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Cell Reports

IL-17 triggers the onset of cognitive and synaptic deficits in early stages of Alzheimer's disease

Graphical abstract



Highlights

- IL-17-producing cells accumulate in the brain and the meninges of 3xTg-AD mice
- The increase of IL-17 producers associates with short-term memory deficits
- Neutralization of IL-17 prevents cognitive impairments and synaptic dysfunction
- IL-17 triggers Alzheimer's disease onset independently of Aβ and tau pathology

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In brief

Using a mouse model of Alzheimer's disease, Brigas et al. demonstrate that the onset of cognitive decline associates with an accumulation of IL-17-producing cells in the brain and the meninges. Targeting IL-17 into the ventricle prevents short-term memory and neuronal synaptic plasticity deficits at early stages of disease.



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Article

IL-17 triggers the onset of cognitive and synaptic deficits in early stages of Alzheimer's disease

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SUMMARY

Neuroinflammation in patients with Alzheimer's disease (AD) and related mouse models has been recognized for decades, but the contribution of the recently described meningeal immune population to AD pathogenesis remains to be addressed. Here, using the 3xTg-AD model, we report an accumulation of interleukin-17 (IL-17)-producing cells, mostly $\gamma\delta$ T cells, in the brain and the meninges of female, but not male, mice, concomitant with the onset of cognitive decline. Critically, IL-17 neutralization into the ventricles is sufficient to prevent short-term memory and synaptic plasticity deficits at early stages of disease. These effects precede blood-brain barrier disruption and amyloid-beta or tau pathology, implying an early involvement of IL-17 in AD pathology. When IL-17 is neutralized at later stages of disease, the onset of short-memory deficits and amyloidosis-related splenomegaly is delayed. Altogether, our data support the idea that cognition relies on a finely regulated balance of "inflammatory" cytokines derived from the meningeal immune system.

INTRODUCTION

Alzheimer's disease (AD) is a highly prevalent neurodegenerative disorder characterized by cognitive deficits due to synaptic dysfunction and sequential formation of amyloid-beta (A β) plaques and hyperphosphorylated tau tangles. There is currently no effective treatment for AD (Dubois et al., 2010; Reitz and Mayeux, 2014), urging the discovery of additional mechanisms underlying the etiology of the disease.

Neuroinflammation, characterized by increased proinflammatory cytokines, infiltrating immune cells, and activated glial cells, is increasingly recognized as a prominent player in AD. However, the role of the immune system is not limited to the brain parenchyma, as it comprises complex interactions between the central nervous system (CNS) and the peripheral systems (Louveau et al., 2015). For instance, T cells, B cells, dendritic cells, innate lymphoid cells, and natural killer (NK) cells reside in the healthy meninges—a direct interface between the brain parenchyma and the peripheral organs (Aspelund et al., 2015; Louveau et al., 2015; Gadani et al., 2017; Korin et al., 2017). Importantly, immune cells and their soluble mediators, namely cytokines, play a key role in CNS function by modulating neuronal connectivity and thus impacting on sensory function (Chen et al., 2017), social behavior (Filiano et al., 2016), as well as learning and memory (Derecki et al., 2010; Monteiro et al., 2016; Ribeiro et al., 2019). It was recently shown that, upon aging and in AD, meningeal lymphatic vasculature deteriorates, thus impairing cognition (Da Mesquita et al., 2018). Furthermore, the ablation of meningeal lymphatic vessels worsened the outcome of anti-A β immunotherapy in the 5xFAD mouse model (Da Mesquita et al., 2021). However, the role of specific meningeal immune populations and their molecular mediators in the context of AD was never addressed.

We have recently reported that $\gamma\delta$ T cells are a major source of interleukin-17A (IL-17) in the healthy meninges, which promotes specifically short-term memory by supporting glutamatergic synaptic plasticity of CA1 hippocampal neurons (Ribeiro et al., 2019). Additionally, IL-17 produced by astrocytes after stroke was also shown to have a beneficial role by promoting tissue repair (Lin et al., 2016). In contrast, murine IL-17-producing cells have been linked to experimental autoimmune encephalomy-elitis (EAE) (Sutton et al., 2009), cerebral ischemia-reperfusion







