## **Blend Times in Stirred Tanks**

## **Reacting Flows - Lecture 9**

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# Evaluation of mixing performance

- Methods to evaluate mixing performance:
  - Characterization of homogeneity.
  - Blending time.
- General methods to characterize homogeneity:
  - Visual uniformity.
  - Quantitative change in local concentration as a function of time.
  - Review instantaneous statistics about the spatial distribution of the species.
    - Average concentration
    - Minimum and maximum
    - Standard deviation in the concentration.
    - Coefficient of variation CoV = standard deviation/average.

# Visual uniformity

- Experimentally measure the time it takes to obtain visual uniformity.
- Can be done with acid-base additions and a pH indicator.
- Offers good comparisons between performance of different mixing systems.
- Not a suitable approach for CFD.









## Visual uniformity example: glass mixing

- Glass exits the glass ovens with variations in temperature and material concentrations.
- As a result, when the glass hardens, there will be visual nonuniformities.
- So, glass needs to be mixed before it is used. Because of the highviscosity and temperature, special mixers are used,
- Optical quality of glass is still often determined visually.



# Quantitative variation in a point

- Measuring the tracer concentration as a function of time c(t) in one or more points in the vessel, is a common experimental method.
- The mixing time is then the time it takes for the measured concentration c(t) to stay within a certain range of the final concentration c<sub>∞</sub>.
- Advantage: easy to use in experiments.
- Disadvantage: uses only one or a few points in the vessel.
- Does not use all information present in a CFD simulation.



# Blend time calculations with CFD

- Transport and mixing of a tracer:
  - Add tracer to the domain.
  - Mass fraction of tracer calculated and monitored as a function of time.
  - Determine blend time based on the mass-fraction field satisfying a pre-specified criterion.
- Flow field required can be steady, frozen unsteady or unsteady.
- Benefit of CFD:
  - The full concentration field is known.
  - Can use more data to determine blend time than what can be measured experimentally using probes.
- Main question: what should be the mixing criterion?

# CFD analysis for blend time

|   |                  | Physical<br>Lab | CFD<br>Lab |
|---|------------------|-----------------|------------|
| Addition of<br>Tracer                     | Volume of Tracer | Controlled      | Exact      |
|   | Delivery Time    | finite          | zero       |
|   | Location         | variable        | fixed      |
| Concentration<br>Measurement<br>of Tracer | Conductivity     | Yes             | No         |
|   | Color            | Yes             | No         |
|   | Mass Fraction    | inferred        | Yes        |

- We will now:
  - Illustrate the blend time analysis using a 2-D Rushton turbine flow field example.
  - Tracer added and its transport and mixing calculated. Mass fractions are monitored as a function of time.
  - Blend time is calculated using different criteria.



Rushton Impeller - 50 RPM - 31.6I Vessel Velocity Vectors Colored By Velocity Magnitude (m/s)



Rushton Impeller - 50 RPM - 31.6l Vessel Contours of Mass fraction of water-liquid-tracer (Time=0.0000e+00) FLUENT 6.2 (axi, segregated, spe, rke, unsteady)



#### Measures of variation

- Variations in Y, the mass fraction of tracer, can be measured in several ways. For all measures, greater numbers indicate a greater variation with no upper bound.
- Coefficient of variation. Ratio between standard deviation  $\sigma_{Y}$  and the average  $\langle Y \rangle$ :

$$CoV = \frac{\sigma_Y}{\langle Y \rangle}$$

- Ratio between maximum and minimum mass fractions  $Y_{max}/Y_{min}$ .
- Largest deviation between extremes in the mass fraction and the average:

$$\Delta_{\max} = \max(Y_{\max} - \langle Y \rangle, \langle Y \rangle - Y_{\min})$$

Can also be normalized over < Y>.

# Variation calculation example

- Mass fraction data:
  - Min-max anywhere: 0.0223-0.539
  - Min-max from probes: 0.0574-0.272
  - Average: 0.0943
  - Standard deviation: 0.0493
- Measures of variation:
  - Max/min = 0.539/0.0223 = 24.2 (anywhere)
  - Max/min = 0.272/0.0574 = 4.7 (from probes)
  - CoV = 0.0493/0.0943 = 0.52
  - $-\Delta_{max} = max(0.539-0.0943, 0.0943-0.0223) = 0.44$
  - $-\Delta_{max}/\langle Y \rangle = 0.44/0.0943 = 4.7$





### Measures of uniformity - absolute

- There is a need to have an absolute measure of uniformity U that is ≤ 1 with 1 (or 100%) indicating perfect uniformity.
- Ratio between the minimum and maximum mass fractions.
  - Bounded between 0 and 1.

$$U_{\min/\max} = \frac{Y_{\min}}{Y_{\max}}$$

 $U_{CoV} = 1 - CoV$ 

- Based on coefficient of variance CoV:
  - Not bounded: can be less than 0.
- Based on largest deviation from the average:
  - Conceptually closer to common experimental techniques.
  - Not bounded: can be less than 0.

