Mitral valve prolapse (MVP), an abnormal displacement into the left atrium of a thickened and redundant mitral valve during systole, is a relatively frequent abnormality in humans and may be associated with serious complications. A recent study implicates fibrillin-1, a component of extracellular matrix microfibrils, in the pathogenesis of a murine model of MVP. This investigation represents an initial step toward understanding the mechanisms involved in human MVP disease and the development of potential treatments.
lar dystrophy, where replacement of specific gene products such as the sarcoglycans could easily be detected. Mesoangioblasts, on the other hand, have been shown to be efficacious in restoring expression of α-sarcoglycan in the α-sarcoglycan–null mouse as well as the expression of the whole dystrophin complex, including δ-sarcoglycan (20). All of these cell types are easy to expand in culture and, once their homing and muscle differentiation activities can be optimized, they may represent a better perspective for the stem cell therapy of striated muscle diseases than BM-SP stem cells.

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Marfan syndrome and mitral valve prolapse

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Mitral valve prolapse (MVP), an abnormal displacement into the left atrium of a thickened and redundant mitral valve during systole, is a relatively frequent abnormality in humans and may be associated with serious complications. A recent study implicates fibrillin-1, a component of extracellular matrix microfibrils, in the pathogenesis of a murine model of MVP (see the related article beginning on page 1586). This investigation represents an initial step toward understanding the mechanisms involved in human MVP disease and the development of potential treatments.

Mitral valve prolapse: scope of the problem and evolution of the defining criteria
Mitral valve prolapse (MVP) is generally understood to be the displacement of abnormally thickened, redundant mitral leaflet(s) into the left atrium during systole (Figure 1) (1). One of the possible consequences of this condition is that the malfunctioning mitral valve allows backflow of blood in the left atrium (mitral regurgitation), which, when severe, leads to left ventricular enlargement and failure. Besides severe mitral regurgitation, MVP has been associated with serious complications such as bacterial endocarditis and sudden death, and primary mitral valve surgery is currently performed most frequently to specifically treat MVP (1, 2). Since the early 1970s, research has suggested that echocardiography is the ideal noninvasive technique to visualize the prolapsing mitral leaflets (3–5). However, continually changing echocardiographic techniques and criteria for the diagnosis of MVP have in many cases further obscured, rather than clarified, our understanding of prolapse in its primary form and in association with other disorders (6). Over the past decade, new echocardiographic criteria for MVP have been established based on an understanding of the 3D structure of the mitral valve (7). Defined according to these criteria, prolapse is the displacement of 1 or both mitral leaflets by more than 2 mm above the high points of the mitral annulus as recorded in either the parasternal or apical long-axis view (Figure 1, detail). This 2 mm displacement derives from studies showing that smaller displacements are not associated with increased leaflet thickness, mitral regurgitation, or valve-related complications (8). In cases where leaflet displacement is greater than 2 mm, prolapse is further subdivided into classic and nonclassic forms based on leaflet thickness (classic, ≤5 mm; and nonclassic, <5 mm), with complications such as endocarditis or severe mitral regurgitation generally occurring in patients with classic prolapse (9).

Using these criteria, a recent population study of 3,491 subjects from the offspring...
cohort of the Framingham Heart Study reported a prevalence of 1.3% for classic MVP and 1.1% for the nonclassic form (10). This overall prevalence of 2.4% is strikingly lower than the extremely high values previously reported elsewhere (21% and 34% when diagnosed using M-mode ultrasound or 2D electrocardiographic views, respectively) and significantly lower than the more generally quoted figures of 4–7% for the general population based on older M-mode studies (1, 11). In the recent study of the offspring cohort of the Framingham Heart Study (10), the incidence of MVP did not differ between the sexes, and no predisposition to prolapse was observed in younger or older adult subjects; this was supported by an even distribution (2–3%) among subjects in each decade from 30 to 80 years. Other data regarding the prevalence of MVP in children suggest that prolapse is uncommon before adolescence, but its prevalence increases after the adolescent growth spurt (12, 13). Nonclassic prolapse appears morphologically to be an intermediate stage between normal mitral valve anatomy and classic prolapse, but studies with up to a 10-year follow-up have shown no evidence of progression from one form to another, and the Framingham Heart Study failed to show an age-related shift from one type to the other, although the patient numbers studied in each age group were small.

