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Genotype–Phenotype Correlations in Pathology Caused by Collagen Type IV alpha 1 and 2 Mutations

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Abstract

COL4A1 and COL4A2 are extracellular matrix proteins that form heterotrimers and are present in nearly all basement membranes in every organ. In the past decade, *COL4A1* and *COL4A2* mutations have been identified to cause a multi-system disorder for which penetrance and severity of constituent phenotypes can greatly vary. Here, we compare the outcomes of more than 100 mutations identified in patients and data from a murine allelic series to explore the presence of genotype-phenotype correlations – many of which are shared among other types of collagen. We find that there is a frequency bias for *COL4A1* over *COL4A2* mutations and that glycine (Gly) substitutions within the triple helical domain are the most common class of mutations. Glycine is most often replaced by a charged amino acid, however the position of the mutation, and not the properties of the substituting amino acid, appear to have a greater influence on disease severity. Moreover, the impact of position is not straightforward. Observations from a murine allelic series suggest that mutations in the NC1 domain may result in relatively mild phenotypes via a ‘quantitative’ mechanism similar to other types of collagens, however, this effect was not apparent in human reports. Importantly, other position-dependent effects had differential impacts depending on the phenotype of interest. For example, the severity of cerebrovascular disease correlated with an amino-to-carboxy severity gradient for triple-helical glycine substitutions whereas the penetrance and severity of myopathy and nephropathy appear to involve a functional sub-domain(s). Greater understanding of genotype-phenotype correlations and the interaction of consequences of different mutations will be important for patient prognosis and care and for developing mechanism-based therapeutics to treat individual components of this emerging syndrome.

1) Introduction

Type IV collagens are network-forming basement membrane collagens¹ and are encoded by six genes; collagen, type IV, alpha 1 (*COL4A1*) through 6 (*COL4A6*) in mammals^{2,3}.

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COL4A1 and COL4A2 are present in almost all basement membranes, while the distributions of COL4A3, COL4A4, COL4A5 and COL4A6 are restricted to a smaller number of basement membranes present in the eye, ear and kidney ^{3, 4, 5}. In humans, *COL4A1* and *COL4A2* share a common locus at 13q34 where they are arranged head to head with a bidirectional promoter ⁶⁻⁹. *COL4A1* (52 exons) and *COL4A2* (48 exons) encode proteins of 1669 and 1712 amino acids, respectively, which share 45% identity. COL4A1 and COL4A2 both contain the same three major domains: The non-collagenous (NC1) domain is a globular domain at the carboxy terminus that is required for the initiation of heterotrimer formation between one COL4A2 and two COL4A1 proteins ($\alpha 1(\text{IV})_2 \alpha 2(\text{IV})$) ^{10, 11}. The majority of each COL4A1 and COL4A2 protein comprises a collagenous triple helical domain characterized by repetition of Gly-Xaa-Yaa motifs (where Xaa and Yaa are variable amino acids). In contrast to fibrillar collagens, COL4A1 and COL4A2 have more than 20 interruptions of the Gly-Xaa-Yaa repeats that encompass protein interaction domains and confer local flexibility ^{12, 13}. Following assembly in the endoplasmic reticulum, heterotrimers are transported to the Golgi where they are packaged for vesicular release from the cell ^{2, 3}. Once they reach the extracellular space, heterotrimers participate in macromolecular network organizations whereby two heterotrimers associate via their NC1 domains, and four heterotrimers form anti-parallel lateral interactions via association of their 7S domains.

A targeted mutation of the *Col4a1/Col4a2* locus on murine Chromosome 8 generated mice with null alleles for both *Col4a1* and *Col4a2* and provided the first insights into their biological roles ¹⁴. This work showed that formation of the collagen IV network is dispensable for initial basement membrane assembly, but critical for its structural integrity. Mice homozygous for the *Col4a1;Col4a2* null allele died between embryonic days (E)10.5–11.5 with disrupted basement membranes including Reichert's membrane ¹⁴. In contrast, heterozygous mice were described as having no overt phenotype. Soon after, heterozygous, semi-dominant pathogenic *COL4A1* and *COL4A2* mutations were identified in humans and in multiple mouse lines that model the pathology in patients ¹⁵⁻¹⁷. Consistent with the broad distribution of $\alpha 1(\text{IV})_2 \alpha 2(\text{IV})$ heterotrimers in nearly all basement membranes, *COL4A1* and *COL4A2* mutations cause multisystem disorders with abnormalities in the vasculature, brain, eyes, kidneys and muscles being the most commonly reported to date ¹⁸⁻²⁰.

Cerebrovascular disease (CVD) is one of the most notable consequences of *COL4A1* and *COL4A2* mutations and comprises a growing constellation of clinical manifestations including porencephaly, small-vessel disease, leukoencephalopathy, intracranial aneurysms and recurrent intracerebral hemorrhages (ICH). Other cerebral defects include calcification and cortical malformations similar to those observed in Walker-Warburg syndrome and muscle-eye-brain disease. Patients also frequently present with ocular disease (microphthalmia, cataracts, anterior segment dysgenesis, Axenfeld Rieger syndrome, glaucoma, optic nerve hypoplasia and retinal vascular tortuosity) kidney disease (hematuria and renal cysts of variable severity) and muscular defects (cramps, elevated serum creatine kinase level and muscular dystrophy). An excellent review of clinical consequences of *COL4A1* and *COL4A2* was recently published ²⁰. It is likely that the present observations are not exhaustive and that pathology in additional tissues and organs will be identified as studies advance.

