**Cellular/Molecular**

**Microglial Expression of the B7 Family Member B7 Homolog 1 Confers Strong Immune Inhibition: Implications for Immune Responses and Autoimmunity in the CNS**

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Inflammation of the CNS is usually locally limited to avoid devastating consequences. Critical players involved in this immune regulatory process are the resident immune cells of the brain, the microglia. Interactions between the growing family of B7 costimulatory ligands and their receptors are increasingly recognized as important pathways for costimulation and/or inhibition of immune responses.

Human and mouse microglial cells constitutively express B7 homolog 1 (B7-H1) in vitro. However, under inflammatory conditions [presence of interferon-γ (IFN-γ) or T-helper 1 supernatants], a significant upregulation of B7-H1 was detectable. Expression levels of B7-H1 protein on microglial cells were substantially higher compared with astrocytes or splenocytes. Coculture experiments of major histocompatibility complex class II-positive antigen-presenting cells (APC) with syngeneic T cells in the presence of antigen demonstrated the functional consequences of B7-H1 expression on T-cell activation. In the presence of a neutralizing anti-B7-H1 antibody, both the production of inflammatory cytokines (IFN-γ and interleukin-2) and the upregulation of activation markers (inducible costimulatory signal) by T cells were markedly enhanced. Interestingly, this effect was clearly more pronounced when microglial cells were used as APC, compared with astrocytes or splenocytes. Furthermore, B7-H1 was highly upregulated during the course of myelin oligodendrocyte glycoprotein-induced and proteolipid protein-induced experimental allergic encephalomyelitis in vivo. Expression was predominantly localized to areas of strongest inflammation and could be colocalized with microglial cells/macrophages as well as T cells.

Together, our data propose microglial B7-H1 as an important immune inhibitory molecule capable of downregulating T-cell activation in the CNS and thus confining immunopathological damage.

**Key words:** encephalomyelitis; glia; immunity; microglia; neuropathology; tolerance; costimulation

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**Introduction**

Antigen presentation in the CNS is thought to play a critical role in the initiation and perpetuation of neuroinflammation. Microglial cells, the resident immune cells, can express major histocompatibility complex (MHC) class II molecules and have the potential to act as effective antigen-presenting cells (APCs) to T cells (Sedgwick et al., 1993). In multiple sclerosis (MS), the prototypic autoimmune inflammatory disorder of the CNS, close interactions of MHC class II-positive microglial cells with T cells are visible within the inflammatory lesions (Bo et al., 1994).

A necessary prerequisite for T-cell activation is T-cell receptor engagement by MHC peptides (signal 1) together with a second antigen-independent signal mediated by costimulatory receptors (signal 2) (Lafferty and Woolnough, 1977). Accumulating evidence suggests that during CNS inflammation, activated microglia express costimulatory molecules such as CD80 (B7.1), CD86 (B7.2), and CD40 in addition to certain adhesion molecules (for review, see Aloisi, 2001). The interaction of microglial cells and T cells subsequently leads to either induction of T-cell proliferation, T-effector functions (e.g., cytokine secretion), or both (Becher et al., 2000) (for review, see Aloisi, 2001). The modulation of T-cell function by microglial cells entails an important dichotomy. Microglial cells are capable of secreting high amounts of proinflammatory cytokines [e.g., interleukin-1 (IL-1), tumor necrosis factor-α (TNF-α), and IL-12] during CNS inflammation. In contrast, they can also participate in the downmodulation of inflammation by producing anti-inflammatory cytokines.
such as TGF-β, IL-10, or IL-1 receptor antagonist (Kreutzberg, 1996). Along this notion, different model systems involving an inflamed CNS environment have characterized microglia as potent inducers of T-cell cytokine production, whereas their capacity to induce T-cell proliferation is much less evident (Carson et al., 1999; Juedes and Ruddle, 2001). The discovery of co-inhibitory molecules of the B7-CD28 family has brought tremendous advancements in understanding the control of T-cell immunity in different immune compartments (Chen, 2004). Programmed death-1 ligand (PD-L1) or B7 homolog 1 (B7-H1) has been attributed to costimulatory and immune regulatory functions (Kobata et al., 2000; Coyle and Gutierrez-Ramos, 2001; Carreno and Collins, 2002; Liang and Sha, 2002; Sharpe and Freeman, 2002). Previous work described that the costimulation of T cells with B7-H1-Fc fusion protein induces T-cell proliferation and the secretion of IL-10 and interferon-γ (IFN-γ) (Dong et al., 1999). However, more recent data suggest that B7-H1 may also negatively regulate T cells by inhibiting cell cycle progression (Freeman et al., 2000; Carter et al., 2002). B7-H1 exhibits a broader tissue distribution than B7.1/2 (CD80/CD86) (Freeman et al., 2000; Epiphimer et al., 2002; Mazanet and Hughes, 2002; Petroff et al., 2002; Wiendl et al., 2003). B7-H1 is upregulated by IFN-γ and regulates immune responses by interacting with programmed death receptor-1 (PD-1) and another yet unidentified receptor on activated T cells, although not CD28/cytotoxic T-lymphocyte-associated antigen receptor 4 (Coyle and Gutierrez-Ramos, 2001; Carreno and Collins, 2002; Dong et al., 2002; Liang and Sha, 2002; Sharpe and Freeman, 2002; Wang et al., 2003). Interestingly, PD-1-deficient mice suffer from autoimmune disorders resulting from inappropriate activation of B and T cells (Nishimura et al., 1999, 2001), suggesting that PD-1 is an important inhibitory signal that acts to prevent uncontrolled proliferation of autoreactive T cells.

