

RESEARCH ARTICLE

Open Access



Brca1 is expressed in human microglia and is dysregulated in human and animal model of ALS

Harun Najib Noristani^{1†}, Jean Charles Sabourin^{2†}, Yannick Nicolas Gerber^{1,2}, Marisa Teigell¹, Andreas Sommacal³, Maria dM Vivanco⁴, Markus Weber³ and Florence Evelyne Perrin^{1,2,5*}

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 hSOD1^{G93A} mice, the most widely used animal model of ALS. We first identified unique hSOD1^{G93A} 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 hSOD1^{G93A} 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, hSOD1^{G93A} 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 hSOD1^{G93A} 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

* Correspondence: florence.perrin@inserm.fr

†Equal contributors

¹Institute for Neurosciences of Montpellier (INM), INSERM U1051, 80, rue Augustin Fliche, 34091 Montpellier, Cedex 5, France

²"Integrative Biology of Neurodegeneration", IKERBASQUE Basque Foundation for Science and Neuroscience Department, University of the Basque Country, Bilbao, Spain

Full list of author information is available at the end of the article

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 motoneurons from the lumbar spinal cord of hSOD1^{G93A} mice [8] and two other mouse models of motoneuron disease [9]. We revealed a unique motoneuron 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 DNA 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^{G93A} microglia gene profiling and comparisons to our previous transcriptomic findings in hSOD1^{G93A} motoneurons. 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^{G93A} lumbar spinal cord

We have previously described early microglial disturbances in hSOD1^{G93A} male mice reflected at P90 by a heterogeneous Iba1⁺ microglial distribution with higher density within the grey matter in hSOD1^{G93A} 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^{G93A} mice (Fig. 1b). To further analyse transcriptomic modification specifically in microglia, we isolated microglia of hSOD1^{G93A} 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⁺ microglia in hSOD1^{G93A} versus controls (26 350; $n = 15$ in hSOD1^{G93A} 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 motoneurons

We had previously identified dysregulated genes in hSOD1^{G93A} motoneurons during disease progression [8, 9]. To unravel potential molecular cross-talk between microglia and motoneurons, we performed a comparative analysis of gene dysregulation in both cell populations. Comparison of dysregulated genes at P90 between motoneurons (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 motoneurons 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 motoneurons 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 motoneurons only group (Additional file 3: Table S2B). Signalling and metabolic pathway analysis revealed immune response and cytoskeleton remodeling as first ranked in the commonly and motoneuron