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injury (Shichita et al., 2009; Sutton et al., 2009; Benakis et al., 2016), and autism-like behavior (Choi et al., 2016). In these cases, IL-17-producing cells, namely CD4⁺ helper type 17 (Th17) and $\gamma\delta$ ($\gamma\delta$ 17) T cells, have been pointed out as critical players in disease progression by promoting a local immune amplification loop in the meninges as well as blood-brain barrier (BBB) disruption (Shichita et al., 2009; Sutton et al., 2009; Gelderblom et al., 2012; Benakis et al., 2016). Moreover, in a rodent model of Parkinson's disease (PD), Th17 cells were described to exacerbate neuroinflammation and neurodegeneration (Liu et al., 2019), which is paralleled by elevated production of IL-17 by CD4⁺ T cells in patients with PD (Sommer et al., 2018). Altogether, these findings suggest that IL-17 may be instrumentally involved in various neurodegenerative diseases. Accordingly, an increased proportion of circulating Th17 cells was observed in subjects with mild cognitive impairment (MCI) (Oberstein et al., 2018), and the rise of IL-17 levels in the serum of AD linked to disease progression (Chen et al., 2014). Th17 cells are reported to infiltrate the brain of AD models, promoting inflammation and neuronal death (Siffrin et al., 2010; Yang et al., 2017). In line with this, neutralization of IL-17 was shown to rescue neuroinflammation and memory impairments induced by direct Aβ administration (Cristiano et al., 2019). However, the impact of IL-17 on brain cognitive function at the onset and during the course of AD remains to be addressed.

Herein, we report a substantial increase of IL-17-producing T cells, mostly $\gamma\delta$ T cells, in the CNS of a transgenic model of AD-3xTg-AD mice-at early stages of the disease. Interestingly, this phenotype associated with cognitive deficits observed in females, but not in males, thus translating the sexual dimorphism observed in patients with AD (Fisher et al., 2018). Importantly, anti-IL-17 monoclonal antibody (mAb) intracerebroventricular (ICV) treatment of females prevented short-term memory deficits and synaptic plasticity impairments, prior to the detection of $A\beta$ or tau pathologies. Our data therefore strongly suggest that exacerbated levels of meningeal IL-17 promote synaptic dysfunction underlying the cognitive decline in early stages of AD.

RESULTS

IL-17⁺ cells accumulate in the CNS at the onset of cognitive decline in 3xTg-AD mouse model

An increase of IL-17 levels in the serum has been linked to disease progression in patients with AD (Chen et al., 2014). To investigate the role of IL-17 in neuropathological changes and



memory deficits associated with the early onset of AD, we used the triple-transgenic mouse model of AD (3xTg-AD), a progressive model of the disease developing both amyloid plaques and neurofibrillary tangles (Oddo et al., 2003b).

Animals were tested at an early stage (2 to 3 months old, i.e., prior to detection of any cognitive deficits), at the onset of disease (5 to 6 months old, when the cognitive deficits start), and at later stages (8 to 9 months old, when pathology and memory deficits are overt; Oddo et al., 2003a; Billings et al., 2005, 2007; Giménez-Llort et al., 2007). Short- and long-term memories were tested using the Y-maze and the Morris water maze (MWM), respectively (Ribeiro et al., 2019). Cognitive deficits evaluated in MWM were only observed in female 3xTg-AD mice starting at 5 months of age, but not in males, even at later stages (8 to 9 months old), when compared with aged- and sex-match B6129/J wild-type (WT) controls (Figures 1A and 1B), as previously reported (Oddo et al., 2003a). According to our previous evidence of IL-17 requirement for short-term memory (Ribeiro et al., 2019), but not long-term, we also assessed Y-maze performance. We found a progressive reduction in the discrimination ratio of females throughout age, starting at 5 to 6 months, whereas males were found only impaired at 8 to 9 months old (Figure 1C). Of note, when compared to age and sex-match WT controls, female 3xTg-AD mice had similar levels of motor and exploratory activity and no signs of anxiety (Figures 1D and 1E), in line with published data (Oddo et al., 2003a).

We hypothesized that IL-17 could trigger AD onset; therefore, we assessed whether changes in cognitive function would be associated with alterations on the provision of IL-17 in the CNS or in the periphery. For this, we analyzed by flow cytometry the brain, meninges, cervical lymph nodes (cLNs), and spleens of WT and 3xTg-AD mice upon disease progression (Figure 2). We found that, at the onset of memory deficits (5 to 6 months of age), the meninges, the brain, and the cLNs of female 3xTg-AD mice, but not the spleen, showed higher percentages and absolute numbers of IL-17A⁺ cells compared with age-matched WT controls (Figures 2A-2C). These differences on IL-17⁺ cells were maintained at later stages (8 to 9 months of old). Interestingly, in contrast to females, 3xTg-AD male mice-that did not display particular cognitive decline at 5 months of age-did not exhibit any accumulation of IL-17⁺ cells (Figures 2B and 2C). This led us to hypothesize that exacerbated IL-17 associates with memory deficits at the onset of AD, selectively in female mice (which were used in all subsequent experiments).

