The genetic landscape and clinical implications of vertebral anomalies in VACTERL association

Yixin Chen, Zhenlei Liu, Jia Chen, Yuzhi Zuo, Sen Liu, Weisheng Chen, Gang Liu, Guixing Qiu, Philip F Giampietro, Nan Wu, Zhihong Wu

ABSTRACT
VACTERL association is a condition comprising multi-system congenital malformations, causing severe physical disability in affected individuals. It is typically defined by the concurrence of at least three of the following component features: vertebral anomalies (V), anal atresia (A), cardiac malformations (C), tracheo-oesophageal fistula (TE), renal dysplasia (R) and limb abnormalities (L). Vertebral anomaly is one of the most important and common defects that has been reported in approximately 60–95% of all VACTERL patients. Recent breakthroughs have suggested that genetic factors play an important role in VACTERL association, especially in those with vertebral phenotypes. In this review, we summarised the genetic studies of the VACTERL association, especially focusing on the genetic aetiology of patients with vertebral anomalies. Furthermore, genetic reports of other syndromes with vertebral phenotypes overlapping with VACTERL association are also included. We aim to provide a further understanding of the genetic aetiology and a better evidence for genetic diagnosis of the association and vertebral anomalies.

OVERVIEW OF VACTERL ASSOCIATION
VACTERL association is a condition with multi-system congenital malformations: Vertebral anomalies (V), anal atresia (A), cardiac malformation (C), tracheo-oesophageal fistula (TE) with or without oesophageal atresia, renal dysplasia (R) and limb abnormalities (L). It was first named as VATER (without ‘C’) and ‘L’ association in 1973. The prevalence of VACTERL/VATER association is between 1/7000 and 1/40 000. As there is no available objective laboratory test for its diagnosis, VACTERL association is diagnosed totally based on the clinical manifestations mentioned above. Most clinicians and researchers require the presence of at least three component features for diagnosis. Besides, due to its heterogeneous phenotype and the abundance of overlapping defects of other syndromes, VACTERL association is typically considered a diagnosis of exclusion with no clear evidence for an alternative or overlapping diagnosis such as Coloboma, Heart anomaly, Atrioventricular canal defects, Retarded mental and somatic development, Genital hypoplasia, Ear anomalies (CHARGE) syndrome, DiGeorge syndrome and Pallister–Hall syndrome. The presence of other features not typically seen in VACTERL association may suggest other disorders. Thus, a physical examination and family history are essential to rule out potentially overlapping diagnoses. It is worth mentioning that 5–10% patients with Fanconi anaemia (FA) have birth defects meeting the diagnosis of VACTERL association with hydrocephalus (VACTERL-H). It is suggested that FA with VACTERL-H should be treated separately from the VACTERL association because of the core characteristics of FA such as haematological anomalies and skin pigmentation changes, the different frequencies of VACTERL-associated phenotypes and the prognosis and therapeutic intervention.

Although the clinical criteria for VACTERL association appear to be straightforward, the overlapping either in clinical manifestation or genetic finding is challenging for clinicians and geneticists. The CHD7 gene mutation, which is proved to be associated with CHARGE syndrome, may also be found in patients diagnosed with VACTERL association, even CHARGE syndrome is clinically excluded. Besides, most of the conditions listed are monogenic disorders. Careful genetic evaluation may help ruling out these conditions. In this review, we listed the related monogenic diseases that share two more overlapping manifestations and their genetic findings (table 1). We propose that (1) these syndromes as well as these candidate genes should be considered in diagnostic and genetic studies in VACTERL association; and (2) VACTERL syndrome remains a diagnosis of exclusion following a thoughtful clinical evaluation and consideration of genetic testing for overlapping syndromes. Prior studies have estimated that 90% of the patients diagnosed with VACTERL association had three or fewer phenotypes (referred to as VACTERL-like association) and <1% of patients had all six anomalies. Although the frequency of the six clinical features (CFs) varies, vertebral anomalies is the most common observation in many cohorts of VACTERL association, which have been reported in approximately 60–95% of affected individuals. Additionally, vertebral anomalies are the most prevalent findings in the first-degree relatives of the probands in some cohorts, thus highlighting the importance of vertebral anomalies as a major diagnostic feature for VACTERL association. In this review, we will summarise the genetic studies of the VACTERL association with an emphasis on vertebral anomalies.