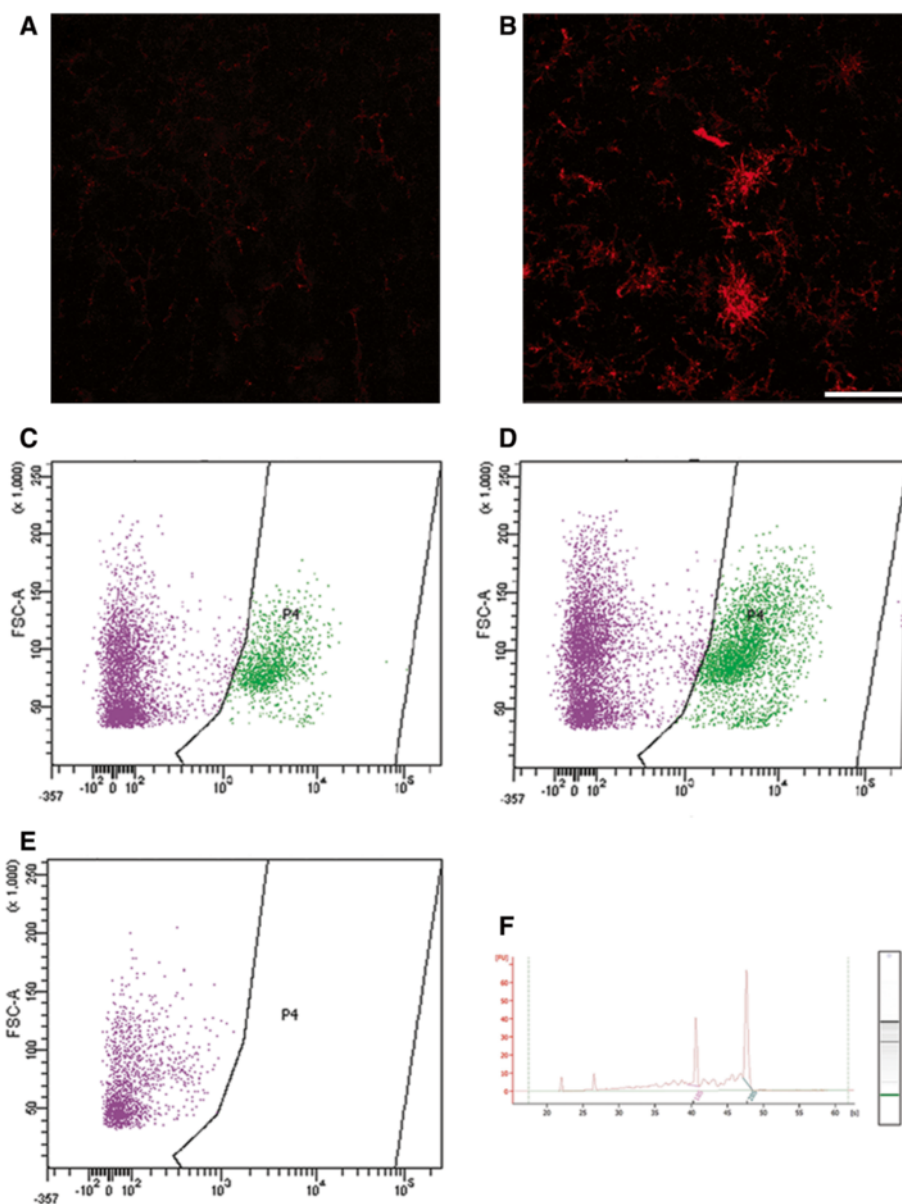


Fig. 1 FACS analysis of CD11b⁺ microglia in control and hSOD1^{G93A} mice from the lumbar segment of the spinal cords at 90 days of age. Microglia were sorted by flow cytometry using the microglia marker CD11b. **a** Confocal images of CD11b expression in spinal cord microglia from control at 90 day of age and **(b)** from transgenic hSOD1^{G93A} mice at early symptomatic age. *Scale bars (a & b):* 50 μ m. **c-d** Representative flow cytometry analysis dot plot displaying microglia profiles. **c** Control and **(d)** hSOD1^{G93A} spinal microglia at P90. In both **(c)** control and **(d)** hSOD1^{G93A} surrounded areas, designed as “P4”, correspond to the labelled cells. **e** Negative control (without CD11b labelling). The X-axis represents the intensity of fluorescence and the Y axis the size of the cells. **f** RNA quality isolated from FACSed microglia

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 motoneurons. Our previous microarrays analysis of microdissected motoneurons at the end stage of the disease (P120)

showed no dysregulation of genes associated with cell death [8], this most likely reflects that dissected motoneurons were at an early demise stage. Indeed, we selected motoneurons that had an identifiable nucleus and a diameter of at least 25 μ m, picking a sub-population of neurons that may resist degeneration. Three hundred twenty genes were uniquely dysregulated in hSOD1^{G93A} motoneurons; 603 uniquely dysregulated in hSOD1^{G93A} microglia; 65 genes were

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 motoneurons at P90, Additional file 4: Table S3). Particularly, inflammation and immune response were ranked top in motoneurons (Additional file 4: Table S3A). Interestingly, signalling and metabolic pathway analysis revealed the involvement of heme metabolism and DNA damage in both motoneurons and microglia (Additional file 4: Table S3C).

Unique transcriptomic profiles of hSOD1^{G93A} microglia

To identify processes and pathways modified in hSOD1^{G93A} 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 %, $p = 2.1E-08$) (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 (*SPP1*) 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 %, $p = 7.3E-04$), with 16 being up-regulated. Inflammation network was also dysregulated in hSOD1^{G93A} microglia (ranked 9th, 9 % of the annotated genes in this process, $p = 1.4E-04$, Additional file 5: Table S4A, Fig. 2d) with 14 up-regulated genes (including a 12.6 fold increase for IGF-1, Fig. 2d). GO cellular processes analysis ranked immune response as first (61/1505 genes, 4 %, $p = 1.3E-18$, Additional file 5: Table S4B, Fig. 2b). Out of the 38 most significantly dysregulated genes 5 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 %, $p = 1.8E-09$, 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 4th dysregulated cellular processes (31/416 genes, 7.45 %, $p = 8.3E-09$, 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 (*Brca1*) pathway is dysregulated in hSOD1^{G93A} mice

Pathways map analysis ranked as the third position DNA damage and specifically the involvement of *Brca1*

as a transcription regulator (Additional file 5: Table S4C, Figs. 2g and 3). Indeed, in hSOD1^{G93A} microglia (7/30 genes, 23 %; $p = 1.6E-05$, 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* (2.18-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 *Brca1* transcript itself was not dysregulated in motoneurons, 4 genes that are involved in *Brca1* pathway were also up-regulated in hSOD1^{G93A} motoneurons 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^{G93A} 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^{G93A} 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*, *Pcna* and *Stat1* as well as down-regulation of *Gadd45a* and *Sp3* in hSOD1^{G93A} microglia (Additional file 6: Figure S2B). It is important to note that dysregulation in *Cdkn1a*, *Myc*, *Pcna*, *Stat1*, *Gadd45a* and *Sp3* transcripts may also be involved in other signalling pathways. These findings confirm *Brca1* involvement in hSOD1^{G93A} microglia is specifically triggered at 90 days of age when the pronounced microgliosis becomes evident.

