Association of Damaging Variants in Genes With Increased Cancer Risk Among Patients With Congenital Heart Disease

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Importance

Patients with congenital heart disease (CHD), the most common birth defect, have increased risks for cancer. Identification of the variables that contribute to cancer risk is essential for recognizing patients with CHD who warrant longitudinal surveillance and early interventions.

Objective

To compare the frequency of damaging variants in cancer risk genes among patients with CHD and control participants and identify associated clinical variables in patients with CHD who have cancer risk variants.

Design, Setting, and Participants

This multicenter case-control study included participants with CHD who had previously been recruited to the Pediatric Cardiac Genomics Consortium based on presence of structural cardiac anomaly without genetic diagnosis at the time of enrollment. Permission to use published sequencing data from unaffected adult participants was obtained from 2 parent studies. Data were collected for this study from December 2010 to April 2019.

Exposures

Presence of rare (allele frequency, \(<1 \times 10^{-5}\)) loss-of-function (LoF) variants in cancer risk genes.

Main Outcomes and Measures

Frequency of LoF variants in cancer risk genes (defined in the Catalogue of Somatic Mutations in Cancer–Cancer Gene Consensus database), were statistically assessed by binomial tests in patients with CHD and control participants.

Results

A total of 4443 individuals with CHD (mean [range] age, 13.0 [0-84] years; 2225 of 3771 with reported sex [59.0%] male) and 9808 control participants (mean [range] age, 52.1 [1-92] years; 4967 of 9808 [50.6%] male) were included. The frequency of LoF variants in regulatory cancer risk genes was significantly higher in patients with CHD than control participants (143 of 4443 [3.2%] vs 166 of 9808 [1.7%]; odds ratio [OR], 1.93 [95% CI, 1.54-2.42]; \(P = 1.38 \times 10^{-12}\)), and among CHD genes previously associated with cancer risk (58 of 4443 [1.3%] vs 18 of 9808 [0.18%]; OR, 7.2 [95% CI, 4.2-12.2]; \(P < 2.2 \times 10^{-16}\)). The LoF variants were also nominally increased in 14 constrained cancer risk genes with high expression in the developing heart. Seven of these genes (\(ARHGEF12\), \(CTNNB1\), \(LPP\), \(MLLT4\), \(PTEN\), \(TCF12\), and \(TFRC\)) harbored LoF variants in multiple patients with unexplained CHD. The highest rates for LoF variants in cancer risk genes occurred in patients with CHD and extracardiac anomalies (248 of 1482 individuals [16.7%]; control: 1099 of 9808 individuals [11.2%]; OR, 1.59 [95% CI, 1.37-1.85]; \(P = 1.3 \times 10^{-10}\)) and/or neurodevelopmental delay (209 of 1393 individuals [15.0%]; control: 1099 of 9808 individuals [11.2%]; OR, 1.40 [95% CI, 1.19-1.64]; \(P = 9.6 \times 10^{-6}\)).

Conclusions and Relevance

Genotypes of CHD may account for increased cancer risks. In this cohort, damaging variants were prominent in the 216 genes that predominantly encode regulatory proteins. Consistent with their fundamental developmental functions, patients with CHD and damaging variants in these genes often had extracardiac manifestations. These data may also implicate cancer risk genes that are repeatedly varied in patients with unexplained CHD as CHD genes.
The growing population of adults with congenital heart disease (CHD) has created increased recognition of additional health issues, including a 1.4-fold to 2-fold higher cancer prevalence than in the general population. While radiation exposure from therapeutic interventions can increase cancer risk (CR), the diversity of malignant conditions outside of radiation fields suggests other risk factors. Additional risks and mechanisms that link CHD to cancer are unknown.

Damaging gene variants contribute to both CHD and cancer, hinting that these disorders share molecular relationships. This model is supported by the increased prevalence of damaging variants in CR genes among children with developmental delays, including autism and intellectual disabilities, which occur in some patients with CHD. To explore potential molecular relationships, we analyzed rare loss-of-function (LoF) variants in CR genes among a large CHD cohort and defined accompanying clinical features. These analyses identify patients with CHD and the highest CR gene burden, who may warrant longitudinal cancer screening.

**Methods**

**Study Participants and Ethical Approval**
The multicenter case-control study was reviewed and approved by the relevant institutional review boards, including at Boston Children’s Hospital. Written informed consent was obtained at the time of enrollment. We studied participants in the Pediatrics Cardiovascular Genetics Consortium with undefined causes for CHD at the time of enrollment and unaffected control participants in studies of autism and schizophrenia (eTable 1 in the Supplement).