Vertebral anomalies
Vertebral anomalies in VACTERL association can be classified as (1) failure of formation, such as hemivertebrae, butterfly or wedge-shaped vertebrae; (2) failure of segmentation such as vertebral bars, fused vertebrae and block vertebrae; and (3) a combination of these two features, resulting in a mixed deformity. Rib anomalies such as...
Table 1  Monogenic diseases overlapping with VACTERL association

<table>
<thead>
<tr>
<th>Syndrome</th>
<th>OMIM</th>
<th>Locus</th>
<th>Gene</th>
<th>Vertebral anomalies</th>
<th>Overlap malformations</th>
<th>Characteristic features beyond VACTERL association</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fanconi anaemia with VACTERL-H</td>
<td>227650;</td>
<td>300514</td>
<td>FANCA; FANCB, etc.*</td>
<td>Same phenotype with VACTERL but lower frequency</td>
<td>V, A, C, TE, R, L</td>
<td>Haematological anomalies; pigmentary changes; hydrocephalus</td>
<td>Holden et al13</td>
</tr>
<tr>
<td>Alagille syndrome</td>
<td>118450</td>
<td></td>
<td>JAG1; NOTCH2</td>
<td>Mostly butterfly vertebra, occasionally hemivertebrae, fusion of vertebrae</td>
<td>V, C, R</td>
<td>Jaundice with conjugated hyperbilirubinemia; dysmorphic facies; posterior embryotoxon and retinal pigmentary changes</td>
<td>Turnpenny and Ellard14</td>
</tr>
<tr>
<td>Basal cell nevus syndrome</td>
<td>109400</td>
<td>9q22;</td>
<td>PTCH1; PTCH2; SUFU</td>
<td>Multiple fusion of vertebral bodies and ribs</td>
<td>V, L</td>
<td>Odontogenic keratocysts of the jaw; palmar or plantar pits; bilamellar calcification of the falx cerebri; basal cell tumours</td>
<td>Oostra and Maas;15 Pino et al16</td>
</tr>
<tr>
<td>Baller–Gerold syndromes</td>
<td>218600</td>
<td>8q24</td>
<td>RECLQ4</td>
<td>Rib fusion and flat vertebrae</td>
<td>V, A, C, R, L</td>
<td>Craniosynostosis; microcephaly</td>
<td>Murthy et al17</td>
</tr>
<tr>
<td>DiGeorge syndrome (22q11.2 deletion syndrome)</td>
<td>188400</td>
<td>22q11</td>
<td>TBX1</td>
<td>Hemivertebrae</td>
<td>V, C, R, L</td>
<td>Thymic abnormality; conotruncal cardiac anomaly; facial dysmorphism; hypocalcaemia</td>
<td>Tsirikos et al;18 Maggadottir and Sullivan19</td>
</tr>
<tr>
<td>Feingold syndrome</td>
<td>164280</td>
<td>2p23-24</td>
<td>N-MYC</td>
<td>Absence of the fifth sacral vertebra and fusion of C5–C7 in a case</td>
<td>V, C, TE, R, L</td>
<td>Microcephaly; brachymesophalangy</td>
<td>Celi et al20</td>
</tr>
<tr>
<td>McKusick–Kaufman syndrome</td>
<td>236700</td>
<td>20p12</td>
<td>MKKS</td>
<td>Vertebral anomalies in one case</td>
<td>V, C, L</td>
<td>Hydrometrocolpos; gastrointestinal malformations</td>
<td>Knowles et al21</td>
</tr>
<tr>
<td>CHARGE syndrome</td>
<td>214800</td>
<td>8q12</td>
<td>CHD7</td>
<td>Idiopathic scoliosis without vertebral anomalies</td>
<td>C, TE, R</td>
<td>Coloboma; choanal atresia/stenosis/cryptophthalmosplasia of semicircular, etc.</td>
<td>Hsu et al;22 Verloes17</td>
</tr>
<tr>
<td>Pallister–Hall syndrome</td>
<td>146510</td>
<td>7p14.1</td>
<td>GLI3</td>
<td>NA</td>
<td>A, C, R, L</td>
<td>Hypothalamic hamartoma; bifid epiglottis; craniofacial abnormalities</td>
<td>Demurger et al23</td>
</tr>
<tr>
<td>Townes–Brocks syndrome</td>
<td>107480</td>
<td>16q21.1</td>
<td>SALL1</td>
<td>NA</td>
<td>A, C, R, L</td>
<td>Dysplastic ears with hearing impairment; intellectual disability</td>
<td>Sudo et al24</td>
</tr>
<tr>
<td>Holt–Oram syndrome</td>
<td>142900</td>
<td>12q24</td>
<td>TBX5</td>
<td>NA</td>
<td>C, L</td>
<td>NA</td>
<td>Goldfarb and Wall201426</td>
</tr>
<tr>
<td>Hemifacial microsomia (DAVS)</td>
<td>164210</td>
<td>14q32</td>
<td>NA</td>
<td>Hemivertebrae, fusion of vertebrae</td>
<td>V, C</td>
<td>Craniofacial anomalies; central nervous system defects: visual and hearing impairment</td>
<td>Beleza-Meireles et al27</td>
</tr>
<tr>
<td>TAR syndrome</td>
<td>274000</td>
<td>1q21</td>
<td>RBMBANA</td>
<td>NA</td>
<td>C, R, L</td>
<td>Thrombocytopenia</td>
<td>Tassanoet al28</td>
</tr>
</tbody>
</table>

