Research Bacteriology -Bldg. 54 - Rm. 16



VETERANS ADMINISTRATION

HOSPITAL
Hines, Illinois

YOUR FILE REFERENCE:

September 19, 1952

IN REPLY REFER TO:

Dr. Joshua Lederberg Department of Genetics College of Agriculture Madison 6. Wisconsin

Dear Dr. Lederberg:

We have used no very precise methods in preparing the McLeod's synthetic medium except for using reagents of highest purity for our work with M N Q. The composition of the medium was as follows:

Ammonium sulfate	4.72 grams
Asparagine	2.0 "
Sodium chloride	5.0 "
Glucose	2.0
$\text{KH}_2\text{PO}_4 \left(\underline{\text{M}}\right)$	
5 5	100. ml.
Va OH (N)	16.4 ml. (or to pH 7.8)
	-

0.5% solutions of MgCl₂, Fe Cl₂ and Ca Cl₂ were freshly prepared separately, then mixed, and 1.0 ml of the mixture was added to the medium.

Distilled water

1000 ml.

Chemicals were weighed only roughly on a triple beam balance accurate to 0.01 gram, even for the M/5 KH₂ PO₄ solution.

pH should be about 7.8 before autoclaving, using the amount of NaOH designated; if not, NaOH was added to this point.

pH after autoclaving at 12 lbs. for 20 minutes, 7.6.

Medium was distributed by hand-pipetting with 25 ml measuring pipettes in 24.5 ml. amounts into $8^{m} \times 1^{m}$ test tubes.

Difco nutrient agar, (B 69), was used for streaking cultures and for slants.

I trust that your efforts will meet with more success this time.

Sincerely colvell