***DACRO (D****isease-****A****ssociated* ***C****hromosomal* ***R****earrangements* ***O****nline)*

34 patients with OFC

PROBLEM: only cytogenetic bands (NO microarray data)

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| **ID** | **Phenotypes** | **Translocation/Inversion** |
| 939 | Cleft lip and palate, Isolated | t(2;8)(q33.1;q24.2) |
| 1047 | Cleft lip and palate, other dysmorphism | t(6;9)(p24.3;q22.33) |
| 460 | Cleft lip and ventricular septal defect | t(2;10)(q53;p13) |
| 461 | Cleft lip and ventricular septal defect | t(10;14)(p13;q24)pat |
| 27 | Cleft palate, isolated | t(2;7)(q33.1;p21.1) |
| 28 | Cleft palate, isolated | t(2;11)(q33.1;p13) |
| 654 | Cleft palate, Isolated | t(6;7)(p25;q31) |
| 1040 | Cleft palate, prominent ears, small chin, tapering fingers | t(11;13)(p15.4;q22.1) |
| 648 | Cleft palate, severe learning disabilities | inv(12)(p11.21q24.31) |
| 645 | Coloboma (bilateral), cleft lip and palate, micropthalmia, mental retardation | inv(4)(q21.22q35) |
| 623 | Congenital glaucoma, cleft lip and palate and mental retardation | t(1;6)(q23;q27) |
| 616 | Dolichocephalic skull, low set abnormal ears, antimongoloid palpebral fissures, epicanthal folds, cleft of soft palate, partial cleft of hard palate, micrognathiaand severe mental retardation | t(2;8;20)(q23;q22;q11.2) |
| 111 | Ectrodactyly, Ectodermal Dysplasia and cleft lip/palate syndrome 1 (EEC1) | t(7;9)(q21.3;p12)pat |
| 112 | Ectrodactyly, Ectodermal Dysplasia and cleft lip/palate syndrome 1 (EEC1) | t(2;7)(q21.1;q22.3) |
| 113 | Ectrodactyly, Ectodermal Dysplasia and cleft lip/palate syndrome 1 (EEC1) | t(7;12)(q21.3;q24.2) |
| 114 | Ectrodactyly, Ectodermal Dysplasia and cleft lip/palate syndrome 1 (EEC1) | t(6;13)(q21;q12) |
| 565 | Epicanthal folds, flat nasal bridge, small mouth, micrognathia, low set ears, cleft palate | t(9;11)(p21.2;p14.2) |
| 399 | Hypertelorism, microtia, and facial clefting syndrome (HMC) | t(1;7)(q31.2;p15.1-p15.3) |
| 827 | Hypogonadotropic hypogonadism (infantile testes), azoospermia, and cleft lip and palate | t(7;8)(p12.3;p11.2) |
| 323 | Median cleft of upper lip/ pedunculated skin masses | t(X;16)(q28;q11.2) |
| 824 | Michelin tyre syndrome with distinctive facial dysmorphia, submucous cleft palate, lateral clefting of the mouth, genital and dental anomalies and moderate developmental delay | inv(7)(q22q31.3)mat |
| 613 | Multiple anomalies including: open metopic suture, midline cleft palate, micrognathia, tetralogy of Fallot variant with high ventricular septal defect, atretic pulmonic valve, hypoplastic pulmonary artery, duplication of vagina and uterus | t(X;17)(q11;q11) |
| 310 | Oblique facial clefts (bilateral), calcaneovarus foot deformity, severe bilateral ocular hypoplasia | t(1;22)(q21;q12) |
| 141 | Orofacial Cleft 1 | t(6;7)(p23;q36.1) |
| 142 | Orofacial Cleft 1 | t(6;9)(p23;q22.3)mat |
| 143 | Orofacial Cleft 1 | t(2;19)(q11.2;q13.3) |
| 145 | Orofacial Cleft 1 | inv(8)(p23q11) |
| 909 | Orofacial Cleft 1 | t(6;9)(p24;p23) |
| 1084 | Orofacial Cleft 1 | t(9;17)(q32;q12) |
| 144 | Orofacial Cleft 1 with hypospadias | inv(4)(p13q21.1)pat |
| 967 | Peters Anomaly associated with multiple midline defects (cranial meningocele, cardiac defects and cleft lip and palate) | inv(4)(q12q13.3) |
| 987 | Severe lower eyelid colobomas, malar and mandibular hypoplasia, bilateral microtia with external auditory canal atreasia, dysplastic ossicles, hearing loss, bilateral choanal stenosis, cleft palate, several oral frenula of the upper lip and micrognathia | t(2;17)(q24.3;q23)  |
| 1077 | Severe mental retardation, muscular hypotonia, seizures, bilateral sensorineural hearing loss, submucous cleft palate, persistent ductus Botalli, unilateral cystic kidney dysplasia and frequent infections | t(11;20)(p15.4;q13.2) |
| 1045 | Ventricular septal defect (VSD), cleft palate, XY sex reversal, hydronephrosis | t(10;11)(q24.2;p12) |