QH301
J83

VOL. 6, No. 2 JUNE 1969

Bio.S.

JOURNAL OF MEDICAL GENETICS

BROWN UNIVERSITY
LIBRARY
JUL 22 1969

Editor: ARNOLD SORSBY

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

Syndrome of Retardation with Urogenital and Skeletal Anomalies (Smith-Lemli-Opitz Syndrome): Clinical Features and Mode of Inheritance 113
L. DALLAIRE

An Association of Rh Type and Gynaecological Disease with Twinning 121
FRANCES V. DE GEORGE

Hypertrophic Pyloric Stenosis: Adult and Congenital Types Occurring in the Same Family 126
CARLOS ZAVALA, ALVARO BOLIO, RAÚL MONTALVO, and RUBÉN LISKER

Inheritance of Coeliac Disease 129
W. M. MCCRAE

Hearing Impairment in Female Carriers of the Sex-linked Syndrome of Deafness with Albinism 132
K. FRIED, M. FEINMESSER, and J. TSITSIANOV

Mongolism and Maternal Menarche 135
J. M. BERG and J. T. R. BAVIN

Consistent and Variable Chromosome Anomalies in Parents of Children with Down's Syndrome 137
J. KAHN and K. ABE

Studies on Glucose-6-phosphate Dehydrogenase Deficient Egyptian Families .. 150
NEMAT HASHEM and ASSMA NOUR

Comparison of Strength of Isoagglutinin Titres in English and Finnish Populations 157
I. J. T. SEPPÄLÄ

XYY Syndrome, and XYY/XXYY Mosaicism also Showing Features of Klinefelter's Syndrome 159
D. A. SPENCER, JULIE W. EYLES, and M. K. MASON

Translocations of D Chromosomes in Two Families: t(13q14q) and t(13q14q) + (13p14p) 166
CATHERINE G. PALMER, PATRICK M. CONNEALLY, and JOE C. CHRISTIAN

Familial Transmission of a Chromosomal Translocation t(2q+;Cp−) 174
CARMEN B. LOZZIO and ANTHONY A. KATTINE

Welshness and Fertility 180
DAVID J. B. ASHLEY

Blood Groups and Lung Cancer 183
DAVID J. B. ASHLEY

A Haptoglobin Johnson Family with Non-hypohaptoglobinaemic Hp^J/Hp^2 187
J. PINTERA, R. DVOŘÁK, J. VACL, and J. FIŠER

Further Evidence for Close Linkage of the Hb^β and Hb^δ Loci in Man 190
MONA K. MISHU and WALTER E. NANCE

Case Reports:

Familial Jaw Cysts in Charcot-Marie-Tooth Disease 193
M. R. SWIFT and S. L. HOROWITZ

Postaxial Polydactyly in Three Indian Families 196
J. MOHAN

Partial Trisomy in a Child with Features Suggesting Mongolism 201
H. K. KUMBNANI and R. A. PFEIFFER

46,XY,21qi/46,XY,21p− Mosaicism in a Child with Down's Syndrome 206
LEONARD ATKINS and MURRAY FEINGOLD

Double Autosomal Trisomy and Mosaicism for Three Cell Lines in Man. Co-existing Trisomy 13–15, Trisomy 17–18, and a Minority Cell Line Trisomic for a Chromosome of Both Groups 209
O. MARGARET GARSON, A. G. BAIKIE, JEAN FERGUSON, and C. H. GREER

A Patient with a Short Arm Deletion of Chromosome 18 (46,XY,18p−) 216
LESTER WEISS and K. MAYEDA

A Patient with 45,XX,G−/46,XX,Gr Mosaicism 220
C. ERIC BLANK and JOHN LORBER

Congenital Malformations Associated with a Ring 4 Chromosome 224
RODNEY CARTER, ELIZABETH BAKER, and DAVID HAYMAN

Book Reviews 228

BRITISH MEDICAL ASSOCIATION · TAVISTOCK SQUARE · LONDON W.C.1

YEARLY SUBSCRIPTION (4 NUMBERS) INLAND £3 15s. ABROAD £4 4s. SINGLE NUMBER 22s. 6d.

J Med Genet: first published as on 1 June 1969. Downloaded from http://jmg.bmj.com/ on April 16, 2024 by guest. Protected by copyright.