Ikaros mediates the DNA methylation-independent silencing of MCJ/DNAJC15 gene expression in macrophages

Nicolás Navasa1,2, Itziar Martin-Ruiz2, Estíbaliz Atondo2, James D. Sutherland2, Miguel Angel Pascual-Itoiz2, Ana Carreras-González2, Hooman Izadi1, Julen Tomás-Cortázar2, Furkan Ayaz3, Natalia Martin-Martin2, Ivana M. Torres1,2, Rosa Barrio2, Arkaitz Carracedo2,3,4, E. R. Olivera5, Mercedes Rincón6 & Juan Anguita1,2,3

MCJ (DNAJC15) is a mitochondrial protein that regulates the mitochondrial metabolic status of macrophages and their response to inflammatory stimuli. CpG island methylation in cancer cells constitutes the only mechanism identified for the regulation of MCJ gene expression. However, whether DNA methylation or transcriptional regulation mechanisms are involved in the physiological control of this gene expression in non-tumor cells remains unknown. We now demonstrate a mechanism of regulation of MCJ expression that is independent of DNA methylation. IFN-γ, a protective cytokine against cardiac inflammation during Lyme borreliosis, represses MCJ transcription in macrophages. The transcriptional regulator, Ikaros, binds to the MCJ promoter in a Casein kinase II-dependent manner, and mediates the repression of MCJ expression. These results identify the MCJ gene as a transcriptional target of IFN-γ and provide evidence of the dynamic adaptation of normal tissues to changes in the environment as a way to adapt metabolically to new conditions.

MCJ (Methylation-Controlled J protein), also known as DNAJC15, is a small protein (147 aa) that contains a highly conserved 70 aa J domain at the C-terminus, an unusual transmembrane domain, and an N-terminal region with no homology to any other known protein1–3. The MCJ gene originated as a gene-duplication from the related gene DnaJC19, already present in flies2. MCJ is located in the inner mitochondrial membrane where it interacts with Complex I of the electron transport chain (ETC), interfering with the formation of supercomplexes composed of complexes I, III and IV4,5. MCJ is the first described endogenous negative regulator of Complex I that has also been associated with the TIM23 translocase and the import of pre-proteins to the mitochondria4. Silencing MCJ expression does not affect cell survival or proliferation5. However, loss of MCJ results in augmented mitochondrial membrane potential, increased oxidative respiration and mitochondrial ATP5. Although MCJ deficiency has no harmful effects under physiological conditions, increased mitochondrial metabolism in the absence of MCJ in vivo prevents the pathological accumulation of lipids in the liver during starvation or high cholesterol diet, and the development of liver steatosis5. MCJ is thus a modulator of mitochondrial

1Department of Veterinary and Animal Sciences, University of Massachusetts Amherst, Amherst, MA 01003. 2CIC bioGUNE, 48160 Derio, Bizkaia, Spain. 3Ikerbasque, Basque Foundation for Science. 4Biochemistry and Molecular Biology Department, University of the Basque Country (UPV/EHU), P. O. Box 644, E-48080 Bilbao, Spain. 5Department of Molecular Biology, Veterinary School, University of León, 24071 León, Spain. 6Department of Medicine, University of Vermont College of Medicine, Burlington, VT 05405. †Present address: Department of Microbiology and Immunology, Dartmouth School of Medicine, Hanover, NH 03755. Correspondence and requests for materials should be addressed to J.A. (email: janguita@cicbiogune.es)
metabolism that acts as a break to attenuate mitochondrial metabolism during adaptation to metabolic stress conditions.

MCJ was initially identified as a gene expressed in some but not all ovarian cancer cell lines and primary ovarian cancer tumors. MCJ is expressed in breast and uterine cancer cells that are sensitive to different chemotherapeutic drugs, but not in those that are multidrug resistant. In normal human and murine tissues, MCJ is highly expressed in heart, liver and kidney and within the immune system, in CD8+ T cells and macrophages. DNA methylation constitutes the only mechanism associated with the regulation of MCJ expression. In ovarian cancer cells, the presence of high levels of CpG island methylation within the first exon of the MCJ gene is associated with loss of expression and correlates with a diminished response to chemotherapy and poor survival. However, the mechanisms that regulate MCJ expression in normal tissues and cells are not known.

