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### Published in:

American Journal of Human Genetics

### Publication status and date:

Published: 01/01/2005

### DOI (link to publisher):

[10.1086/429842](https://doi.org/10.1086/429842)

### Document Version

Publisher's PDF, also known as Version of record

### Citation for the published version (APA):

Klaassens, M., van Dooren, M., Eussen, HJFMM., Douben, H., Dekker, A., van der Lee, C., Donahoe, PK., Galjaard, R.-J., Goemaere, NNT., de Krijger, R., Wouters, C., Wauters, J., Oostra, B., Tibboel, D., & de Klein, A. (2005). Congenital Diaphragmatic hernia and chromosome 15q26: Determination of a candidate region by use of fluorescent in situ hybridization and array-based comparative genomic hybridization. *American Journal of Human Genetics*, 76(5), 877-882.  
<https://doi.org/10.1086/429842>

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## Report

# Congenital Diaphragmatic Hernia and Chromosome 15q26: Determination of a Candidate Region by Use of Fluorescent In Situ Hybridization and Array-Based Comparative Genomic Hybridization

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Congenital diaphragmatic hernia (CDH) has an incidence of 1 in 3,000 births and a high mortality rate (33%–58%). Multifactorial inheritance, teratogenic agents, and genetic abnormalities have all been suggested as possible etiologic factors. To define candidate regions for CDH, we analyzed cytogenetic data collected on 200 CDH cases, of which 7% and 5% showed numerical and structural abnormalities, respectively. This study focused on the most frequent structural anomaly found: a deletion on chromosome 15q. We analyzed material from three of our patients and from four previously published patients with CDH and a 15q deletion. By using array-based comparative genomic hybridization and fluorescent in situ hybridization to determine the boundaries of the deletions and by including data from two individuals with terminal 15q deletions but without CDH, we were able to exclude a substantial portion of the telomeric region from the genetic etiology of this disorder. Moreover, one patient with CDH harbored a small interstitial deletion. Together, these findings allowed us to define a minimal deletion region of ~5 Mb at chromosome 15q26.1–26.2. The region contains four known genes, of which two—*NR2F2* and *CHD2*—are particularly intriguing gene candidates for CDH.

Congenital diaphragmatic hernia (CDH [MIM 142340]), a severe, life-threatening, congenital anomaly characterized by variable defect in the diaphragm, pulmonary hypoplasia, and postnatal pulmonary hypertension, is a relatively common anomaly (Torfs et al. 1992; Beresford and Shaw 2000). CDH can occur as an isolated defect, in combination with multiple congenital anomalies, or as part of a defined syndrome (Fryns et al. 1979; Enns et al. 1998). Little is known about the etiology of CDH. However, there is increasing evidence for a genetic cause of CDH. Various chromosomal anomalies have been described in CDH, with numerical abnormalities (such as

trisomies 13, 18, and 21) as the most common type. Structural chromosomal anomalies, involving almost every chromosome, have been reported (Lurie 2003). Since 1988, clinical data from all Erasmus Medical Centre patients with CDH have been stored in a database. Cytogenetic data from 200 patients with CDH were available, and 24 patients (12%) showed an abnormality. Fourteen patients (7%) showed a numerical abnormality (trisomy 18 or 21). The remaining 10 patients (5%) had a structural anomaly, and 3 of those patients (1.5%) were shown to have a deletion of part of chromosome 15q. The present study focuses on this subset of patients with CDH and chromosome 15q deletions. Data from four previously published patients with CDH and small chromosome 15q deletions were generously made available to us (table 1) (de Jong et al. 1989; Rosenberg et al. 1992; Chen et al. 1998; Schlembach et al. 2001). Clinical evaluation of the seven patients revealed a left-sided diaphragmatic hernia of the Bochdalek type, in-

Received December 9, 2004; accepted for publication February 21, 2005; electronically published March 4, 2005.

