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Butyrophilin, a Milk Protein, Modulates the Encephalitogenic T Cell Response to Myelin Oligodendrocyte Glycoprotein in Experimental Autoimmune Encephalomyelitis

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Experimental autoimmune encephalomyelitis (EAE) induced by sensitization with myelin oligodendrocyte glycoprotein (MOG) is a T cell-dependent autoimmune disease that reproduces the inflammatory demyelinating pathology of multiple sclerosis. We report that an encephalitogenic T cell response to MOG can be either induced or alternatively suppressed as a consequence of immunological cross-reactivity, or “molecular mimicry” with the extracellular IgV-like domain of the milk protein butyrophilin (BTN). In the Dark Agouti rat, active immunization with native BTN triggers an inflammatory response in the CNS characterized by the formation of scattered meningeal and perivascular infiltrates of T cells and macrophages. We demonstrate that this pathology is mediated by a MHC class II-restricted T cell response that cross-reacts with the MOG peptide sequence 76–87, IGEGKVALRIQN (identities underlined). Conversely, molecular mimicry with BTN can be exploited to suppress disease activity in MOG-induced EAE. We demonstrate that not only is EAE mediated by the adoptive transfer of MOG74–90, T cell lines markedly ameliorated by i.v. treatment with the homologous BTN peptide, BTN74–90, but that this protective effect is also seen in actively induced disease following transmucosal (intranasal) administration of the peptide. These results identify a mechanism by which the consumption of milk products may modulate the pathogenic autoimmune response to MOG. The Journal of Immunology, 2000, 165: 2859–2865.

The etiology of multiple sclerosis (MS) is obscure but is believed to involve environmental factors that disrupt immunological self-tolerance to CNS myelin in genetically susceptible individuals (1). The selective activation and expansion of myelin-specific T and B cells in MS patients supports this concept (1, 2), but the identity and mode of action of the environmental factors that initiate or exacerbate this autoimmune response remain unknown.

A major target for the pathogenic autoimmune response in MS and its animal model, experimental autoimmune encephalomyelitis (EAE), is the myelin oligodendrocyte glycoprotein (MOG). MOG is the only myelin autoantigen known to induce both a demyelinating autoantibody response (3, 4) and an encephalitogenic CD4+ T cell response (5–8) in animals with EAE. The demyelinating potential of the MOG-specific Ab response has been studied extensively both in vivo and in vitro (4, 9, 10), and, in MOG-induced EAE, this humoral response acts synergistically with the encephalitogenic T cell response to induce a demyelinating pathology similar to that in MS (4, 11–14). Although the encephalitogenic MOG-specific CD4+ T cell response initiates the recruitment of immune effector cells into the CNS and disrupts the blood brain barrier, demyelination in the rat is dependent on the presence of anti-MOG autoantibodies (4). These Abs bind to MOG exposed on the myelin surface (15) and mediate demyelination by a combination of complement and cell-mediated immune effector mechanisms (16–18). The observation that both MOG-specific T cell (19, 20) and Ab responses (21) are enhanced relative to other myelin Ags in MS suggests that demyelination involves a similar combination of immune effector mechanisms in the human disease. A concept supported by the recent demonstration that anti-MOG-Abs colocalize with myelin debris in actively demyelinating MS lesions (13, 22). However, the mechanism(s) that may initially disrupt self-tolerance to MOG in MS are obscure.

Intriguingly, sequence homologies involving its extracellular Ig-like domain (MOG157) identified MOG as a member of an extended family of “B7-like”proteins (23). Of particular interest was the finding that the highest level of sequence identity, ~50%, was with a homologous extracellular Ig domain of butyrophilin (BTN) (24), a major protein of the milk fat globule membrane (MFGM) (25, 26). This observation led us to speculate that immunological cross-reactivity or “molecular mimicry” with BTN may influence the function of the MOG-reactive autoimmune repertoire.

In the current study, we report that this is indeed the case. In the context of a permissive MHC haplotype, we show that the CD4+ T cell response to BTN and MOG is mutually cross-reactive. Therefore, sensitization with native BTN can initiate an inflammatory response in the CNS mediated by a class II MHC-restricted
MOG-reactive T cell response. Conversely, both intranasal and i.v. treatment with BTN peptide can abrogate MOG-induced EAE. The observation that transmucosal exposure to a BTN peptide can modulate disease activity in MOG-induced EAE suggests that, in the context of an appropriate HLA haplotype, dietary exposure to BTN may modulate the pathogenic autoimmune response to MOG in MS.

