

# IMA Tutorial: Transport & Mixing

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## Lecture 1: Stirring & Mixing

Stirring: mechanical action (cause)  
Mixing: homogenization of a scalar (effect)

$\theta(\underline{x}, t)$  = concentration,  $\underline{u}(\underline{x}, t)$  given

Advection-Diffusion eq.

$$\frac{\partial \theta}{\partial t} + \underline{u} \cdot \nabla \theta = \kappa \nabla^2 \theta, \quad \nabla \cdot \underline{u} = 0 \quad \text{in } \Omega$$

(AD)

$$\text{Boundary conditions: } \left. \begin{array}{l} \hat{n} \cdot \nabla \theta = 0 \\ \hat{n} \cdot \underline{u} = 0 \end{array} \right\} \text{ on boundary } \partial \Omega$$

$$\text{let } \langle \cdot \rangle = \int_{\Omega} \cdot dV$$

Multiply (AD) by  $m \theta^{m-1}$ , integrate:

$$\langle m \theta^{m-1} \partial_t \theta \rangle = \partial_t \langle \theta^m \rangle$$

$$\langle m \theta^{m-1} \underline{u} \cdot \nabla \theta \rangle = \langle \underline{u} \cdot \nabla \theta^m \rangle = \langle \nabla \cdot (\underline{u} \theta^m) \rangle$$

$$= \int_{\partial \Omega} \theta^m \underbrace{\underline{u} \cdot \hat{n}}_{0!} dS = 0$$

$$\begin{aligned} \langle m \theta^{m-1} \kappa \nabla^2 \theta \rangle &= \kappa m \langle \nabla \cdot (\theta^{m-1} \nabla \theta) - \nabla \theta^{m-1} \cdot \nabla \theta \rangle \\ &= \kappa m \int_{\partial \Omega} \theta^{m-1} \nabla \theta \cdot \hat{n} dS - \kappa m(m-1) \langle \theta^{m-2} |\nabla \theta|^2 \rangle \end{aligned}$$

$$\partial_t \langle \theta^m \rangle = -\kappa m(m-1) \langle \theta^{m-2} |\nabla\theta|^2 \rangle$$

$m=0$  is trivial

$m=1$ :  $\partial_t \langle \theta \rangle = 0$  Total amount of  $\theta$  is conserved

$m=2$ :  $\partial_t \langle \theta^2 \rangle = -2\kappa \langle |\nabla\theta|^2 \rangle$   $\langle \theta^2 \rangle$  non-increasing!

Let variance  $\text{Var} = C_2 = \langle \theta^2 \rangle - \langle \theta \rangle^2$

$$\partial_t C_2 = -2\kappa \langle |\nabla\theta|^2 \rangle$$

↑  
constant

Scenario:



- Variance can only decrease.
  - Slows down as  $\langle |\nabla\theta|^2 \rangle \rightarrow 0$
  - But  $\langle |\nabla\theta|^2 \rangle = 0$  iff  $\theta = \text{constant}$ .
- ↑  
in some sense

Hence the system is "driven" towards a homogeneous state where

Assume  $\langle \theta \rangle = 0$  WLOG

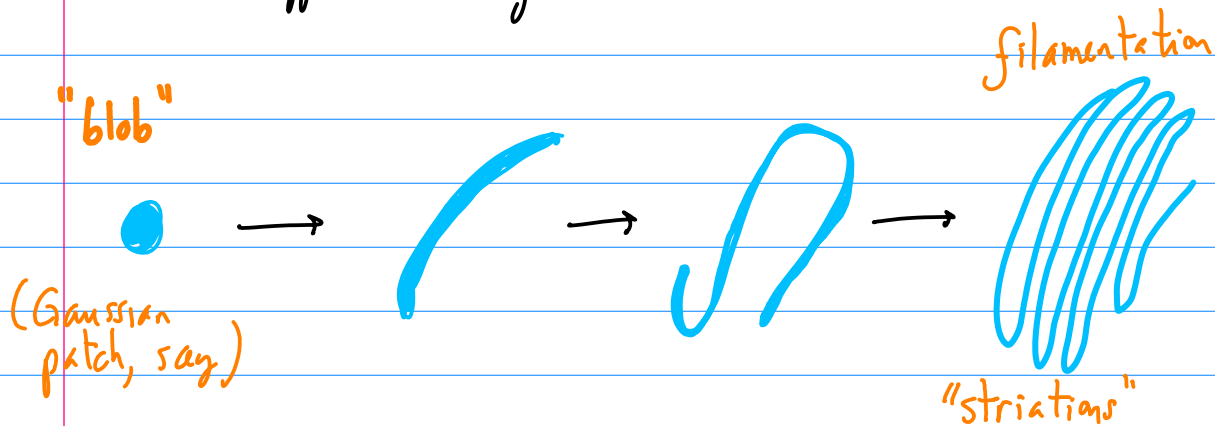
$$\theta(x, t) = \langle \theta \rangle = \text{constant.} \quad (C_2 = 0, \langle \theta^2 \rangle = \langle \theta \rangle^2)$$

