Rayleigh-Taylor instability at a tilted interface

in incompressible laboratory experiments and compressible numerical simulations

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Outline

• Introduction

RT instability at a tilted interface Mixing, available energy and mixing efficiency

Laboratory experiments

At DAMTP, in the Fluid Dynamics Laboratory Incompressible water, NaCl to create density contrast

Numerical simulations

At AWE, using Turmoil3D (with David Youngs) Compressible code, for a mixture of two ideal gases

Conclusions and further work

Introduction

• RT instability

Instability of dense fluid accelerated into less dense fluid

$$\rho_{1} > \rho_{2}$$

$$p_{1}$$

$$p_{2}$$

$$g$$

$$p_{1}$$

$$p_{2}$$

$$P_{1}$$

$$p_{2}$$

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$$P_{2}$$

Much more efficient mixing than other mechanisms (shear instability, mechanical stirring)

An important mixing process within larger-scale flows (3D instability of 2D shear billows)

In environment, R-T instability has non-ideal initial conditions

At a tilted interface, there is competition between local instability and large-scale overturning



DAMTP, University of Cambridge

Rayleigh-Taylor instability

• Definitions of mixing

Distinguish between reversible and irreversible mixing:

Reversible mixing - interleaving of fluid with different properties - "reversible mixing = stirring"

Irreversible mixing - homogenisation of fluid properties at the molecular scale - "irreversible mixing = stirring + diffusion"

Irreversible mixing is important for

- chemical reactions
- removal of available energy when mixing density gradients across a gravitational field

• How do we measure mixing?

Mixing can be measured by a molecular mixing fraction For two fluids, volume fractions f and (1-f): $\vartheta(\mathbf{x},t) = f(\mathbf{x},t)(1-f(\mathbf{x},t))$

Alternatively, for fluids of varying density in a gravitational field, can measure the mixing efficiency η

For a fluid at rest, stirred by an energy input and returning to rest,

 $\eta {=} \frac{increase \ in \ potential \ energy}{amount \ of \ energy \ added}$

fraction of energy lost to fluid motion doing work against gravity

• Mixing in R-T instability

Measurements of η in laboratory experiments high values with some dependence on A Linden & Redondo (1991)

Numerical simulations show sensitivity to initial conditions Linden, Redondo & Youngs (1991), Cook & Dimotakis (2001)

• Diffusion and viscosity in incompressible fluids

Mechanical energy density per unit volume $E_v = \frac{1}{2}\rho|u|^2 + \rho gz$ $\frac{\partial}{\partial t}E_v(x,t) + \nabla f_v(x,t) = -\varepsilon_v(x,t),$ f_v energy flux ε_v energy dissipation

Water/salt system $\nu = 1.0 \times 10^{-2} \text{ cm}^2 \text{s}^{-1}$ kinematic viscosity $\kappa = 1.4 \times 10^{-5} \text{ cm}^2 \text{s}^{-1}$ diffusivityconcentration fluctuations persist at smaller scales than
velocity fluctuations - $Pr = \frac{\nu}{\kappa} = 700$

<u>*Turbulent flows*</u> - eddy viscosity = eddy diffusivity effectively Pr = 1

• Available energy in incompressible flow Mechanical energy in whole fluid $E = \int_{V} E_{v} dV$ decomposes:



Lorenz (1955), Thorpe (1977), Winters et al. (1995)

In unforced, decaying flow

 $\frac{d}{dt} \left(E_{back} + E_{avail} \right) = -\varepsilon$

$$\frac{d}{dt}(E_{back}) = q$$

loss of E due to turbulent dissipation gain in E_{back} due to

molecular mixing

Define cumulative mixing efficiency

$$\eta_{cumulative} = \frac{\int_{t_0}^t q \, dt}{\int_{t_0}^t q + \varepsilon \, dt} = \frac{\Delta P E_{back}}{-\Delta E_{avail}}$$

and instantaneous mixing efficiency

$$\eta_{instantaneous} = \frac{q}{q+\varepsilon} = \frac{\delta P E_{back}}{-\delta E_{avail}}$$

• Available energy in compressible flow

Now concerned with total energy (mechanical + internal) so $E_v = \frac{1}{2}\rho|u|^2 + \rho gz + \rho e$, *e* internal energy. In whole fluid:



Lorenz (1955), Andrews (1981), Shepherd (1993)

In unforced, decaying flow

$$\frac{d}{dt} (E_{back} + E_{avail}) = 0$$
$$\frac{d}{dt} (IE_{back}) = \varepsilon$$

 $\frac{d}{dt} \left(PE_{back} \right) = q$

E is conserved

gain in IE_{back} due to

turbulent dissipation and molecular mixing

gain in PE_{back} due to molecular mixing

Same definitions of mixing efficiency apply, so η is still

fraction of energy lost to fluid motion (reduction in E_{avail}) doing work against gravity (gain in E_{back})

Laboratory Experiments

Configuration



• Initial conditions

A solid barrier introduces significant shear



Dalziel, Linden & Youngs (1999)