Materials and Methods

Animals and Ags

Dark Agouti (DA) and Brown Norway (BN) rats (120–200 g) were obtained from Charles River Breeding Laboratories (Sulzfeld, Germany), and Lewis (LEW) rats were obtained from the animal facility of the Max-Planck Institute for Biochemistry ( Martinsried, Germany). CFA, IFA, and heat-killed Mycobacterium tuberculosis (H37Ra) were purchased from Difco (Detroit, MI). Purified protein derivative was purchased from the State Serum Institute (Copenhagen, Denmark). Recombinant MOG<sup>64</sup> (amino acids 1–120) (12) and BTN<sup>64</sup> (amino acids 27–144 notation including signal sequence) (26) corresponding to the N-terminal Ig-like domains of the two proteins with a C-terminal hexahistidine tag were expressed in *Escherichia coli*. The recombinant proteins were purified by chromatography on Ni-NTA agarose to a purity of >95% as assessed by SDS-PAGE (Qiagen, Chatsworth, CA) and stored at −20°C. MFGM was purified from fresh bovine milk as described previously (25). Synthetic peptides were purchased from Genosys (Cambridge, U.K.).

Immunization protocols and generation of Ag-specific T cell lines

Rats were immunized s.c. at the base of the tail with 100 μg Ag emulsified in CFA containing 225 μg of heat-killed *M. tuberculosis* (H37Ra) in a total volume of 100 μl. Ag-specific T cell lines were generated as described previously (12). Briefly, the draining lymph nodes were removed 10 days postimmunization (d.p.i.), and a single cell suspension was cultured for 72 h at a concentration of 10<sup>7</sup> cells/ml in DMEM supplemented with glutamine, penicillin, streptomycin, sodium pyruvate, essential amino acids (Life Technologies, Rockville, MD), and 1% rat serum in the presence of the selecting Ag (20 μg/ml). T cell blasts were then isolated by density gradient centrifugation and propagated for a further 5–10 days in medium containing IL-2. Ag-specific T cell lines were subsequently maintained by cycles of Ag-specific restimulation using irradiated (5000 rad) syngeneic APCs containing IL-2. Ag-specific T cell lines were subsequently maintained by cycles of Ag-specific re-stimulation using irradiated (5000 rad) syngeneic APCs, followed by expansion in IL-2–containing medium.

Cytofluorometric analysis was performed using samples of 2 × 10<sup>5</sup> viable cells washed with PBS containing 0.2% BSA and 10 mM NaN₃ and incubated with the primary mAb for 1 h on ice. After washing, the cells were stained with fluorescein-conjugated goat anti-mouse IgG (Dianova, Hamburg, Germany) for 1 h on ice. After removing unbound fluorescein-conjugated Ab by washing, the cells were analyzed using a fluorescein-activated cell sorter (FACScan; Becton Dickinson, Heidelberg, Germany). A live gate was obtained by incubating the cells in PBS containing propidium iodide.

Proliferation assays were performed in flat-bottom 96-well tissue culture plates in a total volume of 200 μl containing either 5 × 10<sup>3</sup> lymph node cells, or 2 × 10<sup>5</sup> T cell lines plus 5 × 10<sup>3</sup> syngeneic, irradiated (5000 rad) thymus cells as APCs. Ag-specific proliferation was assessed by the incorporation of [³H]thymidine (10 μCi/well) during the final 16 h of a 72-h culture period using a Packard (Meriden, CT) Matrix 96 Direct β counter.

EAE induction and treatment

Female DA rats (6–8 wk old) were treated intranasally daily for a period of 10 days with a dose of 50 μg of Ag (1 mg/ml in water), which was injected using a micropette (25 μl per nostril). Three days after the last intranasal application of Ag, the animals were injected with 100 μg of MOG<sup>64</sup> in IFA. Adoptive transfer experiments were performed using activated T cell blasts isolated after 72 h restimulation with Ag in vitro. The T cell blasts were suspended in a volume of 1 ml DMEM and injected into the tail vein of female DA rats (6–8 wk old). Treatment with high-dose soluble Ag was performed using 500 μl of Ag (2 mg/ml in DMEM) injected in the tail vein 2 and 4 days after T cell transfer. Animals were weighed and examined daily for clinical signs of EAE that was scored on a scale of 0–5 as described previously.

