

Among these, the condensation product with mono-*t*-butyl-*p*-cresol showed the highest activity, which was superior to commercial 2,6-di-*t*-butyl-*p*-cresol. The condensation products with mono-*t*-butyl-*o*-cresol and mono-*t*-butyl-*m*-cresol, respectively, showed some activities, but other products did not show high activities.

Some of the polynuclear alkylphenols previously reported, that is condensation products of dimethylol-*p*-cresol with *o*-cresol (C-3), thymol (C-5), eugenol (C-6), *p*-hydroxyanisole (C-7), respectively, as well as the condensation product of eugenol and formaldehyde (E-2) were divided into three or four portions with different molecular weights by fractional precipitation with the aid of acetic acid and water. In the case of (C-3), (C-5), (C-6), (C-7), a portion with molecular weight of 300-500 were most effective. An anomaly was observed in the case of (E-2), that is, the monomethylolated product of eugenol, i. e., eugenotine alcohol, m. p. 35°C, was the highest in the activity.

The effects of these compounds to the stability of vitamine A were investigated by G. Katui in the Laboratory of Prof. R. Takata. The results, which coincided with our ones, will be published elsewhere.

16. Polymerisation of Phenyl Glycid Ether and Thioepichlorohydrin

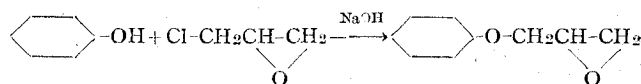
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A. Phenyl Glycid Ether (P.G.E.)

1) Synthesis

P. G. E. was prepared by the method of Boyd and Marle (J. Chem. Soc., 93, 838, (1908)) according to the following reaction:



2) Polymerisation

Exp. No.	Polym. temp. (°C)	Polym. time (hr)	Catalyst (%)	Mol. wt. (Cryoscopic method)
1	120~130	18	NaOH (5)	1357
2	"	79½	" (5)	1844
3	80~90	18½	" (1)	1816
4	120~130	47½	" (1)	1279
5	"	24½	KOH (5)	1531
6	"	14	" (1)	1120
7	"	65	" (1)	923

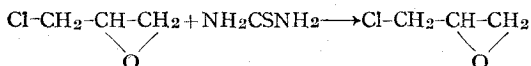
A brown, soft polymer was obtained, when the monomer was treated with molten and powdered KOH or NaOH as catalyst in a sealed tube. It is im-

$$\text{C}_6\text{H}_5\text{OCH}_2\text{CH}_2\text{O} \xrightarrow{\text{Alkali}} \cdots \text{CH}(\text{CH}_2\text{C}_6\text{H}_5)\text{CH}_2\text{OCH}(\text{CH}_2\text{C}_6\text{H}_5)\text{CH}_2\text{OCH}(\text{CH}_2\text{C}_6\text{H}_5)\text{CH}_2\text{O} \cdots$$

As the end group of P.G.E. polymer is thought to be OH-group, we tried to prepare a non-ionic surfactant by reacting ethylene oxide upon the polymer.

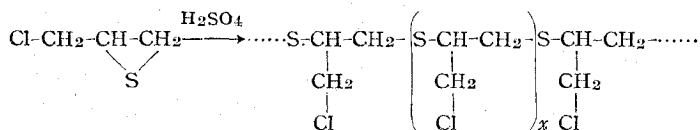
	1%	0.5	0.2	0.1	0.05	0.02	0.01	pure water (16°C)
dyne/cm	51.6	51.9	52.3	53.2	53.9	55.9	58.7	73.34

1) Synthesis



2) Polymerisation

The polymerisation can be considered to proceed as follows:



On the other hand, gradual polymerisation at ordinary temperature resulted a colourless, transparent, soft resin.

Calcd.	32.72%;	Found	32.68%
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(51)

Resin	Aniline	CHCl ₃	React. time.	React. temp.	Product
2.7g.	11g.	20g.	2½ hr.	b. p. of CHCl ₃	3.6g.

According to the above details, light yellow and brittle resin was formed. Theoretical value of N contents calculated from the weight which increased in the product was 6.11%, and analytical value was 6.14%. From this result, it is apparent that 63% of Cl was replaced by 'aniline.

17. Studies on Cobalt Metabolism of *Bacillus subtilis* by Using Co⁶⁰

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Among many microbes, *Streptomyces griseus* and *Bacillus subtilis* are reported to be the good producers of vitamin B₁₂. Microorganisms are expected that they absorb cobaltous ion from the culture media and synthesize this vitamin. These investigations were undertaken with the intention of explaining physiological functions of vitamin B₁₂ and other cobalt containing substances for microorganisms.

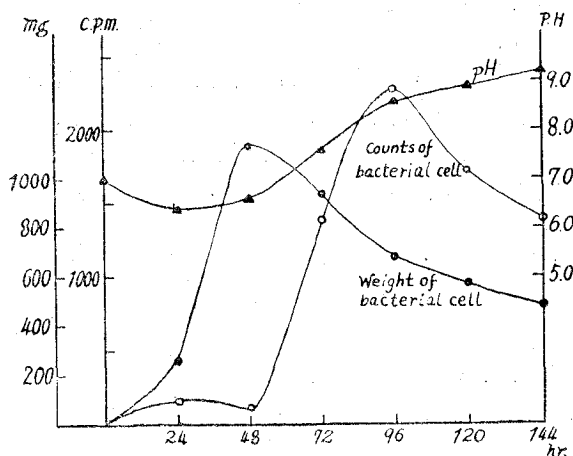


Fig. 1. Absorption of Co⁶⁰ by *bacillus subtilis* (in 250 ml. medium).

The chemical constituents of *bacillus subtilis*, grown on Waksman's media containing 2 p. p. m. radioactive Co⁶⁰, and of the remaining media were precisely analysed during the growth. Total cobalt content in the unit weight of bacterial cells was found to increase with the growth. Co⁶⁰ in the 75 % alcoholic extract, however, showed the maximum after 48 hr. of inoculation. The amount of vitamin B₁₂ in bacterial cells and remaining media were also estimated by means of colorimetric determination of cyanide in cyanoco-