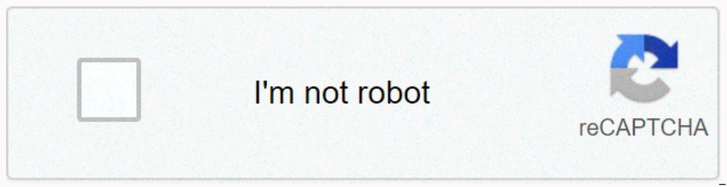

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Unique rare chromosome

Unique is a source of information and support for families and individuals affected by any rare chromosome or newly turned gene disorder, as well as for professionals who work with them. Founded in 1984, Unique is a British charity (1110,661), but welcomes members around the world. Unique's membership is free, but the group receives no government funding and relies heavily on donations and fundraising to continue its work. The unique offers contact with families with the affected member who has the same rare chromosome or gene disorder or that has similar symptoms or practical problems, regardless of specific chromosome or gene disorder. It produces a wide range of family,medically verified information manuals on various rare disorders of the chromosom. Uniquely coordinates families to assist in research and has password-protected discussion forums for registered members, it also has social media sites on Facebook, Twitter, etc. it publishes a magazine three times a year. Unique hosts occasionally study and family weekend conferences. Unique has more than 19,000 affected families and many relevant professionals as members worldwide in more than 100 countries. The network of local contacts around the world contributes to the awareness of chromosomes and gene disorders and is unique to the general public and professionals. This page was last updated on June 13, 2019, Uniqueness is a source of information and support for families and individuals affected by any rare chromosome or gene disorder, as well as for professionals who work with them. Founded in 1984, Unique is a British charity (1110,661), but welcomes members around the world. Unique's membership is free, but the group receives no government funding and relies heavily on donations and fundraising to continue its work. Unique, offers contact with families with the affected member who has the same rare chromosomous disorder or who has similar symptoms or practical problems, regardless of specific chromosomous disorder. It produces a wide range of family,medically verified information manuals on various rare disorders of the chromosom. Uniquely coordinates families to assist in research and has password-protected discussion forums for registered members, it also has social media sites on Facebook, Twitter, etc. it publishes a magazine three times a year. Unique hosts occasionally study and family weekend conferences. Unique has more than 21,500 affected families and many relevant professionals as members worldwide in 100+ countries. A network of local contacts around the world promotes awareness of rare chromosomous disorders and is unique to the general public and professionals. Individually, these chromosomous disorders are rare, but collectively they More than one in every 200 babies are thought to be born with a rare chromosomous disorder, with more than one in every 1,000 babies having symptom symptoms birth or early childhood, the rest suffer when they grow up and try to have children on their own. Contact InformationBay Searle Contact positionCEO Phone 01883 723 356 E-mail info@rarechromo.org Site www.rarechromo.org NameThe address of the Station Road West Oxted Surrey PostcodeRH8 EE Is this information correct? Unique is a source of information and support for families and individuals affected by any rare chromosome or newly turned gene disorder, as well as for professionals who work with