Histopathological analysis

Histological evaluation was performed on paraformaldehyde-fixed, paraffin-embedded sections of brains and spinal cords sampled at various time points of disease. Paraffin sections were stained with hematoxylin-eosin, Luxol fast blue, and Bielschowsky silver impregnation to assess inflammation, demyelination, and axonal pathology. In adjacent serial sections, immunohistochemistry was performed with Abs against macrophages/activated microglia (ED1; Serotec, Oxford, U.K.) or T cells (W3/13; Seralab, Sussex, U.K.). Bound primary Ab was detected with a biotin-avidin technique as previously described in detail (27). Control sections were incubated in the absence of primary Ab or with nonimmune rabbit serum.

Results

Sensitization with BTN induces a strain-specific inflammatory response in the CNS

LEW, BN, and DA rats were immunized with either MFGM, or alternatively BTN<sup>64</sup> in CFA and monitored for the development of clinical and/or histopathological evidence of EAE. Strikingly, although neither immunogen induced clinical disease in these rat strains, histopathological analysis identified a subclinical inflammatory response in the CNS of DA rats that was absent in DA controls immunized with PBS/CFA. This pathologic response was characterized by the formation of scattered meningeal and perivascular infiltrates of T cells and macrophages throughout the CNS (Fig. 1a). This is in contrast to the pathology of MOG-induced EAE in the rat that is dominated by Ab-mediated demyelination (28). In contrast, neither immunogen induced any CNS pathology in the other two rat strains. Moreover, no pathology developed in the CNS of DA rats immunized with PBS/CFA. These observations indicate that BTN<sup>64</sup>, a major component of MFGM, can act as a strain-specific encephalitogenic in the rat.

MOG-specific T cells cross-react with a defined BTN peptide in the DA rat

To test the hypothesis that molecular mimicry with MOG<sup>64</sup> at the level of the T cell response was responsible for the CNS pathology induced by BTN, we investigated the ability of BTN and synthetic BTN peptides to stimulate a panel of encephalitogenic class II MHC-restricted MOG-specific T cell lines (TCL) derived from LEW (MHC RT<sup>1</sup>), BN (MHC RT<sup>1</sup>), and DA (MHC RT<sup>1</sup><sup>Ⅰ</sup>) rats. The encephalitogenic epitopes recognized by these TCLs are strain-specific and together span 70% of the MOG<sup>64</sup> sequence (Table I).

In agreement with the inability of BTN<sup>64</sup> to mediate an encephalitogenic response in either LEW or BN rats, MOG-specific TCLs derived from these strains did not proliferate in response to either BTN<sup>64</sup> or synthetic BTN<sup>64</sup> peptides in vitro. In contrast, DA MOG-specific TCLs proliferated in response to BTN<sup>64</sup> in the presence of syngeneic APCs. Using a panel of overlapping synthetic peptides, we demonstrated that this cross-reactive response was restricted to the overlapping BTN peptides BTN<sub>76–87</sub> and BTN<sub>74–90</sub> (Fig. 2a).

The overlapping sequence of these peptides spans an encephalitogenic MOG T cell epitope for the DA rat, the peptide sequence MOG<sub>76–87</sub>, suggesting that molecular mimicry was restricted to this epitope and did not involve a second epitope located within MOG<sub>66–107</sub> (Fig. 2a; Table I; Stefferl and Schubart, unpublished observations). We investigated this further using synthetic peptides corresponding to amino acids 74–90 of the IgV-like domains of the two Ags. Strikingly both MOG<sub>74–90</sub> and BTN<sub>74–90</sub> stimulated the MOG-specific TCL to a similar extent, inducing a proliferative response similar to that obtained with the recombinant Ag and the longer peptides (Fig. 2a).

