

Fluidization of fungal pellets using miniaturized 3D-printed fluidized bed

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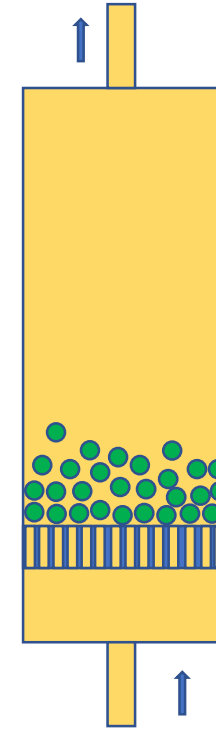
- Poor mixing
- Not scaleable
- Short-term culture



Culture flask



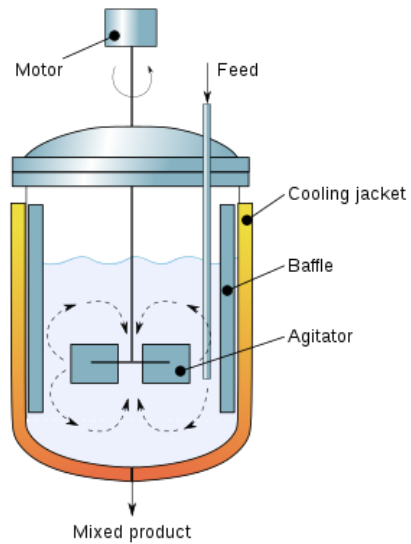
Gas/liquid outlet



- Good mixing
- Low shear stress
- Simplified construction

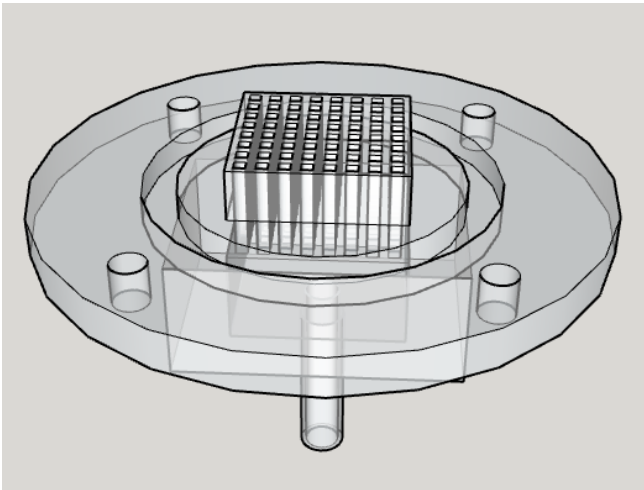
Fluidized bed reactor

- High capital cost
- Shear stress
- Cell damage/death

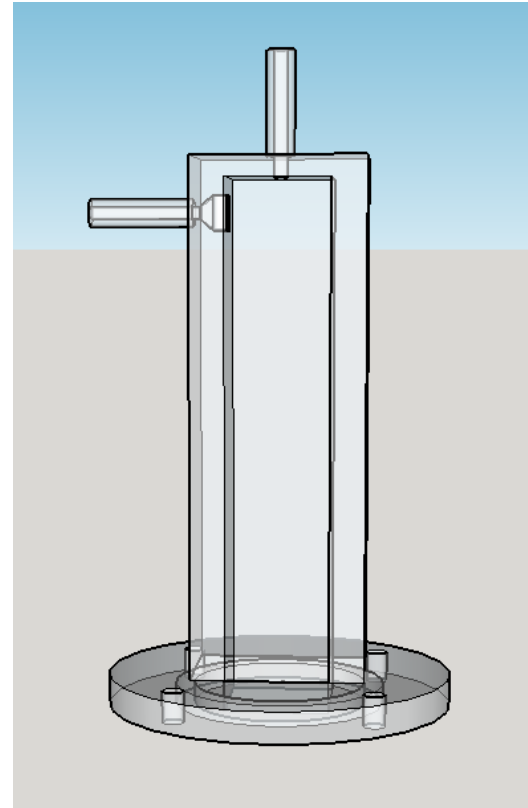