No fluctuations from the mean! When  $C_2$  is small "enough", we say the system is mixed.

Big Q: Where is  $u(x, t)$  !? (stirring)

It doesn't appear in the variance equation!

But of course the variance equation is not closed: it depends on  $\nabla\theta$ .  
 What happens when you stir?



This hints at the answer: stirring increases  $\nabla\theta$

$$\partial_t \langle \theta^2 \rangle = -2\kappa \langle |\nabla\theta|^2 \rangle$$

*this becomes larger as we stir*

By how much are gradients increased? After all, if  $|\nabla\theta|$  becomes too large, then  $\langle \theta^2 \rangle \rightarrow 0$ , so there are no gradients anymore!

Answer: for "good" stirring, the system is driven to a state where

$$\kappa \langle |\nabla\theta|^2 \rangle \rightarrow \text{independent of } \kappa$$

Hence,  $\nabla\theta \sim \kappa^{-1/2}$

This is the chaotic/turbulent mixing scenario:

$\frac{\partial \langle \theta^2 \rangle}{\partial t}$  becomes independent of  $\kappa$  after a "short" transient

*(How short? Typically  $\sim \log \kappa$ )*

*This is the Platonic ideal of mixing*

Furthermore, the smallest scales visible in the concentration field  $\theta(x, t)$  have size  $\sim \sqrt{\kappa}$ . (missing a dimensional factor  $\rightarrow$  see later)

Note that  $\partial_t \langle \theta^2 \rangle$  independent of  $\kappa$  is crucial: in most applications,  $\kappa$  is tiny!

Heat:  $\kappa = 2.2160 \times 10^{-5} \text{ m}^2/\text{s}$  at 300K

10 m room: diffusion time  $\sim \frac{L^2}{\kappa} = \frac{(10\text{m})^2}{(2 \times 10^{-5} \text{ m}^2/\text{s})} \sim 4.5 \times 10^6 \text{ sec}$

So we better stir!  
Even thermal convection  
is often enough.

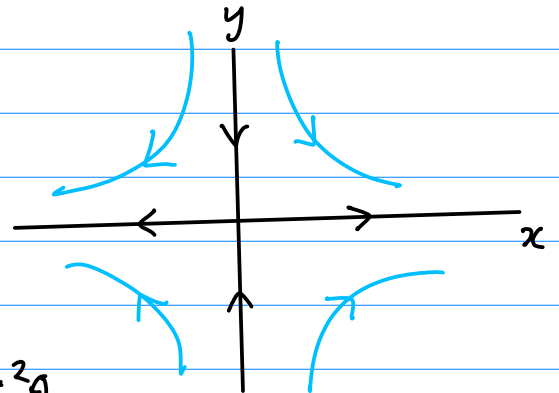
$\sim 1300 \text{ hours}$

$\sim 53 \text{ days!}$

Example of a good mixer:

$$\underline{u}(x, t) = (\lambda x, -\lambda y)$$

"hyperbolic point"



AD:  $\partial_t \theta + \lambda x \partial_x \theta - \lambda y \partial_y \theta = \kappa \nabla^2 \theta$

Can solve this exactly (we'll say more next time), but let's do the simplest thing: look for an  $x$ -independent solution of the form:

