Brca1 is expressed in human microglia and is dysregulated in human and animal model of ALS
Harun Najib Noristani, Jean Charles Sabourin, Yannick Nicolas Gerber, Marisa Teigell, Andreas Sommacal, Maria Dm Vivanco, Markus Weber, Florence Evelyne Perrin

To cite this version:
Harun Najib Noristani, Jean Charles Sabourin, Yannick Nicolas Gerber, Marisa Teigell, Andreas Sommacal, et al.. Brca1 is expressed in human microglia and is dysregulated in human and animal model of ALS. Molecular Neurodegeneration, BioMed Central, 2015, 10 (1), 10.1186/s13024-015-0023-x. hal-02156103

HAL Id: hal-02156103
https://hal.archives-ouvertes.fr/hal-02156103
Submitted on 7 Jun 2021

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L’archive ouverte pluridisciplinaire HAL, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d’enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Distributed under a Creative Commons Attribution 4.0 International License
Brca1 is expressed in human microglia and is dysregulated in human and animal model of ALS

Harun Najib Noristani1, Jean Charles Sabourin2†, Yannick Nicolas Gerber1,2, Marisa Teigell1, Andreas Sommacal3, Maria dM Vivanco4, Markus Weber3 and Florence Evelyne Perrin1,2,5*

Abstract

Background: There is growing evidence that microglia are key players in the pathological process of amyotrophic lateral sclerosis (ALS). It is suggested that microglia have a dual role in motoneurone degeneration through the release of both neuroprotective and neurotoxic factors.

Results: To identify candidate genes that may be involved in ALS pathology we have analysed at early symptomatic age (P90), the molecular signature of microglia from the lumbar region of the spinal cord of hSOD1G93A mice, the most widely used animal model of ALS. We first identified unique hSOD1G93A microglia transcriptomic profile that, in addition to more classical processes such as chemotaxis and immune response, pointed toward the potential involvement of the tumour suppressor gene breast cancer susceptibility gene 1 (Brca1). Secondly, comparison with our previous data on hSOD1G93A motoneurone gene profile substantiated the putative contribution of Brca1 in ALS. Finally, we established that Brca1 protein is specifically expressed in human spinal microglia and is up-regulated in ALS patients.

Conclusions: Overall, our data provide new insights into the pathogenic concept of a non-cell-autonomous disease and the involvement of microglia in ALS. Importantly, the identification of Brca1 as a novel microglial marker and as possible contributor in both human and animal model of ALS may represent a valid therapeutic target. Moreover, our data points toward novel research strategies such as investigating the role of oncogenic proteins in neurodegenerative diseases.

Keywords: Microglia, Transcriptomics, hSOD1G93A mice, ALS patients, Brca1

Background

Amyotrophic lateral sclerosis (ALS) is characterised by selective motoneurones degeneration in the spinal cord, brainstem and motor cortex leading to progressive muscle weakness, atrophy and paralysis. Approximately 90 % of ALS patients are sporadic whilst 10 % are familial cases with genetic mutations in SOD1 (Cu/Zn superoxide dismutase 1), FUS (fused in sarcoma), TARDBP (also known as TDP-43) and C9ORF72, among others [1]. Transgenic mice over-expressing the human mutated gene for SOD1 develop an adult-onset paralysis that closely recapitulates human ALS [2]. Recent studies have established that ALS is a complex multi-factorial disease that involves several cellular partners including glial cells [3].

Microglia, the resident immune cells of the central nervous system (CNS), when activated, release pro-and anti-inflammatory cytokines and chemokines that are generally associated with M1 and M2 phenotypes [4, 5]. Microglia have a dual role in ALS with an early protective effect on motoneurones but also a detrimental effect due to the secretion of neurotoxic factors [6]. It is hypothesised that progressive motoneurone death results from the combination of intrinsic motoneurones vulnerability and toxicity from neighbouring cells such as microglia [6]. In ALS patients and animal models, there is a clear microglia activation [3], in particular we have shown an early involvement of microglia in hSOD1G93A mice [7]. Understanding the contribution of microglia to motoneurone degeneration is of high priority. One means of analysing the role of a cell population in a process network is to study gene expression alterations in this given population. In
addition, an integrative comparison of the specific molecular signatures of several cellular partners is necessary to decipher the crosstalk between these cells. We have previously identified gene dysregulation in pure motoneurones from the lumbar spinal cord of hSOD1<sup>G93A</sup> mice [8] and two other mouse models of motoneurone disease [9]. We revealed a unique motoneurone gene expression profile characterised by an absence of dysregulation of genes associated with cell death and a massive up-regulation of genes involved in cell growth [8].

Growing evidence points toward mitochondrial dysfunction and oxidative damage in ALS [10]. Defence mechanisms, including SOD, counteract excessive accumulation of reactive oxygen species, however in ALS, cellular antioxidant defences are insufficient leading to damage of nucleic acids, proteins and lipids [11]. Inherited mutations in breast cancer susceptibility gene 1 (Brca1), a well-known tumour suppressor implicated in familial breast and ovarian cancers, is one of the best defined risk factor for development of breast and ovarian cancer. Brca1 plays important roles in a broad spectrum of functions including transcription regulation, cell cycle checkpoint activation, apoptosis, chromosomal remodelling, ubiquitination and DNA repair [12]. The role of Brca1 in each of these processes remains to be fully understood but it is hypothesized that it act as a scaffold for the formation of complexes with a wide range of proteins [13]. This ability of Brca1 to interact with different proteins may underlie its involvement in a variety of cellular processes [13]. Brca1 also exerts a protective role against oxidative stress via up-regulation of antioxidant genes and maintenance of the redox balance through up-regulating the expression of heat shock protein HSP27 [14, 15].