Stirred-tank reactor

- Fluidized bed design and fabrication



Lower Part



Upper Part



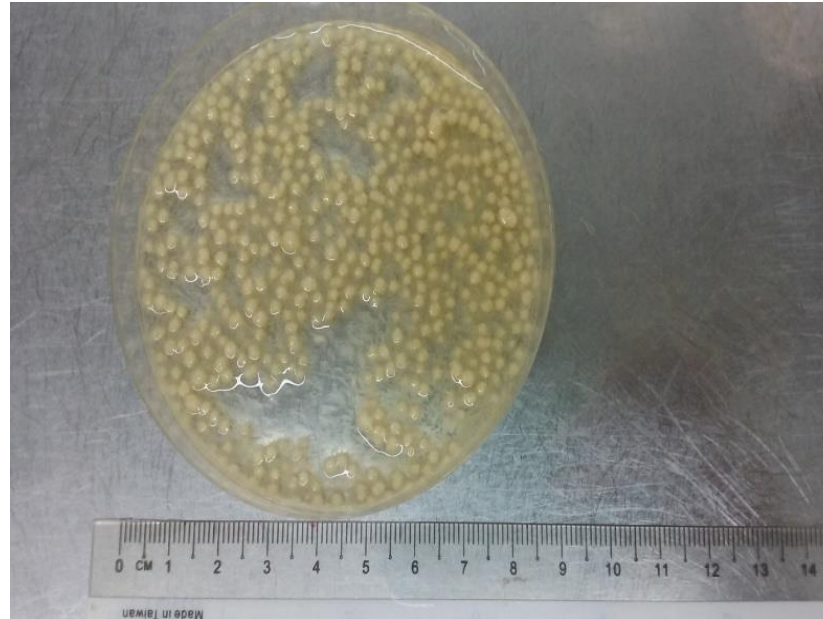
Assembled fluidized bed

- Designed by: Google SketchUp
- Printed by: Form2 SLA printer
- Cross-section: 15×15 mm; Bed height: 15 mm
- Planar distributor (5 mm in height): contains 1mm square holes

- Fungal pellets



Fungi in nature



Fungal pellets in petri dish



Single Pellet in microscope

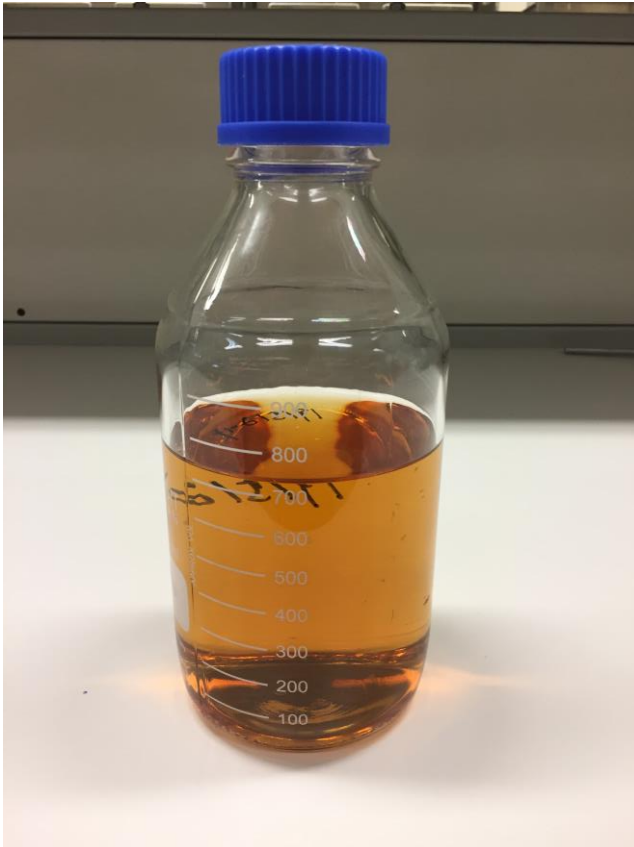
Fungal cells: *Nidula niveo-tomentosa* (Bird's nest fungi)

