Spinal and bulbar muscular atrophy: a motoneuron or muscle disease?
Cynthia L Jordan¹ and Andrew P Lieberman²

Kennedy disease (KD, or spinal and bulbar muscular atrophy) is caused by a CAG/polyglutamine expansion in the androgen receptor (AR) gene. Both motoneurons and muscles are affected by KD, but where mutant ARs act to initiate this disease is not clear. We discuss recent insights into this disease with two main themes. (1) KD is androgen-dependent, suggesting that blocking androgen action may be an effective treatment. (2) Androgens may trigger KD by acting in muscles, which indirectly affects the motoneurons, suggesting that blocking AR function in muscles may rescue motoneurons from disease and provide an effective treatment. Future research will provide a better understanding of how androgens trigger KD and the relative contributions of motoneurons versus muscles in this disease.

Introduction

Many disorders exhibit striking sex-biases in the human population, with some more common in males and showing an X-linked pattern of inheritance. One such disease affecting only men is spinal and bulbar muscular atrophy and Kennedy disease (KD), a so-called ‘lower motoneuron disease’ characterized by an adult-onset, slowly progressive loss of motor function. First described by Kennedy et al. [3], a core feature of KD is a progressive weakness and wasting of muscles, which interferes with the performance of routine motor tasks such as walking, lifting and climbing stairs. Although symptoms of KD typically lead to diagnosis in mid-life (between 30 and 50 years), men with KD often report experiencing frequent muscle twitching and/or cramping for a decade or more before diagnosis, suggesting that the disease process begins well before reaching the threshold of clinical muscle weakness. In addition to affecting limb muscles, bulbar symptoms of KD include difficulty swallowing, chewing and enunciating clearly. While individuals with longer CAG repeats tend to develop the disease earlier, there is marked variation in disease onset even among related individuals with the same CAG repeat size [4,5]. Female carriers with one or two copies of the KD allele commonly experience subclinical symptoms (e.g. muscle twitching) without significant loss in muscle strength [5,6]. Another core feature of KD is reduced androgen-sensitivity, such as gynecomastia and testicular atrophy [3,5,7,8], suggesting a partial loss of normal AR function in individuals with KD. However, it is not this loss of normal function but rather a gain of new and toxic functions conferred by the expanded glutamine tract that causes the neuromuscular pathology of KD, since humans inheriting completely dysfunctional AR do not appear.
prone to such weakness. Finally, blood creatine kinase levels are consistently elevated in KD patients, suggesting that muscle damage is significant in this disease.

**Neurophysiological measures indicate loss of motor and sensory axons and compensatory sprouting in KD**

Nerve conduction and electromyographic studies suggest that the muscle atrophy and weakness in KD are caused by a progressive loss of motor axons, resulting in denervation and atrophy of muscle fibers [5,8]. High amplitude motor unit potentials are also detected, suggesting that muscle fibers are denervated, nearby surviving motor axons sprout to re-innervate them. Such studies also indicate a robust loss of sensory axons in men with KD. Which sensory neurons are affected and the degree of loss within different populations are not well documented, but the consistent absence of tendon reflexes in men with KD suggests that the sensory neurons most affected are the large caliber 1A afferents that relay information to the motoneurons about muscle stretch. Thus, it is likely that losses in both motor and sensory neurons contribute to the loss of motor function in KD.

**Histopathological correlates confirm loss of motor and sensory neurons in KD**

Direct examination of tissues from KD patients reveals a significant loss of motor axons accompanied by a marked and selective depletion of motoneuronal cell bodies in the spinal cord and brain stem [3,8,9]. Moreover, large diameter sensory axons appear selectively depleted from the dorsal roots and the sensory-rich peripheral sural nerve, consistent with the idea that 1A afferents supplying muscle spindles may be selectively lost in this disease. Muscle biopsies, however, suggest that two separate processes may be occurring: (1) ongoing denervation and sprouting indicated by muscle fiber atrophy and fiber-type grouping, and (2) primary myopathy indicated by pathology (e.g. fiber splitting, centralized nuclei, fiber degeneration) typical of diseases that originate in muscle [10]. Another hallmark of KD is an accumulation of mutant AR into insoluble aggregates or inclusions in neuronal and nonneuronal cells [11,12]. While the significance of these microscopically visible aggregates is not clear [12,13**], experimental evidence indicates that toxicity is mediated by soluble forms of the misfolded mutant protein [14*].

