Chromosome 15q24 microdeletion syndrome

Pilar L Magoulas1 and Ayman W El-Hattab2*

Abstract

Chromosome 15q24 microdeletion syndrome is a recently described rare microdeletion syndrome that has been reported in 19 individuals. It is characterized by growth retardation, intellectual disability, and distinct facial features including long face with high anterior hairline, hypertelorism, epicanthal folds, downsloping palpebral fissures, sparse and broad medial eyebrows, broad and/or depressed nasal bridge, small mouth, long smooth philtrum, and full lower lip. Other common findings include skeletal and digital abnormalities, genital abnormalities in males, hypotonia, behavior problems, recurrent infections, and eye problems. Other less frequent findings include hearing loss, growth hormone deficiency, hernias, and obesity. Congenital malformations, while rare, can be severe and include structural brain anomalies, cardiovascular malformations, congenital diaphragmatic hernia, intestinal atresia, imperforate anus, and myelomeningocele. Karyotypes are typically normal, and the deletions were detected in these individuals by array comparative genomic hybridization (aCGH). The deletions range in size from 1.7-6.1 Mb and usually result from nonallelic homologous recombination (NAHR) between paralogous low-copy repeats (LCRs). The majority of 15q24 deletions have breakpoints that localize to one of five LCR clusters labeled LCR15q24A, -B, -C, -D, and -E. The smallest region of overlap (SRO) spans a 1.2 Mb region (Figure 1) [1-10]. Routine high-resolution karyotypes typically do not detect the abnormalities. Additional common features include hypotonia, behavioral problems, and recurrent infections (table 1) [1-10]. Although some of the characteristic features may be non-specific, the combination of these findings should prompt the clinician to suspect this diagnosis. Most of the deletions are between 1.7 to 6.1 megabases (Mb) in size, with the smallest region of overlap (SRO) spanning a 1.2 Mb region (Figure 1) [1-10]. Routine high-resolution karyotypes typically do not detect the

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deletion and all individuals with 15q24 deletion syndrome were diagnosed by using array comparative genomic hybridization (CGH) [1-10].

Epidemiology
The incidence of 15q24 deletion syndrome is unknown. To date, there have been 19 published reports of individuals with 15q24 deletion syndrome in which clinical information and detailed mapping of the genomic breakpoints are available [1-10]. The first case was reported in 2007 by Sharp et al [1]. Prior to this time, the widespread use of array CGH was limited, therefore the diagnosis may have gone undetected and undiagnosed. With the advent of array CGH, we anticipate that the present number of reported cases represents an underestimate of the actual prevalence of the disorder. A recent study by Cooper et al [11] reviewed 15,767 cases with intellectual disability and various congenital defects referred to their laboratory for array CGH. They identified 15q24 deletion in nine patients with intellectual disability, for an incidence of approximately 1 in 1751 in this population. This would make it approximately 1 in 42,000. We would expect the deletion to occur with equal frequency in males and females; however 16 of the 19 reported cases (84%) are males. The reason for the discrepancy in the male to female sex ratio of 15q24 deletion is not clearly understood and may just be representative of the small sample size of cases and ascertainment bias, rather than a true phenomenon. The deletion occurred as a de novo event in the 18 individuals that the parents were tested [1-10].

Clinical description
The clinical characteristics of 15q24 deletion syndrome are based on the clinical features reported in the 19 patients with 15q24 microdeletion and presented in Table 1 [1-10]. At birth, approximately one-third of the individuals had low birth weight consistent with intrauterine growth retardation. Feeding difficulties and failure to thrive were reported in approximately 20%. However, in later childhood, growth parameters were extremely variable with nearly 30% of individuals still exhibiting postnatal growth retardation and short stature, while 17% had obesity. Additionally, almost 20% of individuals were reported to have microcephaly. Two individuals with 15q24 deletion had documented growth hormone (GH) deficiency. This might be an underascertainment of the actual prevalence of GH deficiency in this
growth delay, feeding difficulties, and distinct facial features were often the presenting early symptoms of 15q24 deletion syndrome. Distinct facial features, including long face with high anterior hairline, epicanthal folds, hypertelorism, downslanting palpebral fissures, sparse and broad medial eyebrows, broad and/or depressed nasal bridge, long smooth philtrum, and small mouth with full lower lip were common in all individuals. Ear abnormalities were variable, but present in nearly two-thirds of individuals and included large ears, ear lobe pits, anteverted ear lobes, and protuberant ears. These features may be subtle in the newborn period, but may become more evident with age.