*Numbers of genes been implicated in the pathogenesis associated with Fanconi anaemia.29

A, anal atresia; C, cardiac malformations; CHARGE, Coloboma, Heart anomaly, Atresia of choanae, Retardation of mental and somatic development, Genital hypoplasia, Ear abnormalities; L, limb abnormalities; NA, not available; DAVS, ouzo-austilo-vertebral spectrum; R, renal anomalies; TAR, thrombocytopenia-absent radius; TE, tracheo-oesophageal fistula; V, vertebral anomalies; VACTERL, vertebral anomalies (V), anal atresia (A), cardiac malformations (C), tracheo-oesophageal fistula (TE), renal dysplasia (R) and limb abnormalities (L); VACTERL-H, VACTERL association with hydrocephalus.

GENETIC STUDIES ON VACTERL ASSOCIATION

The aetiology of VACTERL association is not well understood (figure 2). As its phenotypes are too heterogeneous to be defined as a syndrome, and there is no major gene for this condition, thus it is still referred to as an ‘association’. The familial clustering phenomenon suggests a genetic role in its causality.24 41 42

X-linked VACTERL association by ZIC3 mutation

So far, the ZIC3 gene has been demonstrated to cause X-linked VACTERL association. Different types of ZIC3 mutations, including point mutations, deletions and polyalanine expansion, have been reported to be responsible for both VACTERL or VACTERL-like association.43 44 45 Cardiac defects are most commonly found as ZIC3 has important function in cardiac development and mutations in ZIC3 also cause X-linked heterotaxy (MIM#306953);43 44 46 47 anal atresia is present in most patients with ZIC3 mutations; vertebral anomalies are not commonly observed and demonstrated phenotypic variability.45 In animal models, Zic3 knockout mice mimic the human heterotaxy and cardiac phenotype with occasional vertebral/rib anomalies. Zic3 expression was present at all stages of embryonic

as rib fusion and increased or decreased number of ribs are commonly accompanied with vertebral anomalies. In some studies, rib anomalies may occur without vertebral anomalies.7 10 38 19 Although patients with anorectal malformations may have dysplastic sacral vertebra, it is not clear whether these should be regarded as a vertebral anomalies component for diagnosis of VACTERL syndrome.2 Clinical signs of scoliosis or kyphosis may be the first sign of vertebral anomalies when VACTERL association is suspected.40 Radiology is needed for discerning vertebral and rib anomalies.