**CR Genes**
The Catalogue of Somatic Mutations in Cancer–Cancer Gene Consensus database defines 723 CR genes (eTable 2 in the Supplement). This includes 38 CR genes that also cause CHD (eTable 3 in the Supplement and Online Mendelian Inheritance in Man [OMIM]; https://omim.org/), 216 CR genes that regulate RNA transcription or processing, and 227 CR genes with LoF mechanisms, including 107 genes with LoF germ-line mechanisms.

**Variant Calls and Statistical Analyses**
Whole-exome sequences from patients with CHD and control participants were processed using established pipelines to identify rare (allele frequency, $\leq 1 \times 10^{-5}$) heterozygous LoF variants. All $P$ values reflect binomial tests after Bonferroni correction with a $P$ value threshold of $1.67 \times 10^{-3}$ (10 gene lists and 3 comparisons). A false discovery rate $P < .05$ was used as the significant threshold throughout. The eMethods in the Supplement includes further methodological details. Data were collected for this study from December 2010 to April 2019. The software program R version 3.6.0 (R Foundation for Statistical Computing) was used for analysis.

**Results**

**LoF Variants in CR Genes Among Patients With CHD**
Initial analyses of LoF variants in CR genes (eTable 4 and eMethods in the Supplement) and prespecified subsets (Table 1) demonstrated significantly higher frequencies in patients with CHD who were randomly assigned to the discovery group ($n = 2222$) or the replication group ($n = 2221$) in comparisons with independent control cohorts ($n = 3578$ and $n = 6230$; eTables 5 and 6 in the Supplement). As such, we present combined data from 4443 patients with CHD and 9808 control participants (Table 1). Patients with CHD ranged in age from 0 to 84 years, with a mean (SD) age of 13.0 (14.8) years; 2225 of 3771 patients with CHD with a reported sex (59.0%) were male. Control participants ranged in age from 21 to 92 years, with a mean (SD) age of 52.1 years, and 4967 of 9808 participants (50.6%) were male.

The presence of CR variants was not associated with any specific CHD subtype (eTable 7 in the Supplement). Most patients with CHD ($599$ of $642$ [93.3%]) had a single CR variant, while a minority had 2 CR variants ($43$ of $642$ [6.7%]) or 3 CR variants ($3$ of $642$ [0.5%]). No individual had 2 independent CR variants in the same gene. Analyses restricted to participants of European ancestry (CHD: 448 of 3106 individuals [14.4%]; controls: 849 of 9501 individuals [8.9%]) remained significant (odds ratio [OR], 1.72 [95% CI, 1.52-1.94]; $P < 2.2 \times 10^{-16}$; eTable 8 in the Supplement). Because many CR genes are associated with adult-onset malignant conditions, we compared LoF variant frequencies in patients with CHD who were younger than 16 years ($n = 3338$) or older than 16 years ($n = 1105$). A higher proportion of older individuals had LoF variants, albeit with comparable ORs (1.33-1.37; eTable 9 in the Supplement).

Thirty-eight CR genes have dominant patterns of transmission for CHD (denoted OMIM; Table 1); these genes had significantly more LoF variants in patients with CHD than control participants (OR, 7.19 [95% CI, 4.23-12.22]; $P < 2.2 \times 10^{-16}$). The presence of LoF variants was also increased among CR genes with regulatory functions (OR, 1.93 [95% CI, 1.54-2.42]; $P = 1.38 \times 10^{-12}$), a prominent feature of many CR and
CHD genes. Genes found in both categories (OMIM and regulatory; n = 17) had the highest frequency of LoF variants in individuals with CHD (OR, 9.89 [95% CI, 4.80-20.40]; P = 2.2 × 10^{-16}). The CR genes with dominant variants that cause cancer by haploinsufficiency (n = 46) were also enriched in those with CHD (OR, 2.36 [95% CI, 1.60-3.47]; P = 3.45 × 10^{-8}; Table 1).