In breast cancer, Brca1 cellular localisation as well as the significance of its altered localisation, is still a matter of debate. It had been recently shown that in normal breast, Brca1 nuclear expression is strong and uniform in parenchymal cells whereas in malignant cells its expression is reduced if not absent from the nucleus and is, in some cases, observed in the cytoplasm [16]. Interestingly, altered expression of Brca1 was associated with poor prognosis and shortened survival. In the adult rodent CNS, the presence of Brca1 is detected only in neurons [17] whereas a high Brca1 expression is observed in embryonic [17, 18] and adult neural stem cells and is involved in cell proliferation [18].

Here we identify putative Brca1 involvement in ALS via hSOD1<sup>G93A</sup> microglia gene profiling and comparisons to our previous transcriptomic findings in hSOD1<sup>G93A</sup> motoneurones. We then demonstrated that Brca1 is a novel marker of human microglia and is up-regulated in ALS patients.

**Results**

**Transcriptomic analysis of FACS isolated microglia from control and hSOD1<sup>G93A</sup> lumbar spinal cord**

We have previously described early microglial disturbances in hSOD1<sup>G93A</sup> male mice reflected at P90 by a heterogeneous Iba1<sup>+</sup> microglial distribution with higher density within the grey matter in hSOD1<sup>G93A</sup> mice as compared to control [7, 19]. Since activated microglia/macrophages exhibit increased CD11b expression, we carried out CD11b immunostaining (Fig. 1a & b). CD11b-positive microglia displayed enlarged somata with short and thick processes that are typical of a reactive phenotype and were predominantly found in hSOD1<sup>G93A</sup> mice (Fig. 1b). To further analyse transcriptomic modification specifically in microglia, we isolated microglia of hSOD1<sup>G93A</sup> and control littermate males at early symptomatic age (P90) from the lumbar spinal cord (L1-L5) that corresponds to the onset of degeneration. Microglia were isolated by fluorescence-activated cell sorting (FACS) using CD11b (Fig. 1c–e). We observed a 1.65-fold increase in the total number of CD11b<sup>+</sup> microglia in hSOD1<sup>G93A</sup> versus controls (26 350; n = 15 in hSOD1<sup>G93A</sup> and 15 900; n = 26 in control; Fig. 1c & d). RNA extracted from FACS purified microglia was of high quality (Fig. 1f) and microarrays analysis revealed 630 dysregulated genes (260 down-regulated and 370 up-regulated, Additional file 1: Table S1).

**Cross-talk between microglia and motoneurones**

We had previously identified dysregulated genes in hSOD1<sup>G93A</sup> motoneurone during disease progression [8, 9]. To unravel potential molecular cross-talk between microglia and motoneurones, we performed a comparative analysis of gene dysregulation in both cell populations. Comparison of dysregulated genes at P90 between motoneurones (102 genes) and microglia (668 genes) revealed 19 common genes (Additional file 2: Figure S1A). Process network rankings were clearly different in the commonly dysregulated genes (in motoneurones and microglia) and uniquely dysregulated genes. Antigen presentation was classified first in the common group, whilst cytoskeleton and cytoplasmic microtubules genes were top ranked in motoneurones only set (Additional file 3: Table S2A). Similarly, cellular processes analysis ranked first immune response and antigen presentation in the common group whereas response to stress, regulation of immune response, system development and wounding response were the top 4 ranked processes in motoneurones only group (Additional file 3: Table S2B). Signalling and metabolic pathway analysis revealed immune response and cytokine remodelling as first ranked in the commonly and motoneurone
only dysregulated genes, respectively (Additional file 3: Table S2C).

We have previously shown that microglial reactivity precedes neuronal death in hSOD1\(^{G93A}\) mice [7]; to seek for potential modifications in microglia that could trigger motoneurone death, we compared dysregulated genes at P90 in microglia and P120 in motoneurones. Our previous microarrays analysis of microdissected motoneurones at the end stage of the disease (P120) showed no dysregulation of genes associated with cell death [8], this most likely reflects that dissected motoneurones were at an early demise stage. Indeed, we selected motoneurones that had an identifiable nucleus and a diameter of at least 25 \(\mu\)m, picking a sub-population of neurones that may resist degeneration. Three hundred twenty genes were uniquely dysregulated in hSOD1\(^{G93A}\) motoneurones; 603 uniquely dysregulated in hSOD1\(^{G93A}\) microglia; 65 genes were

![Fig. 1](image-url)
common (Additional file 2: Figure S1B). Clear differences were highlighted not only between the genes that were commonly and uniquely dysregulated but also in the ranking as compared to the previous analysis (microglia and motoneurones at P90, Additional file 4: Table S3). Particularly, inflammation and immune response were ranked top in motoneurones (Additional file 4: Table S3A). Interestingly, signalling and metabolic pathway analysis revealed the involvement of heme metabolism and DNA damage in both motoneurones and microglia (Additional file 4: Table S3C).