We have shown that MCJ modulates macrophage responses to a variety of proinflammatory insults. Short-term induction of inflammation by infection with Staphylococcus aureus or injection with LPS prevented TNF production in vivo and the development of acute fulminant hepatitis in mice in the absence of MCJ. MCJ is therefore, a potential therapeutic target under conditions of persistent inflammation.

Results and Discussion

Loss of MCJ expression in heart-infiltrating macrophages during infection with B. burgdorferi. During short-term in vivo inflammatory conditions, MCJ regulates the response of macrophages to Staphylococcus aureus as well as LPS treatment in mice sensitized with galactosamine. In order to determine the role of MCJ on the local macrophage response during an infectious process that requires a more complex and long lasting interaction between the pathogen and the host, we infected MCJ KO and WT mice with Borrelia burgdorferi. After 3 weeks of infection, macrophage infiltration was not significantly different in infected MCJ KO mice and WT animals (Fig. 1A). In addition, the amount of TNF expressed in the cardiac tissue upon infection was not altered in the absence or presence of MCJ (Fig. 1B). We also assessed the level of expression of MCJ in heart-infiltrating macrophages at the peak of infection with the spirochete. Surprisingly, in contrast to bone marrow-derived macrophages (BMMs), real time RT-PCR failed to detect appreciable levels of MCJ mRNA in macrophages infiltrating the hearts (Fig. 1C). The downregulation of MCJ expression during infection was selective of macrophages since total heart MCJ expression levels were readily detected in the infected mice (Fig. 1C). The histological analysis of infected joint and heart tissue showed that the degree of cardiac inflammation was not affected by the lack of the MCJ gene (Fig. S1A,B). Furthermore, the levels of spirochetal DNA were similar in

Figure 1. Heart-infiltrating macrophages do not express MCJ upon infection with B. burgdorferi.

The base of the hearts of 3 week infected and uninfected (UI) mice were used to extract RNA and assess macrophage infiltration and TNF expression levels by qRT-PCR using primers specific for F4/80 (A) or TNF (B). NS; Not significant. (C) Macrophages were purified from the hearts of 3-week infected B6 mice and used to extract RNA. qRT-PCR was then performed to detect MCJ mRNA levels, compared to bone marrow-derived macrophages (BMM). As a control, MCJ mRNA levels were also determined in whole heart tissue of 3-week infected mice. The data shown correspond to 5 mice per group and are presented as the mean ± SE. *; Student’s t test, p < 0.05.
WT and MCJ KO mice (Fig. S1C). These results suggested that upon infection with *B. burgdorferi*, MCJ expression is repressed specifically in macrophages infiltrating the heart.

