

INTAKE FRACTION

- * Intake fraction = IF (previously inhalation transfer factor)
- * Simple but powerful tool (screening calcs; inhalation dose)
- * IF = fraction pollutant mass emitted by single source that is inhaled.
 - inhaled by (a) individual or (b) population.
- * Dose (inhalation) = IF $\times$ E $\times$ Δt

$\uparrow$
 mass emission rate.
- * PIF = population intake fraction = $\sum_{\beta} IF$ (all individuals)
 - β = entire urban area; large building, etc.

Mathematically:

$$IF = \frac{\text{mass inhaled}}{\text{mass emitted}} = \frac{\int_{t_0}^{t_{f2}} C(t) Q_{\beta}(t) dt}{\int_{t_0}^{t_{f1}} E(t) dt}$$

$C(t)$ = mass concentration due only to specific source (mg/m³)

$Q_{\beta}(t)$ = breathing rate (m³/hr)

$E(t)$ = specific source emission rate (mg/hr)

t = time (hr) 0 = initial f = final note t_{f1} vs. t_{f2}

At steady-state

$$IF = \frac{C Q_{\beta} \int dt}{E \int dt} = \frac{C Q_{\beta}}{E} = \frac{\text{mass inhalation rate}}{\text{mass emission rate}}$$

Recall mass balance model for $C(t)$

$$\frac{dC}{dt} = \lambda \phi_0 - \lambda C + E/V \quad \therefore \frac{dC}{\lambda C - E/V} = -dt$$

ϕ (why?)

$$\ln \left\{ \frac{\lambda C - E/V}{\lambda C(t=0) - E/V} \right\} = -\lambda \int dt = -\lambda t$$

$$\ln \left\{ 1 - \frac{\lambda C}{E/V} \right\} = -\lambda t \quad \therefore C = \frac{E}{\lambda V} (1 - e^{-\lambda t})$$

or at steady-state $C = \frac{E}{\lambda V}$

Dynamic Case:

$$IF = \frac{\int C(t) \Phi_B(t) dt}{\int E(t) dt} = \frac{\int \frac{E(t)}{\lambda V} (1 - e^{-\lambda t}) \Phi_B(t) dt}{\int E(t) dt}$$

• Assume E, λ, V, Φ_B constant.

$$IF = \frac{\frac{E}{\lambda V} \Phi_B \int (1 - e^{-\lambda t}) dt}{E(t_f - t_i)} = \boxed{\frac{\Phi_B}{\lambda V \Delta t} \int_{t_i}^{t_f} (1 - e^{-\lambda t}) dt}$$

Steady-state Case:

$$IF = \frac{C \Phi_B}{E} = \frac{\frac{E}{\lambda V} \Phi_B}{E} = \frac{\Phi_B}{\lambda V} = \boxed{\frac{\Phi_B}{\Phi_V}}$$

NOTES:

1. $IF \neq$ function (E) in either dynamic or s.s. cases!
2. IF depends on Φ_B, Φ_V (or λV) and time (if dynamic)
3. Simple, but powerful.

A closer look at IF for steady-state

$$IF = \frac{\Phi_p}{\Phi_v} \approx \frac{0.78 \text{ m}^3/\text{hr}}{\lambda V} \quad \left\{ \begin{array}{l} \lambda \approx 0.2 - 1/\text{hr.} \\ V \approx 350 \text{ m}^3 \text{ (varies a lot!)} \end{array} \right.$$

$$IF \approx 0.002 - 0.01 \quad (0.2\% \text{ to } 1\% \text{ of mass emitted is inhaled})$$

* Assumes well-mixed environment (no personal cloud, etc.)

* Assumes no removal terms.

Comparison w/ outdoor sources

* $IF \approx 10^{-12} - 10^{-10}$ (from literature, e.g., Lai et al.)

* $IF(\text{indoor source}) / IF(\text{outdoor source}) \approx 10^7 - 10^{10}$.

* outdoor source must emit $10^7 - 10^{10}$ more for same dose as indoor source (proximity/activity patterns)

EXAMPLE: $PM_{2.5}$ emissions from candle.

* $E_{\text{candle}} \approx 1 - 100 \text{ mg/hr}$. (use Pagels Candle 1 $\approx 2.4 \text{ mg/hr}$)

* mass inhaled = mass emitted $\times (IF)$

* want $\rightarrow$ outdoor source equivalent of 1 candle. ($PM_{2.5}$)

$$IF_{\text{indoor}} \times E_{\text{candle}} = IF_{\text{outdoor}} \times E_{\text{source}}$$

mass inhaled indoors = mass inhaled outdoors

$$E_{\text{source}} = \frac{IF_{\text{indoor}}}{IF_{\text{outdoor}}} \times E_{\text{candle}} \approx \frac{0.006}{10^{-11}} (2.4 \text{ mg/hr})$$

$$= 1.4 \times 10^9 \text{ mg/hr} = 1.4 \times 10^3 \text{ kg/hr} = 1.4 \text{ metric tons/hr}$$

$$\approx 12,000 \text{ metric tons/year (wow!)}$$

How does this compare w/ all major point sources in Texas?

* PM_{10} (2010) $\approx 55,000 - 60,000$ metric tons/year.

* All sources (major point)! & $PM_{2.5} < PM_{10}$!!

POPULATION INTAKE FRACTION (PIF)INDOOR:

$$\text{Dynamic: } PIF = \frac{\sum_{i=1}^N \int C_i(t) \Phi_{bi}(t) dt}{\int E(t) dt}$$

$$\text{Steady-state: } PIF = \sum_{i=1}^N \frac{C_i \Phi_{bi}}{E}$$

} $N = \#$ of
building
occupants.

Removal of Indoor Pollutants: (indoor source)
w/ Removal

$$V \frac{dC}{dt} = \underbrace{Q \phi_0}_{\text{control}} - QC + E - \underbrace{2Q_c C}_{\text{deposition/reaction on surfaces}} - V_d A C$$

$$\frac{dC}{dt} = -\lambda C + \frac{E}{V} - \frac{2Q_c}{V} C - V_d \frac{A}{V} C$$

↑
losses of ozone & particles.

* Assume steady-state to simplify:

* Assume no control:

$$0 = -\lambda C + \frac{E}{V} - V_d \frac{A}{V} C \quad \therefore \quad 0 = -(\lambda + V_d \frac{A}{V}) C + \frac{E}{V}$$

$$C = \frac{E/V}{(\lambda + V_d \frac{A}{V})} \quad \therefore \quad \frac{1}{(1 + \frac{V_d A/V}{\lambda})} \times \frac{E}{\lambda V}$$

$$IF = \frac{C \phi_\beta}{E} = \frac{1}{(1 + \frac{V_d A/V}{\lambda})} \frac{E}{\lambda V} \phi_\beta = \frac{1}{(1 + \frac{V_d A/V}{\lambda})} \left(\frac{\phi_\beta}{\phi_{\text{vent}}} \right)$$

+ $\frac{2Q_c}{V}$
(If control added back in)

NOTES:

* Effects of removal on IF

- If $V_d A/V \sim \lambda$ IF down by $1/2$.

* For OZONE: $V_d A/V > \lambda$ ($\sim 2-5 \times$ as great)

- IF lower than for inert pollutant.

* For particles: $V_d A/V = f_n(d_p)$

- See results in Lai et al. (don't worry about this $\rightarrow$ I will show in lecture)
- Discuss peak IF & why?

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* Discussion of Nazaroff (2008).