$$U_{\Delta} = 1 - \frac{\Delta_{\max}}{\langle Y \rangle}$$

#### Uniformity



Time (s) 15

# <u>Uniformity</u>

- These measures of uniformity:
  - All indicate perfect uniformity at values of 1.
  - Are not all bounded between 0 and 1.
  - Do not take initial conditions into account.
- Generally, it is most useful to be able to predict the time it takes to reduce concentration variations by a certain amount.
- This is then done by scaling the largest deviation in mass fraction at time t by the largest deviation at time t=0.

$$U(t) = 1 - \frac{\Delta_{\max}(t)}{\Delta_{\max}(t=0)}$$

- E.g. for the example case:
  - − At *t*=0s,  $Y_{max}$ =1 and <*Y*>=0.0943 →  $\Delta_{max}$ (*t*=0) = 0.906.
  - − At *t*=10s,  $\Delta_{max}(10s) = 0.44 \rightarrow U(10s) = 0.51$ .
- Data are often correlated in terms of number of impeller revolutions, at t=10s and 50RPM, there were 10\*RPM/60=8.33 impeller revolutions.



## Comparison between systems

- Let's compare two systems with:
  - The same flow field.
  - The same spatial distribution of species.
  - But different initial mass fractions of species.



Layer with  $Y_{tracer} = 1$  on top of fluid with  $Y_{tracer} = 0. < Y > = 0.094$ .

Layer with  $Y_{tracer}$ =0.4 on top of fluid with  $Y_{tracer}$ =0.1. <Y>=0.13.











## Compare two more systems

- Two systems with approximately the same average mass fraction of tracer <Y> ≅ 0.5.
- The initial distributions are very different: layered vs. blocky pattern.



Layer with  $Y_{tracer} = 1$  on top of fluid with  $Y_{tracer} = 0. < Y > = 0.497$ .



Blocky pattern of fluid with  $Y_{tracer}=0$ . and fluid with  $Y_{tracer}=1. < Y>=0.491$ .











Time (s)

# Compare all four systems

| Initial Y          | U    |  |
|--------------------|------|--|
| Layer (0 to 1)     | 20.3 |  |
| Layer (0.1 to 0.4) | 20.3 |  |
| Blocky Pattern     | 10.7 |  |
| Half-Half          | 26.1 |  |

- Table shows number of impeller revolutions it takes to achieve 99% uniformity for all four systems using the two main criteria:
  - U based on reduction in initial variation.
  - $U_{\Delta}$  based on variation from the average.
- Conclusion: systems with good initial distributions mix faster.
- General recommendation: use U (reduction in initial variation) to correlate results or to compare with literature blend time correlations.

## Continuous systems

- Methods so far are for batch systems.
- Do these methods work for continuous systems?
  - Requires some modification.
  - Looking at mass fraction extremes does not work, because these may be fixed by the inlet mass fractions.
- Various approaches used:
  - Compare batch blend time with average residence time of the material (RT = liquid volume divided by volumetric flow rate). If batch blend time is much smaller than RT, assume there is no mixing problem.
  - Perform particle tracking simulation, similar to shown for static mixers in previous lectures. Analyze residence time distributions.
  - Perform tracer mixing calculation.

## Tracer mixing calculation

- Calculate continuous, steady state flow field.
- Initial mass fraction of tracer is zero everywhere.
- Perform transient calculation for tracer mixing, with non-zero mass fraction tracer at inlet.
- Monitor:
  - Average mass fraction in domain < Y>.
  - Mass fraction at outlet Y<sub>out</sub>.
  - Optional: monitor CoV.
- Definition of perfectly mixed system:  $Y_{out} = \langle Y \rangle$ .
- Mixing time is then the time it takes for the ratio Y<sub>out</sub>/<Y> to be within a specified tolerance of 1.
- Mixing time can be expressed in number of residence times: t/RT.

## Compare two systems

- Rushton turbine flow field.
- Continuous system with two different injections:
  - Low velocity feed (0.01 m/s) distributed across liquid surface.
    - Affects flow in upper part of the vessel only.
  - High speed jet feed (9.6 m/s) entering through bottom shaft.
    - Because of the large momentum contained in the jet, it alters the flow field significantly.
- Outflow at center of bottom.
- Average residence time RT=30s, equivalent to 25 impeller revolutions. The RT is similar to the batch blend time.





Rushton Impeller - 50 RPM - 31.6l Vessel - Continuous flow - Surface Inlet Contours of Mass fraction of water-liquid-tracer (Time=0.0000e+00) FLUENT 6.2 (axi, segregated, spe, rke, unsteady)



Rushton Impeller - 50 RPM - 31.6l Vessel - Continuous flow - Shaft Inlet Contours of Mass fraction of water-liquid-tracer (Time=0.0000e+00) FLUENT 6.2 (axi, segregated, spe, rke, unsteady)





# **Comments**

- The main assumption behind this approach is that the system will eventually reach a steady state where Y<sub>out</sub>=<Y>.
  - Not all industrial systems may have a steady state operating condition which, in general, is an undesirable situation that would need to be addressed.
- CoV can still be used to compare uniformity of different systems under steady state operating conditions with multiple species.