**MCJ expression in macrophages is selectively downregulated by IFNγ.** In order to determine whether the interaction of macrophages with bacterial products results in reduced levels of MCJ, we stimulated RAW cells and BMMs with live *B. burgdorferi* and assessed the levels of MCJ. Stimulation with the spirochete did not affect MCJ protein (Fig. 2A) or mRNA (Fig. 2B) levels. LPS stimulation also failed to alter the levels of MCJ in macrophages (Fig. 2A). These data indicate that the regulation of the expression of MCJ occurs independently of pattern-recognition receptor (PRR) stimulation, including TLR4, TLR1/2 and other PRRs stimulated by the interaction of live *B. burgdorferi* with macrophages11–15. Since IFNγ is a major contributor to macrophage function during cardiac infection with *B. burgdorferi*16,17, we stimulated macrophages with IFNγ. Treatment with IFNγ resulted in lower levels of MCJ protein in both RAW cells and BMMs (Fig. 2C,D). Because MCJ is localized in mitochondria, we examined the effect of IFNγ on mitochondrial mass; however, no difference was observed as determined by levels of the mitochondrial protein, VDAC1 (Fig. 2E). The effect of IFNγ was selective of macrophages, because it did not affect MCJ levels in the murine tumor cell line, Hepa 1–6 or primary CD8+ T cells (Fig. 2C). IL-6 has been shown to downregulate MCJ levels in breast cancer cell lines1. Similarly, we found that IL-6 induced the downregulation of MCJ in Hepa liver cancer cells (Fig. 2F). However, IL-6 failed to downregulate MCJ expression in RAW cells or BMMs (Fig. 2F). These results show that MCJ expression in macrophages is selectively silenced by IFNγ.
IFN\(\gamma\) inhibits \(MCJ\) gene transcription independently of DNA methylation. To determine if the downregulation of \(MCJ\) protein levels by IFN\(\gamma\) in macrophages was due to an effect on \(MCJ\) gene expression, we assessed \(MCJ\) mRNA levels in macrophages stimulated with IFN\(\gamma\). The treatment with IFN\(\gamma\) resulted in a significant decrease in \(MCJ\) mRNA levels in RAW cells and BMMs (Fig. 3A). No previous studies have characterized the human or mouse \(MCJ\) gene promoter region and addressed transcriptional regulation. We identified a 1 kb region upstream of the start initiation site of the murine \(MCJ\) gene (Fig. S2A), that was capable to mediate high levels of transcription in RAW cells in luciferase reporter assays (Fig. 3B). Treatment with IFN\(\gamma\) caused a pronounced decrease in the transcriptional activity of this region of the \(MCJ\) promoter (Fig. 3B).

Figure 3. IFN\(\gamma\) represses \(MCJ\) gene expression independently of DNA methylation. (A) RAW cells and BMMs were stimulated for 20 h with IFN\(\gamma\) and analyzed by qRT-PCR for \(MCJ\) mRNA levels. The results correspond to the average of 3 independent experiments. *, Student’s t test, \(p < 0.05\). (B) RAW cells were co-transfected with plasmids containing the luciferase gene under the influence of the 1 kb proximal promoter region of the \(MCJ\) gene or the Renilla luciferase gene under the influence of the SV40 promoter. After 4 h, the cells were stimulated with 100 ng/mL of IFN\(\gamma\) or left unstimulated. Dual luciferase activity was assessed after 16 h of incubation. The promoterless vector, pGL3 was used as a control. *, Student’s t test, \(p < 0.05\). (C) BMMs were left unstimulated or stimulated with 100 ng/mL of IFN\(\gamma\) in the absence or presence of 1 \(\mu\)M of decitabine (DEC) or Azacitidine (Aza). After 48 h, the cells were tested by Western blotting for the presence of \(MCJ\). GAPDH levels were determined to ensure equal loading. (D) CpG-rich region in the \(MCJ\) gene analyzed by bisulfite sequencing. The primers used for amplification are noted in lower case. The percentage of methylated CpG residues in BMMs stimulated with 100 ng/mL of IFN\(\gamma\) or left untreated is marked in each of the 6 CpG residues. Black circles indicate 100% of the samples contained these residues methylated, while white circles represent 0%. The analysis corresponds to BMMs isolated from 6 mice. (E) CHIP analysis of BMM DNA immunoprecipitated with antibodies against the H3 marks corresponding to trimethylation of Lys 4 (H3K4m3) and 27 (H3K27m3) or H3 pan-acetylation (Pan Ac-H3). The binding levels are relative to total H3. The results correspond to the average \pm SE of 3 independent experiments.
effect of IFN-γ treatment on histone H3 marks associated with the activation and repression of gene expression (ref. 31). The treatment of BMMs with IFN-γ did not affect the binding of trimethylated H3 at Lys 4 and Lys 27 or acetylated H3 to the MCJ promoter (Fig. 3E). These data revealed that transcriptional regulation is an alternative mechanism of MCJ expression modulation that is independent of DNA methylation or alteration on histone marks.

