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Functional Analysis of the Mutations in the Human Cardiac β-Myosin that Are Responsible for Familial Hypertrophic Cardiomyopathy

Implication for the Clinical Outcome

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Abstract

More than 30 missense mutations in the β-cardiac myosin heavy chain gene have been shown to be responsible for familial hypertrophic cardiomyopathy. To clarify the effects of these point mutations on myosin motor function, we expressed wild-type and mutant human β-cardiac myosin heavy chains in insect cells with human cardiac light chains. The wild-type myosin was well purified with similar enzymatic and motor activities to those of the naturally isolated V3 cardiac myosin. Arg^{430}→Gln and Arg^{453}→Cys mutations resulted in decreased actin translocating activity (61 and 23% of the wild-type, respectively) with decreased intrinsic ATPase activity. Arg^{453}→Gln mutation greatly decreased actin translocating activity (27% of wild type) with a 3.3-fold increased dissociation constant for actin, while intrinsic ATPase activity was unchanged. Val^{456}→Met mutation only mildly affected the actin translocating activity as well as ATPase activity of myosin. The degree of deterioration by each mutation was closely correlated with the prognosis of the affected kindreds, indicating that myosin dysfunction caused by the point mutations is responsible for the pathogenesis of the disease. Structure/function relationship of myosin is discussed. (J. Clin. Invest. 1996. 98:2866–2873.)

Key words: cardiomyopathy, hypertrophic • mutagenesis • myosin heavy chains • point mutation • genetics, medical

Introduction

Familial hypertrophic cardiomyopathy (FHC) is an autosomal dominant disease characterized by a hypertrophied left ventricle in the absence of other causes of cardiac hypertrophy. While FHC is one of the causes of sudden and unexpected cardiac death, particularly in healthy young individuals, there is no specific clinical feature which identifies the patients at risk of sudden death. At the present time, > 30 different missense mutations in the β-cardiac myosin heavy chain (MHC) gene, which alter only a single amino acid residue, have been described (3–5). Phenotypic examinations of the genotypically defined kindreds showed that certain β-cardiac myosin heavy chain gene mutations carry greater risk of morbidity than others (6–9), demonstrating that mutations in β-cardiac MHC must be responsible for the pathogenesis of FHC. Since each of the altered amino acids has been highly conserved during evolution (6), it has been suggested that these residues are crucial for myosin function. However, it remained to be clarified how these myosin mutations affect myosin function, particularly its motor activity, thus causing cardiac malfunction.

One of the problems for physiological and/or biochemical studies of FHC is the difficulty of obtaining enough myosin sample from a patient’s heart. To avoid this difficulty, Cuda et al. obtained biopsy samples from patient’s slow skeletal muscle fibers which also express β-MHC along with skeletal muscle myosin and attempted to evaluate the effects of mutations on the β-myosin motor function (10). They selected β-myosin from the mixture of skeletal muscle myosin and β-myosin using a β-MHC specific antibody and showed that β-myosin from the patients translocated actin filaments much slower than normal control. However, their preparation is a mixture of the mutated and nonmutated β-myosins and it is postulated that the ratio of the normal and mutant β-myosins in patient’s heart would be different from that in slow skeletal muscle fiber. Furthermore, since the data represent the activity of β-myosin of unknown ratio of the normal and mutant myosins, it is difficult to quantitatively evaluate the function of mutant β-myosin. Obviously, such an analysis can only be made with pure β-myosin mutant.

On the other hand, Sweeney et al. expressed rat α-cardiac myosin in insect cells and reported that Arg^{453}→Gln mutation resulted in reduction of actin activated ATPase activity and actin translocating activity (11). Problem is, however, only β-MHC but not α-MHC is expressed in human adult left ventricle (12) and V1 cardiac myosin isoform, homodimer of α-MHC, has three to fourfold higher activity than V3 isoform, homodimer of β-MHC, in both actin-translocating velocity and actin-activated ATPase activity (13–16). Divergence between α- and β-MHCs clusters in functionally important regions including ATP, actin, and light chain–binding sites (17). Therefore, the effect of the point mutation on the rat α-cardiac myosin may be different from that on the human β-myosin.

The purpose of this study is to express functionally active recombinant human β-myosin mutants found in the FHC patients and to analyze the effects of each mutation on the motor activity. The active human β-MHCs expressed with human ventricular myosin light chains were successfully purified and the motor and enzymatic activities of myosin mutants were determined. Each β-MHC mutant showed abnormal function ac-
cording to the site and the type of the mutation. Interestingly, the degree of deterioration by the mutation was well correlated with the mortality rate among the affected kindreds supporting the idea that the alteration of β-myosin motor activity by mutation is the critical factor for the prognosis of FHC patients.