**Materials and Methods**

**Antibodies and reagents.** Myelin oligodendrocyte glycoprotein (MOG) peptide 35-MEVWGRSPSRVHHYLRNGK-55 was synthesized by MWG Biotech (Ebersberg, Germany). The proteolipid protein 139–151 (PLP139-151) (HCLGKWLGHPDKF) was synthesized using standard g-fluoromethoxybenzyl chemistry. Protein purified derivate (PDP) of *Mycobacterium tuberculosis* was purchased from Statens Serum Institut (Copenhagen, Denmark). Staphylococcal enterotoxin B (SEB) was provided by Sigma (Deisenhofen, Germany). The following primary antibodies (Abs) were used: anti-mouse B7-H1 (PD-L1), M1H5 (Bio-science, San Diego, CA), 10H5 (L. Chen, Mayo Clinic Rochester, Rochester, MN), BAF1019 (R&D Systems, Minneapolis, MN); anti-mouse inducible costimulatory signal (ICOS), HK5.3 (Ebioscience); anti-mouse MHC II, I-A/E (BD Pharmingen, Heidelberg, Germany); anti-mouse MAC3 (Pharmingen); anti-mouse CD45, anti-mouse CD11b, anti-mouse CD3 (all from BD Biosciences, Heidelberg, Germany); anti-mouse F4/80 (Serotec, Kidlington, UK); and anti-mouse GFAP (Dakocytomation, High Wycombe, UK). Secondary Abs are as follows: goat anti-mouse IgG (H+L) F(ab)2, PE; rat anti-mouse IgG F(ab)2, FITC (Dianova, Hamburg, Germany); anti-rabbit IgG F(ab)2, FITC (Dianova), and donkey anti-rat Cy3 (Jackson ImmunoResearch, West Grove, PA). Mouse IFN-γ and TNF-α were from Sigma. Mouse IL-2 and IFN-γ ELISA were from Pharmingen.

**Isolation of human microglial cells.** The studies were performed in accordance with the guidelines set by the Institutional Review Board of McGill University (Montreal, Canada). Primary adult human glial cells were obtained from surgical resections performed for the treatment of non-tumor-related intractable epilepsy. Tissue was obtained from regions requiring resection to reach the precise epileptic focus and was distant from the main electrically active site. Dissociated cultures of microglia were prepared as described previously (Williams et al., 1992), based on the differential adhesion of the glial cells. Briefly, brain tissue was subjected to enzymatic dissociation with trypsin (0.025%) and DNase I (25 µg/ml) (Boehringer Mannheim, Laval, Quebec, Canada) for 30 min at 37°C, followed by mechanical dissociation by passage through a 132 µm nylon mesh (Industrial Fabrics Corporation, Minneapolis, MN). Cells were further separated on a linear 30% Percoll density gradient (Amersham Biosciences, Baie D’Urfe, Quebec, Canada) and centrifuged at 15,000 rpm at 4°C for 30 min. The cells recovered from the interface contained a mixed glial cell population consisting of ~65% oligodendroglia, 30% microglia, and 5% astrocytes. To enrich for microglia, the mixed cell population was suspended in minimal essential culture medium supplemented with 5% FCS, 2.5 U/ml penicillin, 2.5 µg/ml streptomycin, 2 µM glutamine, and 0.1% glucose (all from Invitrogen, Burlington, Ontario, Canada), and left overnight in 12.5 cm² tissue culture flasks (Falcon; Fisher Scientific, Montreal, Quebec, Canada) in a humid atmosphere at 37°C with 5% CO₂. The less-adherent oligodendroglia were removed by gentle pipetting, and the remaining adherent cells were allowed to develop morphologically for 3 days. Remaining microglia were of 95% purity as assessed by immunocytochemistry and flow cytometry (Williams et al., 1992; Becher and Antel, 1996). Microglia were cultured for ~7 d and then harvested by trypsinization (0.25%). T-helper 1 (Th1) as well as Th2 supernatants from T cells were generated as described previously (Kim et al., 2004).

**Isolation of human monocytes.** For monocyte isolation, blood was obtained by venipuncture, and peripheral blood mononuclear cells were isolated by density gradient centrifugation using lymphocyte separating solution (PAA Laboratories, Linz, Austria). Monocytes were enriched by 1 h of adherence to plastic flasks at 37°C in RPMI 1640 (BioWhittaker, Verviers, Belgium) supplemented with 10% FCS (Biochrom, Berlin, Germany) and penicillin (100 international units (IU)/ml)/streptomycin (10 µg/ml) (Invitrogen). Nonadherent cells were removed. Adherent cells were detached using cell dissociation buffer, and purity was analyzed by flow cytometry (>90% CD14-positive cells). Where indicated, monocytes were stimulated by supernatants from Th1 or Th2 cells as described previously (Kim et al., 2004).

**Isolation of murine microglial cells and astrocytes.** Mouse microglial cells and astrocytes were isolated from primary mixed brain glial cell cultures using a modification of methods described previously (Giulian and Baker, 1986; Magnus et al., 2001, 2002). In brief, cultures were prepared from the brains of newborn C57BL/6 mice [postnatal day 0 (P0) toP2; Charles River, Sulzfeld, Germany], which were freed of their meninges and minced with scissors under a dissecting microscope (Wild, Heerbrugg, Switzerland). Mixed cell cultures were then grown in basal Eagle’s medium supplemented with 10% fetal calf serum (Sigma), 50 IU/ml penicillin, and 50 µg/ml streptomycin at 37°C for 10–14 d. Microglial cells and astrocytes were isolated by shaking the culture flasks for 7 h (Primaria; Falcon, Franklin Lakes, NJ). To check the purity of the cell-culture system, we took a small fraction and performed immunocytochemistry with the monoclonal antibodies (mAbs) F 4/80 (1:100) and GFAP (1:100) as described previously (Giulian and Baker, 1986; Zielasek et al., 1992). A total of 95% or more of the cells were either F 4/80 positive (microglial cells) or GFAP positive (astrocytes). All cells used for our studies came from the same primary glial cell preparation.