Figure 1. Age-dependent sexual dimorphism in 3xTg-AD mice: only females display cognitive deficits starting at 5 months of age

- (D) Animals were placed in a box for 5 min and allowed to explore, and total distance (cm) was evaluated.
- (E) Anxiety was evaluated by the total number of transitions between open and closed arms.

WT and 3xTg-AD females and males 2 to 3 months old (n = 10–15 for female mice; n = 9–10 for male mice), 5 to 6 months old (n = 20–34 for female mice; n = 9–16 for male mice), and 8 to 9 months old (n = 8–9 for female mice; n = 4–5 for male mice) were tested in the (A and B) MWM-maze, (C) Y-maze, (D) open field, and (E) elevated plus maze.

⁽A) The escape latency, i.e., the time required for the mice to find and climb onto the platform, was recorded for 60 s.

⁽B) % time in the quadrant where the platform was evaluated, in the 60-s probe test.

⁽C) The discrimination ratio between novel (N) and the other (O) arm was evaluated.

Results are representative of 1–3 independent experiments. Error bars, mean ± SEM. *p < 0.05, **p < 0.01, and ***p < 0.001 as calculated by one-way ANOVA or t test.







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$\gamma\delta$ T cells are the major source of IL-17 in the CNS of 3xTg-AD mice

IL-17 can be produced by CD4⁺ T cells, CD8⁺ T cells, $\gamma\delta$ T cells, group 3 innate lymphoid cell (ILC3), and (natural killer) NKT cells, among other immune subsets (Hatfield and Brown, 2015; Zenaro et al., 2015; Kwong et al., 2017). In our experimental context, CD3⁺ T cells accounted for most of the IL-17⁺ CD45⁺ cells in the meninges, brain, and cLNs of WT and AD mice (Figure 3A). In the healthy steady state, $\gamma\delta$ T cells are known to be the main source of IL-17 in the meninges (Benakis et al., 2016; Ribeiro et al., 2019). Expectedly, IL-17 was mainly produced by $\gamma\delta$ T cells in both WT and AD mice (Figure 3A). Also worth mentioning are other sources of meningeal IL-17, namely CD4⁺ or CD8⁺ T cells as well as CD3⁻ non-T cell leukocytes, including ILC3s (Hatfield and Brown, 2015; Kwong et al., 2017) and neutrophils (Zenaro et al., 2015; Figures 3A and 3B). Accordingly to the observed accumulation of meningeal IL-17⁺ cells at AD onset (Figures 2A and 2B), the percentages of $\gamma\delta$ T cells, of IL-17⁺ cells among $\gamma\delta$ T cells, as well as the numbers of IL-17-producing $\gamma\delta$ $(\gamma \delta 17)$ T cells were significantly higher in the meninges of 5- to 6month-old AD mice compared with aged-matched WT controls (Figure 3B). Interestingly, this accumulation was associated with an increase of the myeloid compartment in the meninges, brain, and cLNs of AD female mice (Figures S1A and S1B). In fact, $\gamma\delta$ T cell numbers positively correlated with the numbers of macrophages, monocytes, and neutrophils in the meninges (Figure 3C).

Of note, $\gamma\delta$ T cells from WT and AD mice displayed a typical signature of IL-17 producers, being enriched for the expression of CCR6, RORyt, and IL-1R, when compared to their a counterparts, as well as a particularly activated phenotype, enriched for CD44^{hi} CD69⁺ cells (Figures S2A and S2B). Importantly, we observed a similar percentage of $\gamma \delta 17$ T cells expressing Ki67, a marker of cell proliferation (Figure S2B). In the healthy meninges, $\gamma \delta$ T cell receptor (TCR) repertoire is mostly restricted to gamma chain variable region 6 (V γ 6) (that was operationally defined by default as double negative for V γ 1 and V γ 4; Ribeiro et al., 2019). In EAE, CNS-infiltrating $\gamma \delta 17$ T cells are mostly restricted to the Vy4⁺ subset (Sutton et al., 2009), whereas, in models of ischemic stroke, infiltrating $\gamma\delta$ T cells bear the V $\gamma6$ chain (Arunachalam et al., 2017). Here, we show that the TCR repertoire of $\gamma \delta 17$ T cells in the meninges and brain of AD mice remained highly biased for V γ 1⁻V γ 4⁻ (assumed as V γ 6⁺) subset, although V γ 4⁺ represented only a small fraction of total γ δ 17 T cells (Figure S2C).