Brca1 protein is expressed in human microglia and is up-regulated in ALS patients

To investigate *Brca1* protein expression in human microglia, we performed dual immunofluorescence labelling using *Brca1* and CD11b antibodies (Fig. 4). *Brca1* 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 *Brca1* revealed microglial profile that were identical to *Iba1* (the most commonly used

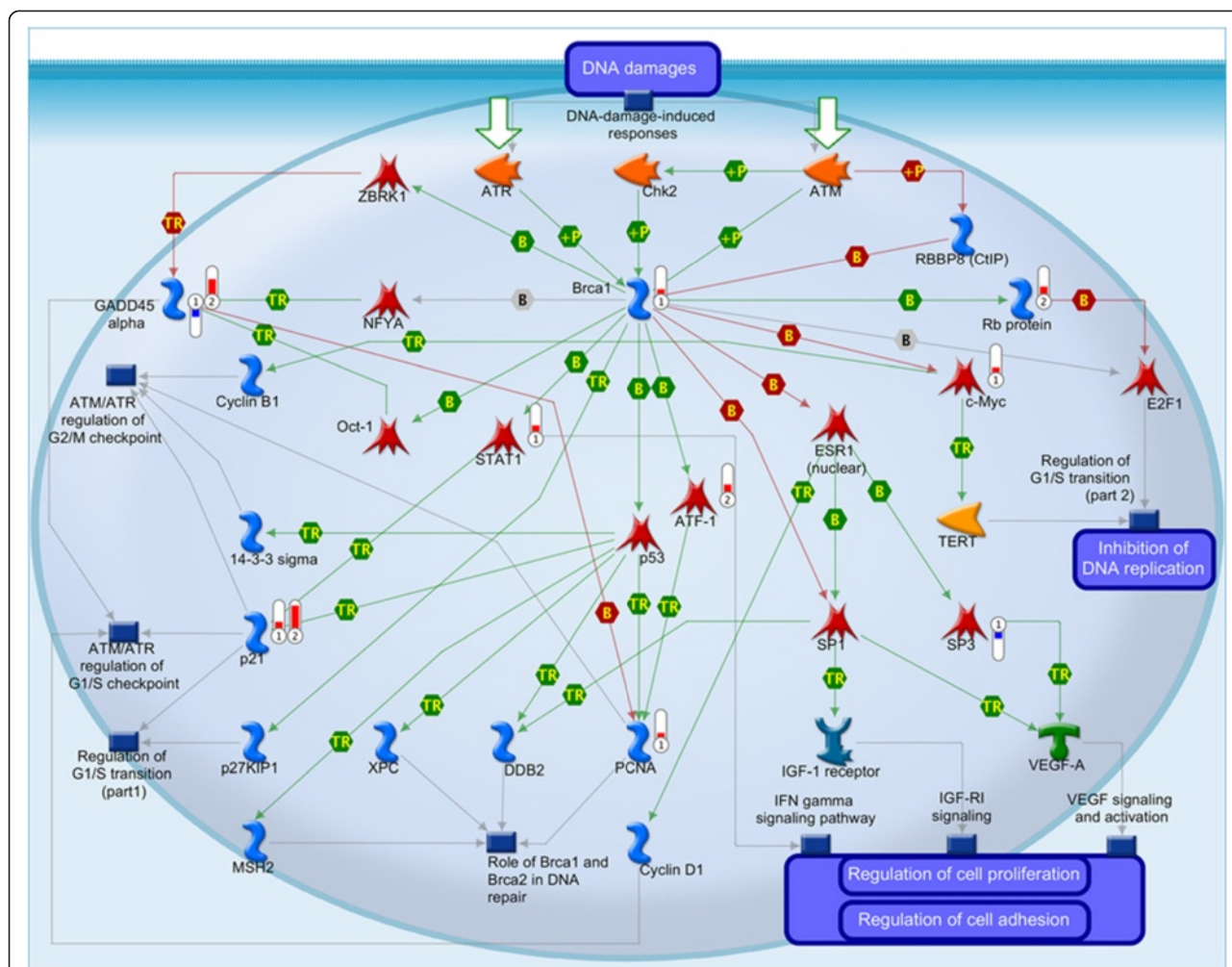


Fig. 3 Brca1 pathway is involved in microglia from hSOD1^{G93A} mice. Gene ontology pathway map analyses of dysregulated genes either in hSOD1^{G93A} microglia at symptomatic age (P90) or in hSOD1^{G93A} motoneurons at the end stage of the disease (P120) demonstrate the involvement of Brca1 pathway. Thermometers indicate gene dysregulation (red: up-regulated; blue: down-regulated), thermometer levels correspond to the level of dysregulation. Thermometers with number 1 represent gene dysregulation in hSOD1^{G93A} microglia and number 2 in hSOD1^{G93A} motoneurons. Interactions between objects: green (positive or activation); red (negative or inhibition); grey (unspecified). B Binding (physical interaction between molecules), TR Transcription regulation (physical binding of a transcription factor to target gene's promoter), +P Phosphorylation (protein activity is altered via addition of a phosphate group). Binding protein Transcription factor Kinase Generic enzyme