For the isolation of splenocytes, the spleen was isolated from syngeneic mice and then cell suspensions were generated and used in the experiments (Jung et al., 2001).

**Coculture experiments with T cells.** Vβ-specific activation of naïve T cells by SEB (5 µg/ml) was measured by coculturing naïve T cells with irradiated syngeneic APC. T cells were mixed with microglial cells or astrocytes in a ratio of 1:5 to achieve good levels of cytokine secretions by the T cells. In the experiments with splenocytes, the ratio was 1:1. The optimal ratio of microglial cells, astrocytes, or splenocytes and T cells for the maximal T-cell cytokine production had been titrated previously. Microglial cells or astrocytes alone produced no IFN-γ or IL-2, and T cells alone (with or without antigens) secreted rather small amounts of IFN-γ or IL-2. A 10- to 100-fold increase in IFN-γ or IL-2 production could be detected after antigen had been added to the cocultures (data not shown). Supernatants were collected at 24 h, centrifuged to remove particulate debris, and stored in aliquots at −70°C.
For the antigen-specific T-cell experiments, T-cell lines were established from inguinal lymph nodes (LNs) or spleens of immunized mice according to standard procedures (Korn et al., 2003). T-cell lines were restimulated with their respective antigen at least two times before the experiments. For in vitro recall, inguinal LN cells or splenocytes were isolated on day 12 after injection and seeded at 75,000 or 150,000 per microtiter-well in restimulation medium. The T-cell lines were CD4 positive and clearly MHC II restricted. PPD (10 μg/ml) for 24 h, which was thoroughly washed off before the interaction with the T cells. B7-H1 or control antibodies were used at a dilution of 1:100. Concentrations had been titrated and tested previously. Of note, IFN-γ, which was used to stimulate microglial cells, astrocytes, or splenocytes, was undetectable in control cultures.

ELISA. OptEIA ELISA kits from Pharmingen were used for the detection of mouse INF-γ and IL-2 in the supernatant.

RNA extraction, cDNA synthesis, and quantitative real-time PCR. Cells were detached by trypsinization, collected and resuspended in 1 ml of TRIzol, and then frozen at −80°C. Total RNA extraction was performed using peqGOLD Tri Fast isolation reagent (peqLab, Erlangen, Germany). For first-strand cDNA synthesis, 2.5 μl of reverse-transcription buffer (10 mM MgCl₂, 500 IU/ml) for 24 h, which was thoroughly washed off before the interaction with the T cells. B7-H1 or control antibodies were used at a dilution of 1:100. Concentrations had been titrated and tested previously. Of note, IFN-γ, which was used to stimulate microglial cells, astrocytes, or splenocytes, was undetectable in control cultures.

For first-strand cDNA synthesis, 2.5 μl of total RNA was dissolved in 21.5 μl of DEPC-treated double-distilled water (H₂O₉₈). A total of 2 μl of random hexamers (200 ng/μl) was added to each sample before incubation at 70°C for 10 min. Samples were cooled on ice and subjected to a mixture consisting of 5X Moloney murine leukemia virus (M-MLV) reverse-transcription buffer (10 μl/sample; Promega, Madison, WI), deoxyNTPs (dNTPs) (10 mM; 10 μl/sample), RNasin (40 μl/μl; 0.25 μl/sample; Promega), M-MLV reverse-transcription buffer (200 μl/μl, 1 μl/sample; Promega), and DEPC-treated H₂O₉₈ (5.25 μl/sample). This mixture of 26.5 μl was added to each RNA/random heteronexomer solution. Samples were mixed and incubated for 10 min at room temperature, for 50 min at 42°C, and finally for 15 min at 70°C.

For quantitative real-time (QRT)-PCR, measurement of gene expression was performed using the ABI prism 7700 Sequence Detection system (Applied Biosystems, Foster City, CA). Primers (BioChip Technologies, Freiburg, Germany) were designed when possible to span exon–exon junctions to prevent amplification of genomic DNA and to result in amplicons <150 bp to enhance efficacy of PCR amplification. Relative quantification of specific gene expression was performed by two-step real-time PCR using cDNA as a template, as described previously (Wiendl et al., 2003). Templates were multiplied using PerkinElmer Life Sciences (Emeryville, CA) SYBR Green PCR Master Mix (containing hot-start AmpliTaqGold, SYBR Green PCR buffer (2×), MgCl₂, and dNTPs). Reverse-transcription PCR of cDNA specimens was conducted in a total volume of 15 μl with 1X TaqMan Master Mix (PerkinElmer Life Sciences) with primers at optimized concentrations. Thermal cycler parameters were 2 min at 50°C, 10 min at 95°C, and 40 cycles of denaturation at 95°C for 15 s followed by annealing/extension at 60°C for 1 min. The fluorescence resulting from binding SYBR Green dye to the double-stranded DNA was measured directly in the PCR tube. Data were analyzed with the ABI PRISM Detection system using the comparative threshold cycle (Ct) method (user bulletin; PerkinElmer Life Sciences). Samples were normalized to 18S rRNA to account for the variability in the initial concentration of the total RNA and conversion efficiency of the RT reaction.

