Beta-2 microglobulin is important for disease progression in a murine model for amyotrophic lateral sclerosis

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INTRODUCTION

Major histocompatibility complex (MHC) class I proteins were originally discovered based on their critical role in the immune system, however immune-independent functions in the nervous system have recently been identified (Huh et al., 2000; Elmer and McAllister, 2012). Beta-2 microglobulin (β2m) is an essential component of MHC class I molecules, being required for expression of all MHC class I on the cell surface. Within the central nervous system β2m has a predominantly motor neuronal expression pattern (Linda et al., 1998, 1999; Thams et al., 2009). This protein is therefore a candidate to contribute to the selective vulnerability of such motor neurons during amyotrophic lateral sclerosis (ALS).

ALS is a progressive neurodegenerative disease, characterized by the selective loss of motor neurons and the denervation of muscle fibers, resulting in muscle weakness and paralysis. In Europe, the disease has an annual incidence of 2.7 cases per 100,000 people (Logroscino et al., 2010) and the disease duration post-diagnosis is 3–5 years. In 10% of patients, ALS is a familial disease and 20% of these familial ALS patients contain mutations in the gene encoding superoxide dismutase 1 (SOD1). Based on these mutations, ALS rodent models have been generated that predictably mimic the patient disease process (Julien and Kriz, 2006). As the disease progression is indistinguishable between familial and sporadic ALS, common disease mechanisms are predicted. One of these mechanisms is decreased (peripheral) neuronal plasticity that can influence the ability of neuronal networks to compensate for a loss of (motor) neurons in the network. β2m is expressed in motor neurons in the lumbar spinal cord (Linda et al., 1999) as well as in motor axons (Thams et al., 2009). Additionally, β2m promotes recovery after axotomy (Linda et al., 1998; Oliveira et al., 2004) and sciatic nerve crush (Oliveira et al., 2004), which implies that it may be of importance in ALS too.

In this study, we investigated the role of β2m in ALS mice. To this end, we assessed the gene expression of β2m and interbred mice genetically lacking β2m with SOD1G93A mice and assessed survival and disease pathology.

MATERIALS AND METHODS

ANIMAL EXPERIMENTS

Mice overexpressing human wild-type SOD1 (SOD1WT) or human SOD1G93A and β2m knockout mice were purchased from The Jackson Laboratories (Bar Harbor, USA) and maintained on a C57BL/6 background. The SOD1G93A and β2m knockout were interbred allowing for approx. 50% of the mice to be littermate controlled in this study. Chow and water were provided ad libitum.
and mice were housed in the specific pathogen free animal facility of the KU Leuven under standard conditions according to the guidelines of the KU Leuven. End stage was defined as the age at which mice could no longer right themselves within 30 s when placed on their back. End stage is used as a measurement of survival and is the condition at which mice are euthanized to prevent further suffering. Disease onset was defined as the age at which mouse weight dropped below 90% of the average day 90–105 weight. The animal caretakers and scientists were blinded to the genotypes of the mice when assessing “end stage”. All animal experiments were performed with the approval of the Animal Ethical Committee of KU Leuven (020/2010).

**LASER DISSECTION MICROSCOPY**

Murine spinal cords were snap-frozen in Tissue-Tec (Sakura Finetek Europe, Alphen aan de Rijn, The Netherlands) to make cryostat sections of 20-μm thickness. Then, cresyl violet–stained motor neurons, located in the ventral horn of the lumbar spinal cord, were collected on membrane slides 1.0 PEN (Carl Zeiss AG, Oberkochen, Germany), using dissection by a laser-dissection microscope (Carl Zeiss AG) and capturing in Adhesive Cap 500 opaque (Carl Zeiss AG). Only motor neurons in which the nucleus was visible and with soma area > 250 μm², were collected. At least 1,500 motor neurons were dissected for each animal.

