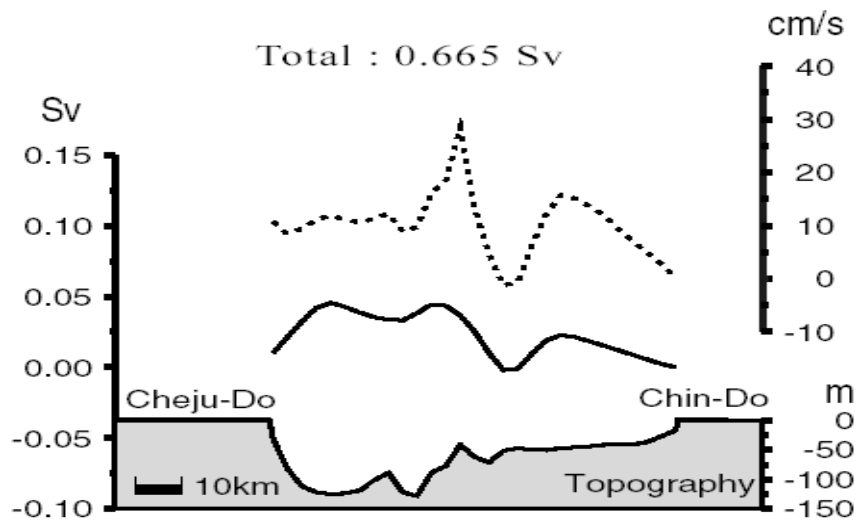




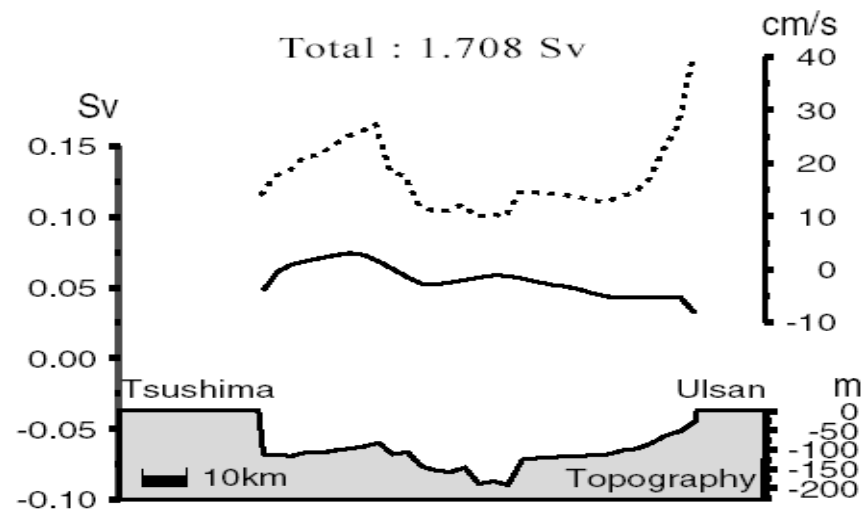


Tracking of germinated cells

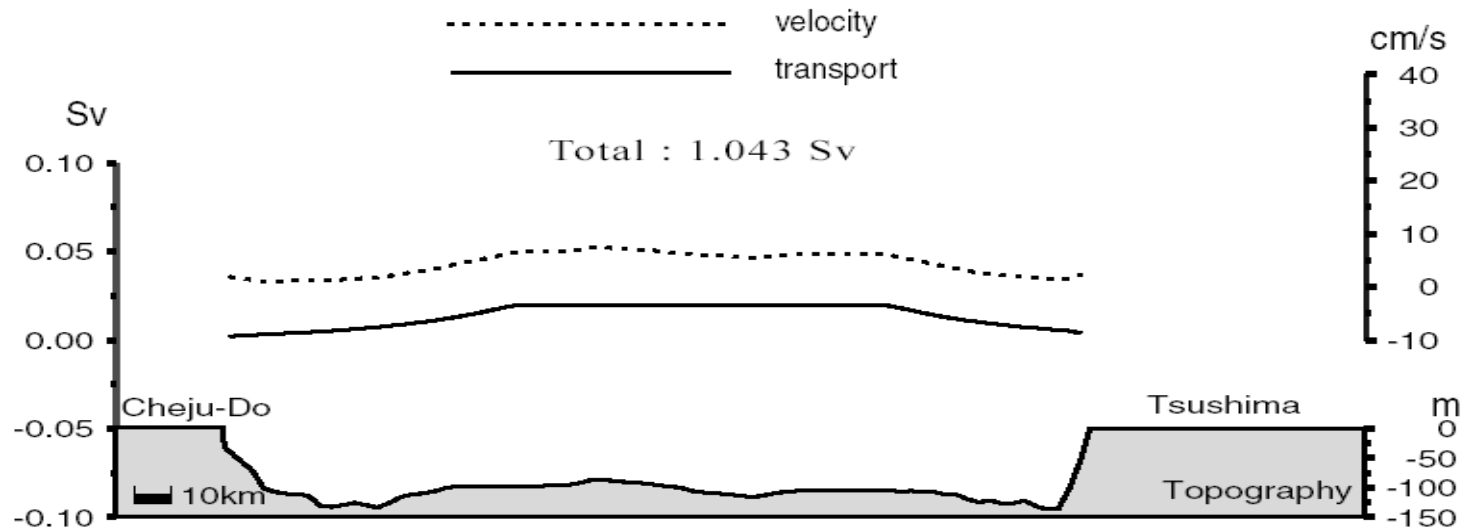




WESTERN BOUNDARY



EASTERN BOUNDARY



SOUTHERN BOUNDARY