**Clinical reports describing KD raise questions**

**What factors besides CAG repeat size determine disease onset and progression?**

Although individuals with longer CAG repeats tend to develop the disease at an earlier age, disease onset can vary by a decade or more for a given CAG repeat size, even within a family [1,5]. Moreover, females having one or both copies of the KD allele show only mild subclinical symptoms and are not deemed as having KD [5,6]. These observations suggest that factors other than the expanded CAG repeat affect expression of this disease.

**Is KD a disease of motoneurons, muscles or both? In other words, where does the mutant AR act to trigger KD?**

Calling KD a ‘motoneuron disease’ carries with it the implicit assumption that the mutant AR acts in motoneurons to cause their death, resulting in denervation-induced atrophy of skeletal muscles and loss of motor function. However, muscles affected by KD undergo deterioration that is not solely attributable to denervation. In particular, men with KD generally show significantly elevated levels of creatine kinase, questioning whether KD involves primary myopathic processes, akin to the muscular dystrophies.

**Experimental models of KD provide some answers and hope for treatment**

Six different mouse models of KD have been described that express full length AR and largely recapitulate the disease phenotype, with the first models described in 2002. Five are transgenic (Tg) models [15,16**,17,18,19**] and one is a knockin (KI) model [20**]. Only one Tg model expresses experimental AR in a cell-specific manner [19**], thus allowing investigators for the first time to address directly questions about which cells AR acts in to cause KD.

**KD phenotype in mice**

Despite various promoters and CAG repeat lengths, the different mouse models show remarkably similar phenotypes, with each exhibiting core features of KD (Supp Table 1). For example, the KD phenotype in mice shows a pronounced sex bias, with experimental males affected more than females, reminiscent of KD in humans. Symptomatic male mice show a marked loss of muscle strength, particularly in their hindlimbs, which impairs their ability to walk normally. Affected males are also typically smaller than controls and are kyphotic (showing a dorsal curvature of the spine), aspects of the KD phenotype that appear unique to rodents. The age of onset of KD symptoms can vary widely among these mouse models, from three weeks to six months, and probably reflects differences in both the size of the CAG repeat and the level of AR expression, but the rise in circulating androgens at puberty probably plays a role in all the models.

**KD models establish that testicular androgens are a key factor triggering disease**

That only men develop KD suggested a role for androgens in this disease, but direct support for this idea came only recently from cell and animal models. For example, castrating KD male mice, either at puberty or as adults, can rescue them from disease [15,16**,19**,20**]. Moreover, testosterone (T) treatment has been shown in some models to ‘turn on’ the disease in asymptomatic mutant females, inducing a KD phenotype comparable to that of gonadally intact mutant males [16**,19**].
Finding that KD is triggered by androgens represents a significant step in understanding disease pathogenesis and demonstrates that an expanded polyQ AR is not sufficient to trigger disease but that male levels of androgens are also required. This explains why females, even when homozygous for expanded CAG AR alleles, do not develop the disease. Moreover, differences in androgen titers may account for some of the variability in the age of onset between individuals that have comparable CAG expansions in their AR gene. Finally and most significantly, the apparent androgen dependence of KD indicates that giving androgens to men with the disease to combat muscle atrophy is contraindicated. Although there are few documented cases of how exogenous androgens affect men with KD, one recent report that androgen treatment worsened symptoms in a patient with KD [21] indicates that the KD phenotype in humans also depends on androgens. A clear implication of these data is that agents which interfere with androgen action may reverse KD symptoms in men (discussed in more detail below).

