Introduction

More than two decades have elapsed since the discovery that sarcomere gene defects cause familial hypertrophic cardiomyopathy (HCM). Since then, genetic testing in HCM has developed and expanded, and is now widely available as a potential clinical service in the Western countries. In the meantime, however, the cross-talk between geneticists and clinicians has developed slowly, and still remains unstandardized, with modalities of interaction and degree of mutual comprehension that vary wildly in various settings. In addition, clinicians often question the clinical utility of genetic testing in HCM patients and their families. The apparent lack of practical benefit, in the face of considerable costs, has long hindered large-scale diffusion of genetic testing particularly in developing countries, and still accounts for understandable (but not always justifiable) resistance on the part of the physicians. However, such resistance is in contrast with considerable evidence supporting a role for molecular diagnosis in tailoring management for HCM patients. We here review several sound clinical reasons in favour of systematic genetic testing in HCM, ranging from identification of complex genotypes, heralding severe disease expression and outcome, to the added benefit of multidisciplinary genetic teamwork, enhancing awareness towards inheritable diseases in the cardiology community. We hope to show that to underestimate the clinical potential of genetic testing in HCM, and to defer its implementation until more advanced knowledge becomes available, is to lose an important opportunity for present improvement in care.

Abstract

More than two decades have elapsed since the discovery that sarcomere gene defects cause familial hypertrophic cardiomyopathy (HCM). Since then, genetic testing in HCM has developed and expanded, and is now widely available as a potential clinical service in the Western countries. In the meantime, however, the cross-talk between geneticists and clinicians has developed slowly, and still remains unstandardized, with modalities of interaction and degree of mutual comprehension that vary wildly in various settings. In addition, clinicians often question the clinical utility of genetic testing in HCM patients and their families. The apparent lack of practical benefit, in the face of considerable costs, has long hindered large-scale diffusion of genetic testing, particularly in developing countries, and still accounts for understandable (but not always justifiable) resistance on the part of the physicians. However, such resistance is in contrast with considerable evidence supporting a role for molecular diagnosis in tailoring management for HCM patients. We here review several sound clinical reasons in favour of systematic genetic testing in HCM, ranging from identification of complex genotypes, heralding severe disease expression and outcome, to the added benefit of multidisciplinary genetic teamwork, enhancing awareness towards inheritable diseases in the cardiology community. We hope to show that to underestimate the clinical potential of genetic testing in HCM, and to defer its implementation until more advanced knowledge becomes available, is to lose an important opportunity for present improvement in care.

Genetic testing for hypertrophic cardiomyopathy: ongoing voyage from exploration to clinical exploitation

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left atrial dilatation, redundant mitral leaflets, may all suggest the presence of a mild phenotype. However, the only gold standard by which such suspicion can be tested is a positive mutational status for HCM, especially in presence of positive family history. Therefore, in the last decade, large-scale genetic screening performed at referral centres has been key to validating the most subtle findings provided by state-of-the-art imaging, and to expand the HCM disease spectrum towards its milder end. Conversely, in HCM families with unavailable or negative genetic test, many individuals remain very difficult to label as affected or non-affected based on clinical assessment alone.

In addition, genetic testing is obviously a sine qua non for the identification of genotype-positive, phenotype-negative individuals, a condition also known as pre-hypertrophic. As previously discussed, the impressive accuracy of current imaging standards is causing a constant shift in HCM phenotype paradigms, so that the definition of truly phenotype-negative individuals is becoming progressively more restrictive. For example, isolated mitral valve and subvalvar abnormalities represent primary expressions of HCM which must be taken into consideration in family members, even in the absence of hypertrophy. Nevertheless, due to the incomplete penetrance of HCM-causing mutations, a significant subset of genotype-positive individuals will be clinically unaffected and exhibit a truly negative phenotype in any age group.

The clinical history of this peculiar pre-hypertrophic subset is still largely undefined, but is likely associated with benign clinical course and outcome, even though some of these individuals will develop a clear-cut HCM phenotype over time. In any case, the identification of a genotype-positive status remains important in this subgroup, because of obvious implications for disease transmission to the offspring, deserving appropriate counselling, as well as implementation of follow-up. Furthermore, young phenotype-negative individuals represent ideal candidates for pharmacological trials aiming at preventing phenotype development, based on encouraging pre-clinical data obtained in transgenic animal models of HCM.