Oligonucleotides used in this study are as follows. 18S: 18S-forward (for) (450–469), 5′-GGTGCAAAATGATACGG, 18S-reverse (rev) (636–619), 5′-GCTGAAATATCCGGGCTT. Human, b7-H1 (PD-L1): b7-H1-forward (for) (441–460), 5′-GTATAAAATGCTGCTGCA; 5′-TCAGGTTAAGACTCCATCTTC, b7-H1-reverse (rev) (646–627), 5′-CACCCTTTTGGAAACCTGGTC, hCD80: CD80-forward (for) (450–469), 5′-ATGTGTTGCAGGCAGTTCTG. mPD-L2: PD-L2-forward (for) (579–598), 5′-AGTGAGTTGGCCCATGGCTG; PD-L2-reverse (rev) (630–383), 5′-GTGATCAATGACATTGACCA; PD-L2-reverse (640–439), 5′-CACCACTTTGGAAACTTGCTG. hCD80: CD80-forward (for) (23–43), 5′-AGTTTAGAAGGAAAAATGCTGC; CD80-reverse (rev) (133–112), 5′-TCAGGTTAAGACTCCATCTTC, b7-H1-forward (for) (712–693), 5′-ATTCTGATCTGAGTCTGGTC, b7-H1-reverse (rev) (400–380), 5′-GGAGGTACATTGGTGATCGG. Murine, b7.1 (mCD80): mB7.1-forward (for) (694–713), 5′-GCCAACCACCCATTAGTG; mB7.1-reverse (rev) (843–823), 5′-GAAGCTTATGATATCACGG, mB7.2 (mCD86): mB7.2-forward (for) (368–387), 5′-ACAAAATGCGACCCACAG; mB7.2-reverse (rev) (507–487), 5′-AGTCTGATCTGACCCATG. Green dye (PD-L1): mB7-H1-forward (for) (579–598), 5′-TGCTTCCAATGTGACCCAGG; mB7-H1-reverse (rev) (712–693), 5′-ATGTTGGTGACCCAGTCTGC, mB7-H1-forward (for) (505–524), 5′-AGTACGGTGTCCTGGTCTAC; mB7-H1-reverse (rev) (646–627), 5′-CTAGGCTTGGCCAGTGGCTG.
Mouse strains and induction of experimental allergic encephalomyelitis.
For active induction of MOG<sub>35–55</sub>-experimental allergic encephalomyelitis (EAE), C57BL/6 mice (Charles River) were immunized with 200 μl of an emulsion of equal volumes of MOG<sub>35–55</sub> in PBS (2 mg/ml) and Freund’s incomplete adjuvant oil (Invitrogen) supplemented with M. tuberculosis H37Ra (2 mg/ml; Difco, Detroit, MI). To enhance the immune response, immunized animals were administered 400 ng of pertussis toxin (Sigma) on the day of immunization and on day 2 after injection. Animals were killed after induction of EAE at times indicated in Results, and tissues were fixed in paraformaldehyde.

For kinetic analysis of B7-H1 expression by flow cytometry, SJL mice (female, 6–12 weeks of age; Harlan Winkelmann, Borchen, Germany) were used according to approved protocols. EAE induction was performed essentially as described previously (Bischof et al., 2004). In brief, 50 nmol of the peptide PLP139–151 in PBS emulsified with an equal amount of CFA containing 200 μg of M. tuberculosis H37RA (Difco) was injected subcutaneously in the back of the foot. In addition, mice received a single intravenous injection of 300 ng of pertussis toxin (List Biologic, Campbell, CA) in PBS. Diseased animals had scores between 2 and 4 on the five-point EAE scale, with a score of 0 being disease free and 5 being moribund or dead.

Flow cytometry. For flow cytometric analysis of inducible surface expression of MHC class II and B7-H1, microglial cells, astrocytes, and splenocytes were preincubated with IFN-γ (500 U/ml) for 48 h. Adherent cells were detached from the surface of the 48-well plastic dishes by incubation with 0.02% EDTA in PBS. Subsequently, samples were treated as described previously (Tabi et al., 1994).

As a marker for T-cell activation in coculture experiments, we measured upregulation of the inducible costimulatory protein ICOS on the PPD-specific T cells after they had been separated from the adherent microglial cells by gentle washing.

Flow cytometric analysis of B7-H1 expression in the CNS in vivo during EAE, cells isolated from the CNS were stained on ice with the indicated antibodies for 20 min and directly before analysis with 1 μg/ml propidium iodide. Flow cytometric analysis was performed on life lymphocytes with a Cyan Cytometer (DakoCytomation) using Summit software (DakoCytomation) for data acquisition and analysis.

Histology. Spinal cords and brains were harvested on days 14–21 after immunization (controls as well as antigen-immunized mice) and snap-frozen in optimal cutting temperature compound. Four- to 10-μm-thick sections were cut, fixed in acetone, and subsequently stained. Single stainings were performed using the avidin–biotin technique (Vector Laboratories, Burlingame, CA) and counterstained with hematoxylin. Flow cytometric analysis was performed with life lymphocytes with a Cyan Cytometer (DakoCytomation) using Summit software (DakoCytomation) for data acquisition and analysis.

Statistical analysis. Data are representative of experiments performed at least three times with similar results. Significance was assessed by two-sided t test (*p < 0.05; **p < 0.01).