All of the individuals have developmental delay of varying degrees, usually in the range of mild to moderate intellectual disability. In reviewing the literature, 6/19 cases were classified as having mild intellectual disability, 2/19 were classified as moderate, 1/19 was classified as moderate-to-severe, 3/19 were classified as severe, and the remaining 7/17 were not further characterized based on severity. Behavior abnormalities, such as autism, hyperactivity, aggression, and attention deficit disorder have been reported in 37% individuals with 15q24 deletion syndrome. McInnes et al performed targeted aCGH across the 15q24 region on ~1500 individuals with autism spectrum disorders and identified a de novo atypical deletion of 3.06 Mb in 15q23-q24.1 in a 5 year old male with autism [8]. In addition to autism, he exhibited other characteristic features of 15q24 deletion syndrome including, moderate intellectual disability, low birth weight, distinct facial features, 2,3 toe syndactyly, mild scoliosis, and recurrent respiratory infections. These features may be subtle in the newborn period, but may become more evident with age.

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Skeletal and digital deformities have been described in the majority of individuals. Digit abnormalities such as proximally implanted and/or hypoplastic thumbs, clinodactyly, brachydactyly, overriding toes, toe syndactyly, and small hands were found in approximately 90% of

Table 1 Summary of clinical manifestation in patients with 15q24 microdeletion syndrome

| Characteristic                  | Cases     |
|--------------------------------|-----------|
| Gender                         | Male - 16/19 (84%) |
| De novo occurrence             | 18/18 (100%) |
| Developmental delay             | 19/19 (100%) |
| Growth abnormalities            | 10/18 (56%) |
| Low birth weight/IUGR           | 6/18 (33%) |
| Short stature                   | 5/18 (28%) |
| Obesity                        | 3/18 (17%) |
| Microcephaly                   | 3/18 (17%) |
| Feeding difficulties            | 4/18 (22%) |
| Distinct Facial features        | 19/19 (100%) |
| Long face                       | 6/18 (33%) |
| Facial asymmetry                | 5/18 (28%) |
| High anterior hairline          | 12/18 (67%) |
| Epicanthal folds                | 9/18 (50%) |
| Hypertelorism                   | 8/18 (44%) |
| Downslanting palpebral fissures | 9/18 (50%) |
| Sparse, broad medial eyebrows   | 9/18 (50%) |
| Strabism                       | 7/18 (39%) |
| Nystagmus                       | 2/18 (11%) |
| Broad nasal base                | 6/18 (33%) |
| Depressed nasal bridge          | 5/18 (28%) |
| High nasal bridge               | 2/18 (11%) |
| Ear abnormalities               | 11/18 (61%) |
| Palate abnormalities            | 4/18 (22%) |
| Long smooth philtrum            | 10/18 (56%) |
| Full lower lip                  | 7/18 (39%) |
| Small mouth                     | 6/18 (33%) |
| Genital abnormalities           | 11/18 (61%) |
| Hypospadias                     | 6/15 (40%) |
| Micropenis                       | 5/15 (33%) |
| Cryptorchidis                   | 3/15 (20%) |
| Digit abnormalities             | 16/18 (89%) |
| Thumb abnormalities             | 7/18 (39%) |
| Brachydactyly/short digits      | 6/18 (33%) |
| Clinodactyly                    | 3/18 (20%) |
| Toe abnormalities               | 6/18 (33%) |
| Skeletal abnormalities          | 11/18 (61%) |
| Joint laxity                    | 10/18 (56%) |
| Scoliosis/kyphosis              | 6/18 (33%) |
| Neurologic findings             | 15/19 (79%) |
| Hypotonia                       | 11/18 (61%) |
| Behavior problems               | 7/19 (37%) |
| MRI abnormalities               | 8/18 (44%) |
| Other features                  |        |
| Recurrent infections            | 7/18 (39%) |
| Hernias                        | 5/19 (26%) |

Table 1 Summary of clinical manifestation in patients with 15q24 microdeletion syndrome (Continued)

| Characteristic                  | Cases     |
|--------------------------------|-----------|
| Congenital heart disease        | 4/18 (22%) |
| Hearing loss                    | 4/18 (22%) |
| Diaphragmatic hernia            | 2/18 (11%) |
| Intestinal atresia              | 2/18 (11%) |
| Imperforate anus                | 1/18 (6%)  |
| Coloboma                        | 1/18 (6%)  |
| Dental problems                 | 1/18 (6%)  |
| Myelomeningocele                | 1/18 (6%)  |
| Congenital heart disease        | 4/18 (22%) |
| Hearing loss                    | 4/18 (22%) |
| Diaphragmatic hernia            | 2/18 (11%) |
| Intestinal atresia              | 2/18 (11%) |
| Imperforate anus                | 1/18 (6%)  |
| Coloboma                        | 1/18 (6%)  |
| Dental problems                 | 1/18 (6%)  |
| Myelomeningocele                | 1/18 (6%)  |
individuals with 15q24 deletion. In addition, many individuals were reported to have joint laxity and scoliosis. Some individuals were reported to have hernias which, in addition to the joint laxity, suggest that there may be an underlying connective tissue abnormality in individuals with 15q24 deletion syndrome [3].