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

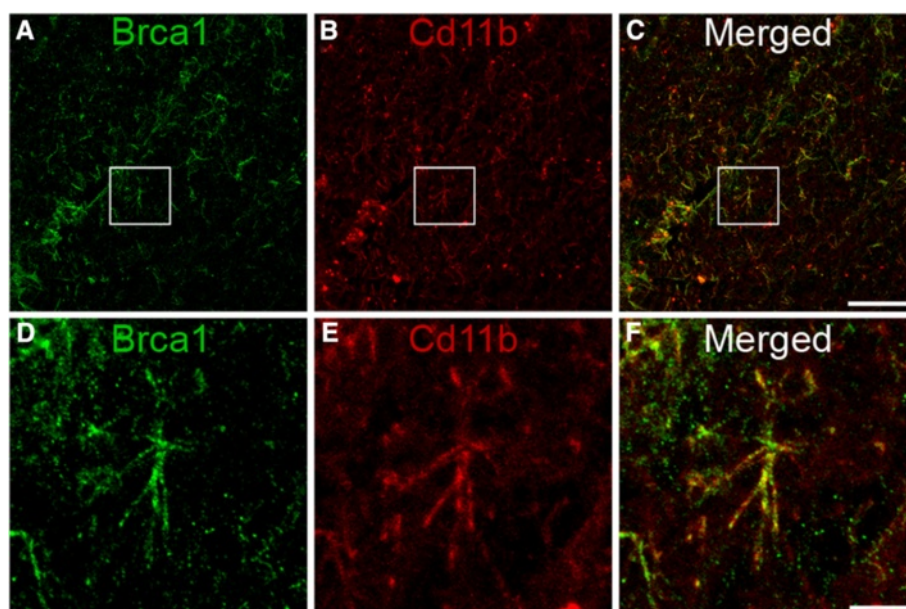


Fig. 4 Microglial expression of Brca1 in human spinal cords. Confocal micrographs displaying Brca1 (a, d), CD11b (b, e) and dual Brca1/CD11b expression (c, f) expression by microglia within the human spinal cord. Brca1 labelled microglia displayed typical ramified morphology with small cell bodies and large and thin processes that completely colocalised with CD11b-positive microglia (c & f). Scale bars (a–c): 50 μ m; (d–f): 10 μ m

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*, *Tnf α* , *Bcl2a1a* 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 hSOD1^{G93A} 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 hSOD1^{G93A} microglia from the lumbar region of the spinal cord over-express *progranulin*, *Igf1* and *osteopontin*, all potential neurotrophic factors, and thus confirmed findings from a previous study [23]. Interestingly, we had previously identified in pure motoneurons of two mouse models of motoneuron disease (hSOD1^{G93A} 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 hSOD1^{G93A} mice [8] and *IGFBP4* mRNA was induced at pre-symptomatic age in *pmm* mice [9]. We also confirmed the up-regulation of potential neurotoxic factors (including *Mmp12*, *tnf- α* and interferon-induced protein with tetratricopeptide repeats) [23]. However, we did not confirm the dysregulation of

the genes coding for *IL-1 β* , *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 hSOD1^{G93A} 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 hSOD1^{G93A} 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 hSOD1^{G93A} 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 IGF1 receptors-mediated cell survival [25, 26]. This simultaneous up-regulation may represent a potential neuroprotective phenotype of microglia in ALS