As of April 2016, there were 93 and 12 mutations identified in human *COL4A1* and *COL4A2*, respectively, and 13 and 3 in the corresponding murine genes (not including three targeted *Col4a1* alleles). This sample size seemed appropriate to explore emerging genotype–phenotype correlations. Identifying genotype–phenotype correlations can be useful for patient management, genetic counseling, cohort stratifications for drug response or adverse events in clinical trials and for providing a better understanding of the biological function of distinct protein sub-domains. Equivalent analyses were performed with similar numbers of patients who have mutations in other types of collagens and these studies have provided valuable insight. The purpose of this review is to draw upon comparisons with other types of collagens and the current literature of human *COL4A1* and *COL4A2* mutations and mouse models to help identify potential genotype–phenotype correlations and understand their implications.

2) Genotype–phenotype correlations in patients with mutations in other types of collagens

Type IV collagens are part of the collagen superfamily comprising 28 members that have the defining feature of a triple-helical domain ¹. Mutations in multiple types of collagens contribute to a diverse constellation of human diseases. There is great a depth of knowledge of disease mechanism for many types of collagens which can be instructive for understanding genotype–phenotype correlations for *COL4A1* and *COL4A2* mutations. However, despite commonalities, members of the collagen superfamily are structurally and functionally diverse and there is no universal disease mechanism. For example, type VI collagen mutations can cause muscular dystrophies that are notable for the breadth of variation in disease severity. The type VI collagens are non-fibrillar collagens and the $\alpha 1(\text{VI})\alpha 2(\text{VI})\alpha 3(\text{VI})$ heterotrimers form beaded microfibrils important for anchoring the basement membrane to other components of the extracellular matrix. Type VI collagen mutations can cause both Ulrich congenital muscular dystrophy (a severe neonatal or early childhood disease) and Bethlem myopathy (where ambulation is often retained into adulthood), which were considered to be distinct diseases until molecular characterization revealed a genotype–phenotype relationship that influences disease expressivity. Dominant missense mutations of Gly residues in the triple helical domain are a common class of type VI collagen mutations and the most severe cases cluster within a small region of ~15 amino acids near the amino terminus of the triple helical domain that is proposed to represent a functional subdomain important for muscle function ²¹.

The type II collagens represent a similar example of phenotypic dichotomy attributable to genotype. Type II collagens are fibril forming $\alpha 1(\text{II})_3$ homotrimers with expression in chondrocytes and in the ocular vitreous. *COL2A1* mutations cause Stickler syndrome type I, characterized by high myopia, vitreoretinal degeneration, retinal detachment and cataracts. On the other end of the disease spectrum resulting from *COL2A1* mutations are severe forms of skeletal dysplasia, chondrodysplasia and achondrogenesis. In a relatively clear example of genotype–phenotype correlations, the preponderance of mutations causing Stickler syndrome are highly disruptive mutations that likely result in functionally null alleles. In a published series of 188 Stickler syndrome type 1 probands, the predominant

cause was loss-of-function mutations comprising splice site alterations, nonsense mutations, or insertion/deletions (indels). Only five of the 77 mutations that caused Stickler syndrome were Gly substitutions in the triple helical domain²². There is further evidence to suggest that the Gly substitutions that cause Stickler syndrome type 1 are near the amino terminal quarter of the protein. In contrast, Gly substitutions nearer the carboxy terminus – especially those for which Gly is replaced by a bulkier amino acid – are more often associated with severe forms of skeletal dysplasia, achondrogenesis or severe short stature phenotypes^{22–25}.

In a departure from distinct clinical entities, vascular Ehlers–Danlos syndrome (vEDS) is caused by dominant mutations in *COL3A1*. Type III collagens are fibril forming $\alpha 1(\text{III})_3$ homotrimers and the triple helical domain consists of 343 strict repeats of the Gly-Xaa-Yaa motif. Typical consequences of vEDS are arterial dissections, bowel perforations and uterine ruptures^{26, 27}. A recent report²⁸ of genotype–phenotype associations in 146 vEDS index patients with molecular diagnosis found 126 distinct mutations that fit into five distinct categories: Gly substitutions in the triple helical domain (56.3%), splice-site and in-frame indels (28.6%), haploinsufficiency (5.5%), non-Gly substitutions in the triple helical domain (3.2%) and mutations outside of the triple-helical domain (6.3%). As a measure of relative severity the age at diagnosis was younger for patients with in-frame splice site mutations (25 years) than for those with Gly substitutions (34 years) and both were younger than for patients with other classes of mutations (45 years). Although there was a frequency bias for substitutions of Gly to more destabilizing amino acids (Aspartic acid, Glutamic acid, Valine), no correlation with disease severity was observed²⁸. Instead, the position of the glycine substitutions appears to play a role with those nearer the carboxy terminus being more severe²⁷. Moreover, the prognosis was better for patients who did not have Gly substitutions or in-frame coding mutations as they were generally spared from digestive complications. In a separate study of 630 index cases for which survival was the outcome measure, patients with triple helical glycine substitutions had significantly shorter survival than patients with null mutations, however, in this case the identity of the substituting amino acid did influence the outcome but the position of the mutation did not²⁹.