Hypotonia has been reported in 60% of individuals with 15q24 deletion syndrome. Approximately half of the individuals had abnormal brain imaging on magnetic resonance imaging (MRI). The features in those with abnormalities are variable and include: dysplastic or abnormal corpus callosum, cortical atrophy, focal cortical dysplasia, and hypoplastic olfactory bulbs. The incidence of seizure activity is relatively low with only one individual having reported seizures [1].

Genital abnormalities are reported in a majority of individuals (60%) and include hypospadias, microphallus, and cryptorchidism. These findings, in addition to other possible malformations or dysmorphisms in the newborn period, may prompt an earlier genetics evaluation and diagnosis when compared to females who have not been reported to have urogenital anomalies. This may possibly explain why the majority of reported patients are males.

Nearly 40% of individuals with 15q24 deletion syndrome had a history of recurrent infections, such as chest, upper airway, and ear infections. The etiology of the recurrent infections is currently unknown. Immunodeficiency could be the underlying cause; however, immunologic work up has not been reported in any of those individuals. Recurrent ear infections may be one of the factors that predispose to hearing loss which has been formally evaluated and diagnosed in nearly 25% of individuals with 15q24 deletion syndrome.

Birth defects have been reported in several individuals including congenital heart defects (such as tetralogy of Fallot and ventricular septal defects) in four (~20%), intestinal atresia in two, imperforate anus in one [5], iris coloboma in one, myelomeningocele in one, and diaphragmatic hernia in two individuals. Other features that have been reported in a minority of cases include strabismus [1-3,6,10] nystagmus [1,5,8] and dental abnormalities [1].

One individual with 15q24 deletion syndrome was reported to have acute lymphoblastic leukemia [4] and another individual was diagnosed with cardiac rhabdomyoma [7]. No additional malignancies or tumors have been reported in this population, therefore it is currently unknown if there is a predisposition to tumors and neoplasia in individuals with 15q24 deletion syndrome.

Two individuals with 15q24 deletion syndrome had another concurrent chromosome abnormality [7,9]. These individuals exhibited additional clinical features with a more severe phenotype than that observed in previously reported patients with isolated 15q24 rearrangements and it was suggested that the number of genomic rearrangements (genomic mutational load) may contribute to the phenotypic severity and variability in patients with 15q24 deletions [7,12,13].

Etiology

15q24 deletion syndrome is a microdeletion syndrome resulting from the interstitial loss of 15q24 ranging in size from ~1.7 to 6.1 Mb. All but one case share the smallest region of overlap (SRO) of ~1.2 Mb in size (see Figure 1) [1-10]. The 15q24 deletions are primarily mediated by nonallelic homologous recombination (NAHR) between paralogous low-copy repeats (LCRs). The majority of 15q24 deletions have breakpoints that localize to LCR clusters. Five LCR clusters have been identified, labeled, LCR15q24A, LCR15q24B, LCR15q24C, LCR15q24D, and LCR15q24E. The SRO spans the 1.2 Mb region between LCR15q24B to LCR15q24C. Some patients have had breakpoints that do not lie within the LCR regions. Breakpoint sequencing for the 15q24 deletion in one of those patients revealed microhomology suggesting that other possible mechanisms include nonhomologous end joining (NHEJ) and fork stalling and template switching (FoSTeS)/microhomology-mediate break-induced replication (MMBIR) [7,14-17].

The SRO is a gene-rich region that includes several genes including: CYP11A1, SEMA7A, CPLX3, ARID3B, STRA6, SIN3A and CSK. CYP11A1 encodes the cholesterol side-chain cleavage enzyme and it has been suggested that haploinsufficiency of CYP11A1 may contribute to the genital abnormalities in patients with 15q24 deletion syndrome [7]. SEMA7A and CPLX3 are two genes that may play an important role in brain development and may contribute to the cognitive disability in individuals with this deletion [7]. Haploinsufficiency of ARID3B and STRA6 may contribute to the congenital malformations observed in a handful of patients with 15q24 deletion syndrome. The ARID3B gene is involved with transcription regulation in embryonic development. Homozygous null mutations in the ARID3B mouse model predispose to an increased incidence of cardiovascular malformations [18]. Similarly, homozygous loss-of-function mutations in STRA6 have also been associated with cardiovascular malformations, microphthalmia, and diaphragmatic hernias [19,20]. SIN3A and CSK both play roles in tumor suppression and their deletion may lead to an increased predisposition to and risk of neoplasia in this patient population [7].