them. Founded in 1984, Unique is a British charity (1110,661), but welcomes members around the world. Unique membership is free - the group receives no government funding and relies heavily on donations and fundraising to continue its work. Unique, offers contact with families with the affected member who has the same rare chromosomous disorder or that has similar symptoms or practical problems, regardless of specific chromosome or gene disorder. It produces a wide range of family,medically verified information manuals on various rare disorders of the chromosom. Unique coordinates families to assist in research and has private Facebook groups for registered members. He publishes the magazine three times a year. Unique hosts occasionally study and family weekend conferences. Unique has more than 21,000 affected members and many relevant professional members worldwide in more than 100 countries. A network of local contacts around the world promotes awareness of chromosomous disorders and is unique to the general public and professionals. Supported Diseases 15q11.2 microdeletion Synonyms: Chromosome 15q11.2 microdeletion, Chromosome 15q11.2 deletion, Del(15)(q11.2), 15q11.2 microdeletion syndrome, Monosomy 15q11.2, 15q11.2 BP1-BP2 microdeletion syndrome 15q13.3 microdeletion syndrome Synonyms: Chromosome 15q13.3 deletion syndrome, Microdeletion 15q13.3 syndrome, Chromosome 15q13.3 microdeletion syndrome, 15q13.3 microdeletion 15q13.3 microduplication syndrome Synonyms: 15q13.3 microduplication, Chromosome 15q13.3 duplication syndrome, Chromosome 15q13.3 microduplication syndrome, Microduplication 15q13.3 syndrome 15q24 microdeletion syndrome Synonyms: Del(15)(q24), Monosomy 15q24 16p11.2 deletion syndrome Synonyms: Chromosome 16p11.2 deletion syndrome, Del(16)(p11.2), Monosomy 16p11.2, Microdeletion 16p11.2 16p11.2 duplication Synonyms: Susceptibility to Autism, 14B, AUTS14B, 16p11.2 duplication syndrome, 16p11.2 microduplication 17q12 deletion syndrome Synonyms: Chromosome 17q12 deletion syndrome, 17q12 microdeletion syndrome, Del(17)(q12), Monosomy 17q12, 17q12 recurrent deletion syndrome 17q12 duplication Synonyms: Chromosome 17q12 duplication syndrome , Microduplication syndrome 17q12, Trisomy 17q12, Recurrent duplication 17q12, microduction 17q12, Dup(17)(q12) 17q23.1q23.2 Synonym microdelece: 17q23.1-q23.2 microdelelyation microdelation Del(17)(q23.1q23.2), Monosomy 17q23.1-q23.2, Monosomy 17q23.1q23.2, Chromosome 17q23.1-q23.2 deletion syndrome 1q duplications Synonyms: Partial trisomy 1q, Partial trisomy of the long arm of chromosome 1, Partial trisomy of chromosome 1q, Partial duplication of chromosome 1q, Partial duplication of the long arm of chromosome 1 20p12.3 microdeletion syndrome Synonyms: Del(20)(p12.3), Monosomy 20p12.3 22q11.2 deletion syndrome Synonyms: Chromosome 22q11.2 deletion syndrome, Velocardiofacial syndrome, VCFS, DiGeorge syndrome, Shprintzen syndrome, Sedlackova syndrome, CATCH22, Autosomal dominant Opitz G/BBB syndrome, Conotruncal anomaly face syndrome, Cayler cardiofacial syndrome 22q13.3 deletion syndrome Synonyms: Phelan-McDermid syndrome, Deletion 22q13.3 syndrome, Chromosome 22q13.3 deletion syndrome, Monosomy 22q13, 22q13 deletion, 22q13.3 deletion, Monosomy 22q13.3 2q37 deletion syndrome Synonyms: Chromosome 2q37 deletion syndrome, Albright hereditary osteodystrophy-like syndrome, Brachydactyly-Intellectual disability syndrome 47 XXX syndrome Synonyms : Trisomy X, Triple X syndrome, Triple-X female, XXX syndrome, Triple-X chromosome