$$\theta(x, t) = e^{-\lambda t} f(y)$$

$$-\lambda f - \lambda y f' = \kappa f''$$

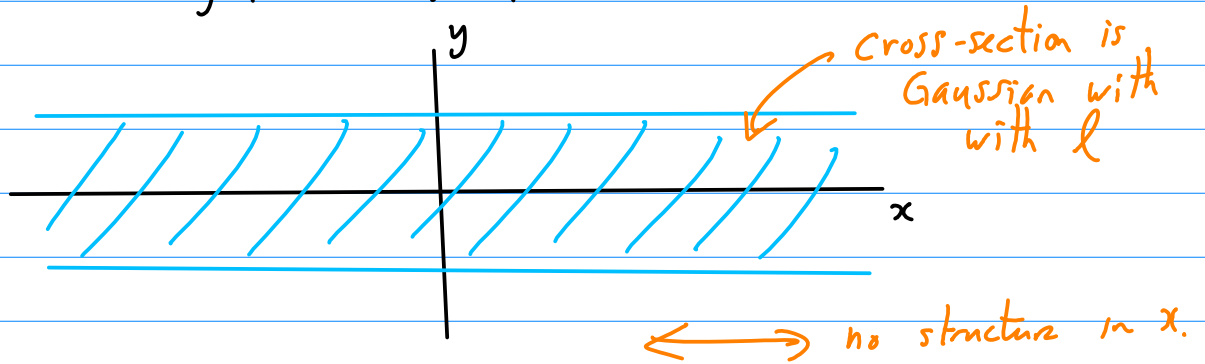
Boundary condition:

$$f \rightarrow 0 \text{ as } y \rightarrow \pm \infty.$$

Solution is:  $f(y) = e^{-y^2/2l^2}$ , where  $l^2 = \frac{\nu}{\lambda}$

Hence,  $\theta(x, t) \sim e^{-\lambda t} e^{-y^2/2l^2}$

This is the "filament" solution:



In fact, this solution tells us about the ultimate state of any compactly-supported initial condition:

"blob"



"filament"



"intensity fading" as  $e^{-\lambda t}$

central part

$\sim$  Gaussian cross-section

For this case, we know the length scale of "striations";

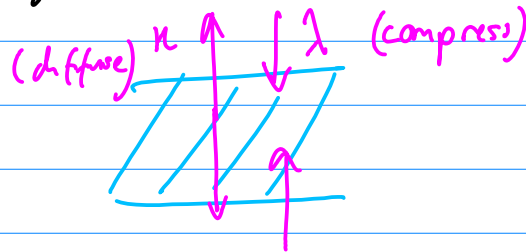
$$l = \sqrt{\frac{\nu}{\lambda}}$$

Batchelor length

Note  $l \sim \sqrt{\nu}$ , as necessary to make decay rate indep. of  $\nu$ !

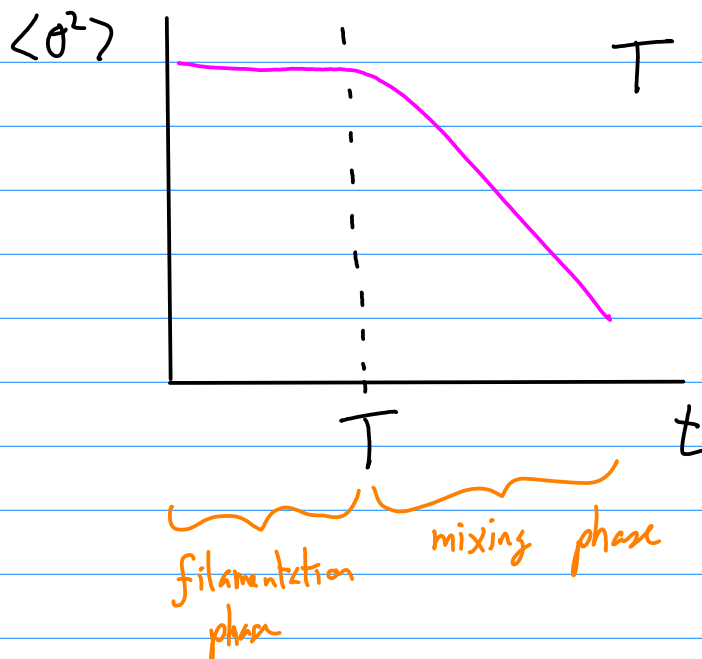
In practical applications,  $\lambda$  is often taken to be the local rate of strain.

$l$  is set by a balance between compression and diffusion



Summary: how mixing proceeds

- A blob is stirred  $\bullet \rightarrow$
- For a while,  $\langle \theta^2 \rangle$  is  $\sim$  constant, since  $\kappa$  is small
- When  $\nabla \theta$  reaches scales of order  $l$ , diffusion takes over
- After that,  $\langle \theta^2 \rangle$  decays at a  $\kappa$ -independent rate



$T$  given by:  $e^{-1} T \sim \sqrt{\kappa}$

$T \sim \lambda^{-1} \log \kappa$