Toy models for Rayleigh-Taylor instability:

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> with thanks to Joanne Holford (DAMTP) David Youngs (AWE)

The growth question:

$$h = \alpha Agt^2$$
, where $A = \frac{\rho_1 - \rho_2}{\rho_1 + \rho_2}$

But what is α ?

◊ 0.10, ... 0.07, 0.06, ... 0.03, 0.02 **?**

Timescale:
$$T = \sqrt{\frac{H}{Ag}}$$

If
$$\delta = h/H$$
, and $\tau = t/T$,

then $\delta = \alpha \tau^2$

Experiments





Appropriate modelling (?)



Growth

Dimensional analysis/similarity theory

 $h = \alpha Ag t^2$.

Single mode

Layzer (1955)

For $\zeta(x, y) = a_0 \cos \frac{2\pi x}{\lambda}$

if
$$\frac{dh}{dt} = w$$
,

then
$$(2+E)\frac{dw}{dt} = Ag(1-E) - C_D \frac{w^2}{\lambda},$$

where

$$E = \exp\left(\frac{-6\pi h}{\lambda}\right).$$

Experimentally $C_D \sim 10$ \diamond Does this make sense?

Early time
$$\rightarrow$$
 linear theory $\frac{d^2 h}{dt^2} = \frac{2\pi Ag}{\lambda} h$
Late time \rightarrow constant velocity $w_{\infty} = \sqrt{\frac{Ag\lambda}{C_D}}$

$$\Rightarrow h \rightarrow w_{\infty}(t-t_0)$$





Often described as 'bubbles'... ...but more like 'thermals' in miscible fluids

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Thermals



Self-similar

 $r = \theta z$ $V = \gamma r^3.$

Buoyancy conserved

$$g'V=g'\gamma r^3=g'_0V_0.$$

Constant Froude number

$$F^2 = \frac{w^2}{g'r}$$

Integrating w = dz/dt

$$\frac{\gamma^{1/2}\theta}{2F(g_0'V_0)^{1/2}}z^2 = t$$

Experimental results $\rightarrow F \approx 1.2$.

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Rayleigh-Taylor as thermals



Froude number ~ 1.2 (aspect ratio 0.72) $\Rightarrow C_{Thermal} \approx 1.3$.

Rayleigh-Taylor bubbles a little like thermals $\rightarrow C_D \approx 1.3$

But in Rayleigh-Taylor environment

• Density field not hydrostatic in ambient

 \diamond Hydrostatic in mean density \Rightarrow halve buoyancy force

 $\rightarrow C_D \approx 2.6$

• Flow around bubble affected by bubble moving in opposite direction

 \diamond Drag due to twice rise speed of bubble $\rightarrow C_D \approx 10.4$

In agreement with single mode experiments

Multi-mode

What happens if λ grows with h?

Let $\lambda = \psi h$

Late times approximation:

$$\frac{dh}{dt} = \left(\frac{Ag}{C_D}(1-E)\psi\right)^{1/2}h^{1/2}$$

 \Rightarrow

$$h = \frac{Ag}{C_D} (1 - E) \psi (t - t_0)^2 = \alpha Ag (t - t_0)^2$$

For $C_D = 10$ and $\psi = 1$, $\alpha = 0.025$.

[Full Layzer growth with $\psi = 1$ gives $\alpha = 0.023$.]

Growth rate maximised with $\psi \sim 10$ giving $\alpha \sim 0.103$



Where do the modes begin? How do they interact?

- ◊ Nonlinear interaction?
- ◊ Initial perturbation?



If modes independent and equal amplitude:

Instantaneous nonlinear mode halving interaction when $h = \lambda$:



Which is it?

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Mixing

See talk by Joanne Holford Energy budget

Can decompose *PE* into **Background PE** and **Available PE**. PE_{back} is the minimum energy state that is achieved by adiabatic rearrangement of fluid parcels.

Mixing increases PE_{back} – it cannot decrease it!

 PE_{avail} is the component of *PE* that can be converted into *KE*, heat (through dissipation) and, if mixing occurs, into PE_{back} .





Mixing efficiency



*

Thermal

Entrainment into a thermal

$$\frac{dV}{dt} = \beta wA$$

•••

$$\beta = 0.18.$$

Energetics of a thermal

Mixing efficiency not well defined: depends on size of domain!