Size: ~2mm

Density: $1.1\text{g}/\text{cm}^3$

Pellet properties unique to fungi are an important consideration

- Culture medium



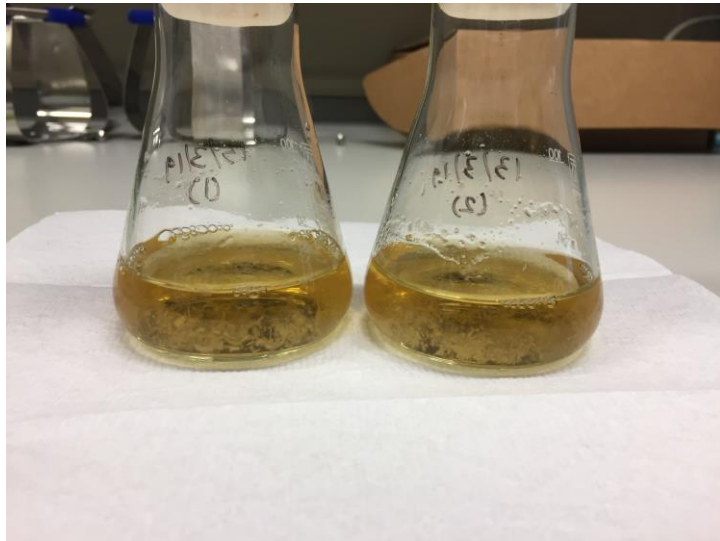
Density: 1.02 g/cm^3

Viscosity: $1.1 \text{ mPa}\cdot\text{s}$

Components of cultivation medium

Components in Media	Concentration (g/L)
glucose monohydrate	50