Apparent reciprocal 15q24 duplications involving the SRO have been reported in 4 individuals [4,7,21]. Two of these individuals shared multiple clinical features
with one another, including developmental delay, hypertonia, joint limitations, digital abnormalities, and distinct facial features such as down-slanting palpebral fissures, epicanthal folds, smooth philtrum, and full lower lip [4,21]. However, other reported individuals with the reciprocal 15q24 duplication did not have the same clinical phenotype, therefore, it is unclear whether 15q24 duplications result in a distinct clinical phenotype [7,21].

Differential diagnoses

15q24 deletion syndrome should be differentiated from other microdeletion/duplication and genetic syndromes that have similar presentations. Individuals with 22q11.2 deletion syndrome (velo-cardio-facial syndrome) can have distinct facial features, asymmetric facies, and developmental delay. However, congenital heart defects and palate abnormalities are much more common in Noonan syndrome. In addition, obesity in older children with Noonan syndrome are more characteristic, whereas the facial features of individuals with 15q24 deletion may be rather subtle, especially in the newborn period. Failure to thrive and feeding difficulties, while common in both conditions, tends to be more severe in Noonan syndrome. Both conditions have a high incidence of skeletal malformations; however, pectus deformities are more common in Noonan syndrome, whereas digit abnormalities are predominant in 15q24 deletion syndrome. Molecular sequencing of genes in Ras-MAPK (mitogen-activated protein kinase) pathway will diagnose Noonan syndrome (and related conditions) in the majority of cases.

Genetic counseling

Variable expressivity exists between individuals with 15q24 deletion syndrome. Some of the variability may be attributed to the specific size and location of the deletion, however, even individuals with similar breakpoints exhibit diverse phenotypes. Offering accurate anticipatory guidance and counseling for individuals with 15q24 deletion syndrome can be challenging especially if the individual has additional genomic rearrangements that may predispose to other clinical consequences. This may interfere with the clinicians’ ability to provide an accurate assessment or outcome in the child. The phenotype, clinical features, and prognosis will depend on the size, location, and genes involved in the additional chromosome abnormality.

The combination of the additional rearrangements with the 15q24 deletion may have an additive effect as the presence of multiple genomic rearrangement in an individual (the genomic mutational load) can result in more severe phenotype and additional features that are not typically observed in individuals carrying single rearrangements [7,12,13,22]. Genetic counseling should take into account this potential effect and be attentive to additional phenotypic variability that may occur.

To date, all of the cases where parents were available for testing (18 out of 19 cases) indicated that the deletion occurred as a de novo event in the child. However, for genetic counseling, reproductive planning, and anticipatory guidance, it is recommended that both parents have FISH or aCGH testing in order to confirm that neither parent also carries the deletion. This is particularly important if the individual has an additional chromosome abnormality or imbalance that may have been inherited from a balanced rearrangement in one of the parents.

Individuals with Noonan syndrome may share some similarities to individuals with 15q24 deletion syndrome, particularly, distinct facial features, skeletal abnormalities, short stature, cryptorchidism, failure to thrive, and developmental delay. However, the facial features of individuals with Noonan syndrome are more characteristic, whereas the facial features of individuals with 15q24 deletion are rather subtle, especially in the newborn period. Failure to thrive and feeding difficulties, while common in both conditions, tends to be more severe in Noonan syndrome. Both conditions have a high incidence of skeletal malformations; however, pectus deformities are more common in Noonan syndrome, whereas digit abnormalities are predominant in 15q24 deletion syndrome. Molecular sequencing of genes in Ras-MAPK (mitogen-activated protein kinase) pathway will diagnose Noonan syndrome (and related conditions) in the majority of cases.
adolescence and adults with 15q24 deletion will help determine if the high incidence of genital abnormalities predisposes to fertility issues later in life.

**Antenatal diagnosis**

Deletion of 15q24 can be detected in amniotic fluid or chorionic villi samples. Since routine karyotyping is not typically sufficient to detect the deletion, aCGH should be offered to parents who carry a structural rearrangement that may predispose to 15q24 deletion or if fetal abnormalities are noted antenatally. Prenatal ultrasonography may detect some cardiovascular malformations, diaphragmatic hernias, and growth retardation in the fetus with 15q24 deletion, however, the most characteristic features of the condition, specifically distinct facial features and digit abnormalities, may be difficult to detect.

