that their failure to detect a complement-fixation antibody response is evidence that an allergic reaction or a previous infection had not taken place. In our own series (Scott and Gardner, 1970), and in a larger series yet to be published, we have shown that infants of 6 weeks and onwards can respond to infection by the rapid production of RS neutralizing antibodies in their secretions in intervals of even less than 7 days. We find this, and the presence of detectable antibody in the secretion at onset of bronchiolitis in approximately 50% of cases, far more convincing evidence of the responsiveness of young infants to infection and consistent with bronchiolitis being a second infection. As children get older, repeated exposure to RS virus could lead to desensitization by the production of large amounts of specific IgG.

We hope Dr. Ross and her colleagues will extend their studies in the first 3 months so that we can increase our knowledge of what we feel is a critical period for our understanding of pathogenesis of bronchiolitis.

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REFERENCES

We showed the above letter to the authors, whose comments follow:

There are two misinterpretations of our paper in the letter by Gardner and his colleagues. Firstly, we did not claim that CF was as sensitive as neutralization but as reliable as and more convenient than neutralization tests for diagnosis of acute respiratory infections due to RS virus. Secondly, our suggestion that bronchiolitis in children under 3 months was probably not a secondary infection was not based on ‘failure to detect’ a complement fixation antibody response but on the lack of ‘boost’ responses in this youngest group.

In the paper cited in their letter (Scott and Gardner, 1970), only 30 patients were studied and from only 10 was a second secretion obtained; in only 4 of these 10 was there a fourfold or greater rise in neutralizing activity. This seems to afford no proof of secondary as opposed to primary infection.

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Growth Pattern in a Boy with Thyroxine-binding Globulin Deficiency

Sir,

We read with interest the article by J. L. Penfold, G. M. Kneebone, M. Wellby, and R. K. Oldfield, on ‘Growth Retardation and Thyroxine-binding Globulin Deficiency’ in the February 1971 issue (p. 115). While they document short stature in terms of a single height measurement, they make no reference to the more important parameter of growth; namely, how fast is the patient growing at this time? We have followed for 6 years the growth pattern in a boy first diagnosed at 13 months of age. No other such longitudinal study has been reported previously.

Case Report

N.F. is the only child in the family. Pregnancy and delivery were normal. Birthweight at term was 2608 g. Jaundice was present for 3 weeks but no cause was found. At 10 months of age, he was found to have dry eczema and no dental eruption but no other abnormal signs. No goitre was present. PBI was 1·1 μg/100 ml. He was, therefore, admitted for further study at the age of 13 months. The bone age at the left wrist was 6 months. Serum cholesterol was 85 mg/100 ml. 132 uptake was 24·4% and 19% at 3 and 5 hours. Total serum plasma protein was 7·5 g/100 ml. Electrophoresis of thyroxine binding proteins showed absence of TBG, slight increase in thyroxine binding albumin (TBA), and marked increase in thyroxine binding prealbumin (TBPA) (Fig. 1.) Because of the normal radioactive iodine uptake and the apparent clinical euthyroid state, no therapy was given. The patient was followed at intervals. His growth in height and weight are shown in Fig. 2. At 2 years 3 months, he had 16 teeth and was saying a few words. Bone age was 16 months. At 4 4/12 years, his bone age was 3½ years. The patient was now shown to be deaf in the right ear only, for which there was no apparent cause. At 4 5/12 years, some thyroid studies were repeated. i21 uptake was 31% and 33%, at 3 and 4½ hours. T3 uptake was 75% (N = 70 to 140%). PBI was 2·3 μg/100 ml (N = 4 to 8). Free thyroxine index was 17·2. These results were considered to be consistent with the earlier diagnosis of low or absent TBG. When last seen at the age of 6 10/12 years, the patient had normal dental age and a bone age of 6½ years. His mental development has remained normal. Though no therapy was instituted, the patient has grown at a constant rate each year and has followed a pattern which is parallel to but just below the third centile line. His bone age though initially delayed is now within normal limits. He did show early clinical features which might have suggested hypothyroidism, but subsequent tests, follow-up, and clinical features have excluded this diagnosis. The other 11 previously reported cases or families (some patients over 50 years