### Comorbidities in Patients With CHD and LoF Variants in CR Genes

Damaging CR variants are increased in individuals with developmental delays.\(^3\) Because these delays can occur with CHD, we partitioned patients into those with extracardiac anomalies (ECA; n = 1482), neurodevelopmental defects (NDD; n = 1393), both ECA and NDD (n = 878), and neither ECA or NDD (isolated CHD; n = 1379). The LoF variants in CR genes were highest among patients with CHD and ECA (CHD: 248 of 1482 individuals [16.7%]; control: 1099 of 9808 individuals [11.2%]; OR, 1.59 [95% CI, 1.37-1.85]; P = 1.3 × 10^{-10}; eTable 9 in the Supplement) and patients with CHD and NDD (CHD: 209 of 1393 individuals [15.0%]; control: 1099 of 9808 individuals [11.2%]; OR, 1.40 [95% CI, 1.19-1.64]; P = 9.6 × 10^{-5}), while the LoF in CR genes in patients with isolated CHD was comparable with that of control participants. Notably, 10 genes had LoF variants in 3 or more patients with CHD and NDD or CHD and ECA (supporting cohort size for statistics: CHD: 1482 individuals with ECA and 1393 with NDD; control: 9808 individuals; P < .05), including 2 CR regulatory genes (catenin beta 1 [CTNNB1]; n = 1393 individuals with CHD and ECA, 2 with CHD and NDD, and 0 control participants) and transcription factor 12 [TCF12]; n = 1393 individuals with CHD and NDD, 2 patients with CHD and ECA, and 2 control participants; eTable 10 in the Supplement). However, we observed no significant functional enrichment in genes with LoF variants within each CHD group (eMethods in the Supplement).

### CR Genes as Candidate CHD Genes

Given the dual role of some genes in both CHD and cancer, we considered if some CR genes might contribute to CHD. When considering only patients with CHD without pathogenic variants in OMIM CHD genes (n = 4293), we identified significantly more LoF variants in affected individuals (576 in 4293 [13.4%]) than control participants (1080 of 9744 [11.1%]; OR, 1.2 × 10^{-6}; OR, 1.24 [95% CI, 1.12-1.39]; eTable 9 in the Supplement). Moreover, 7 CR genes that are LoF intolerant (probability of LoF intolerance by the Genome Aggregation Database, >0.50) and highly expressed in the developing heart had LoF variants in 3 or more patients with unexplained CHD (rho guanine nucleotide exchange factor 12 [ARHGEF12], CTNNB1, lipoma-preferred partner [LPP], afadin [MLLT4], phosphatase and tensin homolog [PTEN], TCF12, and transferrin receptor [TFRC] genes; Table 2).

### Discussion

In a large CHD cohort, we demonstrated an increased prevalence of LoF variants in CR genes, particularly those with established roles in CHD or that regulate gene expression (Table 1). Patients with CHD and variants in the CR genes were more likely to have ECA and/or NDD (eTable 9 in the Supplement), thereby prioritizing patients for prospective studies to determine if clinical outcomes validate CR. Importantly, we identified no increase rates of LoF in CR genes among patients with isolated CHD. Additionally, our data (Table 2) indicate that LoF variants in some CR genes contribute to CHD, prioritizing new molecules for mechanistic studies in heart development.

Recent genetic analyses\(^6\) have demonstrated that many CHD genes function in regulating the epigenome and gene transcription and translation—processes that are critical to orchestrating cardiac progenitor cell proliferation, lineage commitment, and differentiation. These genes broadly participate in human development and harbor the highest rates of damaging variants among patients with CHD and ECA or NDD.\(^6\) Stem cells, with germline and somatic variants in genes with similar key functions, cause cancer.\(^13\) That subsets of patients with CHD (with ECA or NDD; eTable 9 in the Supplement) had the highest burden of LoF variants in these CR genes emphasizes the broad expression patterns and shared molecular mechanisms for some developmental

### Table 1. Patients With Congenital Heart Disease (CHD) and Rare Loss-of-Function (LoF) Variants in Subsets of the Catalogue of Somatic Mutations in Cancer–Cancer Gene Consensus Cancer Risk Genes