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0002-9297/2005/7605-0015\$15.00

**Table 1****Summary of Clinical and Cytogenetic Data for Patients with Deletions Involving Chromosome 15**

Patient	Karyotype	CDH <sup>a</sup>	Other Abnormalities <sup>b</sup>	Deletion Size (Mb)
1	46,XY,t(1;14),inv(6),del(15)(q26)	Yes	Genital anomalies; IUGR	6.3
2	46,XY,r(15)(p11q26)	Yes	Dysmorphic features; cardiac, renal, genital, and limb abnormalities; IUGR	19.9
3	46,XY,r(15)(p11q26)	Yes	Dysmorphic features; cardiac abnormalities; IUGR	23.3
4	46,XX,der(15)t(3;15)(q29;q26.1)	Yes	Dysmorphic features; cardiac and limb abnormalities; two-vessel umbilical cord; IUGR	22.8
5	46,XY,r(15)(p11q26.1)	Yes	Dysmorphic features; genital and limb abnormalities; IUGR	23
6	46,XX,der(15)t(8;15)(q24.1;q26.1)	Yes	Hydrocephaly; dysmorphic features; cardiac, renal, limb, and spine abnormalities; IUGR	16.8
7	46,XX,del(15)(q25q26.3)	Yes	Dysmorphic features; renal and limb abnormalities; IUGR	22.3
8	46,XX,r(15)(p11.1q26.3)	No	Mental retardation; mild dysmorphic features; IUGR	16.3
9	46,XY,del(15)(q26)	No	Mental retardation	15.6

NOTE.—Sources for patients 1–3 and 8, Erasmus Medical Centre, Rotterdam, The Netherlands; patient 4, Rosenberg et al. 1992 (case 2); patient 5, de Jong et al. 1989; patient 6, Chen et al. 1998; patient 7, Schlembach et al. 2001; patient 9, J. Wauters, University Hospital Antwerpen, Antwerpen, Belgium.

<sup>a</sup> Left-sided, Bochdalek-type CDH (if present).

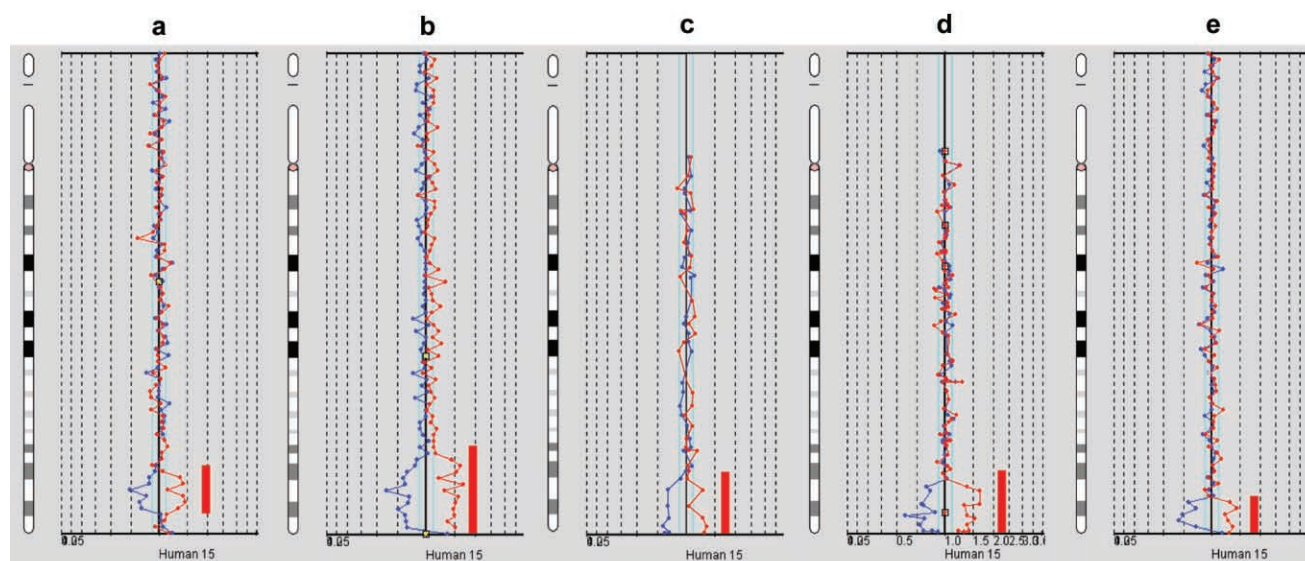
<sup>b</sup> IUGR = intrauterine growth retardation (birth weight <3rd percentile).