syndrome, Triplo X syndrome 47, XYY syndrome Synonyms: YY syndrome, XYY syndrome, XYY Karyotype, Jacobs syndrome, 47,XYY syndrome, Disomy Y, Double Y, Double Y syndrome, Y disomy 48,XXXY syndrome Synonyms: XXXY syndrome 49, XXXXY syndrome Synonyms: XXXXY syndrome, 49,XXXXY 49, XXXYY syndrome Synonyms: XXXYY syndrome 49,XXXXX syndrome Synonyms: Pentasomy X, Chromosome XXXXX syndrome, Penta-X syndrome, Pentasomy X syndrome, Chromosome X pentasomy 5q14.3 microdeletion syndrome Synonyms: Chromosome 5q14.3 deletion syndrome, Monosomy 5q14.3, Del(5)(q14.3), Mental retardation, autosomal dominant 20, 5q14.3 deletion syndrome, Autosomal dominant intellectual disability 20 7q11.23 duplication syndrome Synonyms: Williams-Beuren region duplication syndrome, WBS duplication syndrome, Chromosome 7q11.23 duplication syndrome, Somerville-Van Der AA syndrome , Dup(7)(q11.23), Trisomy 7q11.23, 7q11.23 microduplication syndrome 8p23.1 duplication syndrome Cat eye syndrome Synonyms: Schmid-Fraccaro syndrome, CES, Chromosome 22 partial tetrasomy, INV DUP(22)(Q11) Chromosome 11q duplication Synonyms: Duplication 11q, Trisomy 11q, 11q duplication, 11q trisomy, Partial trisomy 11q, Partial duplication of the long arm of chromosome 11 Chromosome 16p13.3 deletion syndrome Synonyms: 16p13.3 deletion syndrome Chromosome 16p13.3 duplication Synonyms: 16p13.3 duplication, Interstitial 16p13.3 duplication, 16p13.3 microduplication syndrome, Distal duplication 16p, Distal trisomy 16p, Dup(16)(p13.3), Telomeric duplication 16p, Trisomy 16pter Chromosome 18p deletion Synonyms: 18p-, Monosomy 18p Chromosome 19q13.11 deletion syndrome Chromosome 20 trisomy Synonyms: Trisomy chromosome 20, Trisomy 20, Trisomy 20 mosaicism, Mosaic trisomy 20 Chromosome duplication Chromosome 3p- syndrome Synonyms: Del(3p) syndrome, Chromosome 3, monosomy 3p25, Deletion 3p25, Chromosome 3pter-p25 Deletion Syndrome, Telomeric monosomy 3p, Distal 3p deletion, 3p- syndrome, Distal monosomy 3p, Monosomy 3pter Chromosome 8q24.3 deletion syndrome Synonyms: Verheij syndrome Chromosome 9 inversion Synonyms: Inversion 9 Cri du chat syndrome Synonyms: Cat cry syndrome, 5p minus syndrome, Chromosome 5p deletion syndrome, 5p- syndrome, Monosomy 5p, 5p deletion syndrome, Chromosome 5p- syndrome Emanuel syndrome Synonyms: Supernumerary der(22),t(11;22) syndrome, Supernumerary der(22) syndrome Isodicentric chromosome 15 syndrome Synonyms: Duplication/inversion 15q11, Inv dup(15), Non-distal tetrasomy 15q, Non-telomeric tetrasomy 15q, Idic(15), Tetrasomy 15q, Chromosome 15q tetrasomy, Inverted duplication 15 Jacobsen syndrome Synonyms: JBS, Chromosome 11q deletion syndrome, Partial 11q monosomy syndrome Kleefstra syndrome Synonyms : синдром мікроделегування 9q34.3, синдром видалення хромосоми 9q, синдром видалення хромосоми 9q34.3, синдром 9q- синдром синдрому Кулена де Вріса Синоніми: синдром видалення 17q21.31, Моносомія 17q21.31, Синдром мікроделеції 17q21.31, синдром мікроделеції хромосоми 17q21.31, синдром мікроделеції 17q21.31, синдром інтелектуальної інвалідності, пов'язаний з KANSL1, синдром дублювання MECP2 Синоніми: MRXSL, синдром розумової відсталості, пов'язаний з ЛУБсом X (раніше), синдром XLMR, Розумова відсталість, X-пов'язана, Тип Лубса (раніше), Трисомія Xq28 Синдром Міллера-Дікера Синоніми: синдром Міллера-Дікера ліссенцефалії, трисомія МОЗАЇки 7 Синоніми: Мозаїчна трисомія хромосоми 7, Трисомія 7 мозаїчний трисом 8 Синоніми: Мозаїчна трисомна хромосома 8, Трисомна мозаїка 8 мозаїка 8 Мозаїка трисомія 9 Синоніми: Мозаїчна