Active Tracers

We review mixing length theory applied to a set of active scalars (think in terms of biological properties):

$$\frac{D}{Dt}b_i + \nabla \cdot (\mathbf{u}_{bio}b_i) - \nabla \kappa \nabla b_i = \mathcal{B}_i(\mathbf{b}, \mathbf{x}, t)$$

Split the field into an eddy part which varies rapidly in space and time and a mean part which changes over larger (order $1/\epsilon$) horizontal distances and longer (order $1/\epsilon^2$) times:

$$b_i = \overline{b}_i(z|\mathbf{X},T) + b'_i(\mathbf{x},z,t|\mathbf{X},T)$$

We must allow for short vertical scales in both means and fluctuations. Counterbalancing this difficulty is the fact that vertical velocities tend to be weak (order $\frac{U}{fL} \times \frac{UH}{L}$). We assume the mean flows are small $\overline{u} \sim \varepsilon \mathbf{u}'$ and the coefficients in the reaction terms vary rapidly in the vertical but slowly horizontally and in time.

$$\begin{split} \frac{D}{Dt} - \nabla \cdot \kappa \nabla \longrightarrow \\ & \left[\frac{\partial}{\partial t} + u'_m \nabla_m - \nabla \cdot \kappa \nabla - \frac{\partial}{\partial z} \kappa_v \frac{\partial}{\partial z} \right] \\ + \varepsilon \left[u'_m \ \mathbf{\nabla}_m + \overline{u}_m \nabla_m - \mathbf{\nabla}_m \kappa_{mn} \nabla_n - \nabla_m \kappa_{mn} \mathbf{\nabla}_n + \frac{Ro}{\varepsilon} w' \frac{\partial}{\partial z} \right] \\ & + \varepsilon^2 \left[\frac{\partial}{\partial T} + \overline{u}_m \mathbf{\nabla}_m - \mathbf{\nabla}_m \kappa_{mn} \mathbf{\nabla}_n + \frac{Ro}{\varepsilon} \overline{w} \frac{\partial}{\partial z} \right] \end{split}$$

Vertical Structure

1) We assume the case with no flow has a *stable* solution:

$$\frac{\partial}{\partial z} w_{bio} \overline{b}_i = \frac{\partial}{\partial z} \kappa_v \frac{\partial}{\partial z} \overline{b}_i + \mathcal{B}_i(\overline{\mathbf{b}}, z | \mathbf{X}, T)$$

Demos, Page 1: bio dynamics <growth rates>

2) The eddy-induced perturbations satisfy

$$egin{aligned} &\left[rac{\partial}{\partial t}+\mathbf{u}'\cdot
abla-
abla\cdot\kappa
abla
ight]b_i'+rac{\partial}{\partial z}w_{bio}b'=\ &\sum_jrac{\partial\mathcal{B}_i}{\partial b_j}b_j'-\mathbf{u}'\cdot
ablaar{b}_i\equiv\mathcal{B}_{ij}b_j'-\mathbf{u}'\cdot
ablaar{b}_i \end{aligned}$$

with $\mathbf{\nabla} = (\partial/\partial X, \ \partial/\partial Y, \ \partial/\partial z).$

3) The equation for the mean is

$$\begin{split} \left[\frac{\partial}{\partial T} + \overline{\mathbf{u}} \cdot \mathbf{\nabla} - \mathbf{\nabla} \kappa \mathbf{\nabla} \right] \overline{b}_i + \mathbf{\nabla} \cdot (\overline{\mathbf{u}' b'}) + \frac{\partial}{\partial z} w_{bio} \overline{b}_i = \\ \mathcal{B}_i(\overline{\mathbf{b}}, z | \mathbf{X}, T) + \frac{1}{2} \frac{\partial^2 \mathcal{B}_i}{\partial b_j \partial b_k} \overline{b'_j b'_k} \end{split}$$

Summary:

Eddies generate fluctuations by horizontal and vertical advection of large-scale gradients, but the strength and structure depends on the biologically-induced perturbation decay rates.

Perturbations generate eddy fluxes and alter the average values of the nonlinear biological terms.

