

# JOURNAL OF MEDICAL GENETICS

*Editor:* ARNOLD SORSBY

*Assistant Editors:* A. C. STEVENSON and JOHN L. HAMERTON

<b>Genetic Study of Sample of 70 Patients with Myasthenia Gravis .. .. .</b>	<b>257</b>
A. JACOB, E. R. CLACK, and A. E. H. EMERY	
<b>Incidence of Fibrocystic Disease in Wessex .. .. .</b>	<b>262</b>
B. D. HALL and M. J. SIMPKISS	
<b>Multiple Sclerosis: Discordance in Three Pairs of Dizygotic Twins .. .. .</b>	<b>266</b>
W. S. CENDROWSKI	
<b>Anhidrotic Ectodermal Dysplasia: Autosomal Dominant Inheritance with Palate and Lip Anomalies .. .. .</b>	<b>269</b>
R. S. RAPP and W. E. HODGKIN	
<b>Diagnostic and Genetical Aspects of Tuberous Sclerosis .. .. .</b>	<b>273</b>
N. C. NEVIN and W. G. PEARCE	
<b>Amino Acid Patterns in Cystinuric Families .. .. .</b>	<b>281</b>
S. KELLY and W. COPELAND	
<b>Testosterone Excretion Rates in Normal Males and Males with an XYY Complement .. .. .</b>	<b>286</b>
B. T. RUDD, O. M. GALAL, and M. D. CASEY	
<b>Familial Microtia with Meatal Atresia in Two Sibships .. .. .</b>	<b>289</b>
L. C. ELLWOOD, S. T. WINTER, and H. DAR	
<b>Hereditary Non-spherocytic Haemolytic Anaemia with Post-splenectomy Inclusion Bodies and Pigmenturia Caused by an Unstable Haemoglobin Santa Ana-<math>\beta</math>88 (F4) Leucine<math>\rightarrow</math>Proline .. .. .</b>	<b>292</b>
R. W. OPFELL, P. A. LORKIN, and H. LEHMANN	
<b>Combination of Hereditary Elliptocytosis and Heterozygous Beta-thalassaemia: A Family Study .. .. .</b>	<b>298</b>
M. AKSOY and S. ERDEM	
<b>Secretor Status in Asthma and Hay Fever .. .. .</b>	<b>302</b>
M. A. DENBOROUGH and H. J. DOWNING	
<b>Incidence of ABO and Rh Blood Groups in Pulmonary Tuberculosis in Different Ethnic Groups .. .. .</b>	<b>306</b>
N. SAHA and B. BANERJEE	
<b>ABO Blood Groups and Vitiligo .. .. .</b>	<b>308</b>
V. N. SEHGAL and B. DUBE	
<b>Further Observations on Kell Blood Groups in Families Ascertained via a Mongol Propositus .. .. .</b>	<b>310</b>
D. A. PRICE EVANS, P. J. J. WREN, W. T. A. DONOHUE, M. F. BULLEN, M. LEWIS, H. KAITA, B. CHOWN, and I. UCHIDA	
<b>Diverse Chromosomal Anomalies in a Family .. .. .</b>	<b>314</b>
L. ATKINS, C. S. BARTSOCAS, and P. J. PORTER	
<b>Case Reports:</b>	
<b>  Congenital Microcephaly with Hiatus Hernia and Nephrotic Syndrome in Two Sibs .. .. .</b>	<b>319</b>
W. H. GALLOWAY and A. P. MOWAT	
<b>  A Large Deletion of Chromosome No. 1 (46,XY,1?-) .. .. .</b>	<b>322</b>
D. AARSKOG	
<b>  X-linked Hydrocephalus .. .. .</b>	<b>326</b>
M. W. SHANNON and H. L. NADLER	
<b>  Multiple Anomalies Associated with a Small Extra Metacentric Autosome .. .. .</b>	<b>329</b>
A. B. MUKHERJEE, M. W. PARTINGTON, N. E. SIMPSON, and K. A. WALMSLEY	
<b>  An Extra Small Metacentric Chromosome in a Mentally Retarded Boy .. .. .</b>	<b>335</b>
J. ISHMAEL and K. M. LAURENCE	
<b>Review Article:</b>	
<b>  Males with an XYY Sex Chromosome Complement .. .. .</b>	<b>341</b>
W. M. COURT BROWN	
<b>Book Reviews .. .. .</b>	<b>360</b>
<b>Index .. .. .</b>	<b>369</b>