трисомія хромосоми 9, Трисомія 9 мозаїка Наблус масковий синдром Синоніми: синдром мікроделегації 8q22.1 Паллістер-Кілліан мозаїчний синдром Синоніми: Хромосома 12, синдром Ісохромосоми 12 Ніколап, синдром Кілліана , Синдром мозаїки Палістера, синдром Тешлера-Нікола Кілліана, Тетрасомія 12п, мозаїка, синдром Паллістера Кілліана, PKS Часткове видалення синонімів Y: мікроделети Y-хромосоми, Часткове видалення хромосоми Y, Часткове видалення довгої руки Y хромосоми Синдром Потоцьких-Шаффера Синоніми: PSS, Видалення хромосоми 11p11.2, Синдром видалення Проксимальної 11p, видалення 11p11.2 Кільцева хромосома 1 Синоніми: Хромосома 1 кільце, Кільце 1, R1 Кільце хромосоми 10 Синоніми: Хромосома 10 кільце, кільце 10, R10 кільце хромосоми 11 Синоніми: Хромосома 11 кільце, кільце 11, R11 кільце хромосоми 12 Синоніми: Хромосома 12 кільце, кільце 12, R12 Кільце хромосоми 13 Синоніми: Хромосома 13 кільце, кільце 13, R13 Кільце хромосоми 15 Синоніми: Хромосома 15 кільце, кільце 15, R15 кільце хромосоми 16 Синоніми: Хромосома 16 кільце , Кільце 16, R16 Кільцева хромосома Synonyms: Ring chromosom 17, ring 17, P17 Ring chromosom 18 Synonyms: Ring chromosom 18, ring 18, P18 Ring chromosom 19 Synonyms: Chromosom 19 ring, ring 19, R19 Ring chromosom 2 Synonyms: Chromosom 2, ring Ring 2, R2 Chromosom Ring 20 Synonyms: Chromosom 20 Ring, Ring 20, R20, Ring Chromosom 20 Syndrome Ring Chromosom 21 Synonyms: Chromosom 21 Ring, Ring 21, R21 Ring Chromosom 22 Synonyms : Chromosom 22 ring, ring 22, R22 ring chromosom 3 Synonyms: Chromosom 3 ring, ring 3, R3 ring chromosom 4 Synonyms: Chromosom 4 ring, Ring 4, R4 Ring Chromosom 5 Synonyms: Chromosom 5 Ring, Ring 5, R5 Ring Chromosom 6 Synonyms: Chromosom 6 Ring, Ring 6, R6 Ring Chromosom 7 Synonyms: Chromosom 7 Ring, Ring 7, R7 Ring Chromosom 8 Synonyms: Chromosom 8 Ring , Ring 8, R8 Ring Chromosom 9 Synonyms: Chromosom 9 Ring, Ring 9, R9 SATB2 Associated Synonym Syndrome: SATB2 Syndrome, SAS, 2q32q33 Microdelement Syndromes, Monosomy 2q32-q33, Del(2)(q32q33), Del(2)(q32), microdelegate syndrome 2q32-q33, Síndrome de Emanuel Synonyms glass syndrome: Síndrome del der(22) supernumerario, Síndrome der(22) supernumerario, Síndrome der(22)t(11;22) Synonyms Synonymous with Sindrome Smith-Magennis: SMS, Chromosome 17p11.2 Tetrasomy Removal Syndrome 9p Synonyms : Chromosom 9p tetrasomy, Tetrasomy short arm chromosom 9, Mosaic tetrasomy 9p Tetrasomy X Synonyms: 48 XXXX, 48 XXXX syndrome, Tetra X, 48,XXXX syndrome, Four times X Tryploid synonyms: Tryploid syndrome, Tryploidi syndrome, Chromosomic triploid syndrome Trisomy 13 Synonyms: Patau syndrome, Chromosome 13, trisomy 13 full, Complete trisomy 13 syndrome, trisomy syndrome D (formerly) Trisomy 17 mosaic : Chromosome 17 duplication, Trisomy 17 , Chromosom 17 trisomy, chromosom 17, trisomy, chromosom 17, trisomy mosaic Trisomy 2 mosaic Synonyms: Mosaic trisomy 2 Turner Synonymous Syndrome: Ullić-Turner syndrome, Bonvey-Ulrich syndrome, 45 years old, X syndrome, Chromosome X Monosomy X, Gonadal dysgenesis (45,X), Scheleszewkiy Turner syndrome, Turner Varna syndrome Wolf-Hirshhorn Synonyms: WHS, Wolf syndrome, chromosome syndrome 4p, microcephaly, IUGR, hypertension , hooked nose, external ear dysplyase, psychomotor retardation, 4p syndrome, removal distal 4p, 4p- syndrome, distal monosomy 4p, removal of telomerin 4p, Whitver syndrome, Pitt-Rogers-Danks syndrome Y chromosomic infertility Synonymous: Male sterility due to Y-chromosome removal, DAZ, Removed in azoospermia, Y chromosomic microdelity Y chromosomes