Results
Expression of B7-H1 on human microglial cells: strong upregulation by inflammatory stimuli
Human microglial cells were examined for the expression of B7-H1. We analyzed total RNA from cultured human adult microglial cells for different B7 molecule transcripts and HLA-DR by QRT-PCR. Microglial preparations from two different donors were used (HA376 and HA382). We also studied purified human peripheral blood monocytes, which were previously described as constitutively express B7-H1 mRNA and protein (Selenko-Gebauer et al., 2003; Schreiner et al., 2004). mRNA transcripts of B7-H1 (PD-L1) were detectable on monocytes and microglial cells already before the addition of inflammatory stimuli (Fig. 1). Monocytes and microglia were cultured in the presence of supernatants from Th1 or Th2 T-cell lines to assess whether B7-H1 expression on these cell types can be differentially regulated by exposure to a proinflammatory or an anti-inflammatory environment, respectively. Exposure of microglial cells to either Th1 or Th2 supernatants led to a significant upregulation of B7-H1 mRNA, which was clearly higher with Th1 than with Th2 supernatants [relative increases for Th1, 18.5 (HA376) and 9.4 (HA382); for Th2, 10.1 (HA376) and 2.8 (HA382)]. mRNA transcripts for the classical B7 molecules CD80 (B7.1) and CD86 (B7.2) as well as expression of MHC class II were assessed in parallel. Upregulation of CD80 in response to the proinflammatory Th1 supernatants was stronger in monocytes compared with microglia [relative increases for monocytes, 7.7 and 6.2; relative increases for microglia, 2.0 (HA376) and 3.2 (HA382)] (Fig. 1). CD86 was mildly upregulated by Th2 rather than by Th1 supernatants with no substantial differences between monocytes and microglial cells (Fig. 1). Th1 supernatant exposure results in downregulation of CD86 mRNA in monocytes but not microglia. The pattern of regulation of HLA-DR transcripts by T-cell super-
In contrast, TNF-α and the maximum effect was observed at 72 h (data not shown). IFN-γ cultured in the presence of inflammatory stimuli (500 IU/ml on microglial cells was substantially increased when cells were our observations with the Th1 supernatants, B7-H1 expression higher than on astrocytes or splenocytes (Fig. 2). Consistent with Th2 supernatants, we next investigated the expression of B7-H1 and B7 proteins on murine microglial cells in comparison with controls. We measured MHC II expression, which was induced by relative increases and absolute protein expression levels of IFN-γ (500 U/ml) were harvested after 48 h of induction and analyzed for the expression of the indicated mRNA by QRT-PCR. A, B7.1 (CD80); B, B7.2 (CD86); C, B7-H1 (PD-L1); D, PD-L2. Splenocytes and PHA-stimulated splenocytes were used as controls. Bars and numbers represent the relative gene expression of indicated molecules calculated in relation to unstimulated splenocytes (set to 1). Data represent expression analysis from glial cultures from three different animal preparations (mean ± SEM).

**IFN-γ-inducible B7-H1 protein in cultured murine microglial cells: comparison with astrocytes and splenocytes**

To validate our data achieved with RNA from human monocytes or microglial cells cultured in the presence or absence of Th1 or Th2 supernatants, we next investigated the expression of B7-H1 and B7 proteins on murine microglial cells in comparison with astrocytes and splenocytes. In the absence of added cytokines, cultured microglial cells expressed B7-H1 protein, which was higher than on astrocytes or splenocytes (Fig. 2). Consistent with our observations with the Th1 supernatants, B7-H1 expression on microglial cells was substantially increased when cells were cultured in the presence of inflammatory stimuli (500 IU/ml IFN-γ). B7-H1 upregulation was visible 12 h after stimulation, and the maximum effect was observed at 72 h (data not shown).

In contrast, TNF-α had no effect on B7-H1 expression (data not shown). Similar to the observations in microglia, B7-H1 expression was also inducible by IFN-γ in astrocytes and splenocytes. Relative increases and absolute protein expression levels of B7-H1 were higher in glial cells than in splenocytes (Fig. 2). As a control, we measured MHC II expression, which was induced by IFN-γ in all cells (microglia, astrocytes, splenocytes). Four independent experiments of primary cultured glial cells from different animal preparations were examined.

To corroborate the murine protein data, we analyzed total RNA from microglia, astrocytes, and splenocytes for different B7 molecule transcripts by QRT-PCR. According to the literature, murine microglia expressed CD80 and CD86 mRNA, which was upregulated by IFN-γ (Fig. 3A, B). On astrocytes, expression and regulation was lower (Fig. 3A, B). Corresponding to the protein data (Fig. 2), mRNA for B7-H1 (PD-L1) was expressed in unexposed microglia and was substantially induced after exposure to IFN-γ (factor of 6.3–10.5) (Fig. 3C). Astrocytes expressed lower basal levels, which were also highly upregulated after stimulation with IFN-γ. Of note, low levels of PD-L2 were also detected on microglial cells and were upregulated after treatment with IFN-γ (Fig. 3D).

**Functional role of B7-H1 on microglial cells: strong inhibition of naive polyclonal T-cell activation**

To assess the functional consequences of B7-H1 expression on microglial cells, we used a coculture setting in which microglial cells, astrocytes, or splenocytes were used as APC. APC were induced to upregulate B7-H1 and MHC-II by IFN-γ and subsequently cocultured with naive syngeneic T cells and superantigen (SEB) in the presence of neutralizing anti-B7-H1 antibody (MHI5 or 10B5) or the appropriate isotype control antibody. The release of IFN-γ and IL-2 into the supernatant was measured at the times indicated. Both cytokines are produced primarily by T cells and not by microglial cells or astrocytes (data not shown).