**QUANTITATIVE PCR**

Isolation of mRNA was performed using the TriPure (Roche, Basel, Switzerland) method and the RNaseasy kit (Qiagen, Venlo, The Netherlands). Reverse transcriptase polymerase chain reaction (PCR) used random hexamers (Life Technologies, Carlsbad, USA) and Moloney Murine Leukemia Virus Reverse Transcriptase (MMLV RT; Invitrogen, Carlsbad, USA). Quantitative PCR (qPCR) was performed with the StepOnePlus (Life Technologies) and TaqMan Universal PCR Master Mix (Life Technologies). Gene expression assays were purchased from Life Technologies (Carlsbad, USA) and Moloney Murine Leukemia Virus Reverse Transcription (MMLV RT; Invitrogen, Carlsbad, USA). Quantitative PCR (qPCR) was performed with the StepOnePlus (Life Technologies) and TaqMan Universal PCR Master Mix (Life Technologies). Gene expression assays were purchased from Life Technologies and IDT DNA (Coraldale, USA): gapdh (Mm.PT39a.1), β2m (Mm00437762_m1) and cd8b1 (Mm.PT49a.10182911). For this analysis, presymptomatic tissue was collected at 90 days of age and symptomatic at 120 days of age. The scientist performing the qPCR was blinded to the genotypes of the samples.

**NISSL STAINING**

To visualize neurons, Nissl staining was performed on 4% formaldehyde fixed spinal cord sections. Sections were briefly immersed in a cresyl violet solution and subsequently in a 10% acetic acid. Slides were dehydrated by an increased ethanol concentration series and mounted with PerTex® (Histolab AB, Göteborg, Sweden). Images were collected by Zeiss Axio Imager M1 microscope (Carl Zeiss AG) with AxioCam Mrc5 camera (Carl Zeiss AG). The number of (motor) neurons was quantified by measurement of the soma area as visualized by cresyl violet staining in ImageJ (National Institute of Health) on multiple 40 μm thick sections in the ventral horn of the lumbar spinal cord. Characterisation of motor neurons occurred as previously (Fischer et al., 2004) of at least 5–10 ventral horns of the lumbar spinal cord of 2–4 mice per group. The scientist performing the Nissl staining and neuron quantification was blinded to the genotypes of the samples.

**IMMUNOHISTOCHEMISTRY**

Mice were transcardially perfused with phosphate buffered saline (PBS) and subsequently with 4% formaldehyde. Spinal cords were post-fixed with 4% formaldehyde overnight at 4°C and transferred to 30% sucrose for an additional night. After snap freezing, tissue was sectioned by cryostat at 40 μm thickness and stained with a polyclonal antibody directed against ubiquitin (Dako, Glostrup, Denmark). Images were collected by Zeiss Axio Imager M1 microscope (Carl Zeiss AG) with AxioCam Mrc5 camera (Carl Zeiss AG). Ubiquitin immunopositive aggregates were counted per ventral horn using ImageJ of 2–6 ventral horns of the lumbar spinal cord of 2–4 mice per group and presented as the average of the number of aggregates per ventral horn. The scientist performing the immunohistochemistry and aggregate quantification was blinded to the genotypes of the samples.

**STATISTICAL ANALYSIS**

Analysis was performed with the statistical software package Prism Origin (GraphPad Software, La Jolla, USA). Survival was analyzed by Log-Rank testing. Differences between two groups were analyzed using a Student’s t-test. Differences between more than two groups were analyzed by ANOVA with Bonferroni correction for multiple testing. Significance was assumed at p < 0.05. Error bars represent the standard deviation.

**RESULTS**

To assess the potential for β2m to have a functional role in ALS pathogenesis, we assessed gene expression in the spinal cords of non-transgenic, SOD1WT and SOD1G93A mice. We observed a strong increase in β2m gene expression during disease progression in SOD1G93A mice (Figure 1A). This increase is at least partly due to the increased neuron-specific gene expression of β2m, as a greater level of upregulation (10-fold) was identified by qPCR on neurons from SOD1G93A mice compared to neurons from SOD1WT mice isolated by laser dissection microscopy (Figure 1B). As we do not observe an increase of CDA+ T cells in the spinal cord of end stage SOD1G93A mice, as assessed by the gene expression analysis of CD8b1 (Figure 1C), these data indicate potential for a neuronal role for β2m in ALS.