Mitral valve prolapse and mitral regurgitation

The most common complication of MVP is severe regurgitation due to progressive degeneration of the valve and chordae, with myxomatous infiltration (thickening of the mitral layers with glycosaminoglycan accumulation), and fibroelastic and collagen alterations (10, 11, 13). In about 75% of cases, there is sudden deterioration because of chordal rupture (14). The cumulative risk of severe mitral regurgitation and valve rupture is minimal in individuals younger than 50 years of age but then rises steeply, with the risk in men being greater than in women beyond the age of 60. Using current prevalence data, it can be estimated that by the age of 70, approximately 11% of men and 6% of women with classic MVP will need mitral valve replacement (13).

Mitral valve prolapse: a genetic disease?

MVP is generally sporadic but is also associated with a variety of congenital disorders of connective tissue including Marfan syndrome, Ehler-Danlos syndrome, osteogenesis imperfecta, dominant cutis laxa pseudoxanthoma elasticum, and the MASS syndrome (mitral valve prolapse, goroic root dilatation, skeletal changes, and skin changes), among others. It has been estimated that only 0.25% of patients with MVP have Marfan syndrome. This percentage may be somewhat higher if the newer and more stringent criteria for MVP are used, but it is unlikely that more than 1–2% of patients with MVP have an associated connective tissue disorder. Marfan syndrome is associated with mutations in Fibrillin-1 (FBN1) on chromosome 15q21.1 (15, 16). Because MVP is found in many, but certainly not all, patients with Marfan syndrome, it was suggested many years ago that isolated MVP may also be due to a mutation of FBN1 (15). However, despite the availability of literally millions of patients for study, no convincing association has been found to date. Occasional families with MVP have been identified and an underlying gene defect reported. In patients with X-linked myxomatous valvular dystrophy, a rare disorder associated with severe MVP, the defect has been linked to chromosome Xq28 (17). The first locus for nonsyndromic MVP has been mapped to chromosome 16p11.2–p12.2 in 2 of 4 patients at a French surgical center (18). A second locus for autosomal dominant MVP has been mapped to chromosome 11p15.4 (19). Thus, even within families with an autosomal dominant mode of inheritance, there appears to be significant genetic heterogeneity.

The role of mechanical stress in the evolution of MVP is also important. The mitral valve opens and closes more than 3 billion times during the course of the normal human lifespan. During each closure period, it must withstand the full force of ventricular contraction. Because of the normal orientation of the leaflets, this force is unevenly observed that TGF-β–neutralizing antibodies, which confirms a causal relationship between TGF-β dysregulation and this abnormal mitral valve.

These findings are consistent with those from other studies that link TGF-β to the Marfan phenotype (27) and the demonstration that Marfan syndrome can be caused by inactivation mutations in TGFBR2 at 3p24.2–p25 (20) in patients without abnor-

Mitral syndrome and the fibrillin-1/TGF-β pathway

Although abnormalities within FBN1 are responsible for the Marfan phenotype in approximately 80% of patients, Marfan syndrome can also be caused by inactivation mutations in TGF-β receptor 2 (TGFBR2), located at 3p24.2–p25 (20). Fibrillin-I has also been shown to have a biologically important role in the activation of TGF-β in the lung (21). The TGFβs are multistep cytokines that are important modulators of cell growth, proliferation and differentiation, inflammation, extracellular matrix deposition, and apoptosis (22). Virtually every cell in the body produces TGF-β and has receptors for it (23). Their biologic effects are context dependent and vary with tissue type and the activity of other signaling pathways (24). Defects in TGF-β function are associated with a number of pathologic states, including tumor cell growth, fibrosis, and autoimmune disease (23). TGF-β activity is regulated by the conversion of latent TGF-β to active TGF-β. Tissues contain significant quantities of latent TGF-β, and activation of only a small fraction causes a maximal cellular response. There are multiple activators of latent TGF-β (proteases, thrombospondin-1, integrins, reactive oxygen species, and changes in pH), and latent TGF-β has been considered a molecular sensor that responds to specific signals by releasing active TGF-β, which is the effector molecule (22, 25).

In a study reported in this issue of JCI, Ng et al. tested the hypothesis that the fibrillin-1–TGF-β pathway is implicated in the pathogenesis of MVP in a murine model of Marfan syndrome (26). They compared the morphometry of mitral valves in fibrillin-1–deficient mice to those of wild-type mice and report a progressive increase in leaflet length and thickness in heterozygous versus homozygous mice. The leaflets exhibited both increased cell proliferation and decreased apoptosis. Using TGF-β–specific antibodies, they observed that TGF-β activity and signaling were increased in the valve tissue, while the concentration of latent TGF-β remained unchanged. Most interestingly, the mitral valve phenotype was rescued by TGF-β–neutralizing antibodies, which confirms a causal relationship between TGF-β dysregulation and this abnormal mitral valve.