Compared with the isotype control, treatment with anti-B7-H1 mAb strongly augmented the production of IFN-γ and IL-2 by oligoclonally (SEB) stimulated naive T cells (Fig. 4A). In cocultures with microglial cells, in the presence of anti-B7-H1 mAb, the mean amounts of IL-2 and IFN-γ increased from 1789 ± 310 to 2816 ± 776 pg/ml (IL-2) and from 1142 ± 400 to 2110 ± 196 pg/ml (IFN-γ), respectively. For cocultures with astrocytes, IL-2 and IFN-γ were elevated from 1370 ± 60 to 1469 ± 112 pg/ml and from 548 ± 135 to 784 ± 206 pg/ml, respectively, and in the coculture system with splenocytes, IL-2 increased from 2953 ± 324 to 3415 ± 467 pg/ml and IFN-γ from 1285 ± 117 to 1861 ± 349 pg/ml. Albeit in principle, the inhibitory effect of B7-H1 was observed with microglial cells as well as astrocytes and splenocytes, B7-H1 neutralization had its greatest effects in microglial T-cell cocultures, with a mean increase of 57% for IL-2 and 85% for IFN-γ (compared with 7% and 43% in the astrocyte coculture, and 16% and 45% in the splenocyte coculture setting). In general, B7-H1 neutralization had more influence on IFN-γ production than on IL-2 secretion (Fig. 4A). For IL-2 secretion, B7-H1 neutralization only had a significant influence in microglia T-cell cocultures. Together, these data comparing different APC indicate that the inhibitory effect of B7-H1 on T-cell cytokine production is most prominent in microglial T-cell cultures.

**Functional role of B7-H1 on microglial cells: strong inhibition of antigen-specific T-cell activation**

Naive, oligoclonally (SEB) activated T cells may differ considerably from antigen-specific T cells both in their reactivity to TCR-specific stimuli and in their dependence on secondary signals. Therefore, we next assessed the functional significance of B7-H1 expression on the activation of antigen-specific T cells.

Coculture experiments were performed with PPD-specific
T-cell lines, and the modulation of T-cell cytokine production by B7-H1 was assessed. As observed with naive T cells, blocking of B7-H1 on microglial cells resulted in a significant increase in both IFN-γ and IL-2 (for IFN-γ, 17.905 ± 1435 vs 23.054 ± 2880 pg/ml, and for IL-2, 165 ± 96 vs 550 ± 91 pg/ml) (Fig. 4B).

The inhibitory effect of B7-H1 was also demonstrated by assessing the expression pattern of T-cell activation markers following coculture in the presence or absence of a neutralizing B7-H1 antibody. Neutralization of B7-H1 led to increased expression levels of ICOS on PPD-specific T cells, thus demonstrating the inhibitory role of microglia-related B7-H1 for T-cell activation (Fig. 4C). The pooled relative mean fluorescence index from three independent experiments showed an overall elevation in ICOS expression of 39% after inhibition of the B7-H1 signal (p < 0.05).

Expression of B7-H1 in the CNS and in the course of EAE

Our in vitro human and murine experiments suggested that microglia and astrocytes are capable of expressing high levels of inhibitory B7-H1 in the presence of inflammatory conditions. Therefore, we next investigated the expression of B7-H1 during CNS inflammation exemplified by MOG- and PLP-induced EAE. Expression of B7 molecules was assessed by QRT-PCR as well as immunohistochemistry at different time points after immunization in brain specimens and in the spinal cord from animals at an early stage of EAE (score 2) and a late stage of EAE (score 3–4). Under control conditions, B7-H1 mRNA was virtually undetectable in the CNS (Fig. 5). In contrast, B7-H1 expression progressively increased during the course of EAE and seemingly correlated with the disease severity (Fig. 5C). mRNA transcripts for PD-L2, the alternative splice variant of B7-H1, were found in late EAE but to a much lesser extent than B7-H1 (Fig. 5D). Expression levels of CD80 and CD86 mRNA in the brain and spinal cord in principle paralleled this pattern (Fig. 5A, B). According to the mRNA data, B7-H1 protein was virtually undetectable in control animals. However, strong immunoreactivity of B7-H1 was detectable in ongoing EAE, and intensity of staining correlated with the severity of EAE as well as the presence of inflammatory infiltrates in the lesions (Fig. 6). Histochemical analysis of the inflamed spinal cord and the CNS revealed B7-H1 coexpression with MAC3 (microglia/macrophages) but not GFAP (astrocytes) (Fig. 6). Interestingly, B7-H1 also coexisted with CD3 (T cells), corresponding to the notion that T cells are capable of expressing B7-H1 (Dong et al., 2002) (Fig. 6). Thus, our immunohistochemical analysis suggests that B7-H1, although virtually undetectable under physiological conditions, is strongly upregulated in the CNS under inflammatory conditions. As main cellular sources, we found microglia/macrophages as well as invading T cells.

To further substantiate our findings on B7-H1 expression and regulation in vivo, we performed serial flow cytometric analysis of
B7-H1 on CNS cells during the course of PLP-induced EAE (Fig. 7). Using colabeling with CD45 and CD11b, this method also allows differentiation between (resident) microglia (CD45<sub>hi</sub>CD11b<sup>+</sup>) and macrophages (CD45<sub>hi</sub>CD11b<sup>+</sup>) as well as lymphocytes (CD45<sub>lo</sub>CD11b<sup>+</sup>)-negative (Becher et al., 2002). We found a significant increase in microglial B7-H1 expression between baseline, day 14, and day 20 (Fig. 7). Importantly, expression of inhibitory B7-H1 on microglia and macrophages correlated well with the recovery phase of the animals, whereas at maximum disease severity (day 14), 37% of microglia cells were positive, and expression increased up to 84% at the time of recovery (day 20) (Fig. 7). Of note, corresponding to our histochimical results, B7-H1 could also be detected on invading T cells but at a much lower extent (data not shown).

