Haemoglobin Ocho Rios (β52 (D3) Aspartic Acid→Alanine): A New β-Chain Variant of Haemoglobin A Found in Combination with Haemoglobin S

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During a routine study of the parents of a 3-month-old Jamaican child with the sickle-cell trait the father was found to have a haemoglobin electrophoretic pattern identical to that of homozygous sickle-cell disease. Because he was perfectly well and had no haematological abnormalities detailed studies of his haemoglobin constitution were undertaken. These indicate that he is heterozygous for both haemoglobin S and a previously undescribed β-chain haemoglobin variant.

Methods

Haemoglobin analysis was carried out by electrophoresis on cellulose acetate (Kohn, 1969), starch gel using a TRIS-EDTA-borate system, pH 8-5 or phosphate system, pH 7-0 (Weatherall, 1965), agar gel (Robinson et al., 1957; Marder and Conley, 1959) and starch block using a phosphate buffer system, pH 6-5 (Weatherall, 1965). Haemoglobin A2 levels were estimated by the modification of the method of Marengo-Rowe (1965) described by Weatherall et al. (1971). Alkali resistant haemoglobin levels were determined by the methods of Singer, Chernoff, and Singer (1951) and Betke, Marti, and Schlicht (1959). Blood films were also examined for the presence of fetal haemoglobin-containing cells by the method of Kleihauer, Braun, and Betke (1957). Preparation of globin, separation of globin chains on carboxymethyl (CM)-cellulose columns in 8M urea, isolation of purified chains, fingerprinting, and amino-acid analysis followed the methods described by Clegg, Naughton, and Weatherall (1966).

Results

Clinical and Haematological Investigation. The propositus was a 3-month-old Jamaican Negro child with the sickle-cell trait and no haematological abnormalities. The child's mother and father were available for study. The mother's blood picture and haemoglobin electrophoretic pattern were completely normal. The father was a 35-year old Negro who had been born in Ocho Rios on the North Coast of Jamaica. He had no history of illness and clinical examination was completely normal. His haemoglobin level was 15-4 g/100 ml, packed cell volume 45%, and reticulocyte count 2%. The peripheral blood film showed occasional target cells and, on incubation of his blood with coliform organisms, the red cells sickled.

Haemoglobin analysis. Electrophoresis of the father's haemoglobin on cellulose acetate (Kohn, 1969) revealed two components; a major fraction migrating in the position of haemoglobin S and a minor fraction migrating in the position of haemoglobin A2. The haemoglobin A2 value obtained by a quantitative cellulose acetate electrophoresis was 3.3%. This is at the upper limit of the normal adult range (Weatherall et al., 1971). Electrophoresis on agar gel, pH 6-0 (Robinson et al., 1957) or pH 5.9 (Marder and Conley, 1959), revealed two major components; one in the position of haemoglobin S and the other, of almost equal concentration, close to the origin. Starch gel electrophoresis, pH 8-5, of the father's haemolysate gave a pattern identical to that of homozygous sickle-cell anaemia with haemoglobins S and A2 only. No haemoglobins F or A were present. Starch gel electrophoresis, pH 6-5, and starch block electrophoresis, pH 6-5, gave only a single component and the second fraction seen on agar gel electrophoresis was not resolved. There was no increase in alkali resistant haemoglobin by either the method of Singer et al. (1951) or Betke et al. (1959) and none of the red cells showed resistance to acid elution.

Globin was prepared from the father's haemolysate without prior purification and the constituent globin chains examined by CM-cellulose chromatography in 8M urea, pH 6-7. Two globin chain
fractions were isolated, one eluting in the position of the β-chain of haemoglobin S and the other in the position of a normal α-chain. The β-chain was aminoethylated, digested with trypsin, and fingerprinted. When compared with a fingerprint of aminoethylated β-chain three abnormal peptides were noted. One was in the known position of haemoglobin β81 and had the expected amino-acid composition. The normal β1 peptide was also present at a similar concentration to the β81 peptide. The other two abnormal peptides were slightly more basic than, and had very similar chromatographic mobilities to, βA5 and βA5ex, both of which were also seen on the fingerprint. Amino-acid analysis of the two abnormal peptides gave compositions identical to those of βA5 and βA5ex except that there was an additional alanine residue and one less aspartic acid residue in each case. Peptide β5 has one residue of asparagine and two of aspartic acid and since the haemoglobin variant is basic relative to haemoglobin A the alanine substitution must involve one of the two aspartic residues rather than asparagine.

A larger sample of the abnormal β5 peptide was therefore isolated from a trypptic digest of the abnormal β-chains and digested with chymotrypsin. The resulting digest was fractionated by electrophoresis at pH 6.5 and 3.5 and by chromatography in BAWP (Clegg et al., 1966). A peptide was isolated with the following composition: Ser, Thr, Pro, Ala2, Val, Met, corresponding to residues 49–55 of the abnormal β-chain. In the βA-chain, residue 52 is normally aspartic acid. The substitution in the abnormal variant must therefore be β52 Asp→Ala.

The finding in the father of two haemoglobin components on agar gel electrophoresis and of both β81 and β5 and normal and abnormal β5 peptides in the β-chain obtained by urea chromatography indicate that he is heterozygous for both haemoglobin S and a β-chain variant, β52Asp→Ala. Since the latter haemoglobin has not been previously reported it has been designated haemoglobin Ocho Rios after the place of origin of the father.

**Discussion**

The discrepancy between the clinical and laboratory findings in this case led to the identification of a new haemoglobin variant, haemoglobin Ocho Rios. A similar discrepancy led Konotey-Ahulu et al. (1971) to identify the heterozygous state for both haemoglobins S and Osu-Christiansborg (β52 Asp→Asn) which also has the same electrophoretic mobility as haemoglobin S on paper electrophoresis at pH 9 and starch gel electrophoresis at pH 8.6. Thus the two substitutions which have now been

described at this site both result in a haemoglobin variant which has the same mobility as sickle-cell haemoglobin on electrophoresis by routine screening techniques but which is separable by either agar gel electrophoresis or peptide mapping, from sickle haemoglobin. Both these studies underline the importance of a full structural analysis of the ‘haemoglobin S’ from individuals with ‘benign sickle-cell anaemia’. The presence of a second β-chain variant can be deduced by either agar gel electrophoresis or peptide analysis of the ‘β88-chain’ isolated by CM-chromatography in 8M urea even in the absence of critical family members, as in the present case.

Haemoglobin Ocho Rios is a further example of a β-chain variant which does not cause a clinical disorder when inherited in the heterozygous state with haemoglobin S. Like haemoglobin Osu-Christiansborg the substitution is at an external position in the D helix; the gradual acquisition of information of this type may provide information regarding the complementary sites involved in the stacking of haemoglobin S molecules.

**Summary**

A new human haemoglobin variant, haemoglobin Ocho Rios (β52(D3) aspartic acid→alanine) has been characterized. It was found in an individual also heterozygous for the sickle-cell gene and the presence of the two β-chain variants is not associated with any clinical or haematological disorder.

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**References**


