

DEPARTMENT OF HEALTH, EDUCATION, AND WELFARE

CENTER FOR DISEASE CONTROL  
ATLANTA, GEORGIA

SUMMARY MINUTES OF MEETING

October 17-18, 1974

The Immunization Practices Advisory Committee met in Atlanta,  
October 17-18, 1974.

COMMITTEE MEMBERS PRESENT

Dr. David J. Sencer, Chairman  
Dr. H. Bruce Dull, Executive Secretary  
Dr. Elizabeth Barrett-Connor  
Dr. Lonnie S. Burnett  
Dr. William R. Elsea  
Dr. Alexander D. Langmuir  
Dr. E. Charlton Prather  
Dr. Gilbert M. Schiff  
Dr. Eleanor G. Shore

Ex Officio

Dr. Harry Meyer, Bureau of Biologics, FDA, DHEW

Liaison (American Academy of Pediatrics)

Dr. Samuel Katz

OTHERS PRESENT

Dr. Bennett Elisberg, Bureau of Biologics, FDA, DHEW  
Dr. Frank Ennis, Bureau of Biologics, FDA, DHEW  
Dr. Saul Krugman, New York University School of Medicine  
Dr. John Robbins, Bureau of Biologics, FDA, DHEW  
Dr. Morris Schaeffer, Bureau of Biologics, FDA, DHEW

STAFF PRESENT

Bureau of Epidemiology:

Dr. Philip Brachman  
Dr. John Bryan  
Dr. Lawrence Corey  
Dr. Eugene Gangarosa  
Dr. Michael Gregg  
Dr. John Harris  
Dr. Michael Hattwick

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I Bureau of Epidemiology:

(continued)

- Dr. Charles Hoke
- Dr. Jay Jacobson
- Dr. Richard Kaslow
- Mrs. Elizabeth Paz
- Dr. David Snyderman
- Dr. Paul Walter

B Bureau of Laboratories:

- Dr. Walter Dowdle
- Dr. Robert Ellis
- Dr. Gary Noble
- Dr. Morris Suggs

B Bureau of State Services:

- Dr. John Modlin
- Dr. Robert Rosenberg
- Dr. John Witte

The meeting was called to order at 8:30 a.m. by Dr. David J. Sencer, Director, Center for Disease Control, the Committee Chairman. He announced a plan under development for the Center to assemble an expert committee to re-review all cases of oral poliomyelitis vaccine-associated disability. The purpose of such in-depth review will be to reassess the continuing suitability of technical criteria used to judge the possible relationship of poliomyelitis vaccine to paralytic illness.

Dr. Sencer also asked the Committee to prepare to review the data on BCG vaccine effectiveness and the vaccine's utility in tuberculosis prevention and control activities in the United States. Such a review will be placed on the agenda of the next ACIP meeting (winter 1975) and will involve not only the Committee membership but invited consultants as well.

Dr. Sencer asked the Executive Secretary to assume the chair for the remainder of the meeting.

OPEN SESSION

Influenza

Influenza in the United States in 1973-74 was reported fully in the spring meeting and was briefly summarized indicating the type B influenza pre-dominance in the United States with some localized outbreaks of type A influenza later in the winter-spring 1974 season. International influenza B predominated although some considerable type A outbreaks were also observed. No influenza outbreaks have been reported in the United States or in other countries in the recent months.

Influenza viruses isolated from individual cases of illness in various parts of the world during the 1973-74 season and since then have shown some continuing antigenic variation consistent with the expected drifts away from predominant viruses. Type A influenza virus strains continue within the general family of viruses seen in recent years but are less like the Hong Kong, England, and Port Chalmers prototypes than was so in the year ago.

A followup was presented on the live influenza vaccine trials being conducted collaboratively by the Center. General results indicated that the attenuated strain being employed produced minimal adverse effects, namely occasional nasal stuffiness, infrequent virus shedding, non-transmissibility of vaccine virus, and an approximately 70-80 percent antibody response among vaccinees. A continuation of the program is planned in order to investigate specifically the effectiveness of the product.

Dr. Meyer and Dr. Ennis described a recent BoB-sponsored vaccine workshop for purposes of analyzing data on the effects of endotoxin in influenza



vaccine. In general, it appears that endotoxin and influenza antigens may lead to enhancement of vaccine reactogenicity not explainable by the simple additive effects of the two substances. Further studies are underway. New regulations to limit the amount of endotoxin permissible in influenza vaccines are being developed in order to minimize their undesirable effects.