Discussion

Our study sheds new light on the immunobiological role of the novel B7-family molecule B7-H1 in the CNS and during neuroinflammation. We found that murine and human microglial cells express high amounts of B7-H1, especially when exposed to inflammatory conditions simulated by the addition of IFN-γ or Th1 supernatants in vitro (Figs. 1–3). B7-H1 acts as a strong inhibitor of antigen-specific as well as nonspecific polyclonal T-cell activation in that it reduces both the secretion of pro-inflammatory cytokines (IFN-γ and IL-12) and the expression of T-cell activation markers (ICOS) (Fig. 4). As a "proof of concept," we provide data showing that B7-H1 is highly upregulated on microglial cells/macrophages during the course of EAE, especially in regions with the strongest inflammatory response (Figs. 5–7). Thus, our data propose B7-H1, expressed by microglial cells, as a strong immune inhibitory molecule downregulating T-cell activation and thus contributing to the immune homeostasis in the CNS.

The brain had long been considered an immunologically privileged site. This idea is based on the observation that tissue transplants in the CNS are not commonly rejected by the immune system (Medawar, 1948; Barker and Billingham, 1977). An anti-inflammatory and, with regard to invading immune cells, pro-apoptotic environment in the brain, the limited access of brain-derived antigens to the lymphoid organs, the presence of the blood–brain barrier, low MHC expression in the brain parenchyma, and the absence of dendritic cells were used to explain the lack of an effective immune response to antigens in the brain. However, numerous studies in infectious, autoimmune, and tumor models have challenged this view by showing that potent immune reactions can and do occur in the CNS (Hickey, 2001).

The CNS is constantly patrolled by activated T lymphocytes, which may induce profound damage if they identify their specific or a cross-recognized antigen in the context of appropriate MHC restriction elements. The specialized anatomic barriers like the blood–brain barrier (Fabry et al., 1994) and the peculiarities of the lymphatic drainage do not necessarily guarantee the integrity of this organ. However, limiting the local inflammatory response is crucial for an organ as vulnerable as the CNS. For example, inflammation induces a proapoptotic environment mediated by astrocytes via the CD95 pathway (Bechmann et al., 1999). Together, with the lack of costimulatory molecules (Pender, 1999), this pathway is considered to induce high numbers of apoptotic lymphocytes among infiltrating T cells in autoimmune inflammatory conditions (Gold et al., 1996). Another critical player in the immune homeostasis of the CNS environment is the microglia. Although the exact function of these resident immune cells in the intact CNS remains elusive, early insights from studies of peripheral APC suggest that microglia may have major homeostatic and reparative functions in the normal as well as injured CNS. Considering microglial cells to be the major APC in the CNS, microglia–immune cell interactions may have critical impact on the outcome of brain-derived or brain-directed immune reactions. In this context, costimulatory signals provided by the microglia are key elements in this interface. The lack of costimulatory molecules in the healthy brain changes during the course of an inflammatory response (Aloisi, 2001). Therefore, it seems that a good counterbalance is needed to keep inflammatory situations under control.

Thus far, existing studies have elucidated the relevance of stimulatory second signals on microglial cells (such as CD80, CD86, and CD40), including their contribution to modulate or amplify acute or chronic neuroinflammation (Becher and Antel, 1996; Aloisi et al., 1999; Matyszak et al., 1999; Zehntner et al., 2003). Our study is the first to show the importance of a B7 molecule on microglial cells exerting strong coinhibitory properties, thereby providing novel insights into the complex immunobiology of these CNS APC.
B7-H1, or PD-L1, is a type I transmembrane protein with 20% amino acid identity to B7.1 and 15% amino acid identity to B7.2. Ligation of the B7-H1 leads to diminished proliferation and IL-2 production and induction of cell cycle arrest whereby CD8-T cells appear to be more sensitive to this effect than CD4 cells (Carter et al., 2002). B7-H1 interacts with PD-1 and a yet unidentified non-PD-1 receptor on T cells. B7-H1 is expressed not only on hematopoietic APCs but also on parenchymal cells such as muscle cells, microvascular endothelial cells, renal tubular cells, and cancer cell lines (Dong et al., 1999; Latchman et al., 2001; Eppihimer et al., 2002; Wiendl et al., 2003; Wintterle et al., 2003). In contrast, PD-L2 has a more limited expression, predominantly on cytokine-activated macrophages and dendritic cells (Latchman et al., 2001; Yamazaki et al., 2002). These patterns of (parenchymal) expression may allow for the termination of an immune response in inflamed tissues, limiting organ damage (Wiendl et al., 2003), or in tumors allowing for immune evasion (Dong et al., 2002; Wintterle et al., 2003) (for review, see Chen, 2004). B7-H1 expressed on peripheral professional APC has recently been proposed as a candidate contributing to the maintenance of peripheral tolerance. “Weak” APC, such as monocytes, immature or semimature dendritic cells exhibit tolerogenic rather than immunogenic functions, which negatively control spontaneous autoreactive T-cell activation (Lutz and Schuler, 2002). Accordingly, B7-H1 exerts potent negative regulatory functions for T-cell activation while expressed on these “weak APC” (Brown et al., 2003; Selenko-Gebauer et al., 2003; Schreiner et al., 2004). Following this concept, it is interesting to note that B7-H1 plays a major role in microglial cells (and astrocytes), which are considered to be comparably weak APC in the CNS. In our hands, neither unstimulated microglial cells nor astrocytes were able to induce proliferation in antigen-specific or superantigen-stimulated oligoclonal T cells (data not shown). However, they influence cytokine production and T-cell activation, both features critically influenced by inhibitory B7-H1.