**Unique transcriptomic profiles of hSOD1<sup>G93A</sup> microglia**

To identify processes and pathways modified in hSOD1<sup>G93A</sup> microglia, we carried out gene ontology enrichment and network analysis (Additional file 5: Table S4A–C; Fig. 2). Process network analysis ranked as first chemotaxis (Additional file 5: Table S4A, Fig. 2a) with 23 dysregulated transcripts out of 137 annotated genes in this process (17 %, \( p = 2.18 \times 10^{-4} \)) (Additional file 5: Table S4A). Out of the 18 most significantly dysregulated genes, 4 were down-regulated with a maximum of 2-fold whereas 14 were up-regulated (Fig. 2a). The gene coding for osteopontin (SPPI) presents a 16.8-fold increase (Additional file 1: Table S1). Regulation of angiogenesis (ranked 8th, Additional file 5: Table S4A, Fig. 2e) displayed 22 dysregulated genes (223 genes in this process, 9.8 %, \( p = 7.3 \times 10^{-4} \)), with 16 being up-regulated. Inflammation network was also dysregulated in hSOD1<sup>G93A</sup> microglia (ranked 9<sup>th</sup>, 9 % of the annotated genes in this process, \( p = 2.10 \times 10^{-8} \)) (Additional file 5: Table S4A). Out of the 18 most significantly dysregulated genes, 4 were down-regulated with a maximum of 2-fold whereas 14 were up-regulated (Fig. 2a). The gene coding for osteopontin (SPPI) presents a 16.8-fold increase (Additional file 1: Table S1). Regulation of angiogenesis (ranked 8th, Additional file 5: Table S4A, Fig. 2e) displayed 22 dysregulated genes (223 genes in this process, 9.8 %, \( p = 7.3 \times 10^{-4} \)), with 16 being up-regulated. Inflammation network was also dysregulated in hSOD1<sup>G93A</sup> microglia (ranked 9<sup>th</sup>, 9 % of the annotated genes in this process, \( p = 2.10 \times 10^{-8} \)) (Additional file 5: Table S4A).

Amongst the 29 most significantly dysregulated genes 8 were down-regulated with a maximum of 2.97-fold decrease for the gene coding for alpha-synuclein, whereas 33 were up-regulated. Genes coding for CCL5 (5.1-fold change (FC)) and CXCL13 (5.7-FC) were the most up-regulated (Additional file 1: Table S1). Regulation of blood coagulation was ranked 3rd (50/665 genes, 7.5 %, \( p = 1.8 \times 10^{-9} \), Additional file 5: Table S4B, Fig. 2f). Amongst the 29 most significantly dysregulated genes 8 were down-regulated and 21 were up-regulated. Hypoxia was the 4<sup>th</sup> dysregulated cellular processes (31/416 genes, 7.45 %, \( p = 8.3 \times 10^{-9} \), Additional file 5: Table S4B). Out of the 20 most significantly dysregulated genes 4 were down-regulated and 16 were up-regulated (Fig. 2c).

**Breast cancer 1 (Brcal) pathway is dysregulated in hSOD1<sup>G93A</sup> mice**

Pathways map analysis ranked as the third position DNA damage and specifically the involvement of Brcal as a transcription regulator (Additional file 5: Table S4C, Figs. 2g and 3). Indeed, in hSOD1<sup>G93A</sup> microglia (7/30 genes, 23 %; \( p = 1.6 \times 10^{-5} \), Additional file 5: Table S4C) were dysregulated in the canonical Brca1 pathway. GADD45a and SP3 transcription factor were down-regulated with FC of 2.6 and 1.9, respectively. Genes coding for p21 (1.8-FC), PCNA (1.85-FC), STAT1 (1.9-FC), c-Myc (1.8-FC) and Brca1 (1.76-FC) were up-regulated (Additional file 1: Table S1 and Fig. 2g). Concomitant dysregulation of these genes clearly pointed toward a potential involvement of Brca1 as a transcription regulator (Fig. 2g and red and blue thermometers labelled in Fig. 3). Interestingly, even if Brcal transcript itself was not dysregulated in motoneurones, 4 genes that are involved in Brca1 pathway were also up-regulated in hSOD1<sup>G93A</sup> motoneurones namely p21: 7.88-FC; GADD45a: 5.19-FC; Rb protein: 2.44-FC and ATF-1: 2.38-FC, (Fig. 2h and red thermometers labelled 2 in Fig. 3). To confirm microarray findings, we carried out quantitative real-time polymerase chain reaction (qPCR) in pure populations of hSOD1<sup>G93A</sup> and wild type microglia and assessed the expression profiles of all candidate genes involved in Brca1 pathway (Additional file 6: Figure S2). In addition, we have also included microglial samples at 60 days of age to assess the potential involvement of microglial Brca1 at the initial stages of the disease progression in hSOD1<sup>G93A</sup> mice (Additional file 6: Figure S2A). Our qPCR results showed no significant dysregulation of the genes involved in Brca1 pathway at 60 days of age (Additional file 6: Figure S2A). However, at 90 days of age, and similarly to our microarrays results, we found up-regulation of Brca1, Cdkn1a, Myc, Pena and Stat1 as well as down-regulation of Gadd45a and Sp3 in hSOD1<sup>G93A</sup> microglia (Additional file 6: Figure S2B). It is important to note that dysregulation in Cdkn1a, Myc, Pena, Stat1, Gadd45a and Sp3 transcripts may also be involved in other signalling pathways. These findings confirm Brca1 involvement in hSOD1<sup>G93A</sup> microglia is specifically triggered at 90 days of age when the pronounced microgliosis becomes evident.

**Brcal protein is expressed in human microglia and is up-regulated in ALS patients**

To investigate Brcal protein expression in human microglia, we performed dual immunofluorescence labelling using Brcal and CD11b antibodies (Fig. 4). Brcal staining in human control samples revealed ramified microglial population throughout the spinal cord displaying small cell bodies with long and thin processes (Fig. 4a & d) that co-localised with CD11b-positive microglia (Fig. 4b & e, c & f). Similarly, single immunoperoxidase detection of Brcal revealed microglial profile that were identical to Iba1 (the most commonly used
Fig. 2 Enrichment and network analysis of dysregulated gene in pure hSOD1^{G93A} microglia at 90 days of age. Categories networks and gene ontology processes that are most significantly modified in hSOD1^{G93A} microglia at 90 days as compared to control microglia. In each category, the most dysregulated genes are presented.
microglia marker) using adjacent human spinal cord sections (Fig. 5). Ramified microglia were evident in control cases following both Brca1 and Iba1 immunoperoxidase labelling (Fig. 5a–d). On the other hand, Brca1-positive microglia displayed enlarged cell bodies with short/thick processes in ALS cases similar to Iba1 immunostaining (Fig. 5e–h).

