

1. Record Nr.	UNISA990003555620203316
Titolo	Partenze-ritorni : italiani in America Latina / a cura di Fiorenza Tarozzi e Roberto Vecchi
Pubbl/distr/stampa	Bologna, : Clueb, 1996
ISBN	88-8091-469-3
Descrizione fisica	160 p., [12] p. di tav. : ill. ; 24 cm
Disciplina	909.82
Soggetti	Storia contemporanea - Sec. 20
Collocazione	X.3.B. 6143
Lingua di pubblicazione	Italiano
Formato	Materiale a stampa
Livello bibliografico	Monografia
2. Record Nr.	UNINA9910790678803321
Autore	Stevenson Roger E. <1940->
Titolo	Atlas of X-linked intellectual disability syndromes // Roger E. Stevenson, Charles E. Schwartz, and R. Curtis Rogers
Pubbl/distr/stampa	New York : , : Oxford University Press, , [2012] ©2012
ISBN	0-19-997524-8 0-19-981186-5
Edizione	[Second edition.]
Descrizione fisica	1 online resource (363 p.)
Altri autori (Persone)	SchwartzCharles E RogersR. Curtis <1953-> (Richard Curtis) StevensonRoger E. <1940->
Disciplina	616.85/88042
Soggetti	X-linked mental retardation
Lingua di pubblicazione	Inglese
Formato	Materiale a stampa
Livello bibliografico	Monografia
Note generali	Rev. edition of: X-linked mental retardation / Roger E. Stevenson, Charles E. Schwartz, Richard J. Schroer. 2000.
Nota di bibliografia	Includes bibliographical references and index.