peptone from soy	6
potassium dihydrogen phosphate	2.5
magnesium sulfate	0.5
yeast extract	1.5
calcium chloride dihydrate	.0735
Ultrapure H ₂ O	fill up to 1 L

- Fungal pellets fluidization



Flask culture

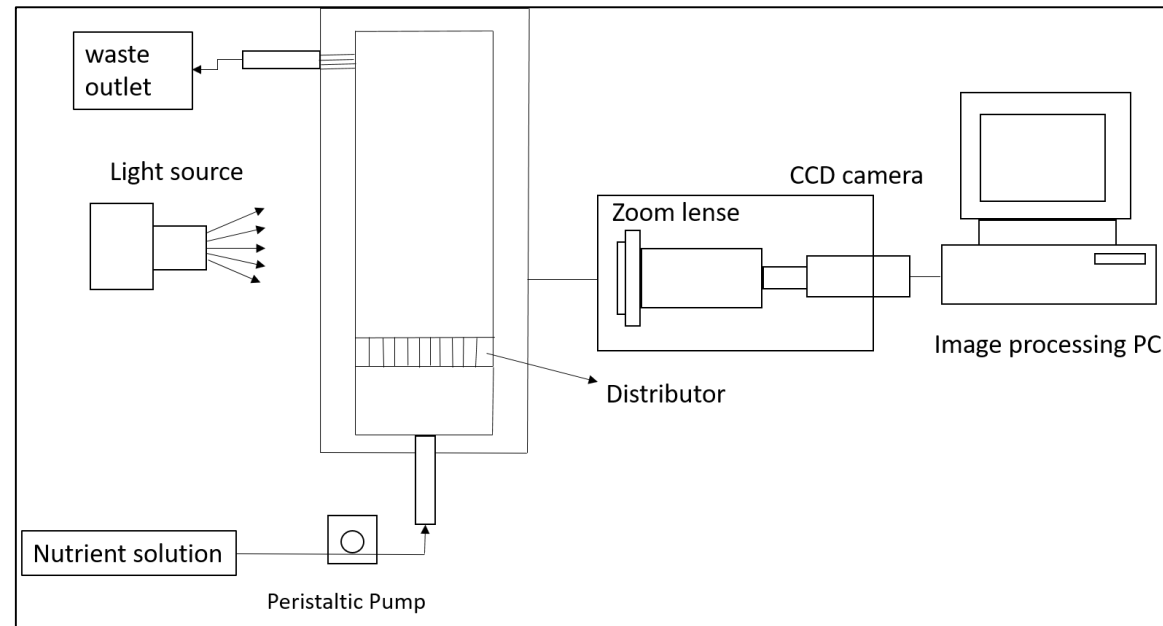
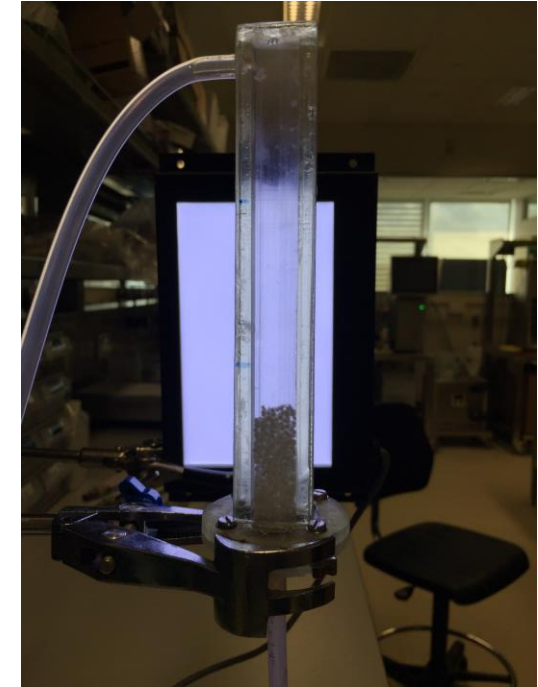
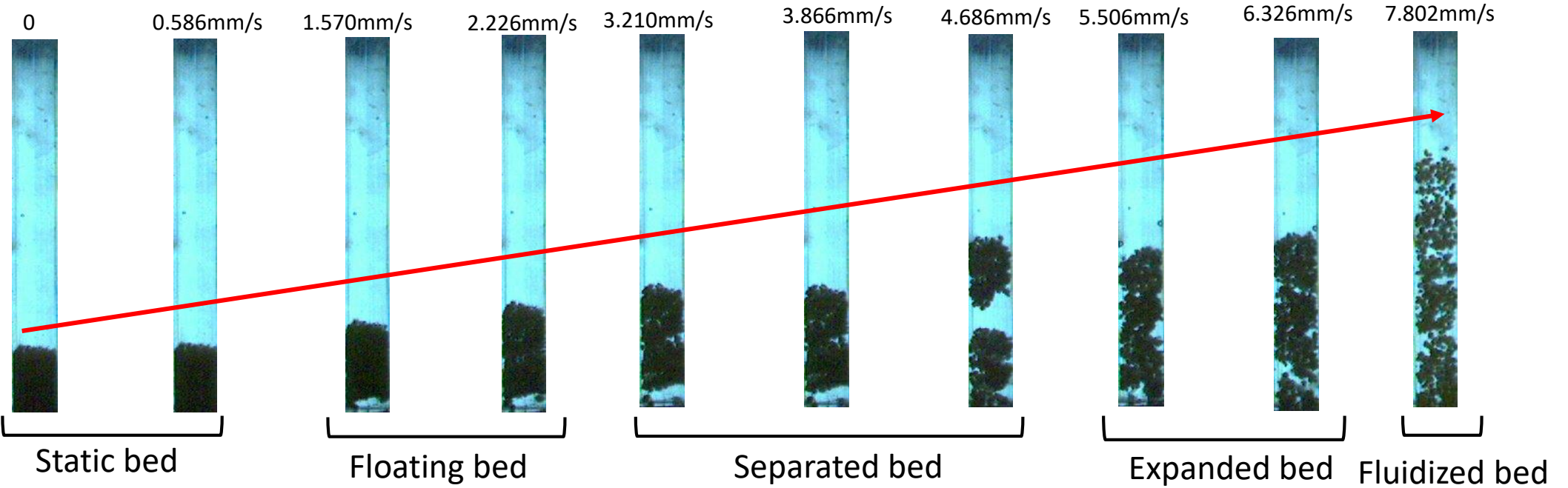