trauterine growth retardation (all patients had birth weights <3rd percentile), and multiple other congenital anomalies, such as cardiac and renal abnormalities. In addition to the seven patients with CDH, we analyzed data from two patients with mental retardation and 15q deletions but without CDH manifestation. Genomic DNA was extracted from cultured cells (from patients 1, 2, 3, 4, and 9) or paraffin-embedded tissue (from patients 5 and 6).

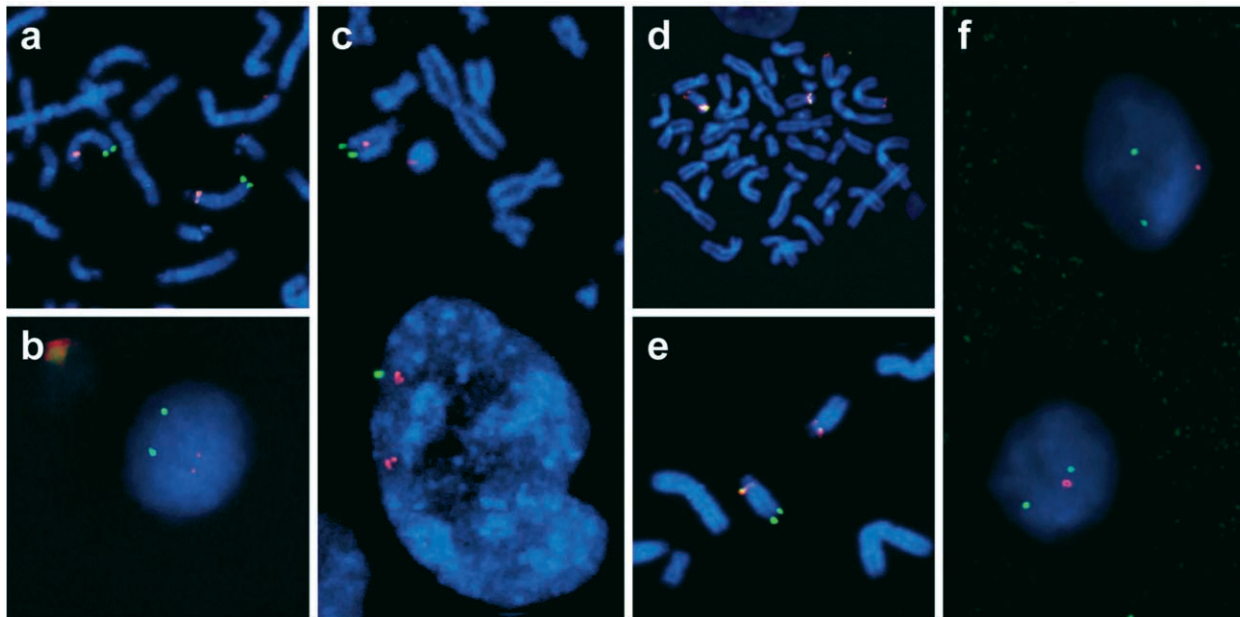
To delineate the possible candidate region, array-

based comparative genomic hybridization (array CGH) was performed using the 1-Mb Human BAC Array (Spectral Genomics) in accordance with the manufacturer's instructions. A dye-swap experimental strategy was used as an additional internal control. Fluorescent signals on the arrays were visualized using the ScanArray Express HT scanner. Images were analyzed with Spectral Ware 2 (Spectral Genomics). Results for patients 1, 3, 4, 7, and 8 are shown in figure 1.