The mechanism by which androgens stimulate toxicity is of great interest. Changes to AR induced by androgens include translocation from the cytoplasm to the nucleus, altered post-translational modifications, and less stable association with molecular chaperones including Hsp70 and Hsp90. Evidence implicates each of these in contributing to pathogenesis [2]. The nucleus appears to be a critical site for triggering KD pathology [22**]. Whether androgen-driven transcriptional activation by the AR is also a requisite step in the pathogenic pathway is less clear, given that AR antagonists fail to block androgens’ disease-promoting effects in some models despite their ability to block normal transcriptional function via classical androgen responsive elements in these same models [22**,23**]. It is possible that nuclear translocation of AR without activation is sufficient to induce toxicity or that transcriptional activity of AR is also required, but is mediated via nonclassical regulatory elements, akin to tamoxifen-bound estrogen receptors acting at AP1 sites [24]. That the length of the polyQ tract affects cofactor recruitment is consistent with this view [25]. Clearly, considerably more work needs to be done to gain a full understanding of androgens’ role in triggering KD.

Evidence from models shifts the focus from cell death to cell dysfunction as the proximate cause of KD

Motoneuronal cell death has long been assumed to crucially mediate the loss of motor function in KD, but data from KD models suggest otherwise. While two of the six KD mouse models show the expected motoneuronal loss [17,18], four do not [15,16*,19**,20**]. Of these, all four models show an androgen-dependent loss of motor function without any apparent loss of motoneurons, with two providing evidence of a distal axonopathy [19**,20**]. Given that castration can rescue KD male mice from the disease well after its onset [15,19**], cell death is probably a late event in KD and cell dysfunction may be at the root of this disorder.

Axonal trafficking defects have been implicated in many neurodegenerative diseases, including KD. In KD Tg mice, abnormal accumulations of neurofilament occur in distal motor axons before the onset of muscle weakness [26**]. Perhaps more tantalizing is finding the same abnormal accumulation of neurofilament in muscle of a KD patient. Katsuno and colleagues also show more directly that axonal trafficking is impaired, demonstrating androgen-dependent deficits in retrograde labeling of motoneurons and dynactin 1 expression in motoneurons. Dynactin 1 is part of the motor protein complex involved in retrograde trafficking in axons. Alterations in anterograde and/or retrograde axonal transport are also suggested by findings in other KD models [15,27,28]. Thus, an androgen-dependent failure in the trafficking of needed cargo along axons, either anterogradely to motor synapses or retrogradely to motoneuronal cell bodies, may be an early and pivotal event triggering the loss of motor function in KD.

Androgens may act in muscles to trigger KD

One great challenge of any chemical intervention is to eliminate the disease-causing action of an endogenous molecule while preserving its health-promoting actions. Thus, for KD, ideal treatments will be aimed at reducing mutant AR action in the cells where the disease begins while sparing the beneficial effects of AR in other tissues, such as those that support reproduction. Knowing where mutant ARs act to trigger KD would greatly facilitate the development of such treatments.

ARs are expressed by many cell types and tissues, including motoneurons and skeletal muscles [29]. Thus, mutant ARs might act in motoneurons, skeletal muscles, or both to cause KD (Figure 1). However, it has long been assumed that ARs act in motoneurons to cause the disease, an idea driven by the fact that men with KD suffer what appears to be ‘neurogenic atrophy’ of skeletal muscles caused by a loss of motoneurons. However, there is little direct evidence to support the idea that mutant ARs act in motoneurons to initiate disease.