These findings are consistent with those from other studies that link TGF-β to the Marfan phenotype (27) and the demonstration that Marfan syndrome can be caused by inactivation mutations in TGFBR2 at 3p24.2–p25 (20) in patients without abnor-
Murine and human MVP: similarities and differences

Similar to human MVP, the murine model pioneered by Ng et al. (26) displays a thickened and redundant mitral valve that prolapses into the left atrium. There are differences, however, among what has been noted in human acquired myxomatous mitral valve disease. Clinically, the MVP that often accompanies Marfan syndrome is not generally considered to be a model of the isolated MVP in the adult for a number of reasons. First, abnormalities in FBN1 have not been demonstrated in sporadic MVP patients or families with the disorder. Second, severe mitral valve disorders in Marfan syndrome are generally manifest in infants, while in adults, the majority of complications are due to aortic aneurysm formation and dissection. This is in contrast to the clinical paradigm of MVP, where complications are typically not seen before the fifth decade of life. Third, the valvular involvement in Marfan syndrome and in these mice affects the whole valve in a relatively homogeneous way. In idiopathic MVP, necropsy studies show that the morphologic abnormalities can involve 1 or both leaflets, and the morphological changes may be heterogeneous within the leaflet itself (28). Fourth, in one recent study (29), prolapse was shown to occur in roughly half of the patients with Marfan syndrome, and in our experience it occurs even less frequently. The mouse model described by Ng et al. (26) differs from human MVP in other ways. The most significant morphologic changes were noted in the homozygous mice. While homozygous and dual heterozygous forms of FBN1 mutations in children have been reported, they appear to be lethal at an early age (30), as was the case in the homozygous mice in the Ng et al. study (26), and thus the homozygous model may be less applicable to the evolution of the human disease. Second, although the histology of the valves in this study is not described in detail, the hypercellularity of the leaflets is striking. An increase in interstitial cells has been noted in surgical specimens and valves from MVP patients; however, hypercellularity is not generally a feature of MVP (31). The difference may lie in the fact that the valves available for study in MVP patients generally come from autopsy or have been surgically removed and thus represent advanced disease, while in the Ng et al. study, the valves were examined at 0.5, 2.5, 4.5, and 6.5 postnatal days. It would be interesting to study the histologic features of the valves in older heterozygous mice to see whether they are similar to those found in clinical MVP. The morphological changes of the mitral valve in this murine model were evident shortly after birth, while idiopathic MVP is uncommon in children and generally becomes manifest after the adolescent growth spurt. It may be that clinically undetectable changes are present at birth in MVP and only become apparent later in life, but this remains to be shown. While an increase in TGF-β activity may be shown to be present in the idiopathic form of MVP (particularly given the multiplicity of activators, including stress), both the presence and primacy of this relationship have yet to be demonstrated.

Future studies using this model would ideally include a more detailed description of the anatomic features of the murine model. A complete echocardiographic assessment of MVP and of the functional consequences of the valve morphology on the left-atrial and left-ventricular size and left-ventricular function at birth and in older mice would be of particular interest, as would echocardiographic assessment of the effects of TGF-β-neutralizing antibodies on the aortic dilation present in the model. Since in humans there is considerable phenotypic variability among individual genotypes, it would also be interesting to see how consistent the phenotypic expression is in the murine model. Importantly, while prolapse is noted at 9 months of age in heterozygous mice, it would be useful to show that the treated heterozygous mice do not also go on to develop prolapse due to incomplete protection by the TGF-β–neutralizing antibodies.

In summary, it appears that MVP is a final common pathway for a variety of genetic and acquired disorders, which presumably weaken connective tissue of the valve and lead to leaflet elongation, thickening, and often degeneration. The finding that TGF-β dysregulation in the connective tissue plays an important role in the development of prolapse in Marfan syndrome (26) raises the question as to whether this cytokine may play a role in other forms of prolapse, including sporadic MVP. It has been suggested that the altered extracellular matrix in fibrillin-1-deficient mice cues cells to
remodel the matrix, and this remodeling is associated with inappropriate and persistent expression of TGF-β. Localized trauma-related disruption of fibrillin and collagen might also trigger TGF-β activation and matrix remodeling. However, a great deal of study remains necessary before the pathogenic mechanisms underlying idiopathic MVP and its relationship to more generalized forms of connective tissue disease are clarified.

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