<table>
<thead>
<tr>
<th>Genes</th>
<th>No.</th>
<th>Patients with CHD</th>
<th>Control participants</th>
<th>Odds ratio (95% CI)</th>
<th>Binomial P valuea</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total participants</td>
<td>14251</td>
<td>4443</td>
<td>9808</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>All cancer risk</td>
<td>723</td>
<td>642</td>
<td>1099</td>
<td>1.34 (1.21-1.49)</td>
<td>2.31 × 10^{-11}</td>
</tr>
<tr>
<td>OMIM CHDa</td>
<td>38</td>
<td>68</td>
<td>18</td>
<td>7.19 (4.23-12.22)</td>
<td>&lt;2.2 × 10^{-16}</td>
</tr>
<tr>
<td>Regulatory</td>
<td>216</td>
<td>143</td>
<td>166</td>
<td>1.93 (1.54-2.42)</td>
<td>1.38 × 10^{-12}</td>
</tr>
<tr>
<td>Regulatory and OMIM CHD</td>
<td>17</td>
<td>40</td>
<td>9</td>
<td>9.89 (4.80-20.40)</td>
<td>&lt;2.2 × 10^{-16}</td>
</tr>
<tr>
<td>Regulatory without OMIM CHD</td>
<td>199</td>
<td>103</td>
<td>157</td>
<td>1.46 (1.13-1.88)</td>
<td>2.01 × 10^{-4}</td>
</tr>
<tr>
<td>Non-OMIM CHD</td>
<td>685</td>
<td>585</td>
<td>1082</td>
<td>1.22 (1.10-1.36)</td>
<td>5.245 × 10^{-6}</td>
</tr>
<tr>
<td>Nonregulatory</td>
<td>507</td>
<td>516</td>
<td>942</td>
<td>1.24 (1.10-1.39)</td>
<td>5.46 × 10^{-6}</td>
</tr>
<tr>
<td>LoF cancer mechanism</td>
<td>227</td>
<td>240</td>
<td>376</td>
<td>1.43 (1.21-1.69)</td>
<td>1.58 × 10^{-7}</td>
</tr>
<tr>
<td>Recessive LoFc</td>
<td>135</td>
<td>158</td>
<td>274</td>
<td>1.28 (1.05-1.57)</td>
<td>1.68 × 10^{-3}</td>
</tr>
<tr>
<td>Dominant LoFd</td>
<td>46</td>
<td>53</td>
<td>50</td>
<td>2.36 (1.60-3.47)</td>
<td>3.45 × 10^{-4b}</td>
</tr>
<tr>
<td>Non-LoF cancer mechanism</td>
<td>496</td>
<td>422</td>
<td>751</td>
<td>1.27 (1.12-1.43)</td>
<td>4.46 × 10^{-6}</td>
</tr>
</tbody>
</table>

Abbreviations: NA, not applicable; OMIM, Online Mendelian Inheritance in Man.

a Bonferroni significance P value threshold: 1.67 × 10^{-4} (10 gene lists by 3 comparisons).

b Significant in both subanalyses.

c The OMIM CHD genes with dominant patterns of transmission.

d Five genes have both dominant and recessive cancer variants; 51 are not characterized.
Table 2. Seven Cancer Risk Genes With Multiple Loss-of-Function (LoF) Variants in Patients With Unexplained Congenital Heart Disease (CHD)

