

Q H301
J83 VOL. 8, No. 3

SEPTEMBER 1971

JOURNAL OF MEDICAL GENETICS

Editor: C. A. CLARKE

Assistant Editors: CEDRIC CARTER and JOHN L. HAMERTON

Polycystic Disease of Kidneys and Liver Presenting in Childhood	257
HELEN BLYTH and BARBARA G. OCKENDEN	
Polycystic Kidneys Associated with Malformations of the Brain, Polydactyly, and Other Birth Defects in Newborn Sibs	285
K. FRIED, E. LIBAN, M. LURIE, S. FRIEDMAN, and S. H. REISNER	
X-linked Spondyloepiphyseal Dysplasia Tarda: Clinical and Linkage Data	291
R. M. BANNERMAN, G. B. INGALL, and J. F. MOHN	
Haemoglobin Osu-Christiansborg: A New β-chain Variant of Haemoglobin A (β52 (D3) Aspartic Acid\rightarrowAsparagine) in Combination with Haemoglobin S	302
F. I. D. KONOTY-AHULU, JUDITH L. KINDERLERER, H. LEHMANN, and B. RINGELHANN	
Differentiation of Two Genetically Specific Types of Depression by the Response to Antidepressant Drugs	306
C. M. B. PARE and J. W. MACK	
Mutation Rate in Duchenne Muscular Dystrophy	310
H. K. GOSWAMI and B. D. CHAURASIA	
Familial Total Anomalous Pulmonary Venous Return	312
J. E. PAZ and E. E. CASTILLA	
Distribution of ABO Blood Groups, G6PD Deficiency, and Abnormal Haemoglobins in Leprosy	315
N. SAHA, H. B. WONG, B. BANERJEE, and M. O. WONG	
Rh Immunization following Incompatible Blood Transfusion and a Possible Long-term Complication of Anti-D Immunoglobulin Therapy	317
M. E. J. BEARD, J. PEMBERTON, J. BLAGDON, and W. J. JENKINS	
Review: Suppression of the Immune Response	321
JOHN BRADLEY and C. J. ELSON	
Group G Deletion Syndromes	341
ROBERT P. KELCH, MARGERY FRANKLIN, and ROY D. SCHMICKEL	
The Inheritance of a Structural Anomaly of One Chromosome No. 16 in a Kindred (46,16-,C+)	346
JUAN CHEMKE and ARTHUR ROBINSON	
The 13q- Deletion Syndrome	351
ELIZABETH GRACE, J. DRENNAN, DOREEN COLVER, and R. R. GORDON	
Somatic Stigmata of Turner's Syndrome in a Patient with 46,XXq-	358
MAUREEN BOCIAN, EVA KRMPOTIC, KATARINA SZEGO, and IRA M. ROSENTHAL	
Case Reports:	
Klinefelter's Syndrome and G Trisomy	364
B. ERDTMANN, A. A. GOMES DE FREITAS, R. P. DE SOUZA, and F. M. SALZANO	
A Case of Cri-du-Chat Associated with Cataracts and Transmitted from a Mother with a 4/5 Translocation	369
HAROLD GROTSKY, LILLIAN Y. F. HSU, and KURT HIRSCHHORN	
A Prisoner with an Unusual Karyotype (46,XY,Dq-)	372
J. KAHN and FREDA REED	
Partial Deletion of a Group-F (19-20) Chromosome in a Physically Handicapped Male Patient	374
PAUL GENEST, MARC BOUCHARD, and JACQUELINE POTY	
An Extra Small Metacentric Autosome in a Mentally Retarded Boy with Multiple Malformations	378
SALVADOR ARMENDARES, LEONOR BUENTELLO, and FABIO SALAMANCA	
An Extra Small Metacentric Chromosome in Association with Multiple Congenital Abnormalities	381
W. H. FINLEY, S. C. FINLEY, and D. MONSKY	
Trisomy D/Trisomy E Mosaicism in an Infant Male	384
RICHARD J. WARREN and JOHN I. KEITH	
X Chromosome Long Arm Deletion in a Patient with Down's Syndrome	387
FRED W. LUTHARDT and CATHERINE G. PALMER	
Trisomy 18 in One of Fraternal Twins	392
DENNIS GERTZER and GERALD NATHENSON	
Book Reviews	395

BRITISH MEDICAL ASSOCIATION LIBRARY
OCT 2 1971

J Med Genet: first published as on 1 September 1971. Downloaded from <http://img.bmj.com/> on January 21, 2021 by guest. Protected by copyright.