**Management**

The proposed management recommendations are based on the current knowledge of the small number of patients reported with this condition in conjunction with the authors' clinical experience, and are meant to serve as a guideline for treating individuals with this condition.

Routine medical management of individuals with 15q24 deletion syndrome and coordination of referrals can be performed by the individual’s primary care physician, however most individuals will require multidisciplinary care from a variety of subspecialists (Table 2). Initial presenting features may include failure to thrive, feeding difficulties, hypotonia, genital anomalies, and distinct facial features. Referral to a clinical geneticist is indicated in any child with a combination of these findings. Detailed physical examination, medical history, and developmental assessment should prompt genetic testing such as aCGH, which will diagnose microdeletions of 15q24. Periodic evaluation by a clinical geneticist is warranted to update the family on new information or symptoms that may be related to 15q24 deletion syndrome.

**Growth and Feeding**

The primary care physician should closely monitor all growth parameters in any individual diagnosed with 15q24 deletion syndrome. Referral to a gastroenterologist and/or feeding and nutrition clinic is warranted in any infant with 15q24 deletion syndrome and feeding difficulties, failure to thrive, or improper weight gain. Because of the risk of childhood obesity in individuals with 15q24 deletion syndrome, weight should be closely monitored even in those subjects with normal growth parameters. Nutritional counseling is recommended in those individuals who experience excessive weight gain. For children with growth retardation, evaluation by an endocrinologist for GH stimulation studies is also recommended. Although no data is available about the effect of GH treatment in individuals with 15q24 deletion, we would except that such treatment would be helpful especially in those with documented GH deficiency.

**Development and Behavior**

Given the high incidence of developmental delay, intellectual disability, and behavior problems in children with 15q24 deletion syndrome, formal neurocognitive and behavior assessment is recommended to determine the child’s specific learning pattern and behavioral

| Specialist                | Evaluation at time of diagnosis                                                                 | Screening and surveillance                                      |
|---------------------------|-------------------------------------------------------------------------------------------------|----------------------------------------------------------------|
| Primary care physician    | Initiate referrals to subspecialist, monitor growth and development                            | Assess growth parameters and development at each visit          |
| Development clinic        | Neurocognitive (developmental) and behavioral evaluation                                       | At time of diagnosis, then annually in childhood               |
| Geneticist                | Confirmation of diagnosis and periodic follow-up evaluations                                   | At time of diagnosis, then annually                           |
| Audiologist               | Routine audiology evaluation                                                                  | At time of diagnosis, then as needed                           |
| Ophthalmology             | Evaluate for coloboma, visual acuity, nystagmus, strabismus                                     | At time of diagnosis, then as needed                           |
| Cardiology                | Echocardiogram to evaluate for congenital heart defects                                        | At time of diagnosis, then as needed                           |
| Allergy and Immunology    | Immune work-up for recurrent and persistent infections                                         | As needed                                                      |
| Endocrine                 | Growth hormone studies for growth retardation                                                  | As needed                                                      |
| Gastroenterology/Fooding and Nutrition | Feeding assessment for failure to thrive and growth retardation                  | As needed                                                      |
| Neurology                 | Brain MRI for structural malformations and EEG for seizures                                   | As needed                                                      |
| Orthopedics               | Spine x-rays and scoliosis evaluation                                                          | As needed                                                      |
| Urology                   | Evaluate for genital abnormalities                                                            | As needed                                                      |
profile. Behavioral therapy may benefit individuals with this syndrome. Therefore periodic follow-up with a developmental and/or behavioral specialist is warranted. Early childhood intervention, encompassing physical therapy, occupational therapy, and speech therapy is needed in most of the cases.

**Neurologic manifestations**
As structural brain anomalies have been reported in approximately half of all cases of 15q24 deletion, brain magnetic resonance imaging (MRI) studies are indicated in any individual with 15q24 deletion and abnormal neurological examination. Referral to neurology is needed in individuals with structural brain malformations or if seizures are suspected.

**Cardiac manifestations**
A baseline echocardiogram is recommended in any individual with the new diagnosis of 15q24 deletion syndrome. Follow-up cardiology evaluations and echocardiograms should be performed based on the results of initial echocardiogram and at the discretion of the patient's cardiologist.

**Ophthalmologic manifestation**
Ophthalmologic abnormalities are common in individuals with 15q24 deletion syndrome; therefore, baseline ophthalmologic evaluations are recommended to screen for structural eye abnormalities and periodic ophthalmologic evaluations should be performed to screen for nystagmus and strabismus.