Of note, besides meninges, brain, cLNs, and spleen, we also observed an inflammatory response of the myeloid, NK, and NKT compartments in the liver of both male and female mice



(Figure S3), which was consistent with the previously described amyloidosis in this organ (Marchese et al., 2014).

IL-17 neutralization prevents short-term cognitive deficits in 3xTg-AD mice

Next, we assessed whether the exacerbated levels of IL-17 in the brain and meninges of female 3xTg-AD mice had a detrimental role on cognitive performances. For this, neutralizing anti-IL-17 mAb (anti-IL-17A; 32.5 µg/day) or isotype control IgG1a (immunoglobulin G [IgG]; 32.5 µg/day) was chronically diffused through a micro-pump delivery system into the right ventricle of 3xTg-AD and WT mice. This chronic delivery of anti-IL-17 was made for a period of 6 weeks at a rate of 0.15 μ L/h starting at 3.5 months of age, prior to any detectable cognitive deficit (3.5 months old; Figure 4A). Of note, treated animals did not show any weight alteration (Figure 4B) or particular signs of locomotor impairments or anxiety, as evaluated in the open-field (OF) test or in the elevated plus maze (EPM) (Figures 4C and 4D). Strikingly, at 5 months of age, we observed that anti-IL-17-treated AD mice (AD+alL-17) were able to discriminate between the novel and the familial arm of the Y-maze test, similarly to WT controls (WT+lgG), whereas control IgG-treated AD mice were not (Figure 4E). These data indicate that anti-IL-17 treatment prevents shortterm cognitive decline observed in 5-month-old AD mice when tested in the Y-maze. Conversely, the ICV administration of 300 ng of IL-17 alone to WT mice was sufficient to induce short-term memory deficits (Figure S4A), which demonstrates the potent detrimental effects of dysregulated and exacerbated levels of IL-17 on brain cognitive functions. This notwithstanding, the long-term cognitive impairments assessed in the MWM could not be rescued by the anti-IL-17 treatment (Figures 4F and 4G). These data indicate that early intervention with neutralizing IL-17 is efficient in preventing short-term reference memory deficits, but not the long-term spatial memory defect observed in AD mice at 5 months of age.

To assess whether manipulation of IL-17 could be efficient in altering the progression of the disease until later stages, a set of 3xTg-AD mice were treated for a prolonged period, from 3.5 to 7 months of age, with either control IgG or anti-IL-17 mAb (Figure S5). We verified that the procedure did not impact on the weight, exploratory behavior, or anxiety of animals (Figures S5A–S5C). Interestingly, the anti-IL-17 treatment in the AD group resulted in preservation of short-term memory performance in the Y-maze, whereas IgG alone did not alter the impaired discrimination of AD mice (Figure S5D). Again, no differences were observed in the MWM between mice treated with either IgG or anti-IL-17 (Figures S5E and S5F). Further histological analysis of the amyloid load and of the phosphoTau levels in

(A) Dot plots represent cell populations from indicated gates.

(B and C) Histograms depict percentages (n = 4–17 mice; B) or absolute numbers (n = 3–5 mice; C) from indicated populations.

Figure 2. Accumulation of IL-17⁺ cells in the CNS at the onset of cognitive deficits

Meningeal (Men), brain, cervical lymph nodes (cLNs), and spleen cell suspensions were prepared from 2- to 3- (n = 4-10 for female mice; n = 4-5 for male mice), 5- to 6- (n = 4-17 for female; n = 4-8 for male mice), and 8- to 9-month-old (n = 5-14 for female; n = 4-6 for male mice) WT and 3xTg-AD female (A and C) and male (B and C) mice. Samples were analyzed for the expression of surface CD45 and intracellular IL-17 markers. Live cells were gated using LiveDead Fixable Viability Dye.

Results are representative of 1–4 independent experiments. Error bars, mean \pm SD. *p < 0.05, **p < 0.01, and ***p < 0.001, as calculated by Mann-Whitney or unpaired t test.