To further delineate the deleted region and to deter-



**Figure 1** Array CGH results. *a*, Patient 1, with CDH and del(15) interstitial deletion. *b*, Patient 3, with CDH and r(15)(p11q26). *c*, Patient 4, with CDH and der(15)t(3;15)(q29;q26.1). *d*, Patient 7, with CDH and del(15)(q25q26.3). *e*, Patient 8, without CDH and with r(15)(p11.1q26.3).



**Figure 2** FISH results. *a*, Patient 1: partial metaphase, probe D15Z4 (red signal) at chromosome 15 centromeric locus and probe RP11-114I24 (green signal) at 15q26.3. *b*, Patient 2: interphase, probe RP11-369K8 (red signal) and RP11-253B9 (green signal) near the chromosome 5 centromeric region at 5p13.2. *c*, Patient 3: partial metaphase and interphase, deletion probe RP11-143C19 (green signal) and normal probe RP11-64K10 (red signal) at 15q23. *d*, Patient 4: metaphase spread, gain of chromosome 3q29; probe RP1-196F4 (red signal) (3qtel) present on der(15) and normal signal probe D15Z4 (yellow/red signal) at the centromeric region of chromosome 15. The der(15) contains both signals. *e*, Patient 4: partial metaphase, deletion probe RP11-183E24 (green signal) at 15q26.2 and normal probe D15Z4 (yellow/red signal). *f*, Patient 6: interphase, deletion probe RP11-57P19 (red signal) and normal probe D15Z4 (green signal). Patients 1–6 all have CDH.

mine the deletions in patients 4, 5, and 9, ~110 BAC clones were selected from the University of California Santa Cruz (UCSC) and Ensembl genome browsers to cover the distal part of chromosome 15. Using the appropriate BAC clones, we performed FISH on metaphase chromosomes from patients 1, 3, and 4 (fig. 2*a* and 2*c–e*). Interphase FISH was performed on nuclei extracted from paraffin-embedded tissues from patients 2, 5, and 6 (fig. 2*b* and 2*f*; data for patient 5 not shown). Only genomic DNA was available from patients 7 and 8, so the size of the deletion in these patients was determined using only array CGH. FISH slides were analyzed using the Axioplan 2 Imaging microscope (Zeiss), and images were collected using the Isis Software System (Metasystems). Combining FISH and array CGH data, we were able to approximately determine the breakpoints in all patients (fig. 3). In patient 1, the interstitial deletion found by array CGH was confirmed and was narrowed to a 6-Mb deletion between BAC clones RP11-79A7 and RP11-616M17. In patient 2, the deletion extended from a region distal to RP11-79A7; in patient 3, it extended from a region distal to RP11-300G22. The proximal breakpoint in patient 4 lies within BAC clone RP11-617F23. In patient 5, the break occurs distal to RP11-300G22. In patient 6, the most distal probe tested that was present on the deleted chromosome 15 was RP11-

79A7. The terminal deletion in patient 7 occurs distal to RP11-360F18. The proximal breakpoints of the deletion interval in the two patients without CDH were similarly determined (fig. 3). In the first patient without CDH, who had a ring chromosome 15, the most distal BAC clone tested that was present on the ring chromosome was RP11-120N1. In the second patient without CDH, the most distal BAC clone present was RP11-262P8. Combining all data, we determined the smallest common deletion interval in patients with CDH to be at 15q26.1–26.2 (which we have termed the “CDH region”). This region is ~5 Mb in size and is bordered by BAC clones RP11-79A7 and RP11-80F4.

CDH is unlike many genetic disorders, for which candidate genes can be determined by using linkage analyses of familial cases, because the vast majority of CDH cases occur de novo. For this type of disorder, the best way to determine which genes are involved is by analyzing a large number of patients for common aberrations by use of different high-resolution genetic methodologies, such as FISH or array CGH. This strategy has already been used successfully to identify genes involved in CHARGE syndrome (MIM 214800) (Vissers et al. 2004) and Cornelia de Lange syndrome (CdLS [MIM 122470]) (Krantz et al. 2004; Tonkin et al. 2004).