**Ikaros is an inducible repressor of MCJ gene transcription.** To identify the specific mechanism by which IFN-γ represses MCJ gene transcription, we performed a search for potential transcription factor binding sites within the 1kb region of the mouse MCJ gene promoter using the tool TFSearch19. Two consensus binding sites for Ikaros (−350 to −361 and −706 to −717) were identified (Fig. S2A). Ikaros is known to act primarily as a repressor of gene expression20. To demonstrate whether Ikaros binds to these putative binding sites in the MCJ promoter and address whether binding was regulated by IFN-γ, we performed chromatin immunoprecipitation (ChIP) assays in BMMs. Binding of Ikaros to both sites was almost undetectable in untreated BMMs (Fig. 4A). However, Ikaros binding to Site 1 (the most proximal
to the transcription start site; Fig. S2A) was highly induced in cells treated with IFN-γ (Fig. 4A). Ikaros binding to Site 2 (Fig. S2A), however, was not induced by IFN-γ (Fig. 4A).

To further analyze the contribution of both putative Ikaros binding sites to the regulation of MCJ gene expression, we generated deletion mutants of both binding sites, as well as a double deletion mutant lacking both binding sites. The deletion of Site 1 resulted in significantly increased transcriptional activation in reporter assays (Fig. 4B) suggesting that this site is bound under basal conditions to a negative gene expression regulator. However, the deletion of Site 2 did not affect the expression activity of the promoter (Fig. 4B). Furthermore, the deletion of both sites resulted in transcriptional activity that was equivalent to that observed with the deletion of Site 1 (Fig. 4B). Overall, these data suggest that Ikaros binds to Site 1 upon induction with IFN-γ and displaces a weaker negative regulator of gene expression.

To demonstrate the role of Ikaros in the IFN-γ-dependent repression of MCJ gene expression, we generated stable lentiviral transductants containing a short hairpin sequence (shRNA) specific for the IKFZ1 gene encoding Ikaros. Transduction with shIKFZ1 in RAW cells caused a prominent reduction of Ikaros levels (Fig. 4C). Importantly, while IFN-γ downregulated MCJ levels in control cells, it did not affect MCJ levels in shIKFZ1-transduced cells (Fig. 4D). These results demonstrate that silencing of MCJ expression by IFN-γ is mediated by Ikaros and reveal this repressor as a key factor in the alternative mechanism regulating MCJ expression.

We then assessed whether IFN-γ upregulates Ikaros expression. The stimulation with IFN-γ did not affect Ikaros levels in macrophages (Fig. 4E), suggesting that the increased binding to the MCJ promoter region was due to post-translational modifications induced by IFN-γ. Ikaros activity is regulated by phosphorylation mediated by Casein Kinase 2 (CK2)21. It has also been reported that IFN-γ regulates the expression of a subset of genes through the activation of CK222,23. To investigate whether IFN-γ promotes Ikaros binding to Site 2 of the MCJ promoter in the presence of a CK2 specific inhibitor, 4,5,6,7-tetrabromobenzotriazole (TBB)24. CHIP analysis demonstrated that the pretreatment with TBB abrogated IFN-γ-induced binding of Ikaros to the MCJ promoter (Fig. 4F).

We then investigated whether silencing of MCJ expression by IFN-γ was mediated by CK2. Pretreatment of macrophages with the CK2 inhibitor prevented downregulation of MCJ expression by IFN-γ (Fig. 4G). In addition, inhibition of CK2 also prevented IFN-γ from suppressing MCJ promoter transcriptional activity (Fig. 4H). Together, these results show that IFN-γ represses MCJ gene transcription in macrophages by promoting CK2-dependent DNA binding of Ikaros to the proximal region of the MCJ promoter.

Our studies identify a novel mechanism of regulation of MCJ gene expression that is independent of the well-established DNA methylation pathway described in several tumors. MCJ/DnaJC15 is emerging as an important regulator of mitochondrial activity and cellular function in vitro and in vivo5,7. Therefore, the control of MCJ transcription constitutes a mechanism to regulate cellular responses to environmental changes. As opposed to DNA methylation, which is considered a long-term mechanism to silence gene expression24, the transcriptional control of MCJ gene expression by Ikaros may allow normal tissues to adapt dynamically to a changing environment. Here, we demonstrate that Ikaros represses MCJ expression in response to IFN-γ in macrophages. Similar mechanisms could be used to alter MCJ levels in other cells or tissues in response to changes in the environment as a way to adapt metabolically to new conditions.