Perhaps the richest source of information for collagen-related genotype–phenotype correlations comes from the type I collagens. Unlike collagen II and III that form homotrimers, collagen I (like collagen IV) form $\alpha 1(\text{I})_2\alpha 2(\text{I})$ heterotrimers. Mutations in *COL1A1* and *COL1A2* cause Osteogenesis imperfecta (OI), a genetically and clinically heterogeneous disorder characterized by skeletal deformities and fragile bones that are susceptible to fracture. An early influential analysis of approximately 70 *COL1A1* and *COL1A2* mutations identified genotype–phenotype correlations that underscored the distinction between the quantitative and qualitative effects of mutations³⁰. As larger sets of patients became available a clearer understanding emerged that appears to have broad applicability across many types of collagens³¹. Mutations were associated with milder forms of OI if they caused quantitative defects (mutations in the carboxyl-terminal domain that preclude the mutant protein from integrating into heterotrimers and frameshifts and null alleles that trigger loss of mRNA leading to a reduction in protein levels). In contrast, qualitative defects (such as Gly substitutions in the triple helical domain, exon skipping and rearrangements) responsible for protein structural defects were associated with more severe forms of OI^{30, 32}. Because mutations within the triple helical domain are common and

associated with severe forms of disease they have been the subjects of greatest interest. A definitive study of more than 800 *COL1A1* and *COL1A2* Gly mutations and altered splice sites in the triple helical domain allowed the discovery and confirmation of a number of genotype–phenotype correlations³¹. Of the 832 independent mutations, 682 resulted in triple helical Gly substitutions (391 in *COL1A1* and 291 in *COL1A2*) and 150 altered splice sites (102 in *COL1A1* and 48 in *COL1A2*)³¹. *COL1A1* splice site mutations often shifted the open reading frame resulting in absence of the mutant chain and manifested as quantitative mutations that were rarely severe (lethal). In contrast, triple helical glycine substitutions produced mutant proteins that incorporate into heterotrimers leading to important qualitative differences. Large-scale analyses of the effects of triple helical Gly substitutions produce a number of interesting genotype–phenotype correlations. Gly is represented by four codons: GGU, GGC, GGA, GGG and transitions or transversions of either of the first two guanine nucleotides results in Gly substitutions to Alanine (Ala), Arginine (Arg), Aspartic acid (Asp), Cysteine (Cys), Glutamic acid (Glu), Serine (Ser), Tryptophan (Trp) or Valine (Val). One third of the Gly substitutions were lethal - especially when the substituting amino acid is charged (Asp and Glu), or has a branched side chain (Arg and Val). When these substitutions were non-lethal they associated with severe outcomes compared to substitutions for Cys, Glu, or Ala. Furthermore, the influence of amino acid substitutions on disease severity was superimposed on the specific location of the mutation. For instance, when comparing only substitutions of Cys for Gly there was a relatively smooth phenotypic gradient from mild to severe in an amino-to-carboxy direction, suggesting a position effect³⁰. While in a larger patient cohort no linear relationship emerged, with few exceptions, Gly substitutions within the first 200 amino acids of *COL1A1* were non-lethal (on the mild end of the OI spectrum) whereas similar mutations closer to the carboxy-terminal end were often severe³¹. Additionally, there were two specific sub-domains associated with lethal phenotypes that align with major ligand binding regions and gap regions without reported mutations that contain putative integrin-binding sites, suggesting critical functions³¹. Mutations causing severe forms of OI were later described in a gap region, re-enforcing the hypothesis that these regions have important functional roles³³.

Notably, compared to *COL1A1* mutations, *COL1A2* mutations were identified much less frequently (291 vs 391 for Gly substitutions and 48 vs 102 for splice site mutations) and there were both similarities and differences in the outcomes³¹. Among patients with *COL1A2* glycine substitutions, those with Asp, Glu and Arg – but not Val – tended to have worse outcomes. *COL1A2* Gly substitutions were predominantly non-lethal, with the exceptions of exon skipping mutations in the carboxyl half of the protein and glycine substitutions that occurred in eight regularly spaced clusters³¹. The lethal clusters do not align with major ligand binding regions but were in regions suggested to be critical for interactions between proteoglycans and the collagen fibrils.

Overall, these studies revealed that mutations causing quantitative differences where the mutant allele is degraded or not incorporated into trimers are found less frequently and are associated with the mildest outcomes. Mutations that cause qualitative differences produce a wide range of phenotypes that depends on many factors including the gene involved and

nature and location of the mutation. Analysis of COL1A1 demonstrated that, in addition to important functional subdomains, worse patient outcomes were associated with Gly substitutions for charged or branched amino acids and for Gly substitutions nearer the carboxy terminus. The position of mutations in COL1A2 appears to have less influence unless the occurred within one of the 8 regularly spaced clusters ³¹.

Mutations in the other type IV collagen isoforms also cause autosomal recessive (*COL4A3* and *COL4A4*) or X-linked (*COL4A5*) Alport Syndrome and may offer insight into to genotype-phenotype correlations ³⁴. The proteins encoded by these three genes form $\alpha3(\text{IV})\alpha4(\text{IV})\alpha5(\text{IV})$ heterotrimers that are present in basement membranes of the glomerulus, cochlea and eye ^{10, 35}. Alport Syndrome is characterized by progressive renal failure, sensorineural hearing loss, lenticonus, cataract and maculopathy ^{36, 37}.