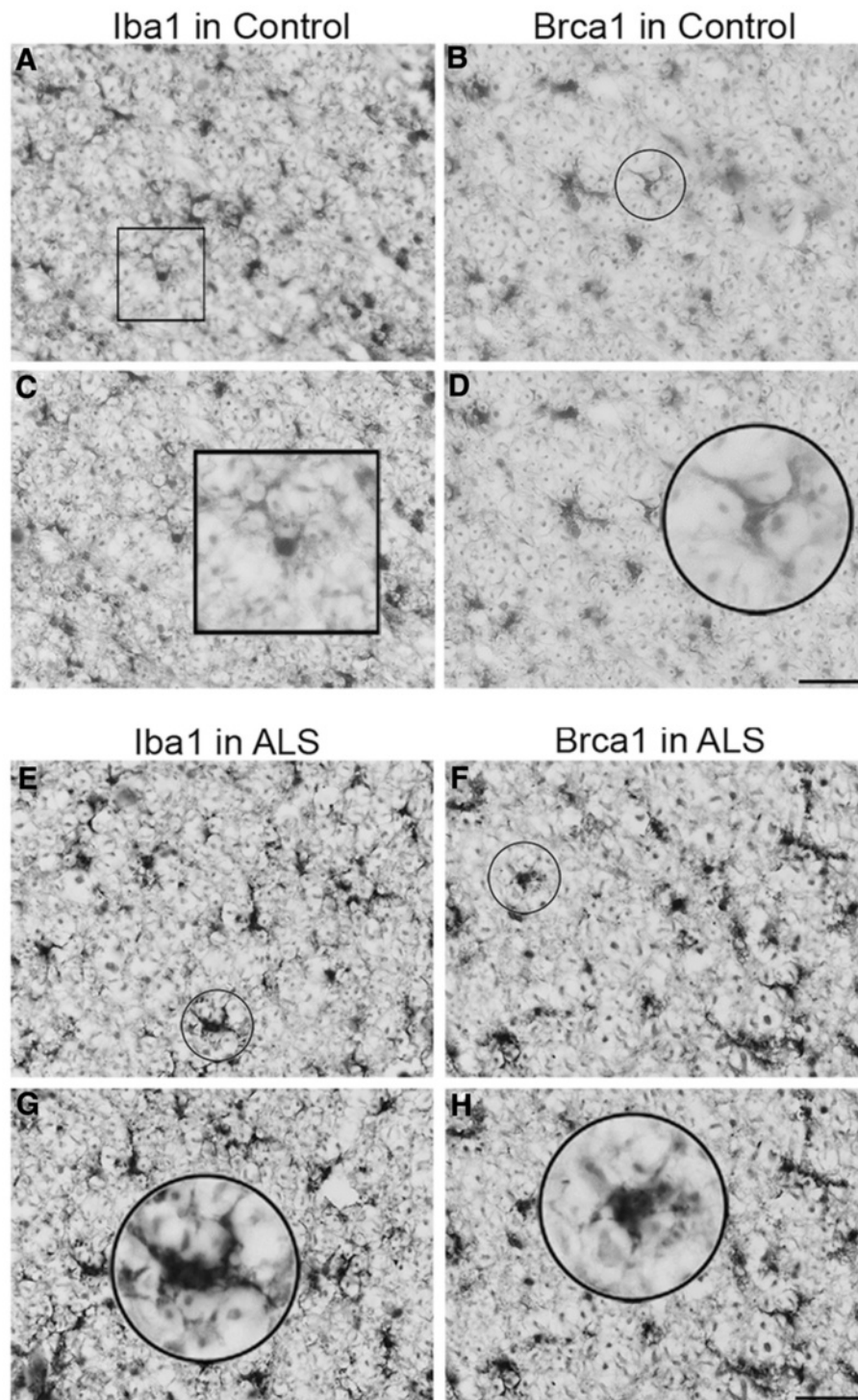


Fig. 5 Microglial Iba1 and Brca1 expression within adjacent human control and ALS spinal cord. Brightfield micrographs indicating Iba1 and Brca1 expression within adjacent sections of control (**a-d**) and ALS (**e-h**) spinal cords. Both Iba1 and Brca1-labelled microglia displayed typical ramified morphology 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 μ m