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3xTg-AD showed no impact of the anti-IL-17 infusion, measured by 6E10 and AT8 antibody staining, respectively (Figure S5G). Regarding neuronal counting and microglial and astrocytic markers (Nissl, Iba1, and GFAP stainings, respectively), we only observed a slight increase in Iba1 staining upon anti-IL-17 treatment and no changes in the remaining markers. As anticipated in our previous study (Ribeiro et al., 2019), this suggests that synaptic function, rather than amyloid-related pathways, is being targeted by the IL-17-mediated regulation of cognitive behavior.

In AD mouse models, splenomegaly is a common manifestation of amyloidosis that is often accompanied by inflammation (Marchese et al., 2014; Yang et al., 2015). Interestingly, the prolonged anti-IL-17 mAb infusion was also associated with a reduced splenomegaly in AD mice (Figure S5H). We therefore propose that prolonged neutralization of IL-17 in 3xTg-AD mice delays the onset of short-term memory deficits and may have a broader impact on the AD-related inflammatory profile in the periphery, beyond the CNS.

IL-17 neutralization prevents synaptic dysfunction independently of $A\beta$ and tau pathology or BBB disruption

To determine whether anti-IL-17 could be beneficial in the prevention of cognitive deficits by affecting AD-related pathological hallmarks, we assessed A_β and tau pathologies by immunohistochemistry and western blot (WB) analysis (Figures 5A-5C). Immunohistochemical studies showed that 5-month-old anti-IL-17- or IgG-treated 3xTgAD mice equally accumulate mild intraneuronal amyloid precursor protein (APP), although they do not yet display Aß plagues (Figure 5A). Also, total APP levels and C-terminal fragments were similar between AD+IgG, AD+alL-17, and WT+lgG mice, as assessed by WB (Figure 5B). Furthermore, biochemical analysis revealed no major changes in total tau expression (N-ter and C-Ter), dephosphorylated Tau (Tau1), or phosphorylation at multiple Tau epitopes (S396, S199, S262, S404, and AT100 [Thr212 and Ser214]) of AD+IgG, AD+alL-17 when compared with WT+lgG mice (Figure 5C). As previously reported (Oddo et al., 2003b), these data further support that disease onset, characterized by mild cognitive impairments, occurs prior to the establishment of $A\beta$ and tau pathologies. Furthermore, we did not find any sign of BBB disruption in AD mice at this age. Upon intraperitoneal injection of Evans blue, the dye was not detected in the parenchyma of the brains of WT or AD mice, indicating that the BBB is structurally and functionally intact (Figure 5D).

We have previously described the involvement of IL-17 in hippocampal glutamatergic transmission and synaptic plasticity in physiological context (Ribeiro et al., 2019). Therefore, we next



sought to address whether the same synaptic targets were deregulated in this context. To investigate basal synaptic transmission, we generated input/output (I/O) curves by recording excitatory postsynaptic potentials (fEPSPs) from the Schaffer/ CA1 synapse at increasing stimulus intensities (0.8-3 mA). The I/O curve from AD+IgG mice attained a significant lower maximum value compared to WT+IgG mice (Figure 5E), as previously described (Oddo et al., 2003b), that was normalized by the anti-IL-17-chronic infusion. We further measured long-term potentiation (LTP) in the CA1 region, as a functional readout for the neural basis of learning and memory (Bliss and Collingridge, 1993). AD-IgG mice display decreased LTP when compared with WT-IgG mice, as expected (Oddo et al., 2003b). Notably, neutralization of IL-17 in AD mice diminished the LTP impairment (Figure 5F). Conversely, we observed that preincubation of IL-17 (30 ng/mL) was sufficient to significantly impair LTP, as opposed to IL-17 (10 ng/mL) that had no effect (Figure S4B). Building on our previous study (Ribeiro et al., 2019), these data suggest that IL-17 action follows a bell-shaped curve, being crucial for synaptic transmission up to a certain level, from which it becomes deleterious.

Altogether, we concluded that exacerbated levels of IL-17 induce glutamatergic synaptic dysfunction and propose that such mechanism would trigger short-term cognitive impairments in AD mice at disease onset.

DISCUSSION

The impact of immune mediators on cognitive loss associated with AD onset remains elusive (Da Mesquita et al., 2018). In the present study, we show that IL-17-producing cells, mostly $\gamma\delta$ T cells, accumulate in the CNS at the onset of cognitive deficits and persist throughout disease progression. IL-17 neutralization was sufficient to prevent short-term memory deficits and hippocampal glutamatergic dysfunction in early stages of disease, in a mechanism that is independent of A β and Tau pathology or BBB disruption. In later stages of disease, prolonged anti-IL-17 infusion resulted in the delay of cognitive impairment, accompanied by a reduction in peripheral inflammation. Therefore, we propose that elevated levels of IL-17 at early stages of disease contribute to synaptic dysfunction and short-term memory deficits in the 3xTg-AD mouse model.