One Tg model of KD expresses wild-type AR only in skeletal muscle fibers, but nonetheless shows an androgen-dependent motor phenotype, including the loss of motor axons akin to other mouse models of KD [19**]. That overexpression of the wild-type AR with 22 CAG repeats triggers disease similar to the expanded polyQ AR suggests that it is mediating toxicity by similar mechanisms and raises the possibility that overexpression of AR in muscle could lead to human disease. Importantly, analysis of these Tg mice led to the conclusion that AR may act in muscle rather than motoneurons to trigger disease. Given that expression of a growth factor, vascular
Schematic showing two possible sites (motoneurons or muscle fibers) where mutant ARs might act to trigger Kennedy disease (KD). Because motoneurons and muscle fibers each express ARs, each represents candidate sites for triggering KD. Mutant ARs could cause the same distal axonopathy (depicted by dashed lines for some motor axons and their terminals) by directly impairing the function of motoneurons or muscle fibers. In one case, distal axonopathy is caused by altered anterograde signaling in the affected motoneuron (shown in red in left panel) while in the other case, axonopathy is induced indirectly via changes in retrograde signals from the affected muscle fibers (shown in red in right panel). On the basis of several mouse models of KD, axonopathy occurs early in the disease process and may involve a perturbation in motoneuronal axonal transport. As this disease progresses, such axonopathy leads to functionally denervated muscle fibers, triggering muscle atrophy and motoneuronal cell death, two key pathological features of KD. It is also possible that AR acts in both cell types to induce the full complement of disease symptoms underlying KD and that AR action at both sites are crucially involved. Studies are needed to fully discern the role of ARs in motoneurons and muscle fibers in causing KD.
endothelial growth factor, is also reduced in muscles of these mice, the disease process may entail a loss of trophic support to motoneurons by muscles. A similar loss of trophic factor expression has been documented in skeletal muscle from a KI mouse model of KD [20**]. While it will be important to test the role of ARs in muscle in other models of KD, these results provide the first direct demonstration that ARs acting in skeletal muscle fibers can trigger a KD phenotype and indirectly cause disease in the motoneurons, including axonal loss and defects in axonal transport (Kemp and Jordan, unpublished data). Such a scenario is in line with the well-established role of muscles in maintaining motoneurons [30] and suggests that therapeutics preventing AR action in muscle may rescue both individuals and their motoneurons from KD.

Molecular targets for therapy emerge from the study of KD models

Hormone dependence

Since the KD phenotype is androgen-dependent, decreasing androgen levels should reduce disease severity. This notion was confirmed by treating KD Tg mice with leuprolerin, a luteinizing hormone-releasing hormone agonist that reduces serum testosterone levels [23**]. Clinical trials to treat KD patients with anti-androgens have been initiated in the United States and Japan, where modest results were reported after a short trial with leuprolerin [26**]. The efficacy of dutasteride, an inhibitor of 5 alpha-reductase, the enzyme that converts testosterone to the more potent dihydrotestosterone, has not yet been reported.

Androgen receptor degradation

The misfolded, polyQ AR protein is the proximal mediator of toxicity in KD, and as such, therapeutic strategies that promote its degradation have attracted considerable attention. ARs associate with the molecular chaperones hsp90 and hsp70, components of the cellular defense against unfolded proteins and possible therapeutic targets. Geldanamycin and its derivatives that inhibit hsp90 impair ligand-dependent trafficking of the receptor, promote its degradation [31] and improve the phenotype of KD Tg mice [32]. Overexpression of hsp70 [33] or the chaperone-dependent E3 ubiquitin ligase CHIP (carboxy terminus of Hsc70-interacting protein) [34] similarly promotes AR degradation to yield phenotypic improvement in KD Tg mice. Additional strategies to promote AR degradation include novel small molecules that disrupt the interaction of the receptor with co-regulators [35], and activation of protein turnover through autophagy [36].

Histone acetylation

Changes in gene expression and chromatin modification accompany cellular dysfunction in KD. These changes may be secondary responses to injury or manifestations of a primary pathway leading to toxicity. In either case, treating cellular [37] and mouse models [38] with histone deacetylase (HDAC) inhibitors, such as sodium butyrate, has proven effective in ameliorating the disease phenotype. Notably, broad spectrum HDAC inhibitors may also influence movement along microtubules that is influenced by HDAC6, a cytoplasmic HDAC. Since this movement machinery is needed for protein degradation, the mechanisms underlying HDACs’ therapeutic effects warrant further study.