Approximately 80–85% of Alport Syndrome cases are X-linked and caused by *COL4A5* mutations ^{38, 39}. Studies of large cohorts of males with X-linked Alport Syndrome reveal genotype-phenotype correlations that are often contradictory to those found in type I collagens. The consensus from studying over 1000 patients with *COL4A5* mutations was that disease severity (age at onset for end stage renal failure) was greatest for truncating mutations, followed by splicing variants and finally missense mutations ^{40–44}. Moreover, truncating or splicing mutations had two-fold greater odds of having ocular pathology compared to missense mutations ⁴¹. This is in contrast to the concept that emerged from type I collagens where outcomes for quantitative mutations are worse than for qualitative mutations. Heterotrimer stoichiometry can offer an explanation for this observation since severe loss of function mutations cannot be compensated for by the homologous chromosome, leaving cells with a virtual absence of COL4A5. Position dependent effects were also reported, however, these too appear to be opposite to those reported in type I collagens. Mutations in the amino terminal signal peptide, regardless of type, led to end stage renal failure at younger ages than mutations in the triple helical or NC1 domains ⁴¹. One study that only looked at triple helical glycine substitutions suggested that mutations in exons 1–20 (amino terminal) might have been less severe than those in exons 21–47 (carboxy terminal) ⁴⁰ while a second, larger study found that mutations nearer the amino terminus were more severe ⁴¹. Finally, a recent study suggested that missense substitutions for bulkier amino acids were associated with more severe outcomes ⁴³.

3) Genotype–phenotype correlations in patients with *COL4A1* or *COL4A2* mutations

This review includes 105 independent occurrences of *COL4A1* or *COL4A2* mutations of which, 93 were in *COL4A1* (at 74 unique sites) and 12 in *COL4A2* (at 10 unique sites) (Tables 1 and 2). All of the mutations described were dominant with equal representation in males (48%) and females (52%) and approximately equal numbers of inherited (47%) or sporadic (53%) incidences. Similar to other types of collagens, triple helical Gly substitutions represented the most prevalent class of mutations (68 out of 93) and the most common substitutions were for charged amino acids; Arg (30), Glu (11), and Asp (10). The remaining substitutions were Ser (7), Val (5), Ala (3), Leu (1), and a termination codon (1). No substitutions for tryptophan or cysteine, which are the only other substitutions possible

with a single nucleotide change of glycine codons, were reported. Splice site mutations within the triple helical domain constitute the second most prevalent class of mutations (8 out of 93). Three of the splice site mutations caused in-frame deletion of an entire exon (exon 21, 25 or 31) and two mutations resulted in nonsense-mediated decay of the transcripts with reduced *COL4A1* expression while the consequences of the other three were not reported. Other classes of mutations were much less frequent including Yaa position substitutions (3), substitutions within the repeat interruptions (3) and frame-shifting indels (3). Eight mutations outside of the triple helical domain have also been reported, seven of which were located in the NC1 domain and one substitution mutation of the start methionine.

The low number of published *COL4A2* mutations decreases the confidence for identifying broad trends, however, triple helical glycine substitutions (6) were also the most prevalent class of mutations where Gly was substituted for Arg (2), Glu (2) and Asp (2). One patient had a frame-shifting deletion and a second patient had a missense mutation, each occurring in the NC1 domain. There were also examples of triple helical Xaa-position substitutions (4), however, three of these were identified independently in the same amino acid and this variant was also identified in controls raising the possibility that this variant might be non-pathogenic. If so, the total number of *COL4A2* mutations could be as low as nine with six being Gly substitutions in the triple helical domain. The failure so far to identify mutations in the repeat interruptions could represent differences between *COL4A1* and *COL4A2* or simply reflect the low number of *COL4A2* mutations discovered to date.

Overall, *COL4A2* mutations appear to associate with milder phenotypes (later age-at-onset of CVD, and fewer system abnormalities reported). However, because the limited number of *COL4A2* mutations, we focus primarily on the 93 *COL4A1* mutations for the investigation of genotype–phenotype correlations in patients. Fourteen *COL4A1* mutations were presumed to reduce the amount of normal $\alpha 1(\text{IV})_2 \alpha 2(\text{IV})$ heterotrimers without producing abnormal heterotrimers and may therefore qualify as quantitative mutations: seven mutations in the NC1 domain, three frame-shifting indels, two frame-shifting splice mutations, one substitution of the start methionine, and one triple helical glycine substitution leading to a premature termination codon. Six out of the seven NC1 mutations and 12 out of the 14 total “quantitative” mutations were reported in patients with porencephaly and the remaining two patients had perinatal ICH. If one considers porencephaly and early age-at-onset as phenotypes on the severe end of the CVD spectrum, quantitative *COL4A1* mutations appear to associate with relatively poor outcomes.

To compare the effects of qualitative measures, we focused on the 68 triple helical Gly substitutions in *COL4A1*. The frequencies of Arg, Glu, and Asp substitutions might suggest that the outcomes of these mutations are more severe, however this conclusion would need to be corrected for codon usage and integrate other potential factors. In four pairs of instances the same glycine residue was independently substituted by distinct amino acids (G498V, G498D; G655R, G655E; G708R, G708V; G990E, G990V). However, in all cases, the substituting amino acids are either branched or charged and so might all be expected to have severe consequences and there were no obvious differences that could be attributed to the identity of the substituting amino acid. To further investigate potential differential effects

of substituting amino acids, we compared the consequences of the 30 Arg substitutions to the Ser (7) and Ala (3) substitutions. No obvious correlations were observed when comparing substitutions that were roughly position-matched. For example, G670R caused porencephaly, whereas G708R caused ocular defects but no CVD and G696S had adult onset ICH. Of four independent occurrences of G755R mutations, none lead to porencephaly and instead all result in childhood or adult onset CVD. In contrast, two independent occurrences for the nearby G749S mutation caused porencephaly. Although these analyses indicate that substitutions to branched amino acids occur more frequently, there is no clear evidence that they are associated with more severe outcomes.