**Orthopedic manifestations**
Physical examination of the curvature of the spine needs to be evaluated with every routine care visit. Spine x-rays are indicated if the child shows any signs of scoliosis and/or kyphosis. If scoliosis and/or kyphosis develop then referral to orthopedics is warranted for follow up and discussing various treatment options.

**Urogenital abnormalities**
A majority of the males with 15q24 have genital abnormalities. Referral to urology for evaluation of the genital abnormalities, if present, is recommended in order to determine if surgical correction or any other treatment is indicated.

**Recurrent infections**
Individuals with 15q24 deletion syndrome are at risk for recurrent chest, upper airway, and ear infections. To date, documented immune deficiency has not been reported in any individual with 15q24 deletion syndrome, however, a referral to immunology may be warranted for recurrent and persistent infections. Referral to otolaryngology is recommended in case of recurrent otitis media and a low threshold is needed for placing pressure equalization (PE) tubes in children with 15q24 deletion syndrome. Recurrent otitis media may also lead to conductive hearing loss, which may affect a child’s expressive and receptive language development; therefore, periodic audiologic evaluation is also indicated.

**Prognosis**
The prognosis for individuals with 15q24 deletion syndrome depends on the severity and extent of congenital malformations. Major birth defects have been reported including, tetralogy of Fallot, myelomeningocele, intestinal atresia, and congenital diaphragmatic hernia (CDH). The prognosis for any individual with CDH is guarded and related to size, location, and visceral involvement of the diaphragmatic hernia [23]. In general, the majority of individuals with 15q24 deletion typically do not have life-threatening organ malformations. The natural history of 15q24 deletion syndrome is currently unknown since there are very few children and even fewer adolescents and adults reported with this deletion. The oldest reported patients, who were 33-years-old at the time of their report, have variable outcome [1,3]. The individual reported by Sharp et al. is a 33-year-old male who had mild developmental delay as a child. He has short stature and relatively mild intellectual disability with good language skills as an adult [1]. Van Esch et al. also reported a 33-year-old male who had severe developmental delay and intellectual disability, hypotonia, and limited speech [3]. He had hyperactive behavior with aggressive outbursts in childhood and adolescence and remained difficult to handle in adulthood despite psychopharmacological therapy. Longitudinal follow-up of individuals with this deletion will clarify the developmental and functional level of independence that individuals may expect to have in adulthood and possible health-related complications as individuals with this deletion age.

**Unresolved questions**
Since 15q24 deletion syndrome is a relatively newly described condition there are still many questions regarding the clinical manifestations that remain unanswered. Salient features that have been reported in a minority of individuals, but have the potential to drastically affect prognosis and anticipatory guidance of individuals with this condition, are the occurrence of neoplasias. At this time, it is unknown if there is a causal relationship between 15q24 deletion and the neoplasias described. Therefore, clinicians should be cognizant of the possible risk of tumors in this population and have low threshold for evaluation if neoplasia is remotely suspected.
The underlying explanation for the discrepancy in the male to female ratio of reported individuals remains unknown. As previously mentioned, this may represent an ascertainment bias rather than a true disparity. A more balanced ratio would be expected as more individuals with 15q24 deletion syndrome are identified.

Lastly, the clinical and phenotypic consequence of the reciprocal duplication of 15q24 remains unknown. Identification of more individuals with 15q24 duplication will enable better classification of clinical features that may be associated with the duplication and a better understanding of the molecular mechanisms that precipitate both the 15q24 deletion and reciprocal duplication.

**Future prospects**

Further characterization of the clinical phenotype, genotype-phenotype correlations, medical complications, and developmental potential of individuals with 15q24 deletion syndrome is imperative in order to fully comprehend the variability and severity of the syndrome. Natural history studies and longitudinal follow-up by the clinician will provide beneficial information on how children, adolescents, and adults with 15q24 deletion age and progress over time. This could lead to the development of practical screening, management, and surveillance guidelines for any individual diagnosed with 15q24 deletion syndrome.

**List of abbreviations**

aCGH: array comparative genomic hybridization; NAHR: non-allelic homologous recombination; LCR: low-copy repeats; SRO: smallest region of overlap; MRI: magnetic resonance imaging; GH: growth hormone; NHEJ: nonhomologous end joining; FoSTeS: fork stalling and template switching; MMBIR: microhomology-mediate break-induced replication; FISH: fluorescent in-situ hybridization; CNV: copy number variant; PWS: Prader-Willi syndrome; MAPK: mitogen-activated protein kinase; EEG: electroencephalogram; PE: pressure equalization; CDH: congenital diaphragmatic hernia.