Isolated reports of distal chromosome 15 deletions



*TFII*"), is a member of the steroid/thyroid hormone receptor subfamily and is involved in retinoic acid metabolism (Kimura et al. 2002). A knockout mouse model of *NR2F2* showed that *Nr2f2*<sup>-/-</sup> mice are not viable and die at E9 in utero because of arrest of cardiac development (Pereira et al. 1999). Heterozygous knockout mice have poor viability in the neonatal period and are smaller than wild-type mice. However, the exact cause of death in these mice is not clear. The second interesting gene is the chromodomain helicase 2 gene (*CHD2* [MIM 602119]), a member of the SNF2/RAD54 helicase gene family. Recently, mutations in another member of this family (*CHD7*) have been found to cause CHARGE syndrome (Vissers et al. 2004). The third gene in the CDH region is the repulsive guidance molecule gene, *RGMA* (MIM 607362), which is involved in the guidance of growth cones in developing neurons (Brinks et al. 2004). This gene is not known to play a role in diaphragm development, nor has it been described in muscle or lung development. The fourth gene located in the smallest region of overlap is the sialyltransferase 8B gene (*SIAT8B* [MIM 602546]), which plays a role in the adhesive properties of neural cell adhesion molecules (Angata et al. 1997).

Haploinsufficiency due to the loss of a copy of one of these genes may be enough to result in a diaphragmatic defect. At the present time, the precise mechanism by which a deletion of or within one of these genes or a related gene mutation causes this developmental defect can only be speculated.

Elsewhere, other genes on chromosome 15q have been suggested as being involved in the pathogenesis of diaphragmatic defects. Biggio et al. (2004) suggested that the myocyte enhancer factor 2A gene, *MEF2A* (MIM 600660), could be involved in the pathogenesis of diaphragmatic defects. *MEF2A* maps to 15q26.3 and is involved in the differentiation of muscle cells from their precursors. Some genes involved in vitamin A metabolism—for example, *RALDH2* (MIM 603687), which maps to 15q21—have also been implicated in the pathogenesis of CDH (Greer et al. 2003). Both *MEF2A* and *RALDH2* are located outside our candidate critical CDH region, which limits their possible role in CDH in our subgroup of patients.

In conclusion, we have mapped a potential critical CDH region to 5 Mb at chromosome 15q26.1-26.2, a region that contains four genes, of which two are especially intriguing candidates in the etiology of diaphragmatic defects. Further research is needed to confirm their exact role in CDH and to determine the pathogenic mechanism. As a first step, we are performing FISH and mutation analyses of a large group of patients with CDH who have normal karyotypes. In the future, prenatal screening for 15q abnormalities when a diaphragmatic hernia is detected could give better clues for

predicting the outcome and could provide more information for genetic counseling.

## Acknowledgments

We thank C. P. Chen, G. de Jong, D. Schlembach, G. Stetten, and B. Ceulemans, for kindly providing us with data from patients with chromosome 15 deletions; T. de Vries Lentsch, for assistance with manuscript preparation; B. Pober, for critical reading of the manuscript; and M. Listewnik, for array CGH technical assistance. This research was funded in part by the Sophia Foundation for Scientific Research, Rotterdam, The Netherlands (SSWO project 441), and National Institute of Health program project grant HD39942 (to P.K.D. and C.L.).

## Electronic-Database Information

The URLs for data presented herein are as follows:

Ensembl Genome Browser, [http://www.ensembl.org/Homo\\_sapiens/](http://www.ensembl.org/Homo_sapiens/)

Online Mendelian Inheritance in Man (OMIM), <http://www.ncbi.nlm.nih.gov/Omim/> (for CDH, CHARGE syndrome, CdLS, Fryns syndrome, *NR2F2*, *CHD2*, *RGMA*, *SIAT8B*, *MEF2A*, and *RALDH2*)

UCSC Genome Browser, <http://genome.cse.ucsc.edu/>

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