Our results identify MCJ gene expression as a transcriptional target of the cytokine IFN-γ, contributing to the regulation of their inflammatory output7. Our data also reinforces the role of IFN-γ as a cytokine that exerts a protective effect during infection with B. burgdorferi26,17,25. Overall, we hypothesize that the combined effect of IFN-γ, including the regulation of MCJ expression, results in a more efficient elimination of the bacteria from the infected tissue without a concomitant increase in the inflammatory damage.

**Methods**

**Mice.** MCJ-deficient mice in a C57Bl/6 (B6) background and wild type B6 mice were bred at UMass Amherst and CIC bioGUNE. The Institutional Animal Care and Use Committees at UMass Amherst and CIC bioGUNE approved all procedures involving animals.

**Infections.** Groups of WT and KO mice were infected by subcutaneous injection with 10⁷ Borrelia burgdorferi 297 in the midline of the back. The mice were sacrificed after 3 weeks of infection and analyzed for inflammatory symptoms in joints and hearts stained with hematoxilin and eosin. Signs of arthritis and carditis were determined blindly as described26. The number of spirochetes in heart tissue was determined by real-time PCR, using primers specific for the recA gene (Table S1) standardized to μg of total DNA with primers corresponding to Glyceraldehyde 3-Phosphate Dehydrogenase, GAPDH, (Table S1)27.

**Cells.** Infiltrating cardiac macrophages were isolated from 3-week infected B6 mice. Hearts were perfused with cold Hank’s balanced salt solution (HBSS, Lonza, Anaheim, CA) and cut into small pieces, followed by digestion with 1 mg/mL of collagenase/dispose (Roche) and homogenization in a Dounce homogenizer. The digest was passed through a 16” gauge syringe to obtain single cell suspensions. The cellular suspension was layered on top of a 3 mL layer of Ficoll (GE Healthcare, Piscataway, NJ) and centrifuged at 400 × g for 40 min without brakes. Monocytes were then purified from the interphasic
cellular fraction using a one-step discontinuous Percoll gradient (46%) under isosmotic conditions. Monocytes were used for RNA extraction.

Bone marrow-derived macrophages were generated as described using 30 ng/mL of M-CSF (Miltenyi Biotec, Bergisch Gladbach, GE). Macrophages were allowed to differentiate in 100 mm x 15 mm petri dishes (Fisher Scientific, Pittsburgh, PA) for 8 days. Non-adherent cells were then eliminated and adherent macrophages were scraped, counted and resuspended in serum-free RPMI medium 2 h prior to use.

CD8+ T cells were purified by positive selection from the spleens of B6 mice using biotinylated anti-CD8 (BD Biosciences, San Diego, CA), anti-biotin microbeads and the MACS system (Miltenyi Biotec, Auburn, CA).

Lentiviral particles containing shRNA targeting Ikaros (Ikzf1 gene, Sigma Chemical Co, St. Louis, MO) were produced as described. Supernatants containing the virus were used to infect RAW 264.7 cells, followed by incubation with puromycin at 2 μg/mL to generate stable lines. Cells containing the empty vector, pLK0.1, were used as a control.

In vitro stimulation. Cells were incubated with 100 ng/mL of murine IFNγ or human IL-6 for the indicated time periods. In some instances, the following inhibitors were used 1 h prior to stimulation: decitabine (DEC, 1 μM), 4,5,6,7-tetramethylumbelliferylazol (TBB, 1 μM; Tocris Bioscience, Bristol, UK), 5-Aza-2-Deoxyctydine (Aza, 1 μM; Sigma Chemical Co.). Stimulation with B. burgdorferi (m.o.i. = 25) or LPS (100 ng/mL) were performed for 4–6 h.

Real-time RT-PCR. RNA from isolated cells or cardiac tissue was extracted by the thioisocyanate method (Amresco, Solon, OH), treated with DNase I (Qiagen), and reverse transcribed using the SuperScript VILO cDNA synthesis kit (Life Technologies). Real-time PCR was then performed using SYBR Green PCR Master Mix (Life Technologies) on a BioRad CFX96 Real-Time System (Bio-Rad, Hercules, CA). Fold induction of the genes was calculated relative to actin, using the 2^(-ΔΔCt) method. The primers used are listed in Table S1.