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**Authors’ contributions**

Both authors participated in writing this review. They read and approved the final version of the manuscript.

**Competing interests**

The authors declare that they have no competing interests.

Received: 22 September 2011 Accepted: 4 January 2012 Published: 4 January 2012

**References**

1. Sharp AJ, Seizcer RR, Velthman JA, Gimelli S, Gimelli G, Striano P, Coppola A, Regan R, Price SM, Knoers NV, Eis PS, Brunner HG, Hennekam RC, Knight SJL, de Vries BBA, Zuffardi O, Eichler EE. Characterization of a recurrent 15q24 microdeletion syndrome. Human Molecular Genetics 2007, 16:567-572.

2. Kloppoki E, Graul-Neumann LM, Greben U, Tonnis H, Ropers H, Horn D, Mundlos S, Ullman R: A further case of the recurrent 15q24 microdeletion syndrome, detected by array CGH. Eur J Pediatr 2008, 167:903-908.

3. Van Esch H, Bacx L, Pijlens L, Fyns J. Congenital diaphragmatic hernia is part of the new 15q24 microdeletion syndrome. European Journal of Medical Genetics 2009, 52:153-156.

4. El-Hattab AW, Smolaérek TA, Walker ME, Schorry EK, Immenk LL, Patel G, Abbott M, Lanpher BC, Ou Z, Kang SL, Patel A, Scaglia F, Lupsiri JK, Cheung S, Tankiewicz P. Redefined genomic architecture in 15q24 directed by patient deletion/duplication breakpoint mapping. Hum Genet 2009, 125:589-602.

5. Andreiux J, Dubourg C, Rio M, Attie-Bitach T, Delaye Y, Mathieu M, Journel H, Copin H, Blondelle E, Doco-Fenzy M, Landais E, Delobel B, Odent S, Manouviour-Hanu S, Horder-Espinasse M: Genotype-phenotype correlation in four 15q24 deleted patients identified by array-CGH. Am J Med Genet Part A 2009, 149A:2813-2819.

6. Massrel-Paulet A, Callier P, Thouvin-Robinet C, Chouchane M, Meiner J, Marie N, Mosca A-L, Ben Salem D, Giroud M, Guiband L, Hueic F, Mugneret F, Fayev L. Multiple cysts of the corpus callosum and psychomotor delay in a patient with a 3.1 Mb 15q24.1q24.2 interstitial deletion identified by array-CGH. Am J Med Genet Part A 2009, 149A:1504-1510.

7. El-Hattab AW, Zhang F, Maxim R, Christensen KM, ward J, Hines-Dowell S, Scaglia F, Lupsiri JK, Cheung S: Deletion and duplication of 15q24: Molecular mechanisms and potential modification by additional copy number variants. Genetics in Medicine 2010, 12:573-586.

8. McInnes LA, Nakamine A, Pilorge M, Brandt T, Jiminez Gonzalez P, Fallas M, Manghi E, Edelmann L, Gleesner J, Hakonanones H, Betancur C, Buxbaum JD. A large-scale survey of the novel 15q24 microdeletion syndrome in autism spectrum disorders identified an atypical deletion that narrows the critical region. Molecular Autism 2010, 1:5.

9. Marshall CR, Noor A, et al: Structural variation of chromosomes in autism spectrum disorder. Am J Hum Genet 2008, 82:477-488.

10. Smith M, Filipek PA, Wu C, Bocian M, Hakim S, Modahl C, Spence MA: Analysis of a 1-megabase deletion in 15q22-23 in an autistic patient: Identification of candidate genes for autism and of homologous DNA segments in 15q22-q23 and 15q11-q13. Am J Hum Genet 2000, 66:765-770.

11. Cooper GM, Coe BP, Gintiraj S, Rosenfeld JA, Vu Th, Baker C, Williams C, Stalker H, Hamid R, Hanning V, Abdel-Hamid H, Bader P, McQueen E, Niyazov D, Leppig K, Thiese H, Hummel M, Alexander N, Gorski J, Kussman J, Shashi V, Johnson K, Rehder C, Ballaf BC, Shaffer LG, Eichler EE. A copy number variation morbidity map of developmental delay. Nature Genetics 2011, 43:838-846.

12. Potocki L, Chen KS, Koeuth T, Killian J, Iannaccone ST, Kashork CD, Stalker H, Hamid R, Hannig V, Abdel-Hamid H, Bader P, McQueen E, Niyazov D, Leppig K, Thiese H, Hummel M, Alexander N, Gorski J, Kussman J, Shashi V, Johnson K, Rehder C, Ballaf BC, Shaffer LG, Eichler EE. A copy number variation morbidity map of developmental delay. Nature Genetics 2011, 43:838-846.