An important caveat regards the issue of genetic testing in children from families with HCM. While regular clinical surveillance remains key in this age group, it is our view that minors should not be offered mutational screening in the absence of reasonable clinical evidence of disease. Because the prognostic relevance of a pre-hypertrophic phenotype remains to be determined, it is important that each individual may freely decide for himself/herself, whether a genotype-positive status should be identified in case of a clinical equivocal diagnosis. In the absence of defined practical advantages (with the only, uncertain and debated exception of reducing risk by restraining sports participation), the burden of anxiety and potential adverse consequences of genetic testing are likely to outweigh benefits in this specific context. Needless to say, this line of reasoning does not apply to minors in whom a clinical diagnosis of HCM can be made based on standard criteria, either as probands or during family screening. In such occurrence, genetic testing is indicated according to the rules employed for adult patients, pending parental approval.
Sarcomere hypertrophic cardiomyopathy as a progressive disease – the case of complex genotypes

Early reports promoted the idea that genotyping of HCM patients may prove useful for risk stratification and prediction of long-term outcome. Subsequent studies, however, have consistently failed to establish meaningful relationships between specific sarcomere myofilament genes, phenotype and outcome. Quite the opposite, the overlap of clinical and pathophysiologic correlates among different genes has shown that genotype-positive, myofilament HCM is a protean but inherently consistent disease, characterized by high prevalence of familial trait and substantial progression towards heart-failure related complications. In this respect, genotype-negative disease appears to behave differently, in that a familial trait is less prevalent, and the long-term course significantly more stable and benign.

Patients with complex genotypes represent an important case in point supporting the concept of myofilament HCM as a progressive disease. A number of independent studies consistently show that multiple sarcomere defects, independent of whether affecting the same or different genes, are associated with earlier onset and more severe clinical phenotype and disease course. In a recent paper describing four HCM patients with triple myofilament gene mutations, representing rare and extreme paradigms of complex genotype, two had a history of resuscitated cardiac arrest or appropriate defibrillator intervention, and three progressed to end-stage HCM by the fourth decade, requiring cardiac biventricular pacing or transplantation (Figure 2). These adverse consequences likely reflect more profound derangement of sarcomere mechanics and cardiomyocyte energetics caused by multiple mutations, but may also be mediated by greater impairment of coronary microvascular function, selectively caused by sarcomere gene defects via unknown molecular pathways.

In the specific HCM patient subsets, genetic screening may thus, provide important additional clues to risk stratification, and potentially indicate the need for differential surveillance strategies based on genotype. Specifically, patients with multiple mutations should be considered for close clinical and imaging follow-up, in order to allow timely recognition of disease progression and increased arrhythmic risk. In the presence of an initial decline in systolic function, the initiation of ACE-inhibition or angiotensin receptor blockade might prevent further LV remodelling and overt systolic dysfunction.

Furthermore, patients with complex genotypes may require heightened attention with regard to primary prophylaxis of sudden cardiac death.

The issue of pre-natal diagnosis

In individual HCM families with particularly severe outcome, mostly related to multiple juvenile sudden deaths, prenatal diagnosis has been successfully employed to terminate pregnancies in which the foetus carried the disease-causing mutation. Nevertheless, because of the uncertain genotype-phenotype relationship, as well as the unpredictable, often benign course of HCM, such an aggressive approach should be considered only in extremely selected instances.

Re-thinking the diagnosis in genotype-negative patients

To date, genotype-negative HCM represents a composite entity, probably comprising a multitude of rare, heterogeneous, and yet-to-be identified HCM-susceptibility genes, as well as, potentially, sporadic disease of non-genetic aetiology. In clinical practice, each patient with unequivocal HCM phenotype and negative first-line genetic screening deserves in-depth individual assessment of alternative molecular diagnoses, including rare HCM-causing genes and phenocopies. In addition, particularly in the young, a negative mutational screening for canonical HCM genes should prompt the search for subtle extra-cardiac manifestations, including abnormal facial and body features and minor renal impairment or neurological deficits. These signs, often overlooked by cardiologists, might point diagnostic efforts in the right direction, highlighting the importance of multidisciplinary approach involving clinical geneticists.

The added value of genetic counselling in clinical practice

An indisputable, although not always appreciated benefit of systematic genetic testing in HCM lies in the cross-fertilization between cardiologists and geneticists. The former, generally show limited expertise and propensity at investigating the hereditary nature of cardiac diseases, and at identifying complex, syndromic phenotypes associated with cardiomyopathies (such as Noonan’s, Leopard’s, mitochondrial disease and Anderson Fabry, just to name a few). Standard protocols for HCM genetic testing routinely include pre-test counselling by a multidisciplinary team involving clinical geneticists (Figure 3). This is a valuable moment for reciprocal education among professionals, ultimately benefiting a wide spectrum of patients with rare conditions. Of note, diseases ranging from idiopathic LV dysfunction to bicuspid aortic valve may have a clear familial background that is all too often neglected in the course of routine clinical evaluations. Implementing systematic protocols for genetic screening of a single entity, in our case HCM, generally proves an important step towards improved recognition and management of inherited heart disorders at large.

Conclusions

Genetic testing for HCM cannot be regarded as an expensive academic gadget, but rather represents a valuable tool in the clinical armamentarium. While major questions remain unanswered, calling for renewed efforts in translational research, what has been achieved so far is sufficient to establish the role of genetic testing in tailored management of HCM patients and their families. To underestimate the actual potential of genetic testing in HCM, and to defer its implementation until more advanced knowledge is available, is to lose an important opportunity for present improvement in care.

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