Emerging targets

While small molecules offer great promise for treating KD, other strategies are under active investigation. Gene silencing by RNA interference can decrease production of toxic proteins and has proven to be effective in animal models of other polyQ diseases. Similarly, replacement of dysfunctional or dead neurons by stem cells is an attractive strategy whose potential remains to be fully explored.

Conclusion

Recent work in KD mouse models has identified a host of molecular pathways that modulate toxicity. Furthermore, the cell types mediating pathogenesis are being defined to include skeletal muscle fibers in addition to motoneurons. Given that skeletal muscles probably contribute to the disease phenotype and may mediate early pathogenic events, they offer a therapeutic target outside the blood brain barrier that is much more accessible to treatment than motoneurons. It is likely that a combination of targets and treatment strategies will prove to be the most effective therapeutic approach for KD and other chronic neurodegenerative disorders.

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Appendix A. Supplementary data


References and recommended reading

Papers of particular interest, published within the period of review, have been highlighted as:

- of special interest
- of outstanding interest


6 Endocrine and metabolic diseases


10. Soraru G, D’Ascenzo C, Polo A, Palmieri L, Vergani L, Lieberman AP, Merry DE: Molecular determinants for the tissue specificity of SERMs in the mouse model of KD. In which gene targeting was used to introduce 113 CAG repeats into the AR gene. This report describes a knockin mouse model of KD, in which gene targeting was used to introduce 113 CAG repeats into the AR gene. Male mice exhibited androgen-dependent weakness associated with the early onset of myogenic and neuropathic changes in muscle and loss of tropomyosin-4 expression. These pathologic changes preceded the occurrence of motoneuron loss and suggested that disease arising in muscle significantly contributes to the KD phenotype.

11. Li M, Miwa S, Kobayashi Y, Merry DE, Yamamoto M, Tanaka F, Nakanishi K, Kobayashi Y, Yamamoto S, Tabata T, Kato S: Androgen-dependent neurodegeneration by polyglutamine-expanded human androgen receptor in Drosophila. Neurogenetics 2002, 35:855-864. These studies show that the drosophila model of the polyglutamine expanded AR causes neurodegeneration in an androgen-dependent manner but that the effect of androgen can be blocked by preventing AR from entering the nucleus. These results suggest that the role of androgens in AR toxicity is to promote translocation of AR to the nucleus where toxicity of AR is then realized.


15. Li M, Chevalier-Larsen ES, Merry DE, Diamond MI: Evidence based on a number of different methodologies (e.g. retrograde labeling of motoneurons, quantitative RT-PCR, and immunocytochemistry), suggesting that androgens induce a loss of motor function by disrupting the normal balance between the intrinsic motor unit pool size and axon field size. These authors describe a series of experiments using a variety of methods including immunoprecipitation, Westerns and atomic force microscopy to show that soluble oligomers form in an androgen-dependent manner well before disease onset and that such oligomers can form proto-fibrillar structures that may underlie cell dysfunction in KD.

16. Chevalier-Larsen ES, O’Brien CJ, Wang H, Jenkins SC, Holder L, Lieberman AP, Merry DE: Castration restores function and neurofilament alterations of aged symptomatic males in a transgenic mouse model of spinal and bulbous muscular atrophy. J Neurosci 2004, 24:4778-4786. These authors describe a series of experiments using a variety of methods including immunoprecipitation, Westerns and atomic force microscopy to show that soluble oligomers form in an androgen-dependent manner well before disease onset and that such oligomers can form proto-fibrillar structures that may underlie cell dysfunction in KD.