13. Cooper GM, Coe BP, Gintiraj S, Rosenfeld JA, Vu Th, Baker C, Williams C, Stalker H, Hamid R, Hanning V, Abdel-Hamid H, Bader P, McQueen E, Niyazov D, Leppig K, Thiese H, Hummel M, Alexander N, Gorski J, Kussman J, Shashi V, Johnson K, Rehder C, Ballaf BC, Shaffer LG, Eichler EE. A copy number variation morbidity map of developmental delay. Nature Genetics 2011, 43:838-846.

14. Zhang F, Khajavi M, Connolly AM, Towne CF, Batish SD, Lupski JR: Analysis of a 1-megabase deletion in 15q22-23 in an autistic patient: Identification of candidate genes for autism and of homologous DNA segments in 15q22-q23 and 15q11-q13. Am J Hum Genet 2000, 66:765-770.

15. Cooper GM, Coe BP, Gintiraj S, Rosenfeld JA, Vu Th, Baker C, Williams C, Stalker H, Hamid R, Hanning V, Abdel-Hamid H, Bader P, McQueen E, Niyazov D, Leppig K, Thiese H, Hummel M, Alexander N, Gorski J, Kussman J, Shashi V, Johnson K, Rehder C, Ballaf BC, Shaffer LG, Eichler EE. A copy number variation morbidity map of developmental delay. Nature Genetics 2011, 43:838-846.

16. Hastings PJ, Ira G, Lupsiri JK: A microhomology-mediated break-induced replication model for the origin of human copy number variation. PLoS Genet 2009, 5:e1000327.

17. Hastings PJ, Lupsiri JK, Rosenberg SM, Ira G: Mechanisms of change in gene copy number. Nat Rev Genet 2009, 10:551-564.

18. Takebe A, Era T, Okada M, Martin J, Luri K, Kuruda Y, Nishikawa S: Microarray analysis of PDGFR alpha + populations in ES cell differentiation culture identifies genes involved in differentiation of mesoderm and mesenchyme including ARID1b that is essential for development of embryonic mesenchymal cells. Dev Biol 2006, 293:35-37.

19. Pasutto F, Sticht H, Hammersen G, Gillessen-Kaesbach G, Fitzpatrick DR, Nürnberg G, Brasch F, Schimer-Zimmermann H, Tolmie J, Chitayat D, Houge G, Fernández-Martinez I, Keating S, Mortier G, Hennekam RC, von der Weense A, Stavrotilak A, Meinecke F, Bitoun P, Becker C, Nürnberg P, Magoulas and El-Hattab Orphanet Journal of Rare Diseases 2012, 7:2 http://www.ojrd.com/content/7/1/2

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Reis A, Rauch A: Mutations in STRA6 cause a broad spectrum of malformations including anophthalmia, congenital heart defects, diaphragmatic hernia, alveolar capillary dysplasia, lung hypoplasia, and mental retardation. Am J Hum Genet 2007, 80:550-560.

20. Golizzi C, Martinovic-Bourel J, Thomas S, Mougou-Zrelli S, Grattagliano-Bessieres B, Bonnere M, Delahaye S, Munnich A, Encha-Razavi F, Lyonnet S, Vekemans M, Attie-Bitach T, Etchevers HC. Matthew-Wood syndrome is caused by truncating mutations in the retinol-binding protein receptor gene STRA6. Am J Hum Genet 2007, 80:1179-1187.

21. Kilholm Lund AB, Hove HD, Kirchhoff M: A 15q24 microduplication, reciprocal to the recently described 15q24 microdeletion, in a boy sharing clinical features with 15q24 microdeletion syndrome patients. Eur J Med Genet 2008, 51:520-526.

22. Girirajan S, Rosenfeld JA, et al: A recurrent 16p12.1 microdeletion supports a two-hit model for severe developmental delay. Nat Genet 2010, 42:203-209.

23. Datin-Dorriere V, Rouzies S, Taupin P, Walter-Nicolet E, Benachi A, Sonigo P, Mitanchet D: Prenatal prognosis in isolated congenital diaphragmatic hernia. Am J Obstet Gynecol 2008, 198:e1-80.e5.

doi:10.1186/1750-1172-7-2
Cite this article as: Magoulas and El-Hattab: Chromosome 15q24 microdeletion syndrome. Orphanet Journal of Rare Diseases 2012 7:2.