NPZ

A simple biological model (mixed layer):

$$\begin{split} \frac{D}{Dt}P &= \frac{\mu PN}{N+k_s} - \frac{g}{\nu}Z[1 - \exp(-\nu P)] - d_P P + \nabla \kappa \nabla P \\ \frac{D}{Dt}Z &= \frac{ag}{\nu}Z[1 - \exp(-\nu P)] - d_Z Z + \nabla \kappa \nabla Z \\ \frac{D}{Dt}N &= -\frac{\mu PN}{N+k_s} + \frac{(1-a)g}{\nu}Z[1 - \exp(-\nu P)] \\ &+ d_P P + d_Z Z + \nabla \kappa \nabla N \\ or \quad N &= N_T - P - Z \end{split}$$

Mean-field approach

We can get a very similar picture using the mean-field approximation: take

$$\begin{split} \frac{\partial}{\partial t}\overline{b}_i + \overline{u}\cdot\nabla\overline{b}_i + \nabla\cdot(\overline{\mathbf{u}'b'}) + \frac{\partial}{\partial z}w_{bio}\overline{b}_i - \nabla\kappa\nabla\overline{b}_i &= \overline{\mathcal{B}_i(\overline{\mathbf{b}} + \mathbf{b}', \mathbf{x}, t)} \\ &\simeq \mathcal{B}_i(\overline{\mathbf{b}}, z | \mathbf{x}, t) + \frac{1}{2}\frac{\partial^2 \mathcal{B}_i}{\partial b_j \partial b_k}\overline{b'_j b'_k} \\ &\frac{\partial}{\partial t}\mathbf{b}'_i + \overline{u}\cdot\nabla\mathbf{b}'_i + \nabla\cdot(\mathbf{u}'b'_i - \overline{\mathbf{u}'b'}) + \frac{\partial}{\partial z}w_{bio}\mathbf{b}'_i - \nabla\kappa\nabla\mathbf{b}'_i = \\ &-\mathbf{u}'\cdot\nabla\overline{b}_i + \mathcal{B}_i(\overline{\mathbf{b}} + \mathbf{b}', \mathbf{x}, t) - \overline{\mathcal{B}_i(\overline{\mathbf{b}} + \mathbf{b}', \mathbf{x}, t)} \end{split}$$

or (dropping the quadratic and higher terms)

$$rac{\partial}{\partial t} \mathbf{b}'_i + \overline{u} \cdot
abla \mathbf{b}'_i + rac{\partial}{\partial z} w_{bio} \mathbf{b}'_i -
abla \kappa
abla \mathbf{b}'_i \simeq - \mathbf{u}' \cdot
abla \overline{b}_i + \mathcal{B}_{ij} b'_j$$

The differences are subtle: the MFA does not presume that the scale of \overline{b}_i is large but linearizes in a way which may not be consistent.

Separable Problems

The mesoscale eddy field has horizontal velocities in the near-surface layer which are nearly independent of z, and the vertical velocity increases linearly with depth $w' = s(\mathbf{x}, t)z$. The stretching satisfies

$$s(\mathbf{x},t) = -\nabla \cdot \mathbf{u}(\mathbf{x},t)$$

For linear (or linearized perturbation) problems in the near-surface layers, we can separate the physics and the biology using Greens' functions.

We define the Greens function for the horizontal flow problem:

$$\left(\frac{\partial}{\partial t} + \mathbf{u}(\mathbf{x}, t) \cdot \nabla - \nabla \kappa \nabla\right) G(\mathbf{x}, \mathbf{x}', t - t') = \delta(\mathbf{x} - \mathbf{x}')\delta(t - t')$$

The perturbation equations can now be solved:

$$b'_{i} = -\int d\mathbf{x}' \int dt' G(\mathbf{x}, t | \mathbf{x}', t') u'_{m}(\mathbf{x}', t') \phi_{m,i}(z, t - t')$$
$$-\int d\mathbf{x}' \int dt' G(\mathbf{x}, t | \mathbf{x}', t') s'(\mathbf{x}', t') \varphi_{i}(z, t - t')$$