17. Kobayashi Y, Doyu M, Sany C, Mike H, Merry DE, Yamamoto M, Kato S, Katsuno M: androgen receptor oligomers underlie pathology in a mouse model of spinal and bulbous muscular atrophy. Proc Natl Acad Sci U S A 2007, 104:18259-18264. These investigators describe a Tg mouse model that expresses normal AR exclusively in skeletal muscle fibers. Unexpectedly, such mice show an androgen-dependent loss of motor function comparable to that shown by other KD mouse models. The authors conclude that AR may act in muscle fibers to trigger KD.

18. Yu Z, Nadgar N, Albertelli M, Gruis K, Jordan C, Robins DM: Lieberman AP: Androgen-dependent pathology demonstrates myopathic contribution to the Kennedy disease phenotype in a mouse knockin model. J Clin Invest 2006, 116:2663-2672. These findings describe a knockin mouse model of KD, in which gene targeting was used to introduce 113 CAG repeats into the AR gene. Male mice exhibited androgen-dependent weakness associated with the early onset of myogenic and neuropathic changes in muscle and loss of tropomyosin expression. These pathologic changes preceded the occurrence of motoneuron loss and suggested that disease arising in muscle significantly contributes to the KD phenotype.

19. Katsuno M, Adachi H, Doyu M, Minamiyama M, Sany C, Kobayashi Y, Inukai A, Soseue G, Leuporelin rescues polyglutamine-related phenotypes in a transgenic mouse model of spinal and bulbous muscular atrophy. Nat Med 2003, 9:768-773. Using a Tg mouse model of KD, this report established that anti-androgen therapy could ameliorate the disease phenotype. The authors showed that gene targeting was used to introduce 113 CAG repeats into the AR gene. Male mice exhibited androgen-dependent weakness associated with the early onset of myogenic and neuropathic changes in muscle and loss of tropomyosin expression. These pathologic changes preceded the occurrence of motoneuron loss and suggested that disease arising in muscle significantly contributes to the KD phenotype.

20. Shang Y, Brown M: Molecular determinants for the tissue specificity of SERMs. Science 2002, 295:2465-2468. These authors describe a series of experiments using a variety of methods including immunoprecipitation, Westerns and atomic force microscopy to show that soluble oligomers form in an androgen-dependent manner well before disease onset and that such oligomers can form proto-fibrillar structures that may underlie cell dysfunction in KD.


Table 1. Characteristics and outcomes of six different mouse models of KD that express a full length AR having variable CAG repeat lengths and controlled by different promoters. Disease characteristics are noted for only males unless specified.