Active immunization with BTN induces an encephalitogenic MOG-reactive T cell response

Although the above observations demonstrate that the DA MOG-specific T cell response exhibits molecular mimicry with BTN in
FIGURE 1. Representative spinal cord histopathology of DA rats induced by BTN. a, Active immunization with BTN\(^{IgV}\); b–d, Adoptive transfer of \(10^7\) BTN \(74-90\)-specific T cell lines. a, Virgin female DA rats immunized with BTN\(^{IgV}\) develop a perivascular inflammatory infiltrate consisting of mononuclear cells (between arrowheads). DA rats immunized with MFGM/CFA developed similar lesions. b, Adoptive transfer of BTN \(74-90\)-specific T cells induced the formation of multiple perivascular inflammatory infiltrates (small arrowheads; the asterisk indicates the infiltrate, which is shown in c and d at higher magnification) consisting of large numbers of T cells (c) but relatively few ED1\(^+\) macrophages (d). Invasion into the parenchyma was minimal. (Magnification: a, c, and d, \(\times 270\); b, \(\times 35\).

vitro, it remained uncertain whether this was responsible for the inflammatory pathology induced by active immunization with BTN in vivo. Therefore, we examined the BTN-specific T cell repertoire for evidence of a cross-reactive and encephalitogenic response to MOG\(74-90\) using short-term TCLs obtained from BTN\(^{IgV}\) immunized donors. Ag-specific TCLs were selected in vitro using a combination of the peptides BTN\(_{63-87}\) and BTN\(_{76-100}\) as the selecting Ag. After two rounds of restimulation in vitro, the encephalitogenic capacity of the TCLs was assessed by adoptive transfer into naive syngeneic recipients.

The i.v. injection of \(5 \times 10^6\) to \(10^7\) T line cells induced an intense inflammatory response in the CNS of all recipients (Fig. 1b), associated with weight loss and hind limb paraparesis. Immunohistochemical analysis of the lesions revealed that they consisted of large numbers of T cells concentrated in the perivascular space. Migration of infiltrating T cells into the parenchyma was restricted and the recruitment of ED1\(^+\) macrophages into the CNS was minimal (Fig. 1, c and d), a pathology similar to that described previously in the Lewis rat following the adoptive transfer of MOG-specific T cells (6).

The observation that BTN\(_{63-87}/BTN_{76-100}\)-selected TCLs proliferate in vitro in response to MOG\(_{74-90}\) and MOG\(^{Igd}\), as well as the selecting peptides and BTN, confirmed that their Ag-specific response to B74–90 and M 74–90 was class II MHC-restricted and the recruitment of ED1\(^+\) macrophages (d). Invasion into the parenchyma was minimal. (Magnification: a, c, and d, \(\times 270\); b, \(\times 35\).

**BTN peptide suppresses adoptively transferred MOG-EAE**

It is now appreciated that molecular mimicry can induce either a pathogenic autoimmune response or, conversely, tolerance, depending on variables such as the route of sensitization, dose, and adjuvant effects (29). This is of particular relevance for peptides derived from ingested Ags, as these will cross the gut mucosal surface and provide tolerogenic signals in the periphery (30, 31). Therefore, could mimicry with BTN be exploited to suppress an autoaggressive MOG-reactive T cell response and abrogate clinical disease in EAE?

The tolerogenic potential of BTN\(_{74-90}\) was first investigated in EAE induced by the adoptive transfer of MOG\(_{74-90}\)-specific T cells. In the DA rat, a dose of \(5 \times 10^6\) MOG\(_{74-90}\)-specific T cells induced a maximal clinical score of 2.5 \(\pm\) 0.3 four days after transfer. Disease severity was dramatically attenuated following i.v. treatment with high-dose BTN\(_{74-90}\) given on days 2 and 4 following T cell transfer (Fig. 3). The same treatment protocol using the nominal MOG peptide ligand MOG\(_{74-90}\) was even more effective and completely abrogated clinical disease (Fig. 3). This differential in vivo effect was not reflected by the in vitro proliferative response of MOG\(_{74-90}\)-specific T cells to MOG\(_{74-90}\) and BTN\(_{74-90}\), which was similar over a wide concentration range (Fig. 2d). We are currently investigating whether this reflects differences in the pharmacological characteristics of the peptides in vivo, or differential effects of the BTN peptide on individual MOG\(_{74-90}\)-specific T cell clones within the TCLs.