Notably, the position of the mutation within the triple helical domain appeared to correlate with CVD severity. There are two Gly substitutions in the first quarter of the triple helical domain suggesting that Gly missense mutations near the amino-terminal end of the triple helical domain might cause milder phenotype. The 30 Arg substitutions occur at 20 unique sites allowing comparison of the effects of position for the same substituting amino acid. If one divides the triple helical domain roughly in half, 14 mutations are in the carboxy-terminal half (amino acid 805 and higher) and, of these, 12 caused porencephaly and two caused schizencephaly. In addition to early age-at-onset, these mutations were more often sporadic or *de novo* cases and were frequently associated with epileptic seizures and motor disorder and/or developmental delay. In contrast, of the 16 Gly to Arg mutations in the amino-terminal half of the protein, only five caused porencephaly, four mutations did not cause ICH/stroke, four mutations were identified in patients who were adults at the time of diagnosis and only four caused epileptic seizures.

In addition to modulating CVD severity, the location of the mutation may also influence the penetrance of other phenotypes. For instance, *COL4A1* mutations cause ocular disease with high penetrance (72%), however the specific ocular defect depends on the position of the mutation. For example, mutations in the amino-terminal one-third of the protein caused retinal arteriolar tortuosity with complete penetrance, whereas mutations causing cataract, anterior segment dysgenesis, glaucoma and microphthalmia tended to occur closer to the carboxy terminus. Of interest, in the amino-terminal one-third of the protein there were mutations reported in six families with a clinical diagnosis of HANAC Syndrome (hereditary angiopathy with nephropathy, aneurysms, and muscle cramps)^{51–53, 112}. The mutations cluster in exons 24 and 25 within a 31 amino acid region (G498–G528) of the triple helical domain that encompasses several integrin-binding domains. While nephropathy, myopathy and aneurysms are not restricted to patients with mutations in this region, these phenotypes may be disproportionately represented in patients from these families. These patients also had retinal arteriolar tortuosity and heart disease but other features including porencephaly, cerebral calcification, cataract, glaucoma, microcephaly or epileptic seizures were not described. Notably, while three Gly substitutions within exon 25 were reported in patients with HANAC syndrome, a splice site mutation causing an in-frame deletion of this exon was found in a patient with bilateral diffuse white matter abnormalities, multiple clinically asymptomatic hemorrhages and finally an acute ICH but without aneurysms, myopathy or renal involvement⁵⁶. Observations from patients with triple helical Gly substitutions support the existence of one or more functional subdomains within this domain that may associate with differential co-morbidities.

There are a number of limitations that demand approaching genotype–phenotype correlations in patients with caution. Most of the mutations occurred only once and to label the ‘severity’ of a mutation based on a sample size of one risks oversimplification of many complex genetic, environmental and stochastic interactions that are integrated in the phenotype. Even in cases of recurrent mutations, the conclusions are not simple. There were eight amino acids that had multiple hits and G755R and G773R were each reported in four independent studies. While the outcomes for G755R were comparable (all patients had ICH, leukoencephalopathy and cataracts with an age of onset between 10 and 21 years, and did not cause porencephaly or epilepsy) those for G773R were less so (three mutations caused porencephaly or severe stroke whereas one mutation cause only very mild white matter changes). Genetic context differences could explain some of the discordance in the expressivity resulting from a given mutation. However, phenotypic differences within families who are more closely matched genetically, including probands with porencephaly born to asymptomatic parents who carried mutations, were also observed. There are multiple possible explanations for these cases including genetic modifiers, mosaicism, biological variation or environmental influences.

4) Genotype–Phenotype correlations in *Col4a1* and *Col4a2* mutant mice

Studies of mouse models with *Col4a1* and *Col4a2* mutations are a powerful way to simplify the genetic and environmental influences and compliment patient studies. In addition to the *Col4a1;Col4a2* null mutation¹⁴, and a *Col4a1* conditional mutation^{101, 110}, there are presently 17 mutations reported in *Col4a1* and *Col4a2* that cause pathology in multiple tissues and organs^{16, 17, 100, 104} (Fig. 1). Because of procedural differences between research groups, notably the focus and depth of analysis, it is difficult to directly compare graded levels of relative severities for each mutation. Importantly, the genetic context in which the mutations are studied also has a significant effect on the penetrance and/or severity of pathology^{18, 80, 101, 111}. For example, *Col4a1* mutant mice maintained on a pure C57BL/6J background have severe ocular dysgenesis, myopathy and ICH that are all suppressed in mutant mice that were crossed for a single generation to the CAST/EiJ inbred strain^{15, 18, 80, 101, 111}. Notably, the 129SvEvTac inbred strain also suppressed anterior segment dysgenesis, but not ICH, indicating that genetic context can have differential effects on distinct phenotypes and suggesting heterogeneity in the pathogenic mechanisms underlying different *Col4a1*-related phenotypes¹⁰¹.