Western blot. Five to 20 μg of protein were run on SDS-PAGE, transferred to nitrocellulose membranes and treated with antibodies specific for MCJ, VDAC1 (D-16) and Ikaros (M-20, Santa Cruz Biotechnology, Dallas, TX). Equal loading was determined using antibodies against GAPDH (6C5) or actin (1-19) from Santa Cruz Biotechnology.

Epifluorescence (Apotome) microscopy. Cells were grown in 8-well chamber slides (Nunc Thermo Scientific, Waltham, MA). Upon incubation with 100 ng/mL of IFNγ (eBioScience, San Diego, CA) for 3 days, the cells were processed as described using anti-MCJ Abs, followed by an anti-rabbit IgG conjugated to Alexa Fluor 594.

Cloning of the proximal 1 kb MCJ promoter and luciferase assays. The proximal 1 kb promoter of the murine MCJ gene was cloned into pGL3 using the primers in Table S1. Deletion mutants corresponding to the putative Ikaros binding sites of the MCJ promoter (Fig. S2) were generated using the QuickChange Site-Directed Mutagenesis kit (Stratagene, La Jolla, CA) and the primers listed in Table S1. 1.9 μg of these constructs plus 0.1 μg of pSVL40 plasmid were cotransfected into RAW cells using the X-TremeGene HP DNA tranfection reagent (Roche). After 6 h, the cells were treated with IFNγ in the presence or absence of the specific inhibitor, TBB. After 20 h incubation, the cells were lysed in lysis buffer (Promega, Madison, WI) and Firefly and Renilla luciferase activities were determined by the Dual Luciferase reporter system (Promega).

Bisulfite sequencing. DNA was extracted from BMMs treated with IFNγ and controls, denatured and subjected to bisulphite conversion as described by Meissner and colleagues. The resultant product was PCR amplified using the primers in Table S1, corresponding to the region in the MCJ gene described by Meissner and colleagues.

Chromatin immunoprecipitation. Fifteen million BMMs were stimulated with 100 ng/mL of IFNγ in the presence or absence of TBB for 16 h. CHIP assays were performed using the SimpleChip Enzymatic Chromatin IP kit-Magnetic beads (Cell Signaling, Beverly, MA) following the manufacturer’s instructions using anti-Ikaros, anti-H3K4m3, anti-H3K27m3, anti-pan acetylated H3 antibodies and anti-H3 (Cell Signaling) or normal rabbit IgG as negative control. The immunoprecipitated DNA was subjected to q-PCR using primers encompassing the two putative Ikaros binding sites (Table S1). The results are presented as fold induction over rabbit IgG immunoprecipitates or total H3 relative to input (percent input method), following the formula: \[ \frac{100 \times 2^{-\Delta\Delta Ct}}{100 \times 2^{-\Delta\Delta Ct}} = 2^{-\Delta Ct_{\text{IgG}} - Ct_{\text{IgG}}} \], where AdjInput = Adjusted input to 100%; CtTEST = Ct of test samples; CtIgG = Ct of samples control.

Statistical Analysis. Results are presented as means ± SE. Significant differences between means were calculated with the Student’s t test. P values of 0.05 or less were considered statistically significant.
References

1. Lindsey, J. C. et al. Epigenetic inactivation of MCJ (DNAJ1) in malignant paediatric brain tumours. *Int J Cancer* **118**, 346–352 (2006).

2. Hatle, K. M. et al. Methylation-controlled J protein promotes c-Jun degradation to prevent ABCB1 transporter expression. *Mol Cell Biol* **27**, 2952–2966 (2007).

3. Schusdziarra, C., Blaminska, M., Azem, A. & Hell, K. Methylation-controlled J-protein MCJ acts in the import of proteins into human mitochondria. *Human molecular genetics* **22**, 1348–1357 (2013).