As an example, we present a 2-year-old Chinese boy with VACTERL association. He was born with oesophageal atresia between L3 and L4, and a right hemivertebra between L5 and S1 (figure 1), which caused evident lumbar scoliosis. He also had an extra thoracic vertebra and an extra pair of ribs without clinical symptoms. Abdominal ultrasound examination revealed horseshoe kidney without impairment of his renal function. He underwent resection of both hemivertebrae with internal fixation and recovered well postoperatively.

development within the anterior pre-somitic mesoderm but not in the developing anal region. Thus, anal atresia was not reported in Zic3-deficient mice, which differs from humans where anal atresia is also prevalent with ZIC3 mutations.

**Sonic hedgehog pathway in VACTERL association**

**SHH** gene has been implicated as the key inductive signal in patterning of the ventral neural tube, the anterior–posterior limb axis and the ventral somites. Studies on animal models indicate that sonic hedgehog (Shh) pathway is important for VACTERL association. Kim *et al.* identified the first animal model that recapitulated the human VACTERL syndrome by knocking out genes (**Shh** and Gli) in Shh pathway. With different genes of the Shh signalling pathway affected, the mutant mice display various combinations, ranges and severity of the VACTERL phenotypes, implying a dosage-dependent effect. Furthermore, a VACTERL-like phenotype was reported in murine with a novel hypomorphic mutation in the **Ift172** gene. The **Ift172** gene encodes a component of the intratflagellar transport, which appears to play an active role in Shh signalling, and Ift proteins are required for both Gli activator and Gli repressor function.

To the best of our knowledge, **SHH** or **GLI3** mutations have not been identified in VACTERL patients. In humans, **SHH** mutation may cause more severe VACTERL phenotypes. Nowaczyk *et al.* reported a patient with holoprosencephaly 3 and **SHH** haploinsufficiency who suffered from sacral anomalies (cleft S1, hemivertebra at S2 and absence of the rest of the sacrum and coccyx), genitourinary abnormality, multiple segments of bowel atresia and limb anomalies. Although this patient has a distinctive diagnosis, the phenotypic features overlap with VACTERL association. There is a possibility that **SHH** mutation causes these overlapping phenotypes.
Some genes that play roles in Shh pathway have been reported to be associated with VACTERL association. A heterozygous de novo 21bp deletion (c.163_183del) in the exon 1 of the HOXD13 gene, which is a downstream target of SHH, was identified in a 17-year-old girl, who was diagnosed with VACTERL association without vertebral anomalies. Another patient with rib anomalies diagnosed with VACTERL association was found with a 451 kb deletion at chromosome 3q28, which contains a single LPP gene. This gene encodes LIM domain containing preferred translocation partner in lipoma that has been shown to bind PEA3, an ETS domain transcription factor that has a role in regulating the SHH pathway. Moreover, CNV (micro-deletions) as well as point mutation in FOXF1 gene have been identified in patients with VACTERL phenotypes. In animal models, Foxf1 has been found to be downregulated in Shh−/− mice and the Foxf1 heterozygotes have been shown to display tracheo-oesophageal atresia and fistulas. Although HOXD13, LPP and FOXF1 mutation were sporadic findings in individuals, these studies argue in favour of that SHH pathway dysfunction is associated with VACTERL association.

### Candidate gene mutations and CNVs

Several candidate gene mutations and CNVs have been reported to be related to VACTERL association (summarised in Table 2). So far, these candidate gene mutations and CNVs listed are found mostly in sporadic cases, which need further large sample verification or functional experiments to confirm their pathogenicity.

Although the genetic aetiology of VACTERL association has been far from established, previous studies did reveal some genetic mutations that can account for one or a few of the six CFs (Table 2). For example, DLL3 gene, which encodes a ligand for the Notch signalling pathway that coordinates somitogenesis, has been found to cause block vertebrae in a Caucasian male VACTERL patient. Saisawat et al. identified recessive mutations in the TNF receptor-associated protein 1 (TRAP1) gene in three families with VACTERL association. They also proved that Trap1 gene is highly expressed in the renal epithelia of 13.5-day-old mouse embryos and its mutations contribute to renal dysplasia.

