ELIZABETH PRINGLE introduced by DR. MARY WILKERS (London). ‘A Study of 112 Children with Coeliac Disease from 1950–69.’ (To be published elsewhere.)

E. RUSSELL (Graz). ‘Precipitins to Aquous Extracts of Flour in Coeliac Disease.’ In certain instances it may be difficult to differentiate between coeliac disease and the ‘coeliac syndrome’. An essential diagnostic criterion in coeliac disease is the disappearance of symptoms on a gluten-free diet, and a relapse caused by reintroduction of gluten into the diet. Normally a long period of observation is necessary to confirm the diagnosis in this way. We have been able to develop a sensitive method of detecting precipitins to aqueous and alcoholic extracts of flour from wheat, barley, oats, and rye in the serum of patients with coeliac disease in 15 patients. The diagnosis of coeliac disease was confirmed by disappearance of the symptoms on a gluten-free diet and by a relapse after reintroduction of gluten into the diet. All of these patients had precipitating antibodies to aqueous extracts of wheat and barley; 13 also showed precipitins to extracts of oats, and 10 to extracts of rye. It was proved that the precipitating antibodies to the aqueous extracts of flour belonged to the IgG group. Sera of 60 children suffering from parenteral diarrhoea, Esch. coli enteritis, disaccharide intolerance, and mucoviscidosis, and pooled serum of 100 healthy adult blood donors all served as controls. In none of these controls could precipitins to aqueous extracts of the cereals be detected. The determination of these antibodies seems to be of value as a diagnostic tool in coeliac disease and could help to unravel the immunological mystery of this disease.

R. A. RISDON introduced by PROFESSOR A. E. CLAIREAUX (London). ‘Renal Dysplasia—A Clinico-pathological Study of 76 Cases.’ (To be published.)

J. MARTIN and P. P. RICKHAM (Liverpool). ‘Pulmonary Metastases in Wilms’ Tumour—Treatment and Prognosis.’ (Archives of Disease in Childhood 45, 805.)


J. A. DUDGEON (London). ‘Prevention of Congenital Rubella Defects. The Need for Long-term Surveillance.’ Several attenuated rubella vaccines have now been developed. Two have already been licensed abroad and one is likely to be licensed in Great Britain early in 1970.

At the autumn meeting of the BPA we presented data of our comparative trials with three of the attenuated vaccines, and outlined a programme for immunization against rubella in schoolgirls aged 10–14 years in the same year tuberculin testing and BCG vaccination are carried out.

The results of vaccine trials now being undertaken on approximately 1,000 schoolgirls will be presented. If rubella vaccine is used on a sufficiently wide scale in the next few years, there should be a decline in the incidence of rubella defects. These can now be fairly readily identified in the first year of life on clinical grounds, and laboratory tests, such as virus isolation, serology, and immunoglobulin levels can provide confirmatory evidence. Defects such as perceptive deafness not recognizable at birth will in many instances be picked up at a school medical examination. A plan for long-term surveillance will be outlined in which it is hoped paediatricians, Medical Officers of Health, and School Medical Officers will co-operate.

L. HOHENAUER (Innsbruck). ‘Calcium and Phosphorus Homeostasis during the First Days of Life.’ (To be published.)

G. J. I. SNODGRASS introduced by DR. L. STIMMLER (London). ‘Interruption of Plasma Magnesium, Calcium, and Phosphorus for 4 Milk Regimens.’ (To be published.)

O. THALHAMMER (Wien). ‘Histidinaemia Detected by Newborn Screening using Guthrie’s Method.’ In 36,462 routine Guthrie tests on newborn infants, 3 cases of histidinaemia have been discovered. Initial tests on day 4, 5, and 6 were positive corresponding to 16, 6, and 6 mg/100 ml. In the same series 62 so-called false positive initial tests were in the same range. There was an excellent concordance between the results of inhibition assay and column chromatography. Histidine blood levels were constantly high in all three infants and histidase activity in stratum corneum was zero. Excretion of histidine in the urine was constantly high, but in spite of this, the ferric chloride test was positive in only two of the three infants. In loading tests urinary histidine output was 27%–47% of loading dose and formiminoglutamic-acid was absent in all samples 24 hours after the loading. Blood levels did not return to starting values less than 24 hours after loading. Histidine-low diet brought histidine blood levels and urine excretion to normal within 1 to 2 weeks.

A Seminar on The Paediatrics of Immigration was held on Saturday morning, 25 April, in the Royal Hotel, Scarborough, with Dr. J. W. Farquhar as Chairman, and introduced by Professor T. E. Oppé. The following speakers took part: Dr. Catriona Hood (London), by invitation,—‘Social and Cultural Factors in the Health of Children of Immigrants.’ Dr. G. S. Prince (London), by invitation,—‘Emotional Deprivation and Pseudo-autism.’ Dr. M. W. Arthurton (Bradford)—‘Immigrant Children and the Day-to-Day Work of a Paediatrician.’ Professor C. E. Stroud (London)—‘Some Important Organic Problems.’

WINDERMERE LECTURE. The Lecture was delivered on 23 April by Professor C. Henry Kempe (University of Colorado Medical Center)—‘Paediatric Implications of the Battered Child Syndrome.’ (To be published in Archives in Disease in Childhood, 1971.)

Members and guests of the Association attended a