However, would transmucosal exposure to the BTN peptide have a significant impact on the course of EAE-induced by active immunization with MOG\(^{Igd}\)? This disease model reproduces the complex immunopathology of MS and is mediated by a combination of both Ab and T cell-dependent effector mechanisms, including an encephalitogenic T cell response directed against two distinct epitopes (Table I; Stefferl and Schubart, unpublished observations). In view of the complex immunopathogenesis of this model and that only one T cell epitope exhibits molecular mimicry with BTN, it was not surprising that intranasal treatment with any of the peptides when given individually (MOG\(_{93-110}\), MOG\(_{74-90}\), or BTN\(_{74-90}\)) was unable to completely suppress clinical disease.

Controls pretreated with OVA developed a biphasic disease characterized by an initial episode of EAE that was rapidly followed by a severe relapse that was normally fatal by day 25 p.i. (Fig. 4a). Nasal
administration of the individual peptides reduced disease severity, but failed to delay disease onset or stop disease progression beyond day 15, as demonstrated in Fig. 4b for animals treated with MOG 93–110. In contrast, the effect of treatment with combinations of either MOG 93–110 and BTN 74–90, or MOG 93–110 and MOG 74–90 was far more effective and both combinations of peptides delayed disease onset, penetrance, and severity (Fig. 4c). This effect was clearly seen 20 d.p.i. when all animals in the control group pretreated with OVA were in relapse with a mean clinical score of 3.5 ± 0.32 (n = 6, Fig. 4c). In contrast, at this time point, only two rats treated with MOG 93–110/BTN 74–90 combination (n = 5) had developed EAE (grades 1 and 2). A similar level of protection was also seen following intranasal pretreatment with the combination MOG93–110/MOG74–90 (data not shown). Therefore, the BTN peptide could be used interchangeably with the corresponding MOG epitope to modify the severity of actively induced MOG-EAE.

Discussion

In this study, we demonstrate that, as a consequence of molecular mimicry, BTN, a common dietary Ag, can modulate the autoimmune T cell response to MOG, a major target in the immunopathogenesis of EAE and MS. This observation extends the concept of

### Table I. Sequence homologies between encephalitogenic MOG T cell epitopes and BTN

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a Data taken from Ref. 6.
b Data for the identity of the BN and DA epitopes was obtained from our unpublished data.

**FIGURE 2.** MOG and BTN provoke a mutually cross-reactive T cell response. T cell lines were derived from DA rats immunized with either (a) MOG1–20 or (b) BTN1–20. Epitope mapping using sets of overlapping synthetic peptides spanning the homologous Ig-domains of BTN or MOG identified a cross-reactive T cell response to an epitope located between amino acids 74 and 90. a. DA MOG-specific T cell lines exhibit a limited proliferative response to BTN1–20 that is similar to that obtained using the peptides BTN74–90 and BTN76–100, their overlapping sequence BTN74–90 and its MOG homologue, MOG74–90. b. A similar cross-reactive response was found when T cells specific for the epitope BTN74–90 were selected from the draining lymph nodes of BTN1–20-immunized DA rats. However, in this case, although both MOG1–20 and the peptides induced a substantial proliferative response, BTN1–20 was relatively poorly processed and presented to the T cells in vitro. c. Recognition of their cognate peptide by MOG74–90-specific (■) and BTN74–90-specific (□) TCLs is restricted by class II MHC (I–A). Addition of the mAb OX6 (anti-I–A), but not mAbs OX17 (anti-I–E) or OX18 (anti-class I MHC), blocks proliferation in vitro (mAb concentration 10 μg/ml). The purified OX mAbs were kindly provided by Dr. R. Weissert, Stockholm. d. The peptides MOG74–90 and BTN74–90 stimulate MOG74–90-specific T cells to a similar extent over a broad range of peptide concentrations.
Intranasal vaccination with BTN peptide ameliorates disease activity in MOG-induced EAE. DA rats were injected with a dose of $5 \times 10^6$ MOG$_{74-90}$-specific T cells that induced severe clinical disease starting from day 4 posttransfer (●). Intravenous administration of 1 mg BTN$_{74-90}$ (■) or MOG$_{74-90}$ (▲) 2 and 4 days after T cell transfer significantly inhibited disease. Injections were performed under light ether anesthesia; control animals were treated with OVA. ∗, $p < 0.05$; Duncan’s post hoc test.