The two functions representing the biological dynamics both satisfy

$$rac{\partial}{\partial au} arphi_i = rac{\partial}{\partial z} \kappa_v rac{\partial}{\partial z} arphi_i + \mathcal{B}_{ij} arphi_j$$

with $\mathcal{B}_{ij} = \partial \mathcal{B}_i / \partial b_j$. These give the diffusive/ biological decay of standardized initial perturbations

$$\phi_{m,i}(z,0) = \mathbf{\nabla}_m \overline{b}_i \quad , \quad \varphi_i(z,0) = z \frac{\partial}{\partial z} \overline{b}_i$$

Demos, Page 3: perturbation structures ${\it <p'z'}$ struct> ${\it <ev}$ of p'> ${\it <ev}$ of z'>

Simple Example

If we ignore vertical diffusion and advection and consider only one component with $\mathcal{B}_{11} = -\lambda$, we have

$$\phi_{m,i} = e^{-\lambda \tau} \boldsymbol{\nabla}_m \overline{b}_i$$

so that

$$b'_{i} = -\left[\int d\mathbf{x}' \int dt' e^{-\lambda(t-t')} G(\mathbf{x},t|\mathbf{x}',t') u'_{n}(\mathbf{x}',t')\right] \mathbf{\nabla}_{n} \overline{b}_{i}$$

The eddy flux takes the form

$$\overline{u'_m b'} = -\left[\int d\mathbf{x}' \int dt' e^{-\lambda(t-t')} \overline{u'_m(\mathbf{x},t)G(\mathbf{x},t|\mathbf{x}',t')u'_n(\mathbf{x}',t')}\right] \mathbf{\nabla}_n \overline{b}_i$$
$$= -\left[\int d\mathbf{x}' \int dt' e^{-\lambda(t-t')} R_{mn}(\mathbf{x},t|\mathbf{x}',t')\right] \mathbf{\nabla}_n \overline{b}_i$$

If we split the right-hand side into symmetric and antisymmetric parts, we find

$$\begin{aligned} \overline{u'_m b'} &= -K_{mn}^{\lambda} \nabla n \overline{b}_i + \epsilon_{mnk} \Psi_k^{\lambda} \nabla n \overline{b}_i \\ &= -K_{mn}^{\lambda} \nabla \overline{b}_i - (\epsilon_{mnk} \nabla n \Psi_k^{\lambda}) \overline{b}_i + \epsilon_{mnk} \nabla n (\Psi_k^{\lambda} \overline{b}_i) \end{aligned}$$

The last term has no divergence and can be dropped. Thus the eddy flux is a mix of diffusion and Stokes' drift:

$$\overline{u'_m b'} = -K^{\lambda}_{mn} \nabla \overline{b}_i + V^{\lambda}_m \overline{b}_i$$

Both coefficients depend on the biological time scale λ^{-1} .

For the random Rossby wave case, the Stokes drift term is

$$\mathbf{V}^{\lambda} = \frac{KE}{(\gamma + \lambda)^2 + \frac{1}{4}} \left(-\cos(2y) \ , \ 0 \right)$$

while the diffusivity tensor is

$$K_{ij}^{\lambda} = 2(\gamma + \lambda) \frac{KE}{(\gamma + \lambda)^2 + \frac{1}{4}} \begin{pmatrix} \cos^2(y) & 0\\ 0 & \sin^2(y) \end{pmatrix}$$

Demos, Page 4: effective coeff <effective k,v>

Not so simple example

"Mixing length" models

$$Flux(b) = -\kappa_e \nabla b$$

even if appropriate for passive tracers are not suitable for biological properties whose time scales may be comparable to those in the physics. Instead, we find

$$\overline{\mathbf{u}_m'b_i'} = -\left[\int d\tau e^{\mathcal{B}_{ij}\tau}R_{mn}(\tau)\right]\nabla_n\overline{b}_j$$

where R_{mn} is the equivalent of Taylor's Lagrangian covariance (but including κ).