To determine whether β2m has an effect in ALS, we interbred β2m−/− mice with SOD1G93A mice and assessed disease progression in β2m−/− SOD1G93A, β2m+/− SOD1G93A and β2m+/+ SOD1G93A mice. The survival of β2m+/− SOD1G93A mice did not differ from β2m+/+ SOD1G93A littermates (data not shown). The complete genetic ablation of β2m did not affect onset of disease (data not shown), but significantly decreased average survival of SOD1G93A mice by 8.9 days (Figure 2A) and reduced disease duration by approximately 50% (Figure 2B). The decrease of survival in β2m−/− SOD1G93A mice demonstrated a protective role for β2m in ALS mice.

To assess whether genetic ablation of β2m alters pathology of SOD1G93A mice, we analyzed pathology in the spinal cords of end stage mice. Decreased numbers of motor neurons were
observed in the end stage spinal cord of SOD1GAS and β2m−/− SOD1GAS mice (Figures 3A–C, quantified in Figure 3D), as were increased ubiquitin-positive aggregates (Figures 3E–G, quantified in Figure 3H). No differences were observed between the end stage pathology of SOD1GAS and β2m−/− SOD1GAS mice for motor neurons (Figures 3B, C) or ubiquitin immunoreactivity (Figures 3F, G). This shows that end stage β2m−/− SOD1GAS mice show the same extent of motor neuron loss and aggregate formation as end stage SOD1GAS mice, although disease progression is faster.

**DISCUSSION**

Here we show that β2m is important in ALS mouse survival and that it is upregulated during disease in the spinal cord and by motor neurons. Upregulation of β2m in neuronal tissues has been reported previously when comparing spinal cord (Edstrom et al., 2004) and brain (VanGuilder Starkey et al., 2012) expression of aged rats to adult controls and in the spinal cord of axotomised rats (Maehlen et al., 1988; Olsson et al., 1989; Linda et al., 1998), which may suggest that stressed neurons increase β2m gene expression to increase plasticity. This concept fits well with reports of the role of β2m in neurons during development and plasticity (Huh et al., 2000; Bilousova et al., 2012), of the hippocampus and visual system (Huh et al., 2000) but not of the cerebellum (Letellier et al., 2008), and the delayed or impaired recovery of β2m knockout mouse post axotomy (Linda et al., 1998; Oliveira et al., 2004) and sciatic nerve crush (Oliveira et al., 2004).

Impaired (peripheral) plasticity by β2m knockout may explain the decrease in survival detected in ALS mice in this study, as increased plasticity is protective in ALS mice and rats (Van Hoecke et al., 2012). A number of plasticity-promoting genetic or pharmacological strategies have proven successful in the past in ALS models, such as EphA4 knockdown and inhibition (Van Hoecke et al., 2012), and vascular endothelial growth factor (VEGF) administration in ALS rodents (Storkebaum et al., 2005).

With the use of a ubiquitous β2m knockout mouse we cannot exclude that the decrease of survival of ALS mice lacking β2m may be due to the effect of removing β2m in the immune system. β2m is necessary for the differentiation of CD8+ T cells and natural killer T (NKT) cells (Koller et al., 1990). The role of these cell types is not yet fully understood in ALS, as varying results are obtained for ALS mouse survival when mature lymphocytes are not present (Beers et al., 2008; Tada et al., 2011). Additionally, NKT cells may be associated to ALS disease pathology as impairments in NKT cells are reported in ALS mice (Finkielstein et al., 2011). That being said, qPCR analysis of CD8+ T cells does not suggest a role for
these cells as CD8b1 gene expression is not increased in ALS spinal cords. Additionally, a role for CD8+ T cells may be predicted to be detrimental in contrast to our data demonstrating a detrimental role for β2m upon removal in ALS mice. Interestingly, β2m has been assessed previously as a biomarker in cerebrospinal fluid or venous blood samples from ALS patients, but with variable results (Brettschneider et al., 2008; Mitchell et al., 2009; Baciu et al., 2012).

The mechanism to which β2m contributes to neuronal plasticity is not fully understood, though it is proposed that it may be through paired-immunoglobulin like receptor-B (PirB) (VanGuilder Starkey et al., 2012). This receptor is located on axons, dendrites and neuronal somata (VanGuilder Starkey et al., 2012) and could thus easily facilitate plasticity. Additionally, β2m is localized at synapses and in synaptosomes (Shatz, 2009). Alternatively, β2m and PirB are associated with decreased plasticity and recovery in other neurodegenerative conditions such as stroke (Adelson et al., 2012), experimental autoimmune encephalomyelitis (EAE; Denic et al., 2012) and ischemia (Wang et al., 2012). These paradoxes are largely affected by the immune system and the beneficial effect of β2m or PirB in these models may be due to the role of the immune system. This notion is supported by work by Linker et al. that show that reconstitution of CD8+ cells in β2m knockout mice delays the effect of EAE compared to β2m knockout littermates (Linker et al., 2005).