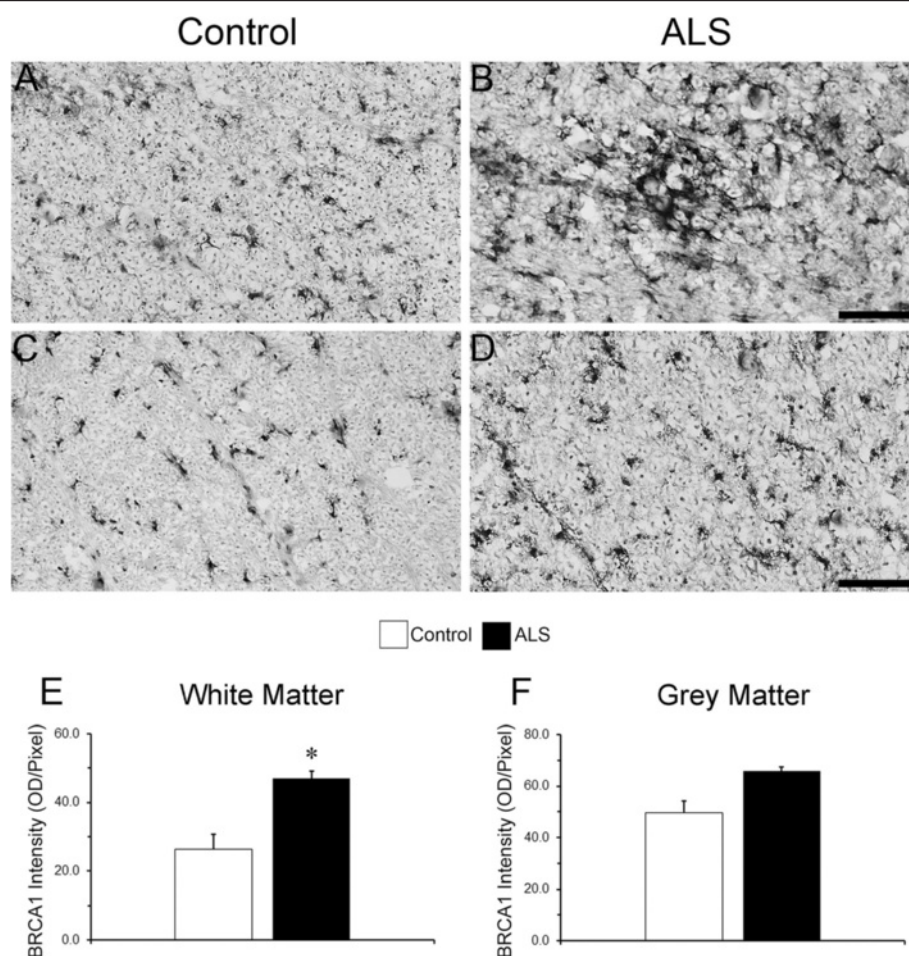


Fig. 6 Up-regulation of microglial Brca1 expression in human ALS spinal cords. Brightfield micrographs displaying Brca1 expression within the spinal cords of control (a & c) and ALS cases (b & d). Bar graphs showing the increase in Brca1 intensity within the white (e) and grey (f) matters of the spinal cord in ALS compared to control cases. Bars represent mean \pm SEM ($n = 5$ for ALS and 14 for controls). * $p < 0.05$. Scale bar (a-d): 100 μ m

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 hSOD1^{G93A} microglia (P90) and up-regulated in motoneurons (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 hSOD1^{G93A} microglia (P90) and motoneurons (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 motoneuron 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

with ALS. *Brcal* is required for normal cerebral cortex size development [30] by preventing apoptosis [31]. Using a neural progenitor-specific driver to delete *Brcal*, Pao et al. demonstrated an important role of *Brcal* in apoptotic and centrosomal functions in neuronal progenitors that may underlie DNA damage and brain size during development [31]. *Brcal* is also associated with lack of spinal cord neural tube closure in *spina bifida* meningocele [32, 33]. Moreover, *Brcal*-deficient embryos presented disorganised neuroepithelium associated with rapid proliferation and enhanced cell death [32].

De-regulation in *Brcal* expression had been reported in Alzheimer's [34, 35] and Huntington's diseases [36]. Even though motor neuron diseases are not typical paraneoplastic 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 *Brcal* involvement in ALS via hSOD1^{G93A} microglia gene profiling and comparisons to our previous transcriptomic findings in hSOD1^{G93A} motoneurons. Nevertheless, mRNA up-regulation of *Brcal* in hSOD1^{G93A} microglia could be simply anecdotal if it were restricted to a mouse model of ALS. This is not the case since we demonstrated that *Brcal* 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 *Brcal* 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 for transcriptomic analysis and immunohistochemistry. Litter-matching between groups were done. 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 SOD^{G93A} 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 DNaseI 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 hSOD1^{G93A}) 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 *Brcal* 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 *Brcal* 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 anaesthetised 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 μ m) 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 % H₂O₂. 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 μ m-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-*Brcal* (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-*Brcal* (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 H₂O₂ 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- μ m-thick section with 330 μ m distance from each other. Statistics: un-paired t-test done with GraphPad Prism version 5.03 (Graph-Pad software, CA, USA). Significance was accepted at $p \leq 0.05$. Results are expressed as mean \pm S.E.M.

Additional files

Additional file 1: Table S1. Database of differential expression comparison of hSOD1^{G93A} microglia microarray data relative to control microglia at 90 days of age. We list information for each dysregulated genes in hSOD1^{G93A} 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 SOD1^{G93A}) with microglia from lumbar spinal cord of at least six pooled mice.