<table>
<thead>
<tr>
<th>Citation describing model</th>
<th>Model type (Tg or KI)/# CAGs/promoter</th>
<th>Sex bias?</th>
<th>Onset of motor symptom(s)</th>
<th>Motor symptoms</th>
<th>Disease-related changes in motoneurons/motor axons</th>
<th>Disease-related changes in muscle</th>
<th>Sensory loss?</th>
<th>Depends on androgens?</th>
</tr>
</thead>
<tbody>
<tr>
<td>(McManamny, Chy et al. 2002)</td>
<td>Tg /120/CMV</td>
<td>Disease severity: Tg♂ = Tg♀&lt;br&gt;Onset age: Tg♂ &lt; Tg♀ (by wks)</td>
<td>~3 wks</td>
<td>Clasping ↓ cage activity ↓ time on hang test</td>
<td>↓ # MNs♂;&lt;br&gt;↓ MN size</td>
<td>mf* type grouping &amp; atrophy ↓ # of mfs</td>
<td>No data</td>
<td>No data</td>
</tr>
<tr>
<td>(Katsuno, Adachi et al. 2002; Katsuno, Adachi et al. 2006)</td>
<td>Tg/97/ chicken β-actin promoter with CMV enhancer</td>
<td>Disease severity: Tg♂ &gt; Tg♀&lt;br&gt;Onset age: Tg♂ &lt; Tg♀ (by wks)</td>
<td>~8 wks</td>
<td>↓ cage activity ↓ time on rotarod impaired gait</td>
<td>marginal loss of MNs ↓ MN size ↓ motor axon size ↓ retrograde axonal transport Accumulation of NF* in distal axons</td>
<td>mf grouped atrophy small angulated mfs ↑ variability of mf size</td>
<td>DRGs intact</td>
<td>Castration prevents KD symptoms in Tg♂ s; Td induces KD symptoms in Tg♀ s; deficits in axonal transport depend on androgens</td>
</tr>
<tr>
<td>(Chevalier-Larsen, O'Brien et al. 2004)</td>
<td>Tg/112/prion</td>
<td>Disease severity: Tg♂ &gt; Tg♀&lt;br&gt;Onset age: Tg♂ &lt; Tg♀ (by mos)</td>
<td>~6 wks</td>
<td>Clasping ↓ time on rotarod ↓ open field activity ↓ grip strength impaired gait</td>
<td>altered NF staining (no change in size or # of MNs);</td>
<td>none</td>
<td>No data</td>
<td>Castration of Tg♂ s reverses deficits in motor function &amp; MN NF staining</td>
</tr>
<tr>
<td>(Sopher, Thomas et al. 2004)</td>
<td>Tg/100/ human AR regulatory elements</td>
<td>Disease severity: Tg♂ &gt; Tg♀&lt;br&gt;Onset age: Tg♂ &lt; Tg♀ (by mos)</td>
<td>~6 mos</td>
<td>↓ time on hang test ↓ stride length impaired gait hindlimb weakness &amp; paralysis</td>
<td>MNs appear smaller, less densely stained and fewer in # ↓ expression of VEGF♂ in spinal cord</td>
<td>hindlimb muscle atrophy ↑ mf staining for NADH* &amp; NCAM* fiber type grouping</td>
<td>No data</td>
<td>No data</td>
</tr>
<tr>
<td>(Yu, Dadgar et al. 2006)</td>
<td>KI/113/mouse AR regulatory elements</td>
<td>Disease severity: Tg ♂ &gt; Tg ♀</td>
<td>~ 10 wks</td>
<td>↓ grip strength</td>
<td>No evidence of cell death based on glial staining Abnormal EMG recordings indicative of neurogenic and primary myopathic processes</td>
<td>↑ levels of AChRα and myogenin mRNA ↓ levels of GDNFk and NT-4l mRNA</td>
<td>No data</td>
<td>Castration of Tg ♂'s partially rescues KI males from muscle weakness and prevents early death</td>
</tr>
<tr>
<td>(Monks, Johansen et al. 2007)</td>
<td>Tg/22/human skeletal actin α</td>
<td>Only Tg ♂ show a motor phenotype</td>
<td>From earliest age examined (peripuberty)</td>
<td>↓ time on hang test ↓ stride length Impaired gait ↓ time on rotarod</td>
<td>↓ motor axons Normal # of MN cell bodies</td>
<td>↓ # of mfs Centralized nuclei split muscle fibers Fiber atrophy ↑ NADH staining ↑ levels of AChRα and myogenin mRNA ↓ level of VEGF mRNA</td>
<td>No data</td>
<td>Castration of Tg males reverses disease symptoms T treatment of asymptomatic Tg females induces disease</td>
</tr>
</tbody>
</table>

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a Tg = transgenic; KI = knock-in  
bMN = motoneuron cell bodies  
cmf = muscle fiber  
dT = testosterone  
eNF = neurofilament  
faltered NF staining = ↓ # of MNs that stain for unphosphorylated neurofilament- heavy chain  
gVEGF = vascular endothelial growth factor  
hNADH = nicotinamide adenine dinucleotide  
iNCAM = neural cell adhesion molecule  
jAChRα = acetylcholine receptor α  
kGDNF = glial-derived neurotrophic factor  
lNT-4 = neurotrophin-4