Finally, to determine Brca1 dysregulation in ALS, we quantified Brca1 immunoreactivity between the control and ALS spinal cords (Fig. 6). Brca1 expression was more evident in ALS compared to control cases (Fig. 6a & b, c & d). Quantitative analysis revealed a significant 78.2 % increase in Brca1 intensity within the white matter in ALS samples compared to controls (26.4 vs 47.1, \( p = 0.015 \); Fig. 6e). Within the grey matter, we observed a 32.8 % increase in Brca1 intensity in ALS samples compared to controls, however data variations in control samples kept them from attaining statistical significance (49.7 vs 66, \( p = 0.0545 \); Fig. 6f).

**Discussion**

Non-cell autonomous toxicity plays a major role in ALS [20] but microglia participation is dual and complex. Microglia reactivity over the course of the disease may be characterised by a continuum of activation states from a M2 neuroprotective state to a deleterious M1 state. In culture, microglia have, at disease onset, a M2 phenotype whereas they are typified by a M1 phenotype.
at disease end-stage [21]. Comparison of our data to previous studies [21, 22] reveals an up-regulation of five M2 priming genes (Clec7a, Igf1, Mmp12, Spp1 and Lgals3) and a down-regulation of Retnla and F13a1. Interestingly, four M1 priming genes are up-regulated CD86, Tnfa, Bcl2a1 and Cxcl10, whilst growth arrest and DNA-damage-inducible alpha gene (Gadd45gip1) is down-regulated. A previous study has reported the gene-expression profile of isolated microglia in hSOD1G93A mice and shown that potentially neuroprotective and neurotoxic factors are induced concurrently during disease progression [23]. The authors have analysed microglia from the entire spinal cord whereas we have restricted our investigations to the lumbar segment where onset of degeneration occurs. We report that hSOD1G93A microglia from the lumbar region of the spinal cord overexpress progranulin, Igf1 and osteopontin, all potential neurotrophic factors, and thus confirmed findings from a previous study [23]. Interestingly, we had previously identified in pure motoneurones of two mouse models of motoneurone disease (hSOD1G93A and pmm) an increase in IGFBP. Also, an IGFBP that binds to IGF-1 and IGF-2 (nephroblastoma over-expressed gene) was up-regulated at all disease stages in hSOD1G93A mice [8] and IGFBP4 mRNA was induced at pre-symptomatic age in pmm mice [9]. We also confirmed the dysregulation of the genes coding for IL-1β, IL-α, IL-10, Ifnar 1 and Ifnar 2 as well as Nox2 at P90. It had been shown that delayed forelimb motor impairment in ALS mice may be partially explained by augmented protective responses in the cervical spinal cords [24], thus gene expression profile of lumbar hSOD1G93A microglia is potentially more homogenous and is more likely to reflect a pathological gene profile than microglia taken from the entire spinal cord. Together, these data confirm that microglia activation states are best characterised as a continuum of M2 and M1 states [21] with a M2 phenotype at early stage of the disease that evolves into a M1 phenotype at disease end-stage.

An unexpected finding was the up-regulation in hSOD1G93A microglia of Brca1 with a 1.76 fold. Using in silico comparison with data from Chiu et al. [23], we found that Brca1 was also deregulated in their study and presented a steady increased with 2.78 and 3.08 fold changes at P100 and P130, respectively. In our study, Brca1 involvement was substantiated by the concomitant dysregulation of a number of other genes. As previously stated, Igf1 was robustly up-regulated in hSOD1G93A microglia; a complex interplay between Brca1 and IGF signalling pathways had been reported in familial cancer, in particular through the convergence of Brca1-mediated tumour protective pathways and IGFl receptors-mediated cell survival [25, 26]. This simultaneous up-regulation may represent a potential neuroprotective phenotype of microglia in ALS.
Fig. 5 Microglial Iba1 and Brca1 expression within adjacent human control and ALS spinal cord. Brightfield micrographs indicating Iba1 and Brca1 expression within adjacent sections of control (a–d) and ALS (e–h) spinal cords. Both Iba1 and Brca1-labelled microglia displayed typical ramified morphology in the control spinal cords with small cell bodies and large and thin processes (c & d). Microglial morphology displayed similar features following Iba1 (a & c) and Brca1 (b & d) immuno staining. In ALS spinal cords, Iba1 and Brca1-labelled microglia displayed both ramified and activated microglia with enlarged cell bodies and short and thick processes (g & h). Microglial morphology were similar using Iba1 (e & g) or Brca1 (f & h) immuno staining. Scale bars (a–h): 50 μm.
at early stage. Converging elements toward the involvement of Brca1 was also pointed through the dysregulation of genes linked to Brca1 and belonging to the DNA damage pathway (Fig. 3). Indeed, GADD45 was down-regulated in hSOD1G93A microglia (P90) and up-regulated in motoneurones (P120) and it had been demonstrated that Brca1 can modulate GADD45 that in turn mediates DNA repair mechanisms and regulates growth arrest [15]. Importantly, we found that the gene coding for cyclin-dependent kinase inhibitor 1A (p21) was up-regulated both in hSOD1G93A microglia (P90) and motoneurones (P120). Indeed, p21 is a downstream target of p53 and regulates several processes such as DNA repair, cell cycle arrest, cell differentiation and apoptosis. Through its antioxidant effects, p21 also protects cells from oxidative damage in vitro and in vivo [15]. Activation of microglial cells and acquisition of deleterious M1 state is associated with an increased generation of reactive oxygen species (ROS) [27] that is likely to participate in motoneurone demise. Polarisation of microglia/macrophages to pro- and anti-inflammatory states is driven by cytokines and other factors such as ROS within the tissue microenvironment [28]. While the functional role of deregulated Brca1 pathway in microglia remains to be determined, one hypothesis is that it may represent an attempt to counteract the detrimental effects of ROS and reflect an antioxidative defence mechanism through modulation of microglia polarisation.