Methods

Protein preparation. Cardiac V3 myosin was prepared from an autopsied human left ventricular muscle as described previously (14). Cardiac heavy meromyosin (HMM) and myosin subfragment 1 (S1) were prepared by α-chymotrypsin digestion (18). Actin was prepared from rabbit skeletal muscle acetone powder (19).

Construction of transfer vectors. Total RNA was prepared from a left ventricular muscle obtained during autopsy of a 44-yr-old patient without clinical and pathological evidence of heart disease (20). With two primers designed based upon the published sequence (21, 22), a truncated human β-cardiac MHC cDNA (3414 bp) encoding Met1-Asp1138 was obtained by reverse transcriptase-coupled PCR (RT-PCR) (23), generating a stop codon at 3415 bp and flanking XbaI sites on both ends (Fig. 1). The forward primer was 5’-CGCCTGGCCTAGAATGGGAGATCCTCGGAGATG-3’, and the reverse primer was 5’-GCTCTCAGACTAGTCTGAGCGGAGCTTCTCCAC-3’. This PCR product was digested with XbaI and subcloned into pBluescript SK II (+) (Stratagene, La Jolla, CA). The sequence of the obtained cDNA was confirmed to be identical to that published previously by dideoxynucleotide termination method using Sequenase 2.0 (United States Biochemical, Cleveland, OH). Four point mutations, 746G→A, 1208G→A, 1357C→T, and 1816G→A were introduced into the β-MHC cDNA individually using a transformer site directed mutagenesis kit (Clontech Inc., Palo Alto, CA). These mutations resulted in substitutions of Arg249 for Gln (Arg249Gln), Arg403 for Gln (Arg403Gln), Arg453 for Cys (Arg453Cys), and Val606 for Met (Val606Met), respectively, which have been shown to be responsible for FHC (6). Wild and mutant human cardiac β-MHC cDNAs thus prepared were excised with XbaI digestion, and subcloned into a baculovirus transfer vector, pBlueBacM (Invitrogen, San Diego, CA) using an unique Nhel site.

A full length cDNA encoding the human ventricular light chain 1 (LC1) (24) was a generous gift from Dr. Yazaki (University of Tokyo, Tokyo, Japan). The cDNA was excised from a eukaryotic expression vector and subcloned into pBlueBacM as a eukaryotic expression vector.

**Figure 1.** Construction of the human β-MHC cDNA transfer vector. A detailed strategy for constructing the transfer vector is described in Methods. The first 3414 bp of the 5’-terminus of human β-MHC cDNA were synthesized by RT-PCR from human left ventricular total RNA. The cDNA fragment is represented as a shaded box. ATG and TAG indicate initiation and termination codons, respectively. The PCR product was excised with XbaI digestion and subcloned into pBluescript SK II (+). After introduction of a missense mutation, the cDNA was excised with XbaI digestion and subcloned into a baculovirus transfer vector, pBlueBacM.
vector, pDR540 (Pharmacia LKB Biotechnology Inc., Uppsala, Sweden), with BamHI digestion and subcloned into pBlueBacM using a BamHI site. A full length cDNA encoding the human ventricular light chain 2 (LC2) was obtained from human left ventricular total RNA by RT-PCR with primers designed based upon the published sequence (25), generating flanking XbaI sites on both ends. The forward primer was 5'-CGCATCTAGAATGGCACCTAAGAAAGCAAAG-3'. The reverse primer was 5'-GACTTCTAGACTAGTCTTCTCTTCCG-3'. This PCR product was digested with XbaI and subcloned into pBlueBacM using a NheI site.

**Purification of the expressed human cardiac myosin.** Recombinant baculovirus was obtained for each cDNA construct by the protocols described by O’Reilly et al. (26). To express a truncated human ventricular myosin fragment, Sf21 cells were coinfected with the recombinant viruses expressing truncated β-MHC, LC1, and LC2 (11, 27). The cells were harvested 84 h after infection. The expressed human cardiac myosin was purified as described previously (27).