Additional file 2: Figure S1. Comparison of gene dysregulation in microglia and motoneurons. (A) Venn diagrams showing that 19 genes are commonly dysregulated in microglia and motoneurons at symptomatic age (P90). (B) Comparison of dysregulated genes in microglia at P90 and motoneurons at disease end stage shows that 65 genes are commonly dysregulated, 603 genes are uniquely dysregulated in hSOD1^{G93A} microglia and 320 uniquely in hSOD1^{G93A} motoneurons. Three chips were used per condition (wild type and SOD1^{G93A}) 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 motoneurons at symptomatic age (P90) and in motoneuron 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 motoneuron 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*, *Cdkn1a*, *Myc*, *Pcna* and *Stat1* as well as down-regulation of *Gadd45a* and *Sp3* in hSOD1^{G93A} 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: Diaminobenzidine; 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 neuropathy; 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 hybridoma 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 I+D+I 2008–2011 and ISCIII-Subdirección General de Evaluación y Fomento de la investigación (P110/00709) [to FEP], the Government of the Basque Country grant (Proyectos de Investigación Sanitaria and Fondo Comun de Cooperación Aquitania-Euskadi) [to FEP], the "Fondation pour la Recherche Médicale" [to FEP] and the French Government, ANR-FNS grant, GIALS (N° ANR-14-CE36-0009-01) [to FEP], the patient organisations "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. ²"Integrative Biology of Neurodegeneration", IKERBASQUE Basque Foundation for Science and Neuroscience Department, University of the Basque Country, Bilbao, Spain. ³Kantonspital St. Gallen. FachMuskelzentrum/ALS clinic, St. Gallen, Switzerland. ⁴CIC bioGUNE, Cell Biology & Stem Cells Unit, Technological Park of Bizkaia, Derio, Spain. ⁵Department "Biologie-Mécanismes du Vivant" Faculty of Science, University of Montpellier, Montpellier, France.

Received: 5 March 2015 Accepted: 25 June 2015

Published online: 01 August 2015

References

- 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.
- Gurney ME, Pu H, Chiu AY, 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.
- Philips T, Rothstein JD. Glial cells in amyotrophic lateral sclerosis. *Exp Neurol*. 2014;262 Pt B:111–20.
- Crain JM, 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. *J Neurosci Res*. 2013;91:1143–51.
- Michelucci A, Heurtaux 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;210:3–12.
- Brites D, Vaz AR. Microglia centered pathogenesis in ALS: insights in cell interconnectivity. *Front Cell Neurosci*. 2014;8:117.