Brca1 is implicated in a broad spectrum of functions; it regulates transcription and cell cycle progression, it is also involved in function that preserve genomic stability such as DNA repair pathways [29] and protection against oxidative damage to DNA. Many of these functions have been associated with CNS development but also with neurodegenerative diseases and in particular

**Fig. 6** Up-regulation of microglial Brca1 expression in human ALS spinal cords. Brightfield micrographs displaying Brca1 expression within the spinal cords of control (a & c) and ALS cases (b & d). Bar graphs showing the increase in Brca1 intensity within the white (e) and grey (f) matters of the spinal cord in ALS compared to control cases. Bars represent mean ± SEM (n = 5 for ALS and 14 for controls). *p < 0.05. Scale bar (a–d): 100 μm
with ALS. Brca1 is required for normal cerebral cortex size development [30] by preventing apoptosis [31]. Using a neural progenitor-specific driver to delete Brca1, Pao et al. demonstrated an important role of Brca1 in apoptotic and centrosomal functions in neuronal progenitors that may underlie DNA damage and brain size during development [31]. Brca1 is also associated with lack of spinal cord neural tube closure in spina bifida meningomyelocele [32, 33]. Moreover, Brca1-deficient embryos presented disorganised neuroepithelium associated with rapid proliferation and enhanced cell death [32].

De-regulation in Brca1 expression had been reported in Alzheimer’s [34, 35] and Huntington’s diseases [36]. Even though motor neuron diseases are not typical para-neoplastic syndromes, association with breast cancer had been regularly reported [37–41]. Moreover, there are occasional reports on improvement of motor neuron syndrome after cancer treatment [42–44].

Conclusion

Here we identify putative Brca1 involvement in ALS via hSOD1G93A microglia gene profiling and comparisons to our previous transcriptomic findings in hSOD1G93A motoneurones. Nevertheless, mRNA up-regulation of Brca1 in hSOD1G93A microglia could be simply anecdotal if it were restricted to a mouse model of ALS. This is not the case since we demonstrated that Brca1 protein is specifically expressed by human microglia and is significantly up-regulated in ALS patients.

These results substantiate that microglia are key non-cell autonomous players in the disease. Thus, the identification of the putative Brca1 involvement in a mouse model and human ALS provides new insights into the pathogenesis of ALS and points towards novel therapeutic targets.

Methods

Animals

Transgenic mice carrying the G93A human SOD1 mutation, B6SJL-Tg (SOD1-G93A)1Gur/J (ALS mice, high copy number) were purchased from The Jackson Laboratory (Bar Harbor, ME, USA) and bred on a B6SJL background. Transgenic mice were housed in controlled conditions (hygrometry, temperature and 12 h light/dark cycle). Ninety days old (P90, symptomatic) males were used. We carried out all animal experiments in accordance with the guidelines approved by the French Ministry of Agriculture and following the European Council directive (2010/63/UE). We minimised the number and suffering of animals.

Flow cytometry sorting of spinal cord microglia from SODG93A and control littermate mice

Mice were deeply anesthetised with tribromoethanol (500 mg/kg) and intracardially perfused with cold RNase-free 0.1 M phosphate base saline (PBS, Invitrogen, Carlsbad, USA); spinal cords were dissected. Only the lumbar (L1 - L5) segment was used and dissociated in 750 µl PBS, 100 µl trypsin 13 mg/ml, 100 µl hyaluronidase 7 mg/ml, 50 µl kinurenic acid 4 mg/ml (Sigma Aldrich, Saint Louis, USA) and 20 µl DNAsel 10 mg/ml (Roche, Rotkreuz, Switzerland) for 30 min at 37 °C. Finally, gentle mechanic dissociation was carried out by pipetting. Cell suspension was sieved on a 40 µm cell strainer (BD Biosciences, Franklin Lakes, USA). To eliminate myelin, cells were re-suspended in PBS-25 % sucrose and centrifuged for 20 min at 750 g. Cells were incubated for 20 min on ice in the primary antibody CD11b-APC 1/100 in PBS (BD biosciences, Franklin Lakes, USA) that specifically labels microglia. Cells were washed with cold PBS and re-suspended in PBS-7-AAD 2 µg/ml (Sigma Aldrich). Cells were sorted with a FACS ARIA (BD Biosciences, Franklin Lakes, USA), equipped with a 488 nm Laser Sapphire 488–20. Size threshold, morphology and 7-AAD were used to eliminate cellular debris and dead cells.

Microarray analysis of gene transcripts

Our data comply with the “Minimal Information About Microarray Experiment (MIAME)” guidelines. Total RNA was isolated using RNeasy Mini Kit, (Qiagen, Maryland, USA) including DNase treatment to remove potential genomic DNA contamination. We tested the quality of the starting RNA and of the amplified cRNA (Agilent 2100 bioanalyzer, RNA 6000 Pico LabChip, Palo Alto, USA) and proceeded only if the RNA quality was satisfactory. A criterion was a cut point for RNA integrity number (RIN) at 7 [45]. Fifty nanograms of RNA per chip were hybridized (three chips per condition) following a T7-based double amplification procedure.