In conclusion, this work shows the detrimental effect of β2m knockout in ALS and identifies β2m signaling as a potential new direction for the development of therapeutic strategies counteracting ALS.

**AUTHOR CONTRIBUTIONS**

Kim A. Staats and Susann Schönefeldt performed the murine behavioral analyses. Kim A. Staats and Marike Van Rillaer conducted the staining experiments. Kim A. Staats and Annelies Van Hoecke analyzed gene expression of motor neurons excised by laser dissection microscopy. Philip Van Damme, Wim Robberecht, Adrian Liston and Ludo Van Den Bosch supervised and designed the experiments. Kim A. Staats, Adrian Liston and Ludo Van Den Bosch wrote the manuscript. All authors approved the final version of the manuscript.

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**REFERENCES**

Adelson, J. D., Barreto, G. E., Xu, L., Kim, T., Brott, B. K., Ouyang, Y. B., et al. (2012). Neuroprotection from stroke in the absence of MHCI or PirB. Neuron 73, 1100–1107. doi: 10.1016/j.neuron.2012.01.020

Baciu, C., Thompson, K. J., Maugeot, J. L., Brooks, B. R., and Weller, J. W. (2012). The LO-BaFL method and ALS microarray expression analysis. BMC Bioinformatics 13:244. doi: 10.1186/1471-2105-13-244

Beers, D. R., Henkel, J. S., Zhao, W., Wang, J., and Appel, S. H. (2008). CD4+ T cells support glial neuroprotection, slow disease progression and modify glial morphology in an animal model of inherited ALS. Proc. Natl. Acad. Sci. U S A 105, 15558–15563. doi: 10.1073/pnas.0807419105

Bilousova, T., Dang, H., Xu, W., Gustafson, S., Jin, Y., Wickramasinghe, L., et al. (2012). Major histocompatibility complex I molecules modulate embryonic neuritogenesis and neuronal polarization. J. Neuroimmunol. 247, 1–8. doi: 10.1016/j.neuroimm.2012.03.008

Brettschneider, J., Mogel, H., Lehnensiek, V., Ahlert, T., Sussmuth, S., Ludolph, A. C., et al. (2008). Proteome analysis of cerebrospinal fluid in amyotrophic
lateral sclerosis (ALS). *Neurochem. Res.* 33, 2358–2365. doi: 10.1007/s11064-008-9742-5

Denic, A., Pirko, I., Wootla, B., Bieber, A., Macura, S., and Rodriguez, M. (2012). Deletion of beta-2-microglobulin ameliorates spinal cord lesion load and promotes recovery of brainstem NAA levels in a murine model of multiple sclerosis. *Brain Pathol.* 22, 698–708. doi: 10.1111/j.1750-3639.2012.00576.x

Edstrom, E., Kullberg, S., Ming, Y., Zheng, H., and Ulbhake, B. (2004). MHC class I, beta2 microglobulin and the INF-gamma receptor are upregulated in aged motoneurons. *J. Neurosci.* 78, 892–900. doi: 10.1002/jnr.20341

Elmer, B. M., and Mcallister, A. K. (2012). Major histocompatibility complex class I proteins in brain development and plasticity. *Trends Neurosci.* 35, 660–670. doi: 10.1016/j.tins.2012.08.001

Finkelstein, A., Kunis, G., Senesyan, A., Ronen, A., Berkutzi, T., Azoulay, D., et al. (2011). Abnormal changes in NKT cells, the IGFr-1 axis and liver pathology in an animal model of ALS. *PLoS One* 6(2):e22374. doi: 10.1371/journal.pone.0022374

Fischer, L. R., Culver, D. G., Tennen, P., Davis, A. A., Wang, M., Castellano-Sanchez, A., et al. (2004). Amyotrophic lateral sclerosis is a distal axonopathy: evidence in mice and man. *Exp. Neurol.* 185, 232–240. doi: 10.1016/j.expneurol.2003.10.004