FIGURE 3. High-dose BTN peptide suppresses MOG$_{74-90}$-induced EAE. DA rats were injected with a dose of $5 \times 10^6$ MOG$_{74-90}$-specific T cells that induced severe clinical disease starting from day 4 posttransfer (●). Intravenous administration of 1 mg BTN$_{74-90}$ (■) or MOG$_{74-90}$ (▲) 2 and 4 days after T cell transfer significantly inhibited disease. The experiment was terminated on day 20 postimmunization, by which time point the entire control group had developed severe neurological deficits. Treatment with a combination of MOG$_{74-90}$ and BTN$_{74-90}$ completely abolished the initial phase of disease and had a significant effect on later disease activity reducing the mean clinical score from 3.5 ± 0.32 to <1.0 on day 20.

FIGURE 4. Intranasal vaccination with BTN peptide ameliorates disease activity in MOG-induced EAE. DA rats were pretreated by intranasal vaccination as described in the text with (a) OVA (n = 6), (b) MOG$_{93-110}$ (n = 6), or (c) a combination of MOG$_{93-110}$ and BTN$_{74-90}$ (n = 5). The experiment was terminated on day 20 postimmunization, by which time point the entire control group had developed severe neurological deficits. Treatment with a combination of MOG$_{93-110}$ and BTN$_{74-90}$ completely abolished the initial phase of disease and had a significant effect on later disease activity reducing the mean clinical score from 3.5 ± 0.32 to <1.0 on day 20.
which case BTN in the diet may not only induce oral tolerance to BTN itself, but also suppress cross-reactive and potentially encephalitogenic MOG-reactive T cell responses. However, such an effect will require a permissive MHC class II haplotype and be further complicated by factors effecting the degradation and uptake of BTN peptides in the gastrointestinal tract before its immunological processing and presentation. However, the mechanisms responsible for oral tolerance are poorly developed at birth, and, in the neonate, feeding Ags may initially prime the immune system rather than inducing tolerance, oral tolerance only developing if feeding is continued beyond a critical age (31). In the case of an autoantigen, this raises the possibility that neonatal exposure may enhance disease susceptibility later in life. This was clearly seen in rats fed MBP for a limited period immediately postpartum and then subsequently immunized with MBP in CFA in adulthood (46). These animals not only exhibited no tolerance to MBP-induced EAE, but also developed more severe disease than control littermates. In contrast, feeding MBP to adult rats induced oral tolerance to MBP, but also developed more severe disease than control littermates (46). These animals not only exhibited no tolerance to MBP-in-induced EAE, but also developed more severe disease than control littermates. In contrast, feeding MBP to adult rats induced oral tolerance and suppressed MBP-induced EAE (46). The role that timing of exposure and the differential effects of syngeneic BTN as opposed to xenogeneic BTN have on the T cell response to MOG is currently under investigation in BTN+/- and wild-type mice.

Intriguingly, the frequency of seropositive responders to milk proteins peaks in childhood and declines as puberty approaches (47), a time frame similar to that reported for the role of the environmental factors in the etiology of MS (1, 48). Moreover, epidemiological studies have identified an association between the consumption of milk and the prevalence of MS (32–34). However, whether or not this is related to molecular mimicry between BTN and MOG must remain a matter of speculation. Indeed, milk in the diet is only one of many environmental factors related to the prevalence of MS (48). Moreover, this simple explanation neglects the existence of multiple BTN homologues that are encoded together with BTN and MOG homologues that are encoded together with BTN telomeric of the nonclassical class 1 MHC locus (23). Mimicry involving the N-terminal domains of these proteins, which are expressed in a wide variety of tissues, may also influence the composition and function of the BTN/MOG repertoire.

In conclusion, we identify BTN as an Ag that can influence the clinical outcome of autoimmune responses to MOG, an important antigenic target in EAE and MS. Modulation of the MOG-specific repertoire as a consequence of molecular mimicry with this dietary Ag BTN may be a significant factor in determining the role MOG plays as a target Ag in the immunopathogenesis of MS. Identification of those MHC haplotypes permissive for a cross-reactive T cell response between MOG and BTN may provide a strategy to identify those at risk of developing encephalitogenic responses to MOG following premature exposure to BTN. 

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References