Our data further suggest an involvement of the B7-H1 pathway in the immune regulatory mechanisms controlling autoreactive T-cell responses relevant for the pathogenesis of CNS autoimmunity. This assumption has recently been fueled by the demonstration of the critical role of B7-H1–PD1 interactions in the regulatory mechanisms of experimental autoimmune encephalomyelitis, in which therapeutic interference had critical impact on the onset and severity of the disease (Salama et al., 2003). Similar to our findings in vivo, B7-H1 was found upregulated in the inflamed CNS. Salama et al. (2003) suggested that the B7-H1–PD-1 pathway is particularly important in the induction phase of a MOG-EAE, an assumption that further emphasizes the strong negative regulatory of B7-H1–PD-1 in the periphery.

What might be the role of B7-H1 in the CNS in the regulation of CNS inflammation, respectively, the effector phase of the host immune response? From our data, it could be argued that the proinflammatory activity of T cells might be regulated via a negative feedback loop by IFN-γ-induced B7-H1 expression on microglia (and possibly invading peripheral monocytes/macrophages). This expression of B7-H1 on local APC could force activated Th1 T cells to reduce their cytokine levels, thus diminishing local levels of inflammatory activity. Because microglia cells are capable of quickly migrating to the site of inflammation or injury (Kreutzberg, 1996), they could participate in very early modulation of inflammation. This scenario is strongly supported by a recent report demonstrating the negative regulatory role of B7-H1 (PD-L1) on antigen-presenting cells, T cells, and host

**Figure 6.** Expression of B7-H1 in the CNS. Fluorescent single or multicolor labeling (top) or enzymatic labeling immunohistochemistry (bottom) in CNS specimens of mice with EAE versus control immunized mice is shown, using antibodies for B7-H1, CD3, MAC3, and GFAP, visualized with secondary fluorochrome reagents (top) or peroxidase (bottom). A–C, B7-H1 (red) is strongly expressed in inflammatory infiltrates (overview in A; higher magnification of a single infiltrate in C) but is absent in the control CNS (no expression in E). B and D show DAPI for staining nuclei. F–I, Triple labeling of one section for DAPI (blue, F), CD3 (G, green), and B7-H1 (H, red) shows partial colocalization of CD3 with B7-H1 (I, yellow). Triple labeling of another section for DAPI (K, blue), GFAP (L, green), and B7-H1 (M, red) shows basically no colocalization of B7-H1 with GFAP-positive astrocytes (M, overlay). Serial cerebellar cryostat sections from EAE mice were stained with antibodies for B7-H1 (O, Q) and MAC3 (P, R). Strong but not completely overlapping staining patterns for B7-H1 and MAC3 were observed. Q and R show a higher magnification focusing on one infiltrate.
tissues by studying PD-L1 knock-out mice (Latchman et al., 2004); experiments using the MOG-EAE model showed that PD-L1 in host tissues and T cells limits responses of self-reactive CD4 T cells in vivo. Importantly, the transfer of encephalitogenic T cells from wild-type mice into PD-L1−/− recipients led to exacerbated disease, demonstrating the importance of host-tissue B7-H1 in negatively regulating the CNS inflammation (Latchman et al., 2004). Because we identified microglial cells as the main cellular source of B7-H1 among the resident CNS cells (Figs. 1, 2, 6, 7), they can be considered key players in mediating the negative regulatory signals in the host tissue, notwithstanding the putative importance of B7-H1 on invading cells (macrophages or T cells). In this view, it is interesting to note that disease was even more severe in PD-L1−/− recipients of PD-L1−/− T cells, therefore suggesting an important negative regulatory role of B7-H1 expressed on invading T cells (Latchman et al., 2004) (Fig. 6). Of note, although we use IFN-γ as a marker of the “Th1 environment,” it is clear that IFN-γ is not the only molecule present within the Th1 milieu that contributes to the observed coinducer molecule induction on monocyte/microglia differentiation. It is likely that multiple molecules in the Th1 environment (including but not limited to IFN-γ) are likely to contribute to the modulation of the human monocyte/microglia (Kim et al., 2004).

The idea of local immunosuppression by B7-H1 possibly affecting disease activity and progression is appealing, from both an immunopathogenic and a therapeutic view. Although purely speculative at present, it is possible that B7-H1 could play a role in direct tolerization of autoreactive cells; a recent study reported tolerance induction within the CNS itself rather than within draining lymphoid tissues. Naive MBP-specific T cells of MBP T-cell receptor transgenic mice readily migrated into the CNS without previous activation but were tolerized within the CNS in situ (Brabb et al., 2000). In a model of autoimmune hepatitis, B7-H1 was critical for determining the accumulation and deletion of intrahepatic CD8+ T lymphocytes as shown in B7-H1 knock-out mice (Dong et al., 2004). It remains to be shown in multiple sclerosis (MS) whether an increase in B7-H1 (on APC or T cells), and a parallel decrease in number or function of pathogenic T cells, could possibly account for the phases of relapses and remissions in MS.

Our demonstration of the expression of B7-H1 in areas of strongest inflammation in EAE further emphasizes our assumption on the importance of this regulatory principle in vivo. This negative regulatory feedback loop should be important for keeping the “anti-inflammatory milieu” in the CNS.

References

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