growth curve batch 3 at 35C

Growth curve of 32 strains

	genome_id exp_id		genome_id exp_id		genome_id exp_id		genome_id exp_id		
1	g2	H2M3R1 9	g11	L2M4R1 17	g22	fp1-2	25	g30	gp1-3
2	g3	H2M3R2 10	g13	L3M5R1 18	g23	fp1-3	26	g31	bg-2
3	g4	H3M1R1 11	g15	L4M2R2 19	g24	fp2-2	27	g32	bg-3
4	g5	H3M3R2 12	g16	L4M3R3 20	g25	crp1-2	28	g33	pms-1
5	g6	H3M4R1 13	g17	L4M4R1 21	g26	crp1-3	29	g34	pms-2
6	g8	H4M5R1 14	g19	L4M7R1 22	g27	crp2-2	30	g35	pms-3
7	g9	L1M2R2 15	g20	L3M1R1 23	g28	gp1-1	31	g36	ppf-1
8	g10	L2M2R1 16	g21	src-2 24	g29	gp1-2	32	g37	40 th- 1

- revive glycerol stock of rhizobia on TY agar
 - prepare labelled plates
 - use a sterilized inoculation loop to streak the glycerol stock on TY agar
 - incubate the petri plates at 30C for 2-3 days until the colony is identifiably large
- prepare ensifer inoculum
 - prepare a DW96 filled with 300uL of PBS
 - pick colonies from the petri plate and dissolve the colonies in the PBS. Use wells in column 1, 5, and 9
 - dispense 100 uL of the dissolved colony to a NUNC96 and measure od at 600 nm
 - standardize the OD600 to 0.1. For target od = 0.1 od_target and target inoculum volume = 100 uL volume_target, and given the measured od of dissolved colonies od_dissolved, use the code below to calculate the needed volume of the dissolved colonies volume_dissolved and the pbs volume volume pbs
 - prepare another NUNC96, follow the table to add required volume of dissolved colonies and pbs to make 100 uL of standardized inoculum
 - measure the od of the standardized inoculum as od_inoculum
- prepare 100 uL of TY medium in NUNC96
 - inoculate 2 uL of the standardized inoculum to each well