**Gel electrophoresis, immunoblot, and ATPase assay.** SDS polyacrylamide gel electrophoresis was carried out on a 7.5–20% polyacrylamide gradient slab gel. Immunoblots were performed as described previously (28) using an anti-human β-MHC monoclonal antibody (HMC14) and an anti-human ventricular LC1 antibody (MLM544) (generous gifts from Dr. Katoh, Yamasa Corp., Chiba, Japan). Nondenaturing gel electrophoresis was performed according to Persechini et al. (29) in the presence of 2 mM MgCl2–ATP. ATPase activity was measured as described previously (30) under conditions described in the legend to Table I.

**In vitro motility assay.** The in vitro motility assay was performed as described previously (14). Each video frame was digitized at a rate of five frames/s into a 320 × 240 pixel array by a video grabber card (Video Blaster; Creative Labs, Milpitas, CA) installed in a personal computer (Compudyne 614295; CompUSA, Dallas, TX). Actin filament velocity was calculated from the movement distance and the elapsed time in successive snapshots.

**Statistical analysis.** For each myosin, measurements were made on three different preparations expressed independently. All data are presented as mean value ± SD. Student’s t test was used for statistical comparison of mean values. A value of P < 0.05 was considered to be significant.

**Results**

**Expression and purification of human β cardiac myosin.** The produced human β-MHC contains Met1–Asp1138 with calcu-
lated molecular mass of 130 kD thus containing the entire S1 and most of the S2 portions of myosin. We chose to produce the heavy chain construct containing both the S1 and S2 portions of the molecule, because the S2 segment is necessary for a stable two-headed structure which is important to represent native motor activity of myosin (27) and because it is technically difficult to express an entire β-MHC (1,935 amino acids) in insect cells. After purification, both wild and mutant myosins contained a 130-kD apparent molecular mass heavy chain, a 25-kD LC1, and a 19-kD LC2 (Fig. 2 A). Approximate stoichiometry of the three polypeptides was 1:1:1 for both wild and mutant β-MHCs based upon gel densitometry, indicating that the recombinant myosins consisted of one heavy chain and one of each class of light chains as is found in naturally isolated cardiac V3 myosin. For both wild-type and mutant myosins, ~ 200 μg of the purified myosins were obtained from 10 grams of S21 cells.

To confirm the authenticity of the expressed proteins, immunoblot analyses were performed using two kinds of antibodies recognizing either human β-myosin heavy chain (HMC14) or human ventricular LC1 (MLM544) (31). The former antibody recognized the expressed truncated β-MHC as well as a naturally isolated human ventricular MHC (Fig. 2 B). The latter antibody reacted with both the expressed LC1 and the naturally isolated one (Fig. 2 C) suggesting that the expressed myosin fragment is the human cardiac β-myosin.

Nondenaturing gel electrophoresis was used to determine whether the expressed β-MHC forms a double-headed structure or a single-headed structure. As shown in Fig. 3, the entire fraction of the expressed myosin migrated at the similar position to that of the naturally isolated cardiac heavy meromyosin (HMM) but much slower than naturally isolated cardiac S1, suggesting that the expressed β-MHC forms a stable dimer, i.e., V3 isoform.

**ATPase activities of the expressed myosin.** The enzymatic function of the expressed myosins was evaluated by measuring K⁺EDTA-activated and Ca²⁺-activated ATPase activities as well as the actin activated Mg²⁺-ATPase activity. The obtained values for the wild-type recombinant β-MHC ATPase are shown in Table I. These values are similar to those measured for naturally isolated V3 cardiac HMM (32) or S1 (33) indicating that the expressed β-myosin is properly folded and retains native enzymatic properties. While the K⁺EDTA-ATPase activity as well as Ca²⁺-ATPase activity of the Arg403Gln and the Val606Met mutant myosins was not significantly different from those of the wild-type myosin, the Arg249Gln and Arg453Cys mutants showed a large difference. The Arg249Gln mutant showed a decreased Ca²⁺-ATPase activity without significant change in K⁺EDTA-ATPase activity, while the Arg453Cys myosin mutation resulted in significant decrease in both K⁺EDTA- and Ca²⁺-ATPase activities, suggesting that these mutations change the conformation of the ATPase active site.

Consistent with these results, the V_max of actin-activated ATPase activity was significantly decreased for both Arg249Gln and Arg453Cys mutants without changing the apparent dissociation constant for actin (K_c). On the other hand, while both Ca²⁺- and K⁺EDTA-ATPase activities were unchanged, the Arg403Gln mutant showed a significantly decreased V_max (31% of the wild-type) and a significantly increased K_c (3.3 times higher than that of the wild-type) for the actin activated ATPase activity suggesting the change in the actin dependent process of myosin ATPase reaction rather than the direct effects on the ATPase active site. The ATPase activities of the Val606Met mutant were not significantly different from those of the wild-type myosin.