4. Roy, A. et al. Mitochondria-dependent reactive oxygen species-mediated programmed cell death induced by 3,3′-diiodomethane through inhibition of F0F1-ATP synthase in unicellular protozoan parasite *Leishmania donovani*. *Mol Pharmacol* **74**, 1292–1307 (2008).

5. Hatle, K. M. et al. *MCJ/DnaJC15*, an endogenous mitochondrial repressor of the respiratory chain that controls metabolic alterations. *Mol Cell Biol* **33**, 2302–2314 (2013).

6. Shridhar, V. et al. Loss of expression of a new member of the DNAJ protein family confers resistance to chemotherapeutic agents used in the treatment of ovarian cancer. *Cancer Res* **61**, 4238–4265 (2001).

7. Navasa, N. et al. Regulation of oxidative stress by methylation-controlled J protein controls macrophage responses to inflammatory insults. *J Infect Dis* **211**, 135–145 (2015).

8. Strathdee, G. et al. Demethylation of the MCJ gene in stage III/IV epithelial ovarian cancer and response to chemotherapy. *Gynecol Oncol* **97**, 898–903 (2005).

9. Strathdee, G., Davies, B. R., Vass, J. K., Siddiqui, N. & Brown, R. Cell type-specific methylation of an intronic CpG island controls expression of the MCJ gene. *Cancerogene* **25**, 693–701 (2004).

10. Ehrlich, M. et al. Hypomethylation and hypermethylation of DNA in Wilms tumors. *Oncogene* **21**, 6694–6702 (2002).

11. Alexopoulou, L. et al. Hyporesponsiveness to vaccination with *Borrelia burgdorferi* OspA in humans and in TLR1- and TLR2-deficient mice. *Nature Medicine* **8**, 878–884 (2002).

12. Behera, A. K. et al. *Borrelia burgdorferi* BBB07 interaction with integrin alpha3beta1 stimulates production of pro-inflammatory mediators in primary human chondrocytes. *Cell Microbiol* **10**, 320–331 (2008).

13. Behera, A. K. et al. Identification of a TLR-independent pathway for *Borrelia burgdorferi*-induced expression of matrix metalloproteinases and inflammatory mediators through binding to integrin alpha 3 beta 1. *J Immunol* **177**, 657–664 (2006).

14. Cervantes, J. L. et al. Phagosomal signaling by *Borrelia burgdorferi* in human monocytes involves Toll-like receptor (TLR) 2 and TLR8 cooperativity and TLR8-mediated induction of IFN-β. *Proc Natl Acad Sci USA* **108**, 3683–3688 (2011).

15. Pétzke, M. M., Brooks, A., Krupna, M. A., Mordue, D. & Schwartz, I. Recognition of *Borrelia burgdorferi*, the Lyme disease spirochete, by TLR7 and TLR9 induces a type I IFN response by human immune cells. *J Immunol* **183**, 5279–5292 (2009).

16. Brown, C. R., Blaho, V. A., Fritsche, K. L. & Lotiacono, C. M. Stat6 deficiency exacerbates carditis but not arthritis during experimental Lyme borreliosis. *J Interferon Cytokine Res.* **26**, 390–399 (2006).

17. Olson, C. et al. Local Production of IFN-gamma by Invariant NKT Cells Modulates Acute Lyme Carditis. *Journal of Immunology* **182**, 3728–3734 (2009).

18. Meissner, A. et al. Genome-scale DNA methylation maps of pluripotent and differentiated cells. *Nature* **454**, 766–770 (2008).

19. Roy, A. C., Wang, M. & Whitesay, M. The ERK/MAPK pathway regulates the activity of the human tissue factor pathway inhibitor-2 promoter. *J Biol Chem* **278**, 6787–6794 (2003).

20. John, L. B. & Ward, A. C. The Ikaros gene family: transcriptional regulators of hematopoiesis and immunity. *Mol Immunol* **48**, 1272–1278 (2011).

21. Cho, S. J., Huh, J. E., Song, J., Rhee, D. K. & Pyo, S. Ikaros negatively regulates inducible nitric oxide synthase expression in macrophages: involvement of Ikaros phosphorylation by casein kinase 2. *Cell Mol Life Sci* **65**, 3290–3303 (2008).

