

THE QUADRANGLE CLUB
CHICAGO

K.O. Kiepschmidt +
Nakum - 1948 or 49

Planta 1951

Brauer

Onervuboro

astroplysed co

Catania

Lively

Freshman i.B.

312 chemostat of Fox B

$\sigma = 6.7$ N limited
has added anthraxite wood to
tank to see if "indol" is
formed and

anthraxite remains unchanged
does hypophane iodide react
turn of conversion from anthraxite to
indol? & it seems, but does anthraxite
get the 2

1) add indol to B nitrogen level
does hypophane or precursor
form and 2

2) add ~~anthraxite~~ ^{anthraxite} to a low nitro-
gen B/it chemostat (at high I)
does precursor form (this chemos-
tat is at high I values hypophane
and)

3) add indol like in No 2)

321 and 321 B and C
low phosphorus B/it

321 at first at $\sigma = 4.7$ hrs it poured
out -

When adding 1.1 mg/l to tank at 3.75 hrs
it does not pour out Beck at

350 & 346

284. 100g/l trypt.

2.3 mg/l arginine

with B/it and arginineless strain D 24/6
to see if one strain steals growth factor
from another.

platings with T1 and T6 (variable counts)

result: at fast flow rates there $\bar{c} = 2 \text{ hrs}$
seemed to be a lower titer ^{of B/it} ~~with T1~~
~~platings~~ indicating that there
may be some stealing of pyrophosphate
(titer falls to $1/2$ at $\bar{c} = 2 \text{ hrs}$)

281 Precursor in glucose medium
at $\bar{c} = 4 \text{ hrs}$ 500g/l trypt.

Beckman at 350: 0.109, 280.437
pours out 0.116/lm

282 auxile mutant precursor exp.

50g/l trypt 2.3 mg/l arginine

Beckman at 350: 0.330 [reverted?]

no conclusions

283 same purpose as 282 no conclusion

286 and 287 low ammonia + 50g/l trypt -
pheme, B/it, precursor as fundan
at T



Think it through!

286^o continued (incomplete) bank)
we added pantamulinic to it (at $T = 3.4$ hr

~~286^o continued~~ if remained
unstrapped, [It does not repress]
this should be repeated at larger values
and also indol should be tried to
see if it helps] 20 mg/ml pantamulinic to bank
but not effect either.

297^a 1 mg/ml of indol was added (to see
if it represses; it does not
repress) for 296 and 297 we ought to have
added at the end excess
305 experiment similar to 284 ^{mycoto}
but this time B10 predominates; ^{phone}

307: Does pyrophosphate prevent precursor
when Chemostat is nitrogen led
30 mg/ml NH₄Cl + 2 mg/ml pyrophosphate +
+ 10 mg/ml of alabine for
congress precursor.
2 B10?

$T = 4.3$ hrs, back at 310: 245

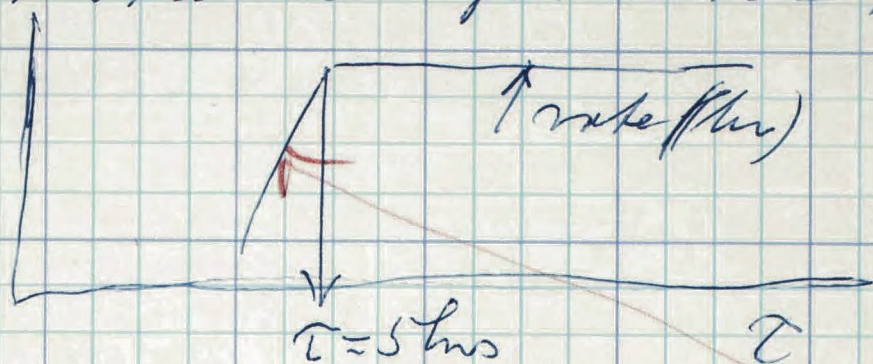
shows signs of autolyses in back-
ground (does not pour out) -

308 same as 307 but no phenylala-
 $T = 4.4$ hrs mine

317 similar result
Bernie Davis strain ~~to~~ 153-11
500 μ l pyrophosphate, $T = 3.25$ hrs
it pours out something like
our precursor

321 confirmed
added P

Best results maximum at $35^\circ = 4 \times 134$
worst τ got curve for precursor



This ought to be repeated with two different absolute conc. of P. ~~fall~~ Question is fall in curve the same at different P. If fall is the same then the growth rate is responsible. If fall is much steeper at higher P that means the tryptophan conc. is controlling the fall. -

321 B and C (about 525 f/l Trypt)
precursor production ~~was~~ determined
as function of τ (See curves)
We don't understand why B ~~doesn't~~ rises
at later times τ does not reach there
curves ~~the~~ unless we assume B has
more P and less Tryptophane

321 C + indol does it pour
out

338 Semi factory
a.) add hypophane
b.) change from $\tau = 2$ hrs to τ of hrs
observe increase in heat density

2 exp Howard

25°C Howard to set up with
1.) $\tau = 9$ hrs 500 g/l to see precursor
and at 37 2.) 500 g/l to see precursor
at $\tau = 3.7$ if suddenly stopped
do lockstep fall

change from small τ to large τ
observe

210

at up 1 mg/l hypophane.

log studies $\frac{-t}{6 \text{ hr}} + \frac{t}{6 \text{ hr}}$

$$\frac{t}{\rho} = \frac{6-2}{12}$$

measures out
with $\tau = 3$

try it and Nitrogen the
Chromostat, add methyl suddenly
every 1/2 hour large growth tube, measure

add
indol

321 B and C - To test this we added
 tryptophane to B. Periodical after came up
 to same value as in C. Therefore both
 should have equal amount of P. - P does not
 pour out now at 8 hours & does it pour
 out at 15.5 hours because tryptophane
 is destroyed by tryptophanase.

increase in precursor and also increase
 in bacterial density

323, 324 sudden change of flow
 rate from start to pour time observe
 1 mg/l trypt. & values & about 2 hrs
 B/14

324 $\tau = 2.03$ hours at 350: 0.212 at 280: 0.637
 system 0.173 and 140 $x_1 = 142$

τ changed by factor 4 and to $\tau = 8$ hrs
 after 1 hour sample poured out
 at 350: 0.244

supernatant gives
 at 280: 0.252 at 250: 0.171 $x_1 = 226$

now on hours $82 + \frac{226 + 142}{8 \times 2}$ it pours out in total

$$\frac{23}{82} = \frac{23}{105}$$

pours out $105 \times \frac{145}{212} = 72$ / hour normalized

$$105 + \frac{173 + 252}{8 \times 2} =$$

072 / hour is really not enough for total
nutriment should be rather 1.2 or 1.4
What does this mean