Huh, G. S., Boulanger, L. M., Du, H., Riquelme, P. A., Brotz, T. M., and Shatz, C. J. (2000). Functional requirement for class I MHC in CNS development and plasticity. *Science* 290, 2155–2159. doi: 10.1126/science.290.5499.2155

Julien, J. P., and Kriz, J. (2006). Transgenic mouse models of amyotrophic lateral sclerosis. *Biochim. Biophys. Acta* 1762, 1013–1024. doi: 10.1016/j.bbadis.2006.03.006

Koller, B. H., Marrack, P., Kappler, J. W., and Smithies, O. (1990). Normal development of mice deficient in beta 2M, MHC class I proteins and CD8+ T cells. *Science* 248, 1227–1230. doi: 10.1126/science.2112266

Letellier, M., Willson, M. L., Gautheron, V., Mariani, J., and Lohof, A. M. (2008). Normal adult climbing fiber monoinnervation of cerebellar Purkinje cells in mice lacking MHC class I molecules. *Dev. Neurobiol.* 68, 997–1006. doi: 10.1002/dneu.20639

Linda, H., Hammarberg, H., Cullheim, S., Levinovitz, A., Khademi, M., and Olsson, T. (1998). Expression of MHC class I and beta2-microglobulin in rat spinal motoneurons: regulatory influences by IFN-gamma and axotomy. *Exp. Neurol.* 150, 282–295. doi: 10.1006/exnr.1997.6768

Linda, H., Hammarberg, H., Piehl, F., Khademi, M., and Olsson, T. (1999). Expression of MHC class I heavy chain and beta2-microglobulin in rat brainstem motoneurons and nigral dopaminergic neurons. *J. Neuroimmunol.* 101, 76–86. doi: 10.1016/s0165-5728(99)00135-6

Linker, R. A., Rott, E., Hofstetter, H. H., Hanke, T., Toyka, K. V., and Gold, R. (2005). EAE in beta2-microglobulin-deficient mice: axonal damage is not dependent on MHC-I restricted immune responses. *Neurobiol. Dis.* 19, 218–228. doi: 10.1016/j.nbd.2004.12.017

Logroscino, G., Traynor, B. J., Hardiman, O., Chio, A., Mitchell, D., Swingler, R. J., et al. (2010). Incidence of amyotrophic lateral sclerosis in Europe. *J. Neurol.* 257(Suppl. 1), 1–18. doi: 10.1007/s00415-010-0334-2

Mitchell, R. M., Freeman, W. M., Randazzo, W. T., Stephens, H. E., Beard, J. L., Simmons, Z., et al. (2009). A CSF biomarker panel for identification of patients with amyotrophic lateral sclerosis. *Neurology* 72, 14–19. doi: 10.1212/WNL.0b013e31819c50b9

Oliveira, A. L., Thams, S., Lidman, O., Piehl, F., Hokfelt, T., Karre, K., et al. (2004). A role for MHC class I molecules in synaptic plasticity and regeneration of neurons after axotomy. *Proc. Natl. Acad. Sci. U S A* 101, 17843–17848. doi: 10.1073/pnas.0408154101

Olsson, T., Kristensson, K., Ljungdahl, A., Maehlen, J., Holmdahl, R., and Klarskog, L. (1989). Gamma-interferon-like immunoreactivity in axotomized rat motor neurons. *J. Neurosci.* 9, 3870–3875.

Shatz, C. J. (2009). MHC class I: an unexpected role in neuronal plasticity. *Neuron* 64, 40–45. doi: 10.1016/j.neuron.2009.09.044

Simmons, Z., et al. (2009). A CSF biomarker panel for identification of patients with amyotrophic lateral sclerosis. *J. Neurol. Neurosurg. Psychiatry* 81, 385–390. doi: 10.1136/jnnp.2008.183525

Mehlen, J., Schroder, H. D., Klarskog, L., Olsson, T., and Kristensson, K. (1988). Axotomy induces MHC class I antigen expression on rat nerve cells. *Neurosci. Lett.* 92, 8–13. doi: 10.1016/0304-3940(88)90733-1

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