**Actin-translocating activity of the expressed myosin fragments.** Fig. 4 shows the sliding velocity of actin filaments on the recombinant human cardiac myosins. More than 90% of actin filaments moved smoothly and unidirectionally on all myosins. There was no significant difference between the wild-type and the mutant myosins in the proportion of the moving actin filaments. The sliding velocity for the wild-type myosin was 1.64±0.14 μm/s at 30°C. This value is similar to that of naturally isolated V3 cardiac myosin (14–16), further confirming the authenticity of the expressed myosin. The sliding veloc-

![Figure 3. Nondenaturing gel electrophoresis of cardiac myosin fragments. (lane 1) Human cardiac HMM; (lane 2) truncated cardiac myosin expressed in insect cells (wild-type); (lane 3) human cardiac myosin S1.](image)

Table I. ATPase Activities of the Expressed Human Cardiac Myosins

<table>
<thead>
<tr>
<th></th>
<th>K⁺EDTA ATPase*</th>
<th>Ca²⁺ ATPase†</th>
<th>V_max (s⁻¹)</th>
<th>K_c (μM)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>nmol Pi/mg per min</td>
<td>nmol Pi/mg per min</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Wild type</td>
<td>198±15</td>
<td>480±34</td>
<td>3.31±0.30</td>
<td>30±4</td>
</tr>
<tr>
<td>Arg249Gln</td>
<td>163±35</td>
<td>111±17</td>
<td>1.88±0.23</td>
<td>35±5</td>
</tr>
<tr>
<td>Arg403Gln</td>
<td>187±11</td>
<td>477±5</td>
<td>1.01±0.15</td>
<td>98±7</td>
</tr>
<tr>
<td>Arg453Cys</td>
<td>74±13</td>
<td>161±8</td>
<td>0.79±0.11</td>
<td>38±5</td>
</tr>
<tr>
<td>Val606Met</td>
<td>168±33</td>
<td>451±11</td>
<td>2.98±0.39</td>
<td>41±6</td>
</tr>
</tbody>
</table>

Measurements were made on three independent preparations of each myosin. All data are presented as mean activity±S.D. *K⁺EDTA-ATPase activity was measured in 0.01 mg/ml myosin, 0.1 mM ATP, 10 mM EDTA, 50 mM Tris-HCl (pH 7.5), 0.6 M KCl at 25°C. The reaction was started by adding ATP and the liberated inorganic phosphate was measured as described previously (30). †Ca²⁺-ATPase activity was measured in 0.01 mg/ml myosin, 0.1 mM ATP, 2 mM CaCl₂, 50 mM Tris-HCl (pH 7.5), and 0.04 M KCl at 25°C. ‡Actin-activated ATPase activity was measured at 25°C in 0.01 mg/ml myosin, 0.1 mM ATP, 50 mM KCl, 20 mM imidazole-HCl (pH 7.5), 6 mM MgCl₂, 0.5 mM EGTA with various concentrations of F-actin (0.1–3.2 mg/ml). The maximum hydrolysis rate (V_max) and the apparent dissociation constant for actin (K_c) for each preparation were determined by using a computed nonlinear least-square curve-fitting program based on a equation, V = V_max/(1 + K_c/[actin]) (42). *P < 0.05 versus wild type.
motor function in terms of ATPase and unloaded actin translocating activity, indicating that these point mutations do not affect proper folding of myosin. Wild-type, indicating that these point mutations do not affect proper folding of myosin.

Particularly, the Arg403Gln and the Arg453Cys mutations most decrease in myosin motor activity by the mutation. Arg403Gln and Arg453Cys, which decrease enzymatic and motor activities severely (70–80% inhibition), are associated with poor prognosis. The Arg249Gln mutation, whose effects on myosin motor function are intermediate (40–50% inhibition), is associated with intermediate prognosis. The Val606Met mutation, which affects myosin function mildly (10–15% inhibition), is associated with good prognosis. The results suggest that myosin dysfunction caused by the point mutations is responsible for the pathogenesis of FHC, although the relative expression level of mutant myosin in patient’s heart might also contribute to the pathogenesis since the expression of the mutant myosin is heterogeneous where the percent mutant expression is unknown from individual to individual. Furthermore, the results of present study imply that the functional analysis of a certain mutation observed in FHC could predict the clinical outcome of the kindreds carrying such a mutation, whose phenotype has not been clinically studied extensively.