Intriguingly, mutations of the same gene may cause variable expressivity among VACTERL patients, even within the same

### Table 2 Candidate genes and CNVs in VACTERL association

<table>
<thead>
<tr>
<th>Chromosome region</th>
<th>Gene</th>
<th>Mutation</th>
<th>Function</th>
<th>Inheritance</th>
<th>Manifestations</th>
<th>Vertebral anomalies</th>
<th>Overlap syndrome</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>1q41</td>
<td>–</td>
<td>–</td>
<td>Duplication</td>
<td>De novo</td>
<td>V, A, C, TE, R</td>
<td>Butterfly vertebrae</td>
<td>–</td>
<td>Hilger et al. 60</td>
</tr>
<tr>
<td>8q24.3</td>
<td>–</td>
<td>–</td>
<td>Duplication</td>
<td>De novo</td>
<td>V, A, C, TE, R</td>
<td>Butterfly vertebrae</td>
<td>–</td>
<td>Hilger et al. 61</td>
</tr>
<tr>
<td>19q13.2</td>
<td>DLL3</td>
<td>p.G269A</td>
<td>Missense</td>
<td>Heterozygous (inherited-mat)</td>
<td>V, C, R, L</td>
<td>Block vertebrae</td>
<td>–</td>
<td>Ueda et al. 64</td>
</tr>
<tr>
<td>13q33.2-qter</td>
<td>–</td>
<td>–</td>
<td>Deletion</td>
<td>De novo</td>
<td>V, A</td>
<td>Block vertebrae</td>
<td>–</td>
<td>Dworschak et al. 65</td>
</tr>
<tr>
<td>22q11.2</td>
<td>–</td>
<td>–</td>
<td>Duplication</td>
<td>De novo</td>
<td>V, A, R</td>
<td>Fusion vertebrae (L4-L5)</td>
<td>22q11.2 duplication syndrome; DiGeorge syndrome</td>
<td>Schramm et al. 66</td>
</tr>
<tr>
<td>Y</td>
<td>–</td>
<td>–</td>
<td>Deletion in Yq and duplication in Yq</td>
<td>NA</td>
<td>V, A, R, L</td>
<td>Block and hemivertebrae in lumbar</td>
<td>–</td>
<td>Bhagat 76</td>
</tr>
<tr>
<td>18q10-q11.2</td>
<td>–</td>
<td>–</td>
<td>Duplication</td>
<td>De novo</td>
<td>V, A, R, L</td>
<td>Dyplastic lumbar and sacral vertebrae, NO detail</td>
<td>–</td>
<td>Felix et al. 77, van der Veken et al. 78</td>
</tr>
<tr>
<td>10q23.31</td>
<td>PTEN</td>
<td>p.H61D</td>
<td>Missense</td>
<td>De novo</td>
<td>V, C, TE, L</td>
<td>Rib anomalies (13 pairs of ribs)</td>
<td>Cowden syndrome</td>
<td>Reardon et al. 69</td>
</tr>
<tr>
<td>3q28</td>
<td>LPP</td>
<td>–</td>
<td>Deletion</td>
<td>De novo</td>
<td>V, C, TE, R</td>
<td>Rib anomalies</td>
<td>–</td>
<td>Arrington et al. 39, Hernandez-Garcia et al. 80</td>
</tr>
<tr>
<td>5q11.2</td>
<td>–</td>
<td>–</td>
<td>Deletion</td>
<td>De novo</td>
<td>V, A, C</td>
<td>No detail</td>
<td>–</td>
<td>de Jong et al. 79</td>
</tr>
<tr>
<td>19p13.3</td>
<td>–</td>
<td>–</td>
<td>Deletion</td>
<td>De novo/inserted-mat</td>
<td>V, A, C, TE, R, L</td>
<td>No detail</td>
<td>–</td>
<td>Peddibhotla et al. 81</td>
</tr>
<tr>
<td>2q31.1</td>
<td>HOXD13</td>
<td>–</td>
<td>Deletion</td>
<td>De novo</td>
<td>A, C, L</td>
<td>Not reported</td>
<td>Brachydactyly-syndactyly syndrome</td>
<td>Garcia-Barcelo et al. 82</td>
</tr>
<tr>
<td>10q24.32</td>
<td>FGFB</td>
<td>p.G29_R34dup; p.P26L</td>
<td>In-frame duplication; missense</td>
<td>Heterozygous</td>
<td>A, C, TE, R, L</td>
<td>Not reported</td>
<td>Kallmann syndrome</td>
<td>Zeidler et al. 83</td>
</tr>
</tbody>
</table>

*Four cases of TRAP1 mutations have been reported and the only case with vertebral anomalies is listed.