Removal of finite thickness barrier causes initial velocity field



• Diagnostic measurements

Image analysis: spatial resolution 1 pixel \cong 0.1cm temporal resolution 25Hz

Assume statistical homogeneity across tank

Add propanol to fresh water to match refractive index

Density measurement

Dense fluid dyed with fluorescent dye

Images corrected for divergence of light sheet and attenuation

Velocity measurement

Fluid seeded with 400µm neutrally-buoyant particles

Lagrangian tracks for particles from tracking a frame sequence Interpolating onto a grid gives Eulerian velocities

Gridded at two scales: 1 cm - resolved velocity 3 cm - mean velocity (overcomes lack of similarity between experiments) Assume isotropy at small scales \Rightarrow estimate of total KE

• Parameters

Atwood number	$0.5 \times 10^{-3} < A < 2.5 \times 10^{-3}$	\Rightarrow Boussinesq
Timescale	$10s > \tau > 4.5s$	
RMS velocity	$0.8 \text{cms}^{-1} < u < 2 \text{cms}^{-1}$	
Integral lengthscale	1.8cm < <i>l</i> < 2.5cm	
Reynolds number	150 < Re < 500	





light source

corrected for by comparing fluorescence pattern of image with uniform dye concentration



(of incident, not fluoresced light)



corrected for by integrating along rays to determine the actual illumination and fluorescence at each point





Energy_evolution 0=5°





Numerical Simulations

• Code type

Semi-Lagrangian finite volume code

(conservation of fluid masses and momentum)

Two ideal gases ($\gamma = 5/3$)

Typical simulation: 3D at resolution 200×160×80

Viscosity and diffusion

Loss of resolution at grid scale ⇒ diffusion-like behaviour for mass fractions, analogous to molecular mixing for KE, analogous to dissipation, and added to IE In some runs, an explicit viscosity was added

• Approximating an incompressible fluid

Normalisation: choose H = 1, Ag = 1, $\rho_1 = 1$

Non-dimensional parameters (ideally small):

Density ratio	$B = \frac{\Delta \rho}{\rho_0} = \frac{2}{g}$	≈0.18	
Mach number	$M = \sqrt{\frac{AgH\rho}{p\gamma}} \approx \sqrt{\frac{3}{5p_0}}$	≈0.08	
Incompressibility ratio	$I = \frac{gH\rho^2}{5p\Delta\rho} \approx \frac{g^2}{10p_0}$	≈0.12	
max_{100}			

Compromise g = 11, $p_0 = 100$

• Initial conditions - basic distribution

Away from interface:

Since $u \approx 0$, require $\frac{\partial p}{\partial z} = -\rho g$. Require neutral stability, buoyancy frequency $N^2 = \frac{g}{T} \left(\frac{\partial T}{\partial z} + \frac{g}{c_p} \right) = 0 \iff \text{isentropic fluid } p = k(s)\rho^{\gamma}.$

At interface:

Choose specific heats at constant volume, c_{v1} and c_{v2} . Require temperature continuous $\Leftrightarrow c_{v1}\rho_1 = c_{v2}\rho_2$.

Everywhere:

Pressure field cannot be entirely hydrostatic.

Require $\frac{\partial}{\partial t}(\nabla . \boldsymbol{u}) = 0$.

Ignoring terms of $O(u^2)$, require

$$\nabla \cdot \left(\frac{1}{\rho} \nabla p\right) \propto \nabla \cdot \left(k^{1/\gamma} \nabla p^{(\gamma-1)/\gamma}\right) = 0,$$

with $\frac{\partial p}{\partial n} = -\rho g \hat{n} \cdot \hat{z}$ on boundaries with outward normal \hat{n} .

• Initial conditions - perturbations

2D velocity field with vorticity at interface models experimental barrier withdrawal.

3D random perturbation to interface position, wavelengths $\frac{L}{40} < \lambda < \frac{L}{20}$, rms amplitude $\sigma = \frac{H}{2500}$.





Energy budget

Numerical results $\eta \approx 0.48$



Typical experimental results $\eta \approx 0.38$



• Why the difference in energy budgets?

<u>Numerical diffusion not $abla^2$ </u>

2D advection test pattern

Numerical viscosity acts preferentially at small scales and is resolution and velocity-dependent

Total dissipation is unaffected by ratio of explicit/numerical viscosity until explicit viscosity dominates



Re of experiments is low

But experiments do not show Re dependence

Energy conservation

Small departures from energy conservation in stable waves

Sensitivity to initial conditions

But there is no change when λ_{random} increased by 4

Different molecular Pr

Does small-scale dynamics adjust to forcing from larger scales?



Cumulative mixing efficiency





Conclusions and further work

• Laboratory experiments

- * $\theta = 0^{\circ}$ $\eta_{cumulative} \approx 0.4$
- * As $\theta \uparrow$, $\eta_{\text{cumulative}} \downarrow$
- * For $\theta \leq 5^{\circ}$, $\eta_{instantaneous} \approx 0.5$

Numerical simulations

- * Models experiments at suitable parameters
- * Good agreement in large-scale overturning

• Further work

- * Investigate sensitivity of mixing to various factors
- * Investigate instability at higher angles up to limiting case:





* Extend study of mixing efficiency to more complex stratifications



Dalziel & Jacobs (2000)