Rayleigh-Taylor





$$h = \alpha A g t^{2} \qquad \qquad \delta = \alpha \tau^{2}$$
$$w = 2 \alpha A g t \qquad \qquad \omega = 2 \alpha \tau$$

 $V = 2L^2h$

Total potential energy

$$PE_{Total}^* = \frac{PE_{Total}}{PE_0} = 1 - 4\alpha^2 \tau^4$$

Background potential energy



Changes due to entrainment between counter-flowing streams. Invoke entrainment hypothesis: $u_e = \beta w$ Area of entrainment independent of *h*

 \Leftrightarrow depth of entrainment comparable with λ

 \Rightarrow entraining area = $\varphi \times$ plan area.

$$PE_{Back}^* = -(1 - \varphi \beta \alpha^2 \tau^4)$$

Available potential energy

$$PE_{Avail}^* = PE_{Tot}^* - PE_{Back}^* = 2 - (4 + \varphi\beta)\alpha^2\tau^4$$





Kinetic energy

 $KE^* = 16\sigma\alpha^3\tau^4$



Available energy changing

$$\frac{dE_{Avail}^{*}}{d\tau} = \frac{dKE^{*}}{d\tau} + \frac{dPE_{Avail}^{*}}{d\tau}$$
$$= -4(4 + \varphi\beta - 16\sigma\alpha)\alpha^{2}\tau^{3}$$

Hence, energy is lost whenever $\alpha < \frac{1}{4}$ (for $\beta = 0, \sigma = 1$).

Instantaneous mixing efficiency

$$\eta_{Inst} = -\frac{\frac{dPE_{Back}^{*}}{d\tau}}{\frac{dPE_{Avail}^{*}}{d\tau} + \frac{dKE^{*}}{d\tau}}{\frac{d\varphi\beta}{4 + \varphi\beta - 16\sigma\alpha}}$$

So for $\varphi = 16$, $\beta = 0.18$, $\sigma = 1$, and $\alpha = 0.06$, then $\eta_{Inst} = 0.49$.



Integral mixing efficiency

If there no mixing after reaching the bottom...



$$\eta_{Integral} = \frac{1}{8} \varphi \beta$$

For $\varphi = 16$ and $\beta = 0.18$, then $\eta_{Integral} = 0.36$.

If there is mixing after reaching the bottom...



$$E_{Avail}^{*(bot)} = \left(1 + 4\sigma\alpha - \frac{1}{4}\varphi\beta\right)$$

If $\Delta E_{Back}^{(After bot)} = \eta_{stab} E_{Avail}^{(bot)}$, then

$$\eta_{Integral} = \frac{1}{8}\varphi\beta + \frac{1}{2}\eta_{stab}\left(1 + 4\sigma\alpha - \frac{1}{4}\varphi\beta\right)$$

For $\eta_{stab} = 0.2$, then $\eta_{Integral} = 0.41$.



Extensions



Let Δ_c be the fractional displacement of the centroid of the *bubble* from z = 0.

$$\eta_{Inst} = -\frac{\delta P E_{Back}^{*}}{\delta P E_{Avail}^{*} + \delta K E^{*}}$$
$$= \frac{\varphi \beta}{4 + \varphi \beta - 4(4 - \varphi \beta) \Delta_{c} - 16\sigma \alpha}$$

Pyramid ($\Delta_c = 1/4$):	$\eta_{Inst} = 0.6.$
Parabolic ($\Delta_c = 1/6$):	$\eta_{Inst} = 0.56.$

(gives linear mean concentration)

 \rightarrow

How can we avoid having to specify C_D ?

Shell model

GOY model (Gledzer–Ohkitani–Yamada):

$$\frac{dU_n}{dt} = \left(ak_n U_{n+1}^* U_{n+2}^* + bk_{n-1} U_{n-1}^* U_{n+1}^* + ck_{n-2} U_{n-1}^* U_{n-2}^*\right)$$
$$- vk_n^2 U_n + F_n$$
$$k_n = \beta^n k_n a = 1, b = -c \text{ and } a = -1 + c$$

with $k_n = \beta^n k_0$, a = 1, $b = -\varepsilon$ and $c = -1 + \varepsilon$.

In Rayleigh-Taylor instability, energy input at all scales.

$$\frac{dU_n}{dt} = (k_n U_{n+1} U_{n+2} - \varepsilon k_{n-1} U_{n-1} U_{n+1} - (1 - \varepsilon) k_{n-2} U_{n-1} U_{n-2}) - \nu k_n^2 U_n + F_n$$

Recall Layzer model: $(2+E)\frac{dw}{dt} = Ag(1-E) - C_D \frac{w^2}{\lambda}$

Hence
$$F_n = A_n g \frac{1 - E_n}{2 + E_n}$$
, where
 $E_n = \exp\left(-\frac{6\pi h_n}{\lambda_n}\right)$ and $A_n = A h_n/h$.

The mode penetrations h_n and total penetration h are obtained from

$$\frac{dh_n}{dt} = U_n$$
 and $h = \max_n h_n$.



- ◊ Approximate quadratic growth
- Ocertificient depends on initial spectrum
- \diamond Possible to replicate $\alpha \sim 0.06$

Conclusions

General

- Initial conditions are important for gross features
- Internal details relatively insensitive to initial conditions
- Appropriate modelling of initial conditions gives close agreement

Thermals model

- Single-mode growth rate consistent with isolated thermal
- Simple model for transfer between modes replicates t^2 growth
- Mixing efficiency consistent with thermal entrainment

Shell model

- Baroclinic input at all scales
- Very simple model replicates t^2 growth
- Growth rate sensitive to initial spectrum

An explanation?

No, but it helps.