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Influenza vaccine prepared for 1974-75 when given to industrial workers in one U.S. locale produced a higher rate of local and systemic reactions than expected. Preliminary data indicate that 25 percent or more of adults receiving the vaccine complained of considerable systemic and local reaction. The implicated lot of vaccine is under continuing re-examination but data from laboratory studies indicate it to be consistent with other lots from the same producer. The Committee concluded that these specific observations on reactogenicity appeared to be atypical for reasons as not yet clearly understood. Additional data are being collected on experiences administering the 1974-75 influenza vaccines to patients, industrial groups, and other groups.

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Traveler Immunization

Committee members were provided with a recently published Supplement to the Morbidity and Mortality Weekly Report dealing with immunization for foreign travel incorporating many ACIP concepts.

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Associated with traveler immunization practices, the Committee asked to consider the present management of yellow fever immunization in the United States through designated centers — a procedure developed primarily to assure the viability of administered vaccine. Requests from individual physicians or clinics for authorization to administer yellow fever vaccine are generally granted on the number of immunizations to be performed.

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After review of current technical issues, the Committee favored facilitation of yellow fever immunization and encouraged the Center to review its practices with respect to authorizing yellow fever vaccination centers. The Committee also recommended that the Bureau of Biologics explore the feasibility of producers' providing single dose vials of yellow fever vaccine. It suggested that efforts be made to assess the suitability of a more liberal policy by conducting periodic surveys of the experience of vaccinees receiving vaccine from large clinics, individual physicians, and other providers.

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Cholera in Europe and Guam

In early spring 1974, cholera was first identified in Portugal. The initial introduction into the country has not been established, although it is likely that it was by means of contaminated shellfish. Cases had been reported from various parts of the country by early May. General spread has been suspected

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to relate variously to contaminated fruits, vegetables, and sh  
Officially, 2,200 cases have been reported and more are continu  
Case identification and therapy have been prompt and effective.  
has been low among hospitalized patients. Cholera has now exte  
Portugal to the Madeira and Cape Verde Islands, the farthest we  
current pandemic of cholera has moved. Serotype of cases has b  
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In Guam, during July 1974, cholera was diagnosed among a small  
construction workers, the initial case, a fatality, diagnosed a  
of postmortem examination. This outbreak appeared to result fr  
source exposure, most likely home-prepared salted, raw fish. T  
cases were generally mild. Environmental studies revealed exte  
tamination.

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CLOSED SESSION

The Committee reviewed with Dr. Morris Schaeffer, Dr. Harry Mey  
Dr. Saul Krugman the progress of the Bureau of Biologics' Viral  
Panel which has been reviewing safety, efficacy, and labeling o  
vaccines. Dr. Krugman, panel chairman, reviewed progress and i  
areas in which regular interaction with the ACIP would be desir  
Questions raised by the panel were discussed; some were schedul  
additional review and discussion.

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OPEN SESSION

CDC Immunobiologics Activities

CDC's Immunobiologics Activity was reviewed, noting the program  
in 1965 and its growth to the present time when 25 different pr  
are provided for special-risk persons or the management of unus  
conditions. Dr. Ellis discussed the mechanics of product distr  
described that the principal emergency products are located in  
airport sites for ready availability.

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Data are being collected on botulinum toxoids following observa  
reduced potency of earlier products. Although antitoxins are a  
for therapy, laboratorians continue to request toxoid immunizat  
to potential exposure. The Committee encouraged continued eval  
the toxoids. No cases of lab-acquired botulism have ever been

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Dr. Ellis reported receiving numerous requests for Schick test  
primarily to study immunologic competency in patients suspected  
deficiency states. The sources of such material are greatly dep  
the Committee generally recommended that efforts be made to stir

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sufficient supply of Schick testing material not only to meet the needs for immunological determinations but also to conduct surveys on the adequacy of diphtheria protection. This latter effort is especially germane in view of localized outbreaks occurring in various parts of the country during the past few years.

Zoster Immune Globulin (ZIG)

The Committee was apprised of continuing efforts by the Center to establish the effectiveness and dosage of ZIG. The Committee noted the differences in judgments on the relative value of ZIG in preventing chickenpox among patients with immunodeficiency states and encouraged that effort to be made to establish objectively the benefits and proper dosages.