The present results also provide fundamentally important information for understanding the structure/function relationship of myosin. The mutation of Arg249Gln or Arg453Cys not only resulted in the decrease in actin translocating activity but also accompanied with a significant reduction of Ca2+-ATPase activity. While the Arg249Gln mutation markedly decreased Ca2+-ATPase activity, the inhibition of K+-ATPase activity was moderate. This might suggest that Arg469 influences the coordination of the metal ion of the ATP substrate to myosin molecule. According to the three-dimensional structure of skeletal muscle myosin S1 (35–37), Arg254 (corresponding to Arg255 of skeletal myosin) is in the β-strand six located at the outer end of the nucleotide binding pocket (Fig. 5 C). This residue is in close proximity of Asp456 (11.6 Å between Arg249 and Asp456 in skeletal myosin) which participates in the coordination of the Mg2+ ion of the ATP substrate (35, 38, 39) which is consistent with the observed change in myosin function. On the other hand, the Arg453Cys mutation decreased both Ca2+- and K+-EDTA ATPase activities to similar extent, suggesting that the mutation influences either ATP binding or the catalysis of ATP hydrolysis. Arg469 is located at the end of the nucleotide binding pocket (37). Although this residue is a little distal from the residues participated in γ-phosphate binding (~ 20 Å), it resides in a loop next to the α-helix containing Asp461, Ala463, Gly464, and Gly466 (corresponding to Asp461, Ser464, Gly465, and Gly466 of Dictyostelium discoideum myosin) which are suggested to participate in the binding of γ-phosphate of ATP (38, 39). One of the residues close to Arg469 is Glu448 (10.8 Å between Arg469 and Asp469 in skeletal myosin) and it would be plausible that charge interaction between Arg469 and an acidic residue contributes to position the α-helix containing the several residues involved in γ-phosphate binding.

Different from above mutations, the Arg403Gln mutation did not change the Ca2+- and K+-EDTA ATPase activities significantly but only decreased the actin dependent ATPase ac-

Figure 4. Sliding velocity of actin filaments on the expressed myosins. Actin movement was observed in 30 mM KCl, 5 mM MgCl2, 25 mM imidazole-HCl (pH 7.5), 1 mM EGTA, 1% 2-mercaptoethanol, 0.5% methylcellulose, 4.5 mg/ml glucose, 216 μg/ml glucose oxidase, 36 μg/ml catalase, and 2 mM ATP at 30°C. Measurements were made on three independent preparations of each myosin. 20–30 actin filaments were measured to obtain an average velocity for each preparation. All values are mean velocities ± SD.
The result suggests that this mutation changes the ATPase reaction steps involved in the actin dependent process (40). Consistent with this finding, the apparent dissociation constant ($K_a$) was significantly increased for Arg403Gln mutant. The observed functional change is well explained by structural point of view that Arg$^{403}$ lies in the loop (Arg$^{403}$-Lys$^{413}$) which forms a close contact with actin (Pro$^{332}$ to Glu$^{334}$ of actin molecule) (Fig. 5 B) (36). The Arg403Gln mutation greatly decreased the actin translocating activity suggesting the importance of this loop for motor activity of myosin.

Val$^{606}$ is in a helix-loop-helix motif just at the upstream of the junction of 50–20 kD subdomains of S1 which contain a part of actin binding interface of myosin molecule (35, 37). The effect of Val606Met mutation on myosin motor and enzymatic activities was mild. On the basis of the structural analysis of acto-S1 complex (36), there are no indications of the importance of this segment for actin–myosin interaction, therefore, the very mild modulation of myosin function by the Val606-Met mutation would be reasonable, although the mild functional modulation would be partly due to the conserved substitution of the amino acid residue, i.e., Val to Met.

It is still unclear how a missense mutation leads to cardiac hypertrophy, myocardial fiber disarray, or fatal arrhythmia, although it may be plausible that myosin motor dysfunction constitutes the primary stimulus for the cardiac hypertrophy as compensation for the reduced contractile function resulting from the mutation.
from the dysfunction of the individual cross-bridge. Although the four mutations tested in this study were different in the degree of the dysfunction of myosin motor, the patients carrying the mutations on the β-MHC tend to produce cardiac hypertrophy to a similar extent of each other (1, 2, 6–9), suggesting that cardiac hypertrophy is not directly related to the prognosis of the disease. Consistent with this notion, it was reported that cardiac hypertrophy is not directly related to the prognosis of the patients carrying the mutations on the σ- and β-form myosin heavy chain complementary DNA clones. J. Clin. Invest. 82:524–531.