We divide the coefficient into symmetric (K) and antisymmetric terms related to the Stokes drift (V)

$$\overline{\mathbf{u}_m'b_i'} = -K_{mn}^{ij} oldsymbol{\nabla}_n \overline{b}_j + V_m^{ij} \overline{b}_j$$

Note that

- Eddy diffusivities and wave drifts mix different components (flux of P depends on gradient of Z).
- If R has a negative lobe, the biological diffusivities can be larger than that of a passive scalar
- The quasi-equilibrium approximation

$$\mathcal{B}_{ij}b'_j = \mathbf{u}' \cdot \boldsymbol{\nabla} \overline{b}_i$$

works reasonably well in the upper water column. In particular

$$\mathcal{B}_{21} = g\overline{Z}\exp(-\nu\overline{P}) > 0$$
 . $\mathcal{B}_{22} = 0$

so that

$$P' = \frac{1}{\mathcal{B}_{21}} \mathbf{u}' \cdot \nabla \overline{Z}$$
 unlike $C' = -\boldsymbol{\xi} \cdot \nabla \overline{C}$

Demos, Page 5: complex diffusion <transport coeff: display -geometry +0+0 -bordercolor white -border 20x20 -rotate 90 ~glenn/12.822t/graphics/t0.ps> up Z grad flux of Pt1.ps <quasiequilibrium fluxes: display -geometry +0+0 bordercolor white -border 20x20 -rotate 90 ~glenn/12.822t/graphics/t1a.ps> downgradient Kpp,KZZt2.ps

Eulerian-Lagrangian

If $\kappa = 0$, we can relate the relevant form of the Eulerian covariance

$$R_{mn}(\mathbf{x},t|\mathbf{x}',t') = \overline{u'_m(\mathbf{x},t)G(\mathbf{x},t|\mathbf{x}',t')u_n(\mathbf{x}',t')}$$

to Taylor's form. The Greens' function equation

$$\frac{\partial}{\partial t}G + \mathbf{u}(\mathbf{x}, t) \cdot \nabla G = \delta(\mathbf{x} - \mathbf{x}')\delta(t - t')$$

has a solution

$$G(\mathbf{x}, t | \mathbf{x}', t') = \delta \left(\mathbf{x} - \mathbf{X}(t | \mathbf{x}', t') \right)$$

where

$$\frac{\partial}{\partial t} \mathbf{X}(t|\mathbf{x}',t') = \mathbf{u}(\mathbf{X},t) \quad , \qquad \mathbf{X}(t'|\mathbf{x}',t') = \mathbf{x}'$$

gives the Lagrangian position of the particle initially at \mathbf{x}' at time t'. But it is more convenient to back up along the trajectory and let

$$G(\mathbf{x}, t | \mathbf{x}', t') = \delta(\mathbf{x}' - \boldsymbol{\xi}(t - t' | \mathbf{x}, t))$$

where the particle at $\boldsymbol{\xi}$ at time t' passes \mathbf{x} at time t (and takes a time τ for this tranistion). Thus the $\boldsymbol{\xi}$'s give the starting position, which, for stochastic flows varies from realization to realization. We can solve

$$rac{\partial}{\partial au} \boldsymbol{\xi}(\tau | \mathbf{x}, t) = -\mathbf{u}(\boldsymbol{\xi}(\tau | \mathbf{x}, t), t - \tau) \quad , \qquad \boldsymbol{\xi}(0 | \mathbf{x}, t) = \mathbf{x}$$

for $\tau = 0$ to $\tau = t - t'$ to find $\boldsymbol{\xi}$.

We can now define the generalization of the Lagrangian correlation function used by Taylor

$$R_{mn}(t - t', \mathbf{x}) = \int d\mathbf{x}' \overline{u'(\mathbf{x}, t)G(\mathbf{x}, t | \mathbf{x}', t')u'(\mathbf{x}', t')}$$
$$= \overline{u'_m(\mathbf{x}, t)u'_n(\boldsymbol{\xi}(t - t' | \mathbf{x}, t), t - (t - t'))}$$
or
$$R_{mn}(\tau, \mathbf{x}) = \overline{u'_m(\mathbf{x}, t)u'_n(\boldsymbol{\xi}(\tau | \mathbf{x}, t), t - \tau)}$$

For homogeneous, stationary turbulence (on the scales intermediate between the eddies and the mean), this will be equivalent to Taylor's

$$R_{mn}(\tau) = \overline{u'_m(\mathbf{X}(t'+\tau|\mathbf{x}',t'),t'+\tau)u'_n(\mathbf{x}',t')}$$

but we include inhomogeneity and (for, general G, diffusion).