To systematically investigate potential genotype–phenotype correlations, nine *Col4a1* and one *Col4a2* mutations¹⁰⁰ were uniformly backcrossed to the C57BL/6J genetic background^{101, 102}. The allelic series comprised seven missense mutations of glycine residues in the triple helical domain (six in COL4A1, one in COL4A2), one missense mutation in the NC1 domain of COL4A1, and a point mutation of splice acceptor site that results in the deletion of exon 41 (ex41) but retains the open reading frame (Fig. 2A). By controlling environmental factors and the genetic context, studies using this allelic series demonstrated allelic heterogeneity between mutations. The severity of ICH was compared between strains of mutant mice aged for 7–9 months and different classes of mutations that implicated three potential genotype–phenotype correlations were identified^{101, 102}. First was a domain effect whereby the NC1 domain mutation (*Col4a1*^{S1582P}), which presumably

reflects a quantitative mutation (Fig. 2B), caused less severe ICH than the triple helical domain mutations (Fig. 2C). Second, for point mutations within the triple helical domain, there was a position effect whereby mutations nearer the carboxy termini (where heterotrimer winding is initiated) caused more severe ICH than mutations nearer the amino termini. Third, there appeared to be a ‘class effect’ whereby *Col4a1*^{ex41} was more severe than missense mutations even when compared to a glycine point mutation within exon 41 (*Col4a1*^{G1180D}). The pathogenicity resulting from *COL4A1* and *COL4A2* mutations is generally attributed to impaired secretion and concomitant intracellular accumulation and extracellular deficiency of mutant heterotrimers. Notably, ICH severity in this allelic series correlated with the relative levels of heterotrimer accumulation between mutations (Fig. 2B and C)^{101, 102}.

An interesting contrast was observed when myopathy was quantified in this same allelic series¹⁰². Similar to the previous trends, mice with a presumed qualitative mutation in the NC1 domain (*Col4a1*^{S1582P}) had little or no phenotype and myopathy resulting from the *Col4a1*^{ex41} mutation was more severe than that resulting from the ‘position-matched’ point mutation (*Col4a1*^{G1180D}) (Fig. 2D). However, in contrast to the trends observed in ICH, there was no apparent position effect for mutations within the triple helical domain. Instead, *Col4a1*^{G394V}, which was among the mutations with the least intracellular accumulation and mildest ICH, had the most severe myopathy demonstrating an interesting dichotomy for this mutation. *Col4a1*^{G394V} occurs near a putative integrin-binding site adjacent to the cluster of HANAC mutations, in a region that may represent a functional sub-domain with an important role in muscle development and function¹⁰².

Kidney abnormalities (albuminuria and thin or split glomerular basement membranes) were previously reported in *Col4a1*^{+/- ex41} mice¹⁵ but had not been examined in other strains from the allelic series. Here we show that hematuria shares a profile that resembles ICH for the *Col4a1* mutations whereby the *Col4a1*^{S1582P} mutation had almost no phenotype, the *Col4a1*^{ex41} mutation was worse than *Col4a1*^{G1180D} and mutations nearer the carboxy termini were more severe than those nearer the amino termini (Fig. 3A). These data may suggest that hematuria has a vascular origin or that hematuria and ICH share pathogenic mechanisms. When we measured albuminuria the results were similar with the notable exception that the *Col4a2* mutation (*Col4a2*^{G646D}) was disproportionately severe compared to *Col4a1* mutations (Fig. 3B). This observation might suggest that *COL4A2* is more important for nephropathy or that this mutation occurs within a functional sub-domain important for renal pathology. In support of the latter alternative, the *Col4a1*^{G658D} mutation, which is nearby, lead to the most severe albuminuria amongst the *Col4a1* point mutations.

Another potential genotype–phenotype correlation was reported in an allelic series of three mutations: *Col4a1*^{G627W}, *Col4a1*^{K950E} and *Col4a1*^{G1046D} (also called Bru, Raw and Svc, respectively)¹⁶. The authors reported that the Gly substitutions (*Col4a1*^{G627W} and *Col4a1*^{G1046D}) caused relatively more severe ocular and renal phenotypes than a Lys substitution affecting the Yaa-position amino acid on the Gly-Xaa-Yaa motif (*Col4a1*^{K950E}), which only causes a retinal vascular phenotype. *Col4a1*^{G627W} was identified in a different genetic background, nonetheless, *Col4a1*^{K950E} and *Col4a1*^{G1046D} were studied in similar genetic contexts and the position of the mutation within the Gly-Xaa-Yaa repeat is a

plausible explanation for the mild phenotype(s) in *Col4a1*^{K950E} mice. An initial report of these alleles suggested that the glomerular basement membrane was unremarkable but Bowman's capsule had basement membrane defects and hypertrophic cells¹⁶ which differed from the ultrastructural glomerular basement membrane abnormalities¹⁵, albuminuria and hematuria described for the other alleles. More recently, a detailed characterization of renal pathology cause by *Col4a1*^{K950E} and *Col4a1*^{G1046D} mutations was described extensive pathology and potentially distinct involvement of glomerular and tubular defects¹⁰⁷. Notably, differences in focus, techniques and study depth underscore the challenges of comparing genotype–phenotype data between research groups. A targeted allele that is analogous to a human HANAC-causing mutation (*Col4a1*^{G498V}) was also recently published¹⁰⁴ and while the relative severities of the phenotypes are difficult to compare in isolation, this report also includes a thorough characterization of renal pathology and supports the existence of one or more amino-terminal sub-domain(s) that disproportionately influence the severity of myopathy and nephropathy.

Despite a limited number of available mutations, findings from studies using murine models of *Col4a1* mutations, including two allelic series, provide insights into disease mechanisms and support the conclusion that there are genotype–phenotype correlations. The data suggest that 1) mutations in the NC1 domain (and likely other 'quantitative' mutations) may associate with milder outcomes than those in the triple-helical domain; 2) Gly substitutions in the triple helical domain have position-dependent effects on biosynthesis that correlate with CVD, but not other phenotypes; 3) in frame, exon-skipping mutations within the triple helical domain may be disproportionately severe compared to Gly substitutions 4) Yaa-position substitutions may be less severe than Gly substitutions 5) one or more functional sub-domains near the amino-termini may be involved in myopathy and possibly nephropathy. The fact that genetic background and genotype–phenotype correlations had differential effects on different phenotypes suggest that there is mechanistic heterogeneity.