Hybridization targets were obtained following a double amplification procedure according to the protocol developed by Affymetrix (GeneChip® Eukaryotic Small Sample Target Labeling Assay Version II, Affymetrix, Santa Clara, USA) and previously used [8, 9]. A hybridization mixture containing 5.5 µg of biotinylated cRNA was generated. The biotinylated cRNA was hybridized to Affymetrix GeneChip® MOE 430 2.0. Three chips per group (wild type and hSOD1G93A) were hybridized, each corresponding to microglia from at least six pooled mice. Chips were visualised on a 3000 gene scanner (Affymetrix, Santa Clara, USA). We selected the differentially expressed transcripts using the Affymetrix software MAS 5.0 and carried out pair-wise comparison analyses where each of the mutant samples was
compared to each of their respective control samples. This analysis is based on the Mann–Whitney pair-wise comparison test and allows the ranking of the results by concordance as well as the calculation of significance ($p$ value) of each identified gene expression [46, 47]. A gene must exhibit 50 % or more of the “present” calls in all samples to be considered “expressed” and has two or more “present” calls among the three sets of samples. Fold differences were calculated as the ratio between the average values within each condition. Signal values and detection calls (present or absent) for all samples were determined using Affymetrix MAS5.0. Based on power analysis, we had selected a cut off threshold of 1.75 ($p$ (α) 0.05, β 0.80) to identify transcripts that are differentially expressed between the controls and hSOD1$^{G93A}$ mutant mice. Statistics: t-test with un-equal variance. Pathway analysis was done with MetaCore (Thomson Reuters).

Quantitative real-time polymerase chain reaction
Candidate genes involved in Brca1 pathway were validated using qPCR. Similar to microarray, total RNA was extracted as described above from CD11-positive microglia isolated using FACS and used as a template in real time PCR. At least two animals were used for each analysis. To assess the involvement of microglia Brca1 at initial stages of the disease progression, we carried out qPCR at 60 and 90 days of age in hSOD1$^{G93A}$ and wild type mice. One round of amplification was done following the first cycle (first cDNA and cRNA synthesis) of the Affymetrix double amplification procedure before undertaking reverse transcription with random hexamers (Superscript II, Invitrogen, Carlsbad, CA). Real time PCR using Syber Green PCR Master Mix and Abi Prism SDS 7900 HT (Applied Biosystems, Foster City, CA) was done according to the manufacturer’s protocol. All amplicons were designed within the 3’ end of the cDNA using Primer Express Software 2.0 (Applied Biosystems, Foster City, CA) and when possible, overlapped exon-exon junctions. For the sequences of the primers, see Additional file 7: Table S5. All samples were analysed in triplicate and the values were normalised to four reference genes mitochondrial ribosomal protein S9 (RPS9), TATA box binding protein (TBP), actin β and eukaryotic translation elongation factor 1 (EEF1).

Human spinal cord samples
Human low thoracic and lumbar (T11-L5) spinal cords were obtained from 14 controls (males and females; 23 to 74 years of age; mean age: 52.4 years) and five ALS patients (males and females; 66 to 79 years of age; mean age: 71 years) from the Kantonsspital St. Gallen Fachbereichsleiter Muskelzentrum/ALS clinic under the approval of the Swiss legislation and from the New York Brain Bank–Taub Institute, Columbia University (NYBB), New York, USA. All donors had given their written consent for the autopsy and we followed the Declaration of Helsinki.

Immunohistochemistry
Mice were anesthetised with tribromoethanol (500 mg/kg) and perfused intracardially with cold PBS followed by cold 4 % paraformaldehyde (PFA, Sigma Aldrich). Spinal cords were removed and post fixed for 2 h in 4 % PFA. Samples were cryoprotected in sucrose 30 %, included in Tissue Teck (Sakura, Alphen aan den Rijn, The Netherlands), frozen and kept at −80 °C until processing.

For mice, free floating spinal cord transverse sections (20 μm) were washed twice in PBS (5 min), treated for 30 min in PBS containing lysine (20 mM, pH 7.2) and for 15 min in 1 % H$_2$O$_2$. Sections were blocked for 1 h with PBS containing bovine serum albumin (BSA, 1 %, Sigma Aldrich) and Triton X-100 (0.1 %, Fisher Scientific, Illkirch, France) and then incubated 48 h at 4 °C with CD11b (1/200, Developmental Studies Hybridoma Bank, Iowa, USA) primary antibody. Alexa-conjugated 594 secondary antibody was used (1/1000; Molecular Probes, Eugene, OR, USA).

For human spinal cord 22 μm-thick cryosections of lumbar and lower thoracic segments were collected on super frost plus slides and were processed as described above. For dual fluorescence labelling, sections were placed in a cocktail of rat anti CD11b (1/100, Hybridoma Bank, University of Iowa, USA) and rabbit anti-Brca1 (1/100, Santa Cruz Biotechnology, Dallas, USA) primary antibodies for 48 h at 4 °C. Sections were washed in 0.1 M PBS followed by incubation in corresponding secondary antibodies conjugated to Alexa 488 and 594 (1/1000; Molecular Probes, Eugene, OR, USA). For peroxidase labelling, sections were placed for 48 h at 4 °C in either rabbit anti Iba1 (macrophage/microglia-specific calcium-binding protein) (1/1000, Wako Pure Chemical Industries, Osaka, Japan) or rabbit anti-Brca1 (1/100, Santa Cruz Biotechnology, Dallas, USA) primary antibodies. Spinal cord sections were then incubated in donkey anti-rabbit (1/500, Jackson ImmunoResearch, Carlsbad, USA) antibody for 2 h at 4 °C. Sections were then washed in TRIS buffer and enzymatic revelation was done with nickel enhanced DAB and H$_2$O$_2$ 0.1 % as a substrate. Sections were then dehydrated in ascending concentration of ethanol and finally xylene. Coverslips were applied using Entellan (Merck KGaA, Darmstadt, Germany).