A, anal atresia; ACD/MPV, alveolar capillary dysplasia with misalignment of pulmonary veins; C, cardiac malformations; L, limb abnormalities; NA, not available; R, renal anomalies; TE, tracheo-oesophageal fistula; V, vertebral anomalies; VACTERL, vertebral anomalies (V), anal atresia (A), cardiac malformations (C), tracheo-oesophageal fistula (TE), renal dysplasia (R) and limb abnormalities (L).
family. Dworschak et al\textsuperscript{69} identified chromosome 13q deletions in two patients with VACTERL phenotypes. The girl was born with perineal fistula, renal hypoplasia, bilateral triphalangeal thumbs and oligodactyly, butterfly vertebrae and cerebral anomalies, and died at 10 months of age. The second patient, a male child, suffered from perineal fistula, block vertebrae at C2–C3 and C4–C5–C6 and bilateral hearing loss. \textit{Pcsk5} gene has been identified as a candidate gene of VACTERL association in mice.\textsuperscript{70} Nakamura et al\textsuperscript{71} reported a Japanese VACTERL boy with eighth thoracic hemivertebra having a frameshift mutation of \textit{PCSK5}, while his healthy father also shared the same mutation. Peddibhotla et al\textsuperscript{72} reported eight patients with chromosome 19p13.3 microdeletions and six of them fulfilled the diagnostic criteria for VACTERL association. Among the six VACTERL patients, one patient has vertebral anomalies while her two children, although with VACTERL association, are free from vertebral anomalies. These phenomena imply other modification factors desperate for further investigation in this condition.

**Chromosomal aberrations**

Chromosomal aberrations also contribute to VACTERL association. Several case reports have been published that describe chromosomal anomalies in VACTERL patients as Felix et al\textsuperscript{73} and Brosens et al\textsuperscript{74} reviewed previously. However, chromosomal aberrations are not included here as they also contribute to the occurrence of congenital malformations beyond what is typically observed in VACTERL association.

**Mitochondrial dysfunction**

Damian et al\textsuperscript{82} first reported an A to G transversion in the mitochondrial NP3243 mutation in cystic kidney of a VACTERL child. Spinal radiograph showed multiple cervical and thoracic vertebral wedging, fusion and fission. She also had limb abnormalities, cardiac malformations and renal anomalies. This child belonged to a family in which other members had mitochondrial encephalomyopathy, lactic acidosis, and stroke-like episodes syndrome and chronic progressive external ophthalmoplegia, which suggests mitochondrial dysfunction may contribute to VACTERL syndrome.\textsuperscript{83} Stone et al\textsuperscript{84} studied a cohort of 62 patients with VACTERL association and none of the affected children had measurable levels of the NP 3243 mutation. A few authors have previously reported an association of VACTERL association in patients with mitochondrial disorders known as complex IV respiratory chain deficiency.\textsuperscript{85–87} Overall, four of the five individuals presented with vertebral anomalies; three showed oesophageal involvement; two had anal atresia and two patients presented with additional minor dysmorphic features. Different combinations of other multiple congenital malformations have also been reported in a series of children with respiratory chain deficiency, leading to the hypothesis that in these patients congenital anomalies might result from an abnormal development during embryogenesis through either a lack of ATP or an alteration of apoptosis controlled by the mitochondrial machinery. However, it is also possible that mitochondrial dysfunction and congenital malformations in the patient described here are both secondary to an as yet unidentified process.\textsuperscript{88} In conclusion, whether mutation of mitochondrial dysfunction causes VACTERL association is still controversial. Some clinical signs and symptoms that may be not common in patients with VACTERL association, including progressive muscle weakness, characteristic patterns of cardiac, neurologic and exocrine dysfunction,\textsuperscript{89} may suggest a potential existence of mitochondrial dysfunction.