Rabies Immune Globulin

Rabies Immune Globulin was licensed in the summer of 1974 and has become generally available since September. It appears that the supplies sufficient and that additional lots of the globulin will become available earlier this fall. Plans have been made for evaluating need in the event that demand exceeds supply. Dr. Hattwick reviewed data previously presented on the use of HRIG and duck embryo vaccine (DEV) which indicates the need for 21 doses of y vaccine plus 2 boosters to produce antibody response among patients who receive vaccine and globulin.

Meningococcal Meningitis

Dr. Jacobson reviewed the epidemic of meningococcal meningitis which has been occurring predominantly among children in Sao Paulo and other areas of Brazil since 1971. Although in the early epidemic, cases were caused by sulfamycin resistant serogroup C meningococci, since spring 1974, the majority of cases have been serogroup A. The overall rate of illness based on international information provided in the summer of 1974 is an incidence of 65 cases per 100,000 population per month.

This is the most serious urban outbreak of meningococcal meningitis ever reported.

As previously reported to the Committee, the Brazilian Ministry of Health has instituted a polysaccharide vaccine campaign for purposes of disease control and also of evaluating its relative effectiveness. In general, data on the serogroup C polysaccharide vaccine administered to school children indicates poor antibody response and lack of protective efficacy. Serogroup A vaccine tested in children in Africa and other parts of the world shows approximately 90 percent effectiveness from one dose. Its use in Brazil is anticipated to provide an opportunity for further evaluation.

Dr. Jacobson presented a theoretical analysis of the cost/benefits of using an effective meningococcal polysaccharide vaccine in countries with only endemic meningococcal disease. The Committee encourages refinement of this approach as part of a general exploration of planning for the appropriate use of polysaccharide vaccines in routine preventive medical practice.

Dr. John Robbins reviewed the general status of polysaccharide vaccines and current production capabilities for such chemically specific products. He pointed out that in order to provide a sufficient supply of meningococcal polysaccharide vaccines would require substantial effort among commercial laboratories capable of carrying out the chemical processes. Such production efforts would likely occur only with a specified national policy

#### Hepatitis-B in Dentists

Although not yet described in the public health literature, there have been several clusters of hepatitis-B cases occurring in association with dental practice. Mechanisms of spread among dental patients or between dentists and patients have not been clearly established. Methods of control, therefore, remain obscure. Efforts are being made to study intensively the epidemiological importance of hepatitis-B antigen carriers in order to apply the data to the potential risk of infection in dentistry. The Committee urges that such data be collected and that management of potential risks be related to known hazards and not to hypothetical risks.

#### Hospital Personnel Immunization

Dr. Kaslow reviewed the status of a CDC report for hospital personnel incorporating general recommendations of the ACIP on immunization. He reviewed specific questions of the need for and suitability of influenza vaccine, mumps vaccine, polio vaccine, and rubella vaccine for susceptible hospital employees. Committee comments and general reactions will be incorporated into a further draft of the report to be distributed for review.

#### Other Business

As a special item of business, Dr. Gangarosa presented data on the comparability of intradermal cholera vaccine to that subcutaneously administered. In general, findings were of approximately equivalent protective value but with substantially fewer local or systemic reactions. Antibody levels and duration of immunity information suggest that the intradermal route is not as good, possibly because of antigen mass. It was pointed out that although the intradermal vaccination could have merit in terms of lowered bacteriogenicity, it might not be acceptable to the quarantine authorities of countries requiring

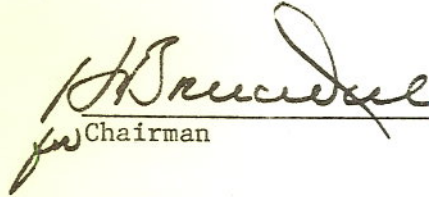


cholera vaccine certification from travelers. Furthermore, the committee felt that if dosage were to be changed, additional information would be required. It recommended that no action be taken at the present time.

The Committee concluded its meeting by selecting potential dates for its winter session. Either January 15-16 or 29-30 appeared to be suitable. Committee members intend to check calendars and designate preference.

The meeting was adjourned at approximately 4:30 p.m.

I hereby certify that, to the best of my knowledge, the foregoing summary of minutes is accurate and complete.

  
Chairman

11/8/74  
Date