Diagram of experimental setup

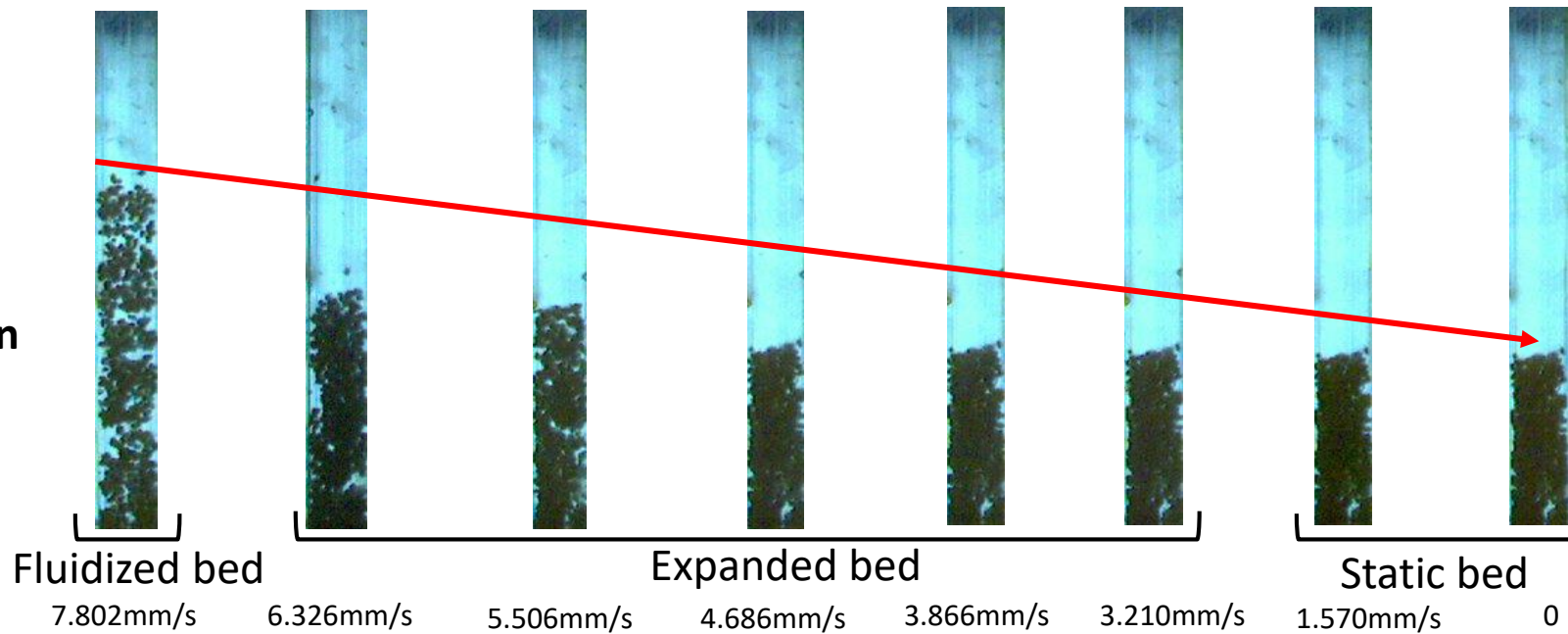


Fluidization process

Fluidization



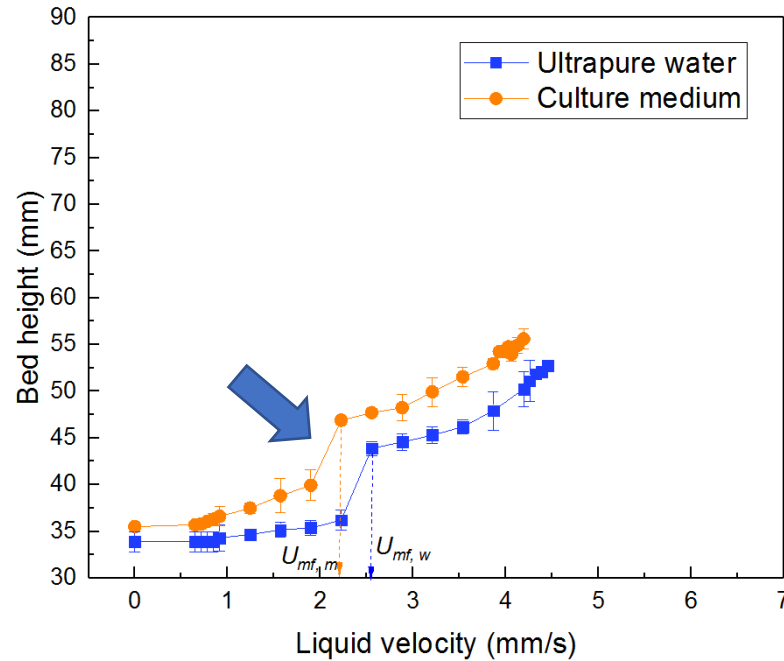
Defluidization



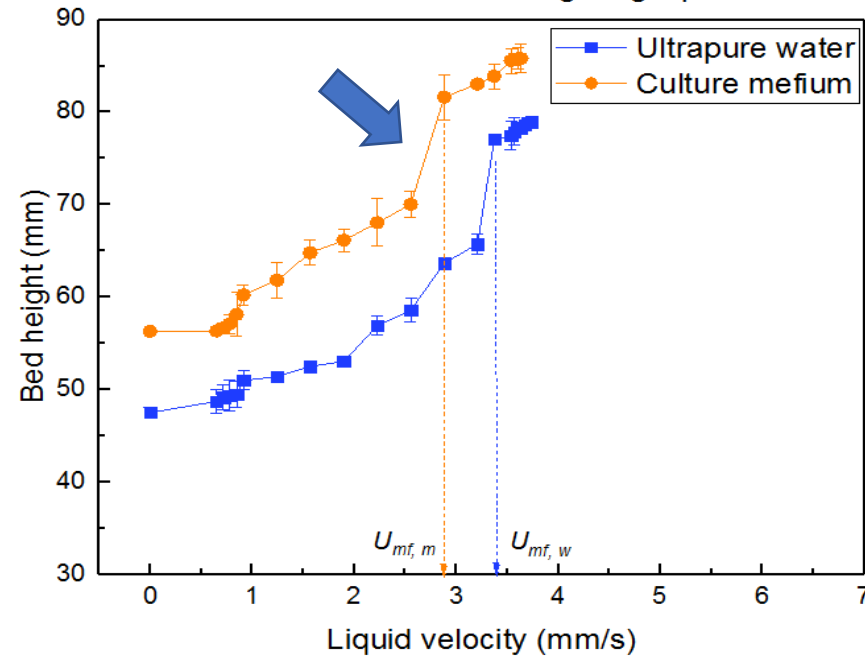
Pellets fluidization



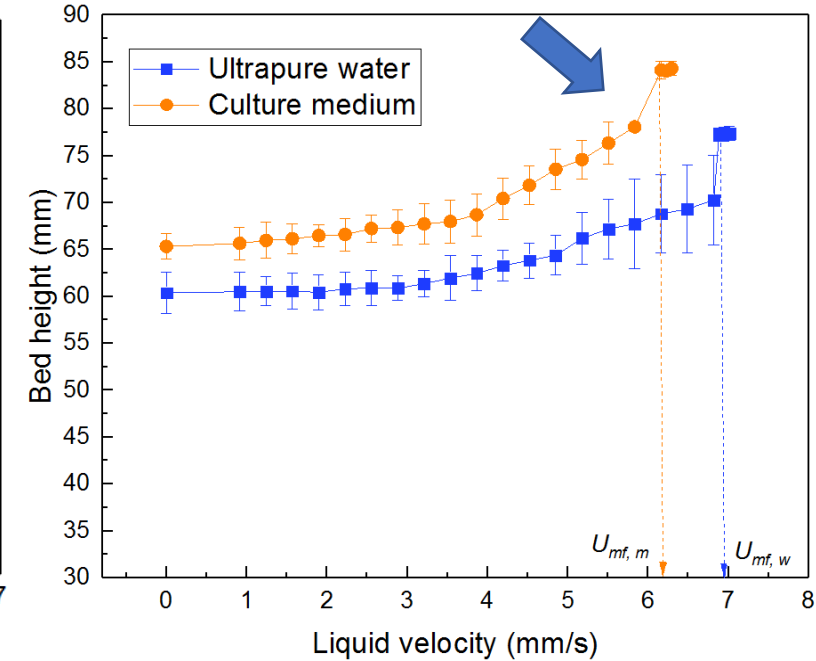
Defluidization curve of 0.5g fungal pellets



Defluidization curve of 1.0g fungal pellets



Defluidization curve of 1.5g fungal pellets



- The bed height jump occurs when agglomerated pellets become dispersed and fully fluidized, when liquid velocity reaches U_{mf} ;
- The U_{mf} is increasing with mass of fungal cells as the voidage of initial packed bed is increasing with overall mass.

- Ergun equation

$$U_{mf} = \frac{d_p^2 (\rho_P - \rho_L) g}{150 \mu_L} \left(\frac{\varepsilon^3 \phi_P^2}{1 - \varepsilon} \right)$$

Pellets mass (g)	0.5		1.0		1.5	
Initial bed voidage, ε_0	0.38		0.43		0.51	
Liquid type	Ultrapure water	Culture medium	Ultrapure water	Culture medium	Ultrapure water	Culture medium
Experimental Minimum fluidization velocity (mm/s) $U_{mf,e}$	2.226	1.898	3.210	2.554	6.818	6.162
Theoretical minimum fluidization velocity (mm/s) $U_{mf,th}$	1.869	1.669	2.698	2.409	5.71	5.105

- There are some transitional regime (separated bed, expanded bed) before actual fluidization regime start usually not present in liquid fluidization.
- The agglomerated fungal pellets are separated into dispersed pellets and homogenously fluidized by the liquid flow when liquid velocity reaches the U_{mf} .
- The homogenous fluidization and suspension of pellets exhibit the feasibility of fungal fermentation using fluidized bed bioreactor.
- The experimental value of $U_{mf,e}$ is larger than $U_{mf,th}$ calculated from Ergun equation, mostly due to increase of initial voidage (wall effects) but possible due to surface forces (cohesive forces between pellets).

Future work:

- Fungal pellets fluidization characterization
- Fermentation in gas-liquid-solid fluidized bed (aeration)

Questions?

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