In summary, the aetiology of VACTERL association appears to be heterogeneous, suggesting that it may be a complex condition. Besides the gene mutations and CNVs mentioned above, some other factors such as intrinsic mutations or epigenetic factors may also play important roles in this condition. Environmental factors including maternal diabetes\textsuperscript{90} and exposure to statins,\textsuperscript{91} which may associated with congenital anomalies, may play a significant role in the pathogenesis of VACTERL syndrome.

**CONCLUSION**

VACTERL association is a rare and complex condition with highly heterogeneous aetiology and manifestations. At the present time, there appears to be evidence for genetic factors contributing to VACTERL syndrome including single-gene mutations, CNVs and structure variants to mitochondrial dysfunction. Future studies are needed to identify epigenetics and environmental causes for VACTERL syndrome. Targeted genetic testing can contribute to eliminating overlapping diagnoses from further consideration in an affected individual. Notably, a given variant may explain a particular CF of VACTERL association, so it may be worth trying to investigate this sophisticated association by focusing on one of the six component features. ‘Vertebral anomalies’ is one of the core component features of VACTERL association, including formation and segmentation vertebral. Wu et al\textsuperscript{85} recently described a compound heterozygous model in which a null allele mutation in combination with a common haplotype of \textit{TBX6} causes congenital scoliosis, suggesting that genetic factors play an important role in vertebral anomalies. Additionally, we suggest that the genetic mutations may contribute to vertebral anomalies in a certain syndrome. Alternatively, VACTERL association may be caused by a ‘two-hit’ model in which two genes or one gene in combination with an epigenetic factor may elicit all associated features.\textsuperscript{93} In the future, combination of new genomic technologies such as whole-exome sequencing, whole-genome sequencing, comparative genomic hybridisation array and whole-genome bisulphite sequencing may well reveal a surprising number of additional contributing loci, delineating the entire spectrum of the VACTERL association in humans.

**Acknowledgements** The authors thank Dr Pengfei Liu from the Department of Molecular and Human Genetics, Baylor College of Medicine, and Dr Xiaoyue Wang from the State Key Laboratory of Medical Molecular Biology, Chinese Academy of Medical Sciences, for their comments on the manuscript. They also express gratitude to the patient described in this article for his willingness to take part in this study.

**Contributors** YC, ZL and JC contributed equally to this article. WC and NW conceived and designed the review. YC interpreted data and contributed to the manuscript preparation. YC, ZL and JC drafted the main manuscript. SL, WC and GL contributed to data acquisition. All authors approved the final manuscript.

**Funding** The research was supported by National Natural Sciences Foundation of China (81501852, 81472046, 81271942, 81130034, 81472045), Distinguished Young Scholars of Peking Union Medical College Hospital (U2010506), Beijing nova program (2016) and The Central Level Public Interest Program for Science Research Institute (No 13, 2015).

**Competing interests** None declared.

**Ethics approval** Ethics Committee of Peking Union Medical College Hospital.

**Provenance and peer review** Commissioned; externally peer reviewed.

**Open Access** This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/4.0/

**REFERENCES**


Hosie S, Holland-Cunz S, Wijers CH, Marcelis CL, van Rooij IA, Hildebrandt F, Whole-exome resequencing reveals recessive mutations in TRAP1 in individuals with CAKUT and VACTERL association.


Belloni E. Holoprosencephaly, sacral anomalies, and situs ambiguus in an infant with mutation in the transcription factor PEA3.


The genetic landscape and clinical implications of vertebral anomalies in VACTERL association

Yixin Chen, Zhenlei Liu, Jia Chen, Yuzhi Zuo, Sen Liu, Weisheng Chen, Gang Liu, Guixing Qiu, Philip F Giampietro, Nan Wu and Zhihong Wu

J Med Genet published online April 15, 2016

Updated information and services can be found at:
http://jmg.bmj.com/content/early/2016/04/15/jmedgenet-2015-103554

These include:

References
This article cites 93 articles, 11 of which you can access for free at:
http://jmg.bmj.com/content/early/2016/04/15/jmedgenet-2015-103554
#BIBL

Open Access
This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/4.0/

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Topic Collections
Articles on similar topics can be found in the following collections

Open access (191)
Oesophagus (28)

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/