# SCIENCE

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## MEDALS OF THE NATIONAL ACADEMY OF SCIENCES<sup>1,2</sup>

## MEDAL PRESENTATIONS: REMARKS BY THE PRESIDENT

ANNUALLY at this time presentation of medals and awards is made by the academy. These are bestowed by formal vote of the academy acting on the recommendations of special committees charged with the responsibility of administering the several trust funds.

Normally the occasion of the bestowal is at a dinner to which the academy invites a number of distinguished men and women to be guests. This year, because of the exigencies of war, the dinner is necessarily confined to the members and the medalists. Although, therefore, the setting for the presentations this evening lacks the brilliance of former occasions, I can assure our medalists that it lacks nothing in the esteem in which the academy holds them and their work.

<sup>1</sup> Meeting at Washington, D. C., April 26 and 27. <sup>2</sup> The presentation of medals was made at the annual dinner, April 26. To-night we have five medals to bestow. Two of them are governed by the same trust fund and under its terms are in recognition of distinguished work published in specific years. With some of the medals go monetary awards and diplomas. The medalists will be presented by the chairman or a representative of the trust fund committee which made the nomination to the academy.

The medals will be awarded in the order of establishment of the trust funds.

The first award is that of the Henry Draper Gold Medal to Ira Sprague Bowen, and the citation is: "in recognition of his contributions to astronomical physics; more especially his researches on the spectra and chemical composition of the gaseous nebulae." Dr. S. A. Mitchell will present the medalist.

The Draper Fund was established on April 13, 1883, by deed of gift from Mary Anna Palmer Draper (\$10,000 and die), and the deed of establishment provides interest and income to be used for the medal to

	cmm CO <sub>2</sub> liberated in 1 hour
Serum + Ringer solution + Acetylcholine	250
Serum + $NaHCO_8$ + Acetylcholine	325
Serum + Ringer solution + Acetylcholine	
$+ \operatorname{CaCl}_2(1/500 \text{ molar})$	146
Serum + NaHCO <sub>8</sub> + Acetylcholine	
$+ \operatorname{CaCl}_2(1/500 \text{ molar})$	234

It appears that the addition of calcium chloride prevents some of the carbon dioxide from leaving the solution. To prove, that the smaller amount of  $CO_2$ found in the presence of calcium chloride was not due to an inhibitory effect of calcium chloride upon the choline-esterase, but to a methodical error, the acetylcholine was replaced by acetic acid in some determinations:

n 10 minut	tes
Serum + NaHCO <sub>3</sub> + N/100 acetic acid 129	
$Serum + NaHCO_{3} + N/100$ acetic acid	
+0,1 cc CaCl <sub>2</sub> molar 120	
$\text{Serum} + \text{NaHCO}_{a} + \text{N}/100$ acetic acid	
+0.2 cc CaCl <sub>2</sub> molar 95	
$\text{Serum} + \text{NaHCO}_{3} + \text{N}/100$ acetic acid	
+0,2 cc CaCl <sub>2</sub> 1/100 molar	
$Serum + NaHCO_{a} + N/100$ acetic acid	
+0,2 cc CaCl <sub>2</sub> 1/1000 molar	

It follows from the above experiments that the titration procedure gives more reliable results, but it has the disadvantage that it can only be used in clear and almost colorless solutions.

Since the manometric method requires a Warburg apparatus we attempted to develop a more rapid procedure for the determination of the activity of cholineesterase in colored solutions. The following was found to be satisfactory: The serum, a NaHCO<sub>3</sub> solution and the acetylcholine solution were placed in the outer chamber of a Conway-jar<sup>3</sup> and a N/10 Ba(OH)<sub>2</sub>solution in the inner part. The total Ba(OH)<sub>2</sub> solution, or 0,5 cc was taken out 40 minutes after the Conway-jar was covered with the glass lid, and the quantity of unreacted Ba(OH)<sub>2</sub> was titrated with N/100 acetic acid. The values were satisfactory and in agreement with the results obtained by the titrimetric method mentioned above.

А.	SACK	
E.	А.	Zeller

BÜRGERSPITAL, BASEL, SWITZERLAND

## CHEMICAL STUDIES ON CRYSTALLINE BARIUM ACID HEPARINATE

WE have obtained analytical data for the crystalline barium acid salt of heparin (barium acid heparinate)

<sup>§</sup> E. J. Conway, "Micro-Diffusion Analysis and Volumetric Error." in comparison with those obtained for the neutral and acid salts of mucoitinsulfuric acid and chondroitinsulfuric acid. The following molar ratio was found for barium acid heparinate. Anhydrohexosamine : anhydrohexuronic  $acid : SO_3 : Ba = 2.0 : 1.9 : 6.0 : 3.0.$ Further, N:S:Ba=2:6:3. Thus all barium is attached to ester sulfate and the carboxyl group of the uronic acid component is free. Summation (89 per cent.) of the above data, in comparison with the high summation (96 per cent.) obtained for the neutral sodium salt of chondroitinsulfuric acid, does not exclude the possible presence of another constituent. d-Glucosamine was identified (as d-glucosamine hydrochloride) in the hydrolyzate of the crystalline barium acid heparinate. Sodium heparinate (purified through the crystalline barium acid salt) consumes one mole (per 1,200 equivalent weight) of periodic acid.

The amino group of the *d*-glucosamine component of barium acid heparinate is not acetylated and is not free. Barium acid heparinate loses its anticoagulant potency on repeated crystallization from warm, dilute acetic acid. This change is accompanied by the appearance of a free amino group in the molecule, no sulfate is lost and the material is still stained with toluidine blue. Thus neither sulfate content nor toluidine blue staining power are true criteria of heparin activity.

Crystalline barium acid heparinate is also biologically inactivated by prolonged drying at elevated temperatures and by treatment ("Roche heparin" used in this experiment) with weakly ammoniacal hydrogen peroxide, the latter reaction resulting in appreciable sulfate loss.

Full details will be communicated at a later date.

M. L. WOLFROM D. I. WEISBLAT<sup>1</sup> R. J. MORRIS C. D. DEWALT J. V. KARABINOS JAY MCLEAN

DEPARTMENT OF CHEMISTRY,

THE OHIO STATE UNIVERSITY

<sup>1</sup> Hoffmann-La Roche post-doctoral fellow of The Ohio State University Research Foundation.

### BOOKS RECEIVED

- BLATT, A. H. Organic Syntheses. Collective Volume II. Illustrated. Pp. ix + 654. John Wiley and Sons. \$6.50.
- HILL, GERALD F. Termites (Isoptera) from the Australian Region. Illustrated. Pp. 479. Commonwealth of Australia.
- MACHOVER, SOLOMON. Cultural and Racial Variations in Patterns of Intellect. Pp. 91. Bureau of Publications, Teachers College, Columbia University. \$1.60.
- REYNIERS, JAMES A. Micrurgical and Germ-Free Methods, Their Application to Experimental Biology and Medicine. Illustrated. Pp. xiv + 274. Charles C Thomas. \$5.00.

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