The role of IL-17 in early stages of Alzheimer's pathology is controversial and poorly explored. If, on one hand, neutralization of IL-17 was shown to rescue A β -induced neuroinflammation and memory impairments (Cristiano et al., 2019), on the other side, it is reported that IL-17A overexpression in AD mouse brain is neuroprotective, decreasing amyloid angiopathy (Yang et al.,

Figure 3. $\gamma\delta$ T cells are a major source of IL-17 in the CNS of 3xTg-AD mice

Meningeal, brain, and cLNs cell suspensions were prepared from 5- to 6- and 8- to 9-month-old WT and 3xTg-AD female and male mice. Samples were analyzed for the expression of surface CD45, CD3, CD4, TCRô, CD11b, F4/80, Ly6C, and Ly6G.

(C) Linear regression plots of the number of $\gamma\delta 17$ T cells and macrophages, monocytes, or neutrophils.

⁽A) Live cells were gated using LiveDead Fixable Viability Dye. Pie charts represent indicated cell populations from indicated gates (n = 4–11 mice).

⁽B) Histograms depict percentages of $\gamma\delta$ T cell (top panels) or IL-17-producing $\gamma\delta$ T cells (n = 5–12 mice; middle panels) and absolute number of IL-17-producing $\gamma\delta$ T cells ($\gamma\delta$ 17 T cells; n = 3–4 mice; lower panels).

Results are representative of 1–3 independent experiments (n = 3–5 mice). Error bars, mean \pm SD. *p < 0.05 and **p < 0.01, as calculated by Mann-Whitney, unpaired t test, and Pearson correlation.







Figure 4. Early IL-17 neutralization prevents short-term cognitive deficits in 3xTg-AD mice

(A) Experimental design: WT and 3xTgAD mice were treated with either neutralizing anti-IL-17 antibody (anti-IL-17A; $32.5 \mu g/day$) or isotype control IgG1a (IgG; $32.5 \mu g/day$), starting at an average age of 3.5 months and continued for 6 weeks, through a micro-pump delivery system (Alzet). Cognitive assays, AD-related hallmarks (amyloid-beta [A β] and tau pathologies), and electrophysiology were performed after this period, in 5- to 6-month-old mice.

(B) Mouse weight in grams (g) after treatment (n = 4–5 mice).

(C) Exploratory behavior evaluated by total distance (cm) in the open field (n = 8–9 mice).

(D) Anxiety in the elevated plus evaluated by the total number of transitions between open and closed arms (n = 8-9 mice).

(E) The discrimination ratio between novel (N) and the other (O) arm was evaluated in the Y-maze test (n = 37-39 mice).

(F) Escape latency during the training phase.

(G) % time in the test quadrant where the platform was in the MWM test (n = 23-29 mice).

Results are representative of 1–6 independent experiments. Error bars, mean ± SEM. **p < 0.01 and ****p < 0.0001, as calculated by one-way ANOVA or t test.

2017). To tackle this question, we took advantage of the tripletransgenic mouse model of AD (3xTg-AD), a progressive model that allows individualizing the different stages of the disease, regarding both cognitive deficits and AD hallmarks. Importantly, it also translates the sexual dimorphism observed in humans, where women are more affected than men in both disease prevalence and symptom progression (reviewed in Fisher et al., 2018).

IL-17 is a recent player in neurophysiology linked to cognitive behavior and anxiety in mice (Ribeiro et al., 2019; Alves de Lima et al., 2020) and described as a neuromodulator of sensory responses in worms (Chen et al., 2017). Here, we show that IL-17-producing cells accumulate in the meninges and in the brain at early stages of the disease, suggesting that AD-related immune alterations might initiate at the onset of cognitive symptoms. In pathology, IL-17 is largely described to mediate inflammation of the CNS (Shichita et al., 2009; Sutton et al., 2009; Gelderblom et al., 2012; Benakis et al., 2016) and was suggested to play a role in AD and PD pathology (Zenaro et al., 2015). More recently, IL-17⁺ CD4⁺ (Th17) cells were described to exacerbate neuroinflammation and neurodegeneration in rodent models of PD. In the same line, elevated production of IL