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, Boisset G, Docquier M, Schaad O, Descombes P, Kato AC. No widespread induction of cell death genes occurs in pure motoneurons in an amyotrophic lateral sclerosis mouse model. *Hum Mol Genet*. 2005;14:3309–20.
9. Perrin FE, Boisset G, Lathuiliere A, Kato AC. Cell death pathways differ in several mouse models with motoneuron disease: analysis of pure motoneuron 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;1:1679–91.
11. Caestecker KW, Van de Walle GR. The role of BRCA1 in DNA double-strand repair: past and present. *Exp Cell Res*. 2013;319:575–87.
12. Somasundaram K. Breast cancer gene 1 (BRCA1): role in cell cycle regulation and DNA repair—perhaps through transcription. *J Cell Biochem*. 2003;88:1084–91.
13. Savage KI, Harkin DP. BRCA1, a 'complex' protein involved in the maintenance of genomic stability. *FEBS J*. 2015;282:630–46.
14. Bae I, Fan S, Meng Q, Rih JK, Kim HJ, Kang HJ, et al. BRCA1 induces antioxidant gene expression and resistance to oxidative stress. *Cancer Res*. 2004;64:7893–909.
15. Vurusaner B, Poli G, Basaga H. Tumor suppressor genes and ROS: complex networks of interactions. *Free Radic Biol Med*. 2012;52:7–18.
16. Rakha EA, El-Sheikh 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.
17. Bernard-Gallon DJ, De Latour MP, Sylvain 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.
18. 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.
19. 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.
20. Ilieva H, Polymenidou M, Cleveland DW. Non-cell autonomous toxicity in neurodegenerative disorders: ALS and beyond. *J Cell Biol*. 2009;187:761–72.
21. Liao B, Zhao W, Beers DR, Henkel JS, Appel SH. Transformation from a neuroprotective to a neurotoxic microglial phenotype in a mouse model of ALS. *Exp Neurol*. 2012;237:147–52.
22. Hickman SE, Kingery ND, Ohsumi TK, Borowsky ML, Wang LC, Means TK, et al. The microglial sensome revealed by direct RNA sequencing. *Nat Neurosci*. 2013;16:1896–905.
23. Chiu IM, Morimoto ET, Goodarzi H, Liao JT, O'Keeffe S, Phatnani HP, et al. A neurodegeneration-specific gene-expression signature of acutely isolated microglia from an amyotrophic lateral sclerosis mouse model. *Cell Rep*. 2013;4:385–401.
24. Beers DR, Zhao W, Liao B, Kano O, Wang J, Huang A, et al. Neuroinflammation modulates distinct regional and temporal clinical responses in ALS mice. *Brain Behav Immun*. 2011;25:1025–35.
25. Liu B, Li D, Guan YF. BRCA1 regulates insulin-like growth factor 1 receptor levels in ovarian cancer. *Oncol Lett*. 2014;7:1733–7.
26. Werner H, Bruchim I. IGF-1 and BRCA1 signalling pathways in familial cancer. *Lancet Oncol*. 2012;13:e537–44.
27. Henkel JS, Beers DR, Zhao W, Appel SH. Microglia in ALS: the good, the bad, and the resting. *J Neuroimmune Pharmacol*. 2009;4:389–98.
28. David S, Kroner A. Repertoire of microglial and macrophage responses after spinal cord injury. *Nat Rev Neurosci*. 2011;12:388–99.
29. Mantha AK, Sarkar B, Tell G. A short review on the implications of base excision repair pathway for neurons: relevance to neurodegenerative diseases. *Mitochondrion*. 2014;16:38–49.
30. Pulvers JN, Huttner WB. Brca1 is required for embryonic development of the mouse cerebral cortex to normal size by preventing apoptosis of early neural progenitors. *Development*. 2009;136:1859–68.
31. Pao GM, Zhu Q, Perez-Garcia CG, Chou SJ, Suh H, Gage FH, et al. Role of BRCA1 in brain development. *Proc Natl Acad Sci U S A*. 2014;111:E1240–8.
32. Gowen LC, Johnson BL, Latour AM, Sulik KK, Koller BH. Brca1 deficiency results in early embryonic lethality characterized by neuroepithelial abnormalities. *Nat Genet*. 1996;12:191–4.
33. King TM, Au KS, Kirkpatrick TJ, Davidson C, Fletcher JM, Townsend I, et al. The impact of BRCA1 on spina bifida meningomyelocele lesions. *Ann Hum Genet*. 2007;71:719–28.
34. Evans TA, Raina AK, Delacourte A, Aprelikova O, Lee HG, Zhu X, et al. BRCA1 may modulate neuronal cell cycle re-entry in Alzheimer disease. *Int J Med Sci*. 2007;4:140–5.
35. Nakanishi A, Minami A, Kitagishi Y, Ogura Y, Matsuda S. BRCA1 and p53 tumor suppressor molecules in Alzheimer's disease. *Int J Mol Sci*. 2015;16:2879–92.
36. Jeon GS, Kim KY, Hwang YJ, Jung MK, An S, Ouchi M, et al. Deregulation of BRCA1 leads to impaired spatiotemporal dynamics of gamma-H2AX and DNA damage responses in Huntington's disease. *Mol Neurobiol*. 2012;45:550–63.
37. Brain L, Croft PB, Wilkinson M. Motor neurone disease as a manifestation of neoplasm (with a note on the course of classical motor neurone disease). *Brain*. 1965;88:479–500.
38. Chio A, Brignolio F, Meineri P, Rosso MG, Tribolo A, Schiffer D. Motor neuron disease and malignancies: results of a population-based study. *J Neurol*. 1988;235:374–5.
39. Forsyth PA, Dalmau J, Graus F, Cwik V, Rosenblum MK, Posner JB. Motor neuron syndromes in cancer patients. *Ann Neurol*. 1997;41:722–30.
40. Rojas-Marcos I, Rousseau A, Keime-Guibert F, Rene R, Cartalat-Carel S, Delattre JY, et al. Spectrum of paraneoplastic neurologic disorders in women with breast and gynecologic cancer. *Medicine*. 2003;82:216–23.
41. Sadot E, Carlier L, Corcia P, Delozier Y, Levy C, Viader F. Breast cancer and motor neuron disease: clinical study of seven cases. *Amyotroph Lateral Scler*. 2007;8:288–91.
42. Berghs S, Ferracci F, Maksimova E, Gleason S, Leszczynski N, Butler M, et al. Autoimmunity to beta IV spectrin in paraneoplastic lower motor neuron syndrome. *Proc Natl Acad Sci U S A*. 2001;98:6945–50.
43. Evans BK, Fagan C, Arnold T, Dropcho EJ, Oh SJ. Paraneoplastic motor neuron disease and renal cell carcinoma: improvement after nephrectomy. *Neurology*. 1990;40:960–2.
44. Forman D, Rae-Grant AD, Matchett SC, Cowen JS. A reversible cause of hypercapnic respiratory failure: lower motor neuropathy associated with renal cell carcinoma. *Chest*. 1999;115:899–901.
45. Kiewe P, Gueller S, Komor M, Stroux A, Thiel E, Hofmann WK. Prediction of qualitative outcome of oligonucleotide microarray hybridization by measurement of RNA integrity using the 2100 Bioanalyzer capillary electrophoresis system. *Ann Hematol*. 2009;88:1177–83.
46. Hubbell E, Liu WM, Mei R. Robust estimators for expression analysis. *Bioinformatics*. 2002;18:1585–92.
47. 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.
48. Noristani HN, Olabarria M, Verkhatsky A, Rodriguez JJ. Serotonin fibre sprouting and increase in serotonin transporter immunoreactivity in the CA1 area of hippocampus in a triple transgenic mouse model of Alzheimer's disease. *Eur J Neurosci*. 2010;32:71–9.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at
www.biomedcentral.com/submit