5) Conclusions and perspectives

Mutations in *COL4A1* and *COL4A2* have been studied for nearly a decade and, compared to type I collagen mutations underlying OI and *COL4A5* mutations underlying Alport Syndrome, the number of patients from which to draw genotype-phenotype correlations is relatively limited. However, this retrospective analysis suggests the existence of genotype–phenotype correlations for *COL4A1* and *COL4A2* mutations that are consistent with observations for some other types of collagens. In both humans and mice, there is a frequency bias for mutations in *COL4A1* over *COL4A2*. This inequity likely reflects a combination of ascertainment bias and true biological differences. Mutations in *COL4A1* were reported several years before mutations in *COL4A2* and routine *COL4A2* screening in patients lagged. As mutation detection shifts from candidate gene sequencing to exome analyses this bias should disappear. However, ascertainment bias alone does not explain the discrepancies in the numbers of mutations in each gene. The genes are roughly equal in size and would be expected to acquire mutations at equivalent rates in patients and during random chemical mutagenesis. It is possible that milder outcomes for *COL4A2* mutations may explain the frequency difference in mutations identified. It is interesting to note however that in the murine allelic series, the *Col4a2* mutation essentially behaved like a

position-matched *Col4a1* mutation with the exception of one phenotype (albuminuria) for which this mutation was *more* severe.

As is the case for other types of collagens, Gly substitutions were the most common class of mutation in patients. Mutations in 50 out of 437 triple helical Gly residues in COL4A1 and six out of 428 triple helical Gly residues for COL4A2 have been identified. While substitutions for branched amino acids occurred more frequently, the identity of the substituting amino acid did not appear to have a strong influence on disease severity. Instead, there appear to be important positional influences; mutations located near the amino terminus of COL4A1 are associated with milder pathology, and the severity of CVD tends to be worse for Gly substitutions that are closer to the carboxyl end of the triple helical domain. Finally, mutations that clustered near the region containing integrin-binding domains appear to be associated with a greater likelihood for patients to have myopathy and nephropathy suggesting the existence of a functional subdomain influencing the penetrance of distinct phenotypes.

In findings from the murine allelic series the mutation with presumed quantitative effects (S1582P in the NC1 domain) had milder effects on heterotrimer biosynthesis, ICH and myopathy than most mutations with presumed qualitative effects (mutations in the triple helical domain). In contrast, in patients, qualitative *COL4A1* mutations do not appear to be associated with more severe outcomes than quantitative mutations. As there are biases toward identification of mutations with severe outcomes in early studies, it is possible that these classes of mutations will diverge as more patients are reported with a broader spectrum of mutations. An additional caveat is that the molecular consequences of most mutations were inferred and not tested, which could lead to erroneous classification. Importantly, although porencephaly and early age-at-onset CVD were considered to be more severe than adult age-at-onset CVD determining the relative outcome severities is subjective and remains a pervasive challenge. There is the additional challenge in integrating multiple co-morbid phenotypes, some of which appear to be mechanistically distinct. In contrast, characterizations of OI severity relies on well-established criteria for bone fragility that are more quantitative and focused on a relatively defined outcome. Finally, the roles of different environmental factors (including anti-coagulants, surgical delivery, and exercise)^{15, 101} must all be considered as they may contribute to misleading conclusions.

In contrast to patient studies, the allelic series of mutant mice comprises less saturation of mutations across the protein or for each class of mutation. For example, there is only one presumed quantitative mutation in the NC1 domain. Although the effect was consistent across phenotypes (and consistent with the absence of phenotypes in mice heterozygous for *Col4a1;Col4a2* null alleles) these findings are based upon observations for a single allele and it would be overly simplistic to presume that mutations across the entire domain will behave equally. However, these resources are powered by standardized environments, genetic homogeneity and large sample sizes for an individual mutation which all improve reproducibility. Continued efforts to characterize pathology in *Col4a1* and *Col4a2* mutant mice will improve our understanding of the phenotypic spectrum of *Col4a1*- and *Col4a2*-related disease. Better definition of the genotype-phenotype correlations and patient outcomes will help to understand risks including co-morbidities or age-at-onset and will

ultimately impact therapeutic approaches, genetic counseling and patient care. This information will ideally come from large, prospective, longitudinal studies to detail the natural histories of patients for whom there are biochemical data, and quantitative outcome measures for multiple phenotypes. Together, these and other advances may reveal pathways that can be targeted therapeutically to reduce, delay or prevent aspects of this syndrome.

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Abbreviations

COL4A1	collagen type IV alpha 1
COL4A2	collagen type IV alpha 2
CVD	cerebrovascular disease
HANAC	hereditary angiopathy with nephropathy, aneurysms and muscle cramps
ICH	intracerebral hemorrhages
OI	Osteogenesis imperfecta
NC1	non collagenous 1
vEDS	vascular Ehlers-Danlos syndrome
Ala	Alanine
Arg	Arginine
Asp	Aspartic acid
Cys	Cysteine
Glu	Glutamic acid
Gly	Glycine
Ser	Serine
Trp	Tryptophan
Val	Valine

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Highlights

We discuss genotype-phenotype correlations emerging from 105 *COL4A1* and *COL4A2* mutations in patients and 17 *Col4a1* and *Col4a2* mutations in mice. The results have similarities and differences when compared to genotype-phenotype correlations for other types of collagens. Triple helical glycine substitutions are the most frequent class of mutation. Glycine is most often replaced by a charged amino acid however the position of the mutation appears to be more important than the identity of the substituting amino acid. Data from mice, but not humans, suggest that NC1 domain mutations might be less damaging than triple helical domain mutations. Moreover, for cerebrovascular disease there appears to be a position-dependent effect whereby triple helical mutations nearer the carboxy terminus may be more damaging than those nearer the amino terminus. However, this association may not hold true for all phenotypes and there appears to be at least one functional subdomain near the amino terminal end of the triple helical domain in which mutations have divergent outcome severities for different sub-phenotypes. The involvement of multiple organs and a lack of quantitative phenotypic measures make it difficult to draw firm conclusions and the different components of the multisystem disorder likely involve different pathogenic mechanisms, which must be considered.