Morphometric bright field photographs had been obtained and analysed using NanoZoomer RS slide scanner (NanoZoomer Digital Pathology System and NDP view software, Hamamatsu, Japan). For immunofluorescence
images, we used laser scanning inverted confocal microscopy (Leica SP5, Mannheim, Germany). Laser intensity and detector sensitivity settings were kept constant for all image acquisitions within a given experiment. Brcal staining intensity measurement was done by measuring their optical density (OD) using ImageJ (National Institutes of Health, USA), as described previously [48]. For each given sample we analysed at least three 22-µm-thick section with 330 µm distance from each other. Statistics: un-paired t-test done with GraphPad Prism version 5.03 (GraphPad software, CA, USA). Significance was accepted at p ≤ 0.05. Results are expressed as mean ± S.E.M.

Additional files

Additional file 1: Table S1. Database of differential expression comparison of hSOD1G93A microglia microarray data relative to control microglia at 90 days of age. We list information for each dysregulated genes in hSOD1G93A microglia as compared to control microglia. With both the p-value and the step-up p-value that is the false discovery rate (FDR) analogue of the p-value. Three chips were used per condition (wild type and SOD1G93A) with microglia from lumbar spinal cord of at least six pooled mice.

Additional file 2: Figure S1. Comparison of gene dysregulation in microglia and motoneurones. (A) Venn diagrams showing that 19 genes are commonly dysregulated in microglia and motoneurones at symptomatic age (P90). (B) Comparison of dysregulated genes in microglia at P90 and motoneurones at disease end stage shows that 65 genes are commonly dysregulated, 603 genes are uniquely dysregulated in hSOD1G93A microglia and 320 uniquely in hSOD1G93A motoneurones. Three chips were used per condition (wild type and SOD1G93A) with microglia from lumbar spinal cord of at least six pooled mice.

Additional file 3: Table S2. Comparison of gene dysregulation in both microglia and motoneurones at symptomatic age (P90) and in motoneurone only (P90) using gene ontology enrichment and network analysis. In all tables the top scored categories have the lowest p-value. Table S2A: Process networks ranking. Table S2B: Gene ontology processes ranking and Table S2C: Pathway maps ranking.

Additional file 4: Table S3. Comparison of gene dysregulation in microglia at symptomatic age (P90) and in motoneurone at the end stage of the disease (P120) using gene ontology enrichment and network analysis. In all tables the top scored categories have the lowest p-value. Table S3A: Process networks ranking. Table S3B: Gene ontology processes ranking and Table S3C: Pathway maps ranking.

Additional file 5: Table S4. Gene ontology enrichment and network analysis of gene dysregulation in microglia at symptomatic age (P90). In all tables the top scored categories have the lowest p-value. Table S4A: Process networks. Table S4B: Gene ontology processes and Table S4C: Pathway maps. Percentage of dysregulated genes corresponds to the ratio of dysregulated genes in our data out of annotated genes in the given category (Gene Ontology).

Additional file 6: Figure S2. Quantitative real-time polymerase chain reaction (qPCR) validation of microarray findings related to candidate genes involved in Brcal pathway. To confirm the microarray results, the seven identified genes involved in Brcal pathway were analysed by real time qPCR. Bar graphs showing up-regulation of Brcal, Cdn1na, Myc, Ptna and Sta1 as well as down-regulation of Gadd45a and Sp3 in hSOD1G93A microglia at 90 (B) but not 60 days (A) as compared to control microglia. For each sample, real time PCR was done in triplicate.

Additional file 7: Table S5. Candidate genes involved in Brcal pathway selected for qPCR to validate the microarray data.

Abbreviations

ALS: Amyotrophic lateral sclerosis; CNS: Central nervous system; DAB: Diamobenzidine; ER: Endoplasmic reticulum; FACS: Fluorescence-activated cell sorting; FALS: Familial amyotrophic lateral sclerosis; FC: Fold change; GO: Gene ontology; OD: Optical density; PBS: Phosphate base saline; PFA: Paraformaldehyde; pmn: Progressive motor neuronopathy; RIN: RNA integrity number; ROS: Reactive oxygen species; SOD: Super oxide dismutase.

Competing interests

The authors declare that they have no competing interests.

Authors’ contributions

HNN participated in the design of the study, performed immunohistology, participated to the analysis and interpretation of data and helped to draft the manuscript. JCS participated in the design of the study, carried out FACS and participated to immunohistology. YNG participated to acquisition of FACS data. MT participated to immunohistology. AS performed autopsy. MdmV participated to acquisition of FACS data. MW performed patient selection and obtained patients consent. FEP: conception, design of the work, analysis and interpretation of data, drafting the work and final approval. All authors read and approved the final manuscript.

Acknowledgments

We are grateful to ALS patients and their relatives that donate their tissues. We acknowledge the New York Brain Bank–The Taub Institute, Columbia University (NYBB). The hyridomada CD11b antibody developed by Timothy A. Springer was obtained from the Developmental Studies Hybridoma Bank developed under the auspices of the NICHD and maintained by the University of Iowa, Department of Biology, Iowa city, IA 52242. We thank the IGE3Genomics Platform, University of Geneva Switzerland for their assistance in transcriptomic and qPCR analysis.

This work was supported by the Spanish Government, Plan Nacional de H+D+I 2008–2011 and ISCIII-Subdirección General de Evaluación y Fomento de la investigación (PI10/07079) [to FEP], the Government of the Basque Country grant (Proyectos de Investigacion Sanitaria y Fondo Comun de Cooperacion Aquatania-Euskadi) [to FEP], the “Fondation pour la Recherche Medicale” [to FEP] and the French Government, ANR-FNS grant, GIALS (N° ANR-14-CE36-0009-01) [to FEP], the patient organizations “Demain Debout Aquitaine” [to YNG and HNN] and “Verticale” [to FEP and HNN].