<table>
<thead>
<tr>
<th>Gene</th>
<th>CHD LoF</th>
<th>Non-CHD LoF</th>
<th>pLI</th>
<th>Heart expression rank</th>
<th>Human CHD gene</th>
<th>CR regulatory gene</th>
<th>Known gene syndrome/phenotype</th>
<th>No. of patients with CHD and ECA phenotype data</th>
<th>CHD phenotypes (No. of patients)</th>
<th>No. of patients with CHD and ECA phenotypes</th>
<th>Proportion of patients with CHD and NDD, No./total No.</th>
</tr>
</thead>
<tbody>
<tr>
<td>KDR</td>
<td>7</td>
<td>0</td>
<td>0.98</td>
<td>87</td>
<td>No</td>
<td>No</td>
<td>Hemangioma</td>
<td>1 LVOTO (1) and ToF (2)</td>
<td>1 With microcephaly, micrognathia, inguinal hernia, cryptorchidism, and hydrocephalus</td>
<td>1/1</td>
<td></td>
</tr>
<tr>
<td>TCF12</td>
<td>5</td>
<td>2</td>
<td>0.97</td>
<td>89</td>
<td>No</td>
<td>Yes</td>
<td>Craniosynostosis</td>
<td>5 ASD (1), Ebstein anomaly (1), and LVOTO (3)</td>
<td>2 With bitemporal narrowing (1), abdominal heterotaxy (1), absent corpus callosum and seizure disorder (1), and/or neonatal AML (1)</td>
<td>3/3</td>
<td></td>
</tr>
<tr>
<td>CTNNB1</td>
<td>3</td>
<td>0</td>
<td>1</td>
<td>99</td>
<td>No</td>
<td>Yes</td>
<td>Neurodevelopmental disorder</td>
<td>3 ASD (1), dilated tricuspid valve (1), and ToF (1)</td>
<td>3 With microcephaly (2), strabismus (2), asthma (1), micrognathia (1), congenital scoliosis (1), and/or hypotonia (2)</td>
<td>2/2</td>
<td></td>
</tr>
<tr>
<td>ARHGEF12</td>
<td>3</td>
<td>0</td>
<td>1</td>
<td>91</td>
<td>No</td>
<td>No</td>
<td>None</td>
<td>3 ASD (2) and truncus arteriosus (1)</td>
<td>1 With hearing loss</td>
<td>3/3</td>
<td></td>
</tr>
<tr>
<td>MLLT10</td>
<td>3</td>
<td>0</td>
<td>1</td>
<td>85</td>
<td>No</td>
<td>Yes</td>
<td>None</td>
<td>2 ASD or VSD (1) and LVOTO (1)</td>
<td>None with ECA</td>
<td>1/1</td>
<td></td>
</tr>
<tr>
<td>PTEN</td>
<td>3</td>
<td>1</td>
<td>0.98</td>
<td>78</td>
<td>No</td>
<td>No</td>
<td>Cowden</td>
<td>3 ASD (2) and VSD (1)</td>
<td>2 With macrocephaly (2), airway malacia (1), congenital scoliosis and abnormal vertebrae (1), Chiari malformation and hydrocephalus (1), and/or Cowden features (1)</td>
<td>2/2</td>
<td></td>
</tr>
<tr>
<td>TFRC</td>
<td>3</td>
<td>1</td>
<td>0.78</td>
<td>82</td>
<td>No</td>
<td>No</td>
<td>Immunodeficiency</td>
<td>3 Pulmonary atresia/stenosis (2) and ToF (1)</td>
<td>2 With frontal bossing (1), and/or intestinal atresia and VATER association (1)</td>
<td>1/1</td>
<td></td>
</tr>
</tbody>
</table>

Abbreviations: AML, acute myelogenous leukemia; ARHGEF12, rho guanine nucleotide exchange factor 12 gene; ASD, atrial septal defect; CTNNB1, catenin beta 1 gene; ECA, extracardiac anomaly; KDR, kinase insert domain receptor gene; LVOTO, left ventricular outflow tract obstruction; MLLT10, histone lysine methyltransferase DOT1L cofactor gene; NDD, neurodevelopmental delay; pLI, probability of loss-of-function intolerance; PTEN, phosphatase and tensin homolog gene; TCF12, transcription factor 12 gene; TFRC, transferrin receptor gene; ToF, tetralogy of Fallot; VATER, vertebral, anal, tracheal, esophageal, and abdominal heterotaxy.

Biallelic variants are often required for oncogenesis, whereas 1 damaging variant is often sufficient to perturb cardiac morphogenesis. This raises the possibility that damaging variants in patients with CHD set the stage but do not directly initiate oncogenesis. Increased CR likely reflects additional, amplifying factors, including radiation and other CHD-associated exposures. Consistent with this model, we note that dominant MLL2/KMT2D variants cause CHD, while biallelic loss occurs in many cancers. We propose that CR is increased because the germline variant provides the first of 2 hits needed for cancer to emerge. Across this CHD cohort, 3.6% had LoF variants in recessive CR genes, while 1.8% had LoF variants in dominant CR genes. We speculate that high lifetime radiation doses might increase somatic variants that complement CHD LoF variants.
tients with CHD and help determine causal associations, as well as whether associations are modified by nongenetic factors. Control participants were considerably older, potentially introducing survivor bias that would enhance the burden of CR variants in patients with CHD. Depth of sequencing precluded differentiation of variants as germline or high-level mosaic. We analyzed only LoF variants since defining missense variants as damaging requires detailed functional assessments, and thus our data provide only a conservative estimate of CR variants in patients with CHD. This also precluded assessment of burden for CR genes that operate by gain-of-function mechanisms in cancer, such as protein tyrosine phosphatase non-receptor type 11 (PTPN11) and other RASopathy genes.14

Conclusions

Decades of therapeutic progress enable long-term survival for newborns with CHD, and current estimates indicate 6 in 1000 adults are survivors of CHD.1 The recognition that CHD genotypes influence CR can promote clinical surveillance and early interventions and further promote lifelong health in adult patients with CHD. Additionally, we suggest that mechanistic studies into the molecular and cellular processes that are disrupted by damaging variants in CHD and cancer genes may uncover insights that inform new treatment strategies for both disorders.

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