Figure 1. Summary of reported phenotypes for murine Col4a1 and Col4a2 mutations
(A) Schematic diagram illustrating murine mutations reported in the literature. (B) List of *Col4a1* and *Col4a2* mutations and the phenotypes that have been reported for each.

Figure 2. Allelic heterogeneity for heterotrimer biosynthesis, intracerebral hemorrhage and myopathy

(A) Schematic diagram illustrating the mutations in the allelic series. All mutations are in the triple helical domain, except COL4A1^{S1582P}, which is in the C-terminal NC1 domain. COL4A1^{ex41} is a splice site mutation causing deletion of exon 41. (B) Western blot quantification demonstrating that the mutation in the NC1 domain (COL4A1^{S1582P}) had relatively low levels intracellular heterotrimer accumulation and that there is a position-dependent graded severity of heterotrimer accumulation for mutations within the triple helical domain whereby mutations nearer the NC1 domain had higher levels of heterotrimer accumulation. (C) Quantification of intracerebral hemorrhage revealed that COL4A1^{ex41} mutation leads to the most severe phenotype and that point mutations in the triple helical domain nearer the C-terminus tended to cause more (or more severe) hemorrhages. (D)

Quantification of non-peripheral nuclei revealed that the Col4a1^{G394V} mutation, which is in an integrin-binding domain, causes the most severe myopathy. For all groups, n>5. Comparisons between wild-type (Col4a1^{+/+}) and mutant mice were performed using Student's t-test; *, p<0.05. **, p<0.01. For comparison among the different strains, a one-way ANOVA followed by a Tukey's post hoc test was performed; †, p<0.05. Panels A, B and D are modified from Kuo *et al.* (2014) ¹⁰² and Panel C is modified from Jeanne *et al.* (2015) ¹⁰¹.

Figure 3. Relative measures of hematuria and albuminuria in mice from an allelic series of Col4a1 and Col4a2 mutations

We collected urine and measured levels of hematuria and albuminuria using colorimetric dipsticks and a clinical chemistry autoanalyzer, respectively (described in Gould *et al.* (2006) ¹⁵). **A**). The proportion of animals of each genotype with no, mild, moderate or severe hematuria are indicated as percentages. Sample sizes for both each genotype are indicated at the top. **B**) We measured albuminuria by calculating the ratio of albumin to creatinine (mg/g) and indicated the proportion of animals of each genotype with no, mild (<30), moderate (30–60) or severe (>60) albuminuria as a percentage. Sample sizes for both each genotype are indicated at the top.

Table 1

		COL4A1	COL4A2		
		Occurrences	Locations	Occurrences	Locations
Start codon	Met substitution	1	1	0	0
Triple helical domain	Gly of Gly-Xaa-Yaa motif	68	50	6	6
		Arg (30), Glu (11), Asp (10), Ser (7), Val (5), Ala (3), Leu (1), Ter (1)		Arg (2), Glu (2), Asp (2)	
	Yaa of Gly-Xaa-Yaa motif	3	3	0	0
		Pro to Leu, Met to Val, Gln to Glu			
	Xaa of Gly-Xaa-Yaa motif	0	0	4	2
				Glu to Gly (3), Gln to Lys	
	Repeat Interruptions	3	3	0	0
	Frameshift	3	3	1	1
		2 deletions, 1 duplication		1 deletion	
	Splice-site	8	8	0	0
NC1 domain	Missense	4	3	1	1
	Nonsense	1	1	0	0
	Duplication	1	1	0	0
	Insertion	1	1	0	0
Totals		93	74	12	10

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Table 3

	Protein	Reference	Notes	Porencephaly	ICN	Hypotension	Anemia	Cortical Lamination Defects	Ocular Dysgenesis/ Glaucoma	Retinal Vascular Tortuosity	Retinal Hemorrhages	Optic Nerve Hypoplasia	Myopathy	Nephropathy	Lung	Hearing loss	Tendon Adhesions	
COL4A1	p.G394V	100-103																Legend: Reported present Reported absent Not reported
	p.G498V	104	Targeted															
	p.G627W	16, 105	called <i>Bru</i>															
	p.G658D	100-102																
	p.G912V	100-102																
	p.K950E	16, 106-108	called <i>Raw</i> (Yaa-position)															
	p.G954R	100																
	p.G1038S	100-102																
	p.G1064D	16, 107-109	called <i>Syc</i>															
	p.1180D	100-102																
	Δ ex41	*, 17, 18, 80, 101-103, 110, 1*	Δ ex41															
	p.1341R	100																
	p.1344D	100-102																
	p.S1582P	100-102																
COL4A1	p.V31F	100	Signal peptide cleavage site															
	p.G646D	100-103																
	p.G646D	100-103																
	Col4a1/a2 null	14	Targeted															
	Col4a1 conditional	101, 103, 110	Targeted (called <i>Flex41</i>)															