Author details

1Institute for Neurosciences of Montpellier (INM), INSERM U1051, 80, rue Augustin Fliche, 34091 Montpellier, Cedex 5, France. “Integrative Biology of Neurodegeneration”, IEREBASQUE Basque Foundation for Science and Neuroscience Department, University of the Basque Country, Bilbao, Spain.
2Kantonspital St. Gallen, FachMuskelzentrum/ALS clini, St. Gallen, Switzerland. 3CIC bioGUNE, Cell Biology & Stem Cells Unit, Technological Park of Bizkaia, Derio, Spain. 4Department “Biologie-Mécanismes du Vivant” Faculty of Science, University of Montpellier, Montpellier, France.

Received: 5 March 2015 Accepted: 25 June 2015

References

1. Leblond CS, Kaneb HM, Dion PA, Rouleau GA. Dissection of genetic factors associated with amyotrophic lateral sclerosis. Exp Neurol. 2014/262 Pt B 91–101.
2. Gurney ME, Pu H, Chiu AI, Dal Canto MC, Polchow CY, Alexander DD, et al. Motor neuron degeneration in mice that express a human Cu, Zn superoxide dismutase mutation. Science. 1994/264/1772–5.
3. Philips T, Rothstein JD. Glial cells in amyotrophic lateral sclerosis. Exp Neurol. 2014/262 Pt B 111–20.
4. Crain JW, Nikodemova M, Watters JJ. Microglia express distinct M1 and M2 phenotypic markers in the postnatal and adult central nervous system in male and female mice. Eur J Neurosci. 2013/91/1143–51.
5. Michelucci A, Heurtiaux T, Grandbarbe L, Morga E, Heuschling P. Characterization of the microglial phenotype under specific pro-inflammatory and anti-inflammatory conditions: effects of oligomeric and fibrillar amyloid-beta. J Neuroimmunol. 2009/210–13.
6. Bites D, Vaz AR. Microglia centered pathogenesis in ALS. Insights in cell interconnectivity. Front Cell Neurosci. 2014/8117.
7. Gerber YN, Sabourin JC, Rabano M, Vivanco M, Perrin FE. Early functional deficit and microglial disturbances in a mouse model of amyotrophic lateral sclerosis. PLoS One. 2012;7:e36000.
8. Perrin FE, Boisvert G, Doquier M, Schaad O, Descombes P, Kato AC. No widespread induction of cell death genes occurs in pure moron neurons in an amyotrophic lateral sclerosis mouse model. Hum Mol Genet. 2005;14:309–20.
9. Perrin FE, Boisvert G, Lathuiere A, Kato AC. Cell death pathways differ in several mouse models with moronone disease: analysis of pure moronone populations at a presymptomatic age. J Neurochem. 2006;98:1959–72.
10. Coppede F. An overview of DNA repair in amyotrophic lateral sclerosis. ScientificWorldJournal. 2011;11:1679–91.
11. Caestecker KW, Van de Walle GR. The role of BRCA1: role in cell cycle regulation and the maintenance of genomic stability. FEBS J. 2011;278:630–46.
12. Bae I, Fan S, Meng Q, Rih JK, Kim HJ, Kang HU, et al. BRCA1 induces antioxidant gene expression and resistance to oxidative stress. Cancer Res. 2003;64:7893–909.
13. Vurasuner B, Poli G, Basaga H. Tumor suppressor genes and ROS: complex networks of interactions. Free Radic Biol Med. 2012;52:7–18.
14. Rakha EA, El-Seikh SE, Kandil MA, El-Sayed ME, Green AR, Ellis IO. Expression of BRCA1 protein in breast cancer and its prognostic significance. Hum Pathol. 2008;39:857–65.
15. Bernard-Gallon DJ, De Latour MP, Syliva V, Vissac C, Aunoble B, Chassagne J, et al. Brca1 and Brca2 protein expression patterns in different tissues of murine origin. Int J Oncol. 2001;18:271–80.
16. Korhonen L, Brannvall K, Skoglosa Y, Lindholm D. Tumor suppressor gene BRCA-1 is expressed by embryonic and adult neural stem cells and involved in cell proliferation. J Neurosci Res. 2003;71:769–76.
17. Gerber YN, Sabourin JC, Hugnot JP, Perrin FE. Unlike physical exercise, modified environment increases the lifespan of SOD1G93A mice however both conditions induce cellular changes. PLoS One. 2012;7:e45503.
18. Ilieva H, Polymenidou M, Cleveland DW. Non-cell autonomous toxicity in neurodegenerative disorders: ALS and beyond. J Cell Biol. 2009;187:761–72.
19. Liu WM, Mei R, Di X, Ryder TB, Hubbell E, Dee S, et al. Analysis of high measurement of RNA integrity using the 2100 Bioanalyzer capillary electrophoresis system. Ann Hematol. 2009;88:1177–83.
20. Hubbell E, Liu WM, Mei R. Robust estimators for expression analysis. Bioinformatics. 2002;18:1585–92.
21. Liu WM, Mei R, Di X, Ryder TB, Hubbell E, Dee S, et al. Analysis of high density expression microarrays with signed-rank call algorithms. Bioinformatics. 2002;18:1593–9.
22. Noristani NH, Olabarria M, Vehrkratsy A, Rodriguez JJ. Serotonin fibre sprouting and increase in serotonin transporter immunoactivity in the CA1 area of hippocampus in a triple transgenic mouse model of Alzheimer’s disease. Eur J Neurosci. 2010;32:71–9.