22. Harris, S. M., Harvey, E. J., Hughes, T. R. & Ramji, D. P. The interferon-gamma-mediated inhibition of lipoprotein lipase gene transcription in macrophages involves casein kinase 2- and phosphoinositide-3-kinase-mediated regulation of transcription factors Sp1 and Sp3. *Cell Signal* **20**, 2296–2301 (2008).

23. Quisner, F. et al. Protein kinase CK2 inhibition induces cell death via early impact on mitochondrial function. *J Cell Biochem* **115**, 2103–2115 (2014).

24. Raynal, N. J. et al. DNA methylation does not stably gene expression but instead serves as a molecular mark for gene silencing memory. *Cancer Res* **72**, 1170–1181 (2012).

25. Bockenstedt, L. K. et al. CD4+ T helper 1 cells facilitate regression of murine Lyme carditis. *Infect Immun* **69**, 5264–5269 (2001).

26. Hawley, K. et al. Macrophage p38 Mitogen-Activated Protein Kinase Activity Regulates Invariant Natural Killer T-Cell Responses During *Borrelia burgdorferi* Infection. *Journal of Infectious Diseases* **206**, 283–291 (2012).

27. Fincato, G., Polentarutti, N., Sica, A., Mantovani, A. & Colotta, F. Expression of a heat-inducible gene of the HSP70 family in human myelomonocytic cells: regulation by bacterial products and cytokines. *Blood* **97**, 579–586 (1991).

28. Garcia-Cao, I. et al. Systemic elevation of PTEN induces a tumor-suppressive metabolic state. *Cell* **149**, 49–62 (2012).

29. Hawley, K. L. et al. CD14 cooperates with complement receptor 3 to mediate MyD88-independent phagocytosis of *Borrelia burgdorferi*. *Proceedings of the National Academy of Sciences of the United States of America* **109**, 1228–1232 (2012).

30. Clark, S. J., Statham, A., Stirzaker, C., Molloy, P. L. & Frommer, M. DNA methylation: bisulphite modification and analysis. *Nat Protoc* **1**, 2353–2364 (2006).

31. LB Rashkov et al. Epigenetic regulation of macrophage polarization and function. *Trends in Immunology* **34**: 216–222.

Acknowledgements

We thank J.M Iglesias-Pedraz for technical support. Supported by grants from the National Institutes of Health (AI-078277 to JA, and CA-127099 to MR), the Spanish Ministry of Economy Plan Nacional (SAF2012-34610 to JA, BFU2011-25986 to RB and JDS), Basque Government Depts. of Industry, Tourism and Trade (Etorreko to AC), Health (2012111086 to AC) and Education (P12012-03 to AC, and P12012/42 to RB and JDS), Instituto de Salud Carlos III (PI10/01484 and PI13/00031 to AC), Marie Curie (277043 to AC), and ERC (336343 to AC). AC is supported by a Ramon y Cajal Award. N.M.M. is supported by the Spanish Association Against Cancer (AEC). 

Author Contributions

N.N., I.M.M., E.A., J.D.S., M.A.P., A.C.G., H.I., J.T.C., F.A. and I.M.T. performed the experiments. N.M.M. and A.C. helped with lentiviral production and transduction experiments. R.B., A.C., E.R.O. and
M.R. provided reagents and helped in the analysis of the data. N.N., M.R. and J.A. wrote the manuscript. J.A. designed the study.

**Additional Information**

**Supplementary information** accompanies this paper at http://www.nature.com/srep

**Competing financial interests:** The authors declare no competing financial interests.

**How to cite this article:** Navasa, N. et al. Ikaros mediates the DNA methylation-independent silencing of MCJ/DNAJC15 gene expression in macrophages. *Sci. Rep.* 5, 14692; doi: 10.1038/srep14692 (2015).

This work is licensed under a Creative Commons Attribution 4.0 International License. The images or other third party material in this article are included in the article’s Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit http://creativecommons.org/licenses/by/4.0/