Cover; TABLE OF CONTENTS; FOREWORD; PREFACE; AARSKOG SYNDROME; ABIDI SYNDROME; ADRENOLEUKODYSTROPHY; AGENESIS OF THE CORPUS CALLOSUM, X-LINKED; AHMAD SYNDROME; AICARDI SYNDROME; ALLAN-HERNDON-DUDLEY SYNDROME; ALPHA-THALASSEMIA INTELLECTUAL DISABILITY (SEE ALSO ATRX-ASSOCIATED XLID); AP1S2-ASSOCIATED XLID; APAK ATAXIA-SPASTIC DIPLEGIA SYNDROME; ARMFIELD SYNDROME; ARTS SYNDROME; ARX-ASSOCIATED XLID; ATAXIA-DEAFNESS-DEMENTIA, X-LINKED; ATKIN-FLAITZ SYNDROME; ATRX-ASSOCIATED XLID; BERGIA CARDIOMYOPATHY; BERTINI SYNDROME; BORJESON-FORSSMAN-LEHMANN SYNDROME; BRANCHIAL ARCH SYNDROME, X-LINKED
CANTU SYNDROME
CARPENTER-WAZIRI SYNDROME (SEE ALSO ATRX-ASSOCIATED XLID); CEREBRO-CEREBELLO-COLOBOMA SYNDROME; CEREBRO-OCULO-GENITAL SYNDROME; CEREBRO-PALATO-CARDIAC SYNDROME (SEE ALSO RENPENNING SYNDROME); CHARCOT-MARIE-TOOTH NEUROPATHY, COWCHOCK VARIANT; CHARCOT-MARIE-TOOTH NEUROPATHY, IONASESCU VARIANT; CHASSAING-LACOMBE CHONDRODYSPLASIA; CHRISTIAN SYNDROME; CHRISTIANSON SYNDROME; CHUDLEY-LOWRY SYNDROME (SEE ALSO ATRX-ASSOCIATED XLID); CK SYNDROME; CLARK-BARAITSER SYNDROME; COFFIN-LOWRY SYNDROME; CORNELIA DE LANGE SYNDROME, X-LINKED; CRANIOFACIOSKELETAL SYNDROME
CREATINE TRANSPORTER DEFICIENCY
DUCHENNE MUSCULAR DYSTROPHY; DYSKERATOSIS CONGENITA; EPILEPSY-INTELLECTUAL DISABILITY IN FEMALES (EIDF); FITZSIMMONS SYNDROME; FLNA-ASSOCIATED XLID; FRAGILE X SYNDROME; GIUFFRE-TSUKAHARA SYNDROME; GLYCEROL KINASE DEFICIENCY; GOLABI-ITO-HALL SYNDROME: (SEE ALSO RENPENNING SYNDROME); GOLDBLATT SPASTIC PARAPLEGIA SYNDROME; GOLTZ SYNDROME; GRAHAM ANOPHTHALMIA SYNDROME; GUSTAVSON SYNDROME; HALL OROFACIAL SYNDROME; HEREDITARY BULLOUS DYSTROPHY, X-LINKED; HOLMES-GANG SYNDROME (SEE ALSO ATRX-ASSOCIATED XLID); HOMFRAY SEIZURES-CONTRACTURES; HYDE-FORSTER SYNDROME
HYDRANENCEPHALY WITH ABNORMAL GENITALIA (SEE ALSO ARX-ASSOCIATED XLID)
HYDROCEPHALY-CEREBELLAR AGENESIS SYNDROME; HYDROCEPHALY-MASA SPECTRUM; HYPOPARATHYROIDISM, X-LINKED; INCONTINENTIA PIGMENTI; JUBERG-MARSIDI-BROOKS SYNDROME; KANG SYNDROME; LENZ MICROPHTHALMIA SYNDROME; LESCH-NYHAN SYNDROME; LISSENCEPHALY AND ABNORMAL GENITALIA, X-LINKED (SEE ALSO ARX-ASSOCIATED XLID); LISSENCEPHALY, X-LINKED; LOWE SYNDROME; LUJAN SYNDROME; MARTIN-PROBST SYNDROME; MEHMO SYNDROME; MENKES SYNDROME; MIDAS SYNDROME; MILES-CARPENTER SYNDROME; MOHR-TRANENBJAERG SYNDROME; MONOAMINE OXIDASE-A DEFICIENCY
MUCOPOLYSACCHARIDOSIS II
AMYOTUBULAR MYOPATHY; N-ALPHA-ACETYLTRANSFERASE DEFICIENCY; NANCE-HORAN SYNDROME; NORRIE DISEASE; OPITZ FG SYNDROME; OPTIC ATROPHY, X-LINKED; ORAL-FACIAL-DIGITAL SYNDROME I; ORNITHINE TRANSCARBAMOYLASE DEFICIENCY; OTOPALATODIGITAL SYNDROME I (SEE ALSO FLNA-ASSOCIATED XLID); OTOPALATODIGITAL SYNDROME II (SEE ALSO FLNA-ASSOCIATED XLID); PAINE SYNDROME; PALLISTER W SYNDROME; PARTINGTON SYNDROME (SEE ALSO ARX-ASSOCIATED XLID); PELIZAEUS-MERZBACHER SYNDROME; PERIVENTRICULAR NODULAR HETEROTOPIA (SEE ALSO FLNA-ASSOCIATED XLID); PETTIGREW SYNDROME
PHOSPHOGLYCERATE KINASE DEFICIENCY

disorders caused by genes on the X chromosome. Clinical and laboratory data on 150 syndromes are presented in a concise and consistent manner. Each syndrome is defined and information is provided on somatic features, growth and development, neurological signs, cognitive performance, imaging and other laboratory findings, and when possible, the nature and localization of the responsible gene. Craniofacial and other somatic findings are extensively illustrated. A differ

3. Record Nr.	UNINA9910689551703321
Titolo	Assessing the Assistive Technology Act of 1998 : hearing before the Subcommittee on 21st Century Competitiveness of the Committee on Education and the Workforce, House of Representatives, One Hundred Seventh Congress, second session, hearing held in Washington, DC, March 21, 2002
Descrizione fisica	1 online resource (iv, 203 p.) : ill
Soggetti	Self-help devices for people with disabilities - Law and legislation - United States Rehabilitation technology - United States Human engineering - United States Telecommunication in higher education - United States
Lingua di pubblicazione	Inglese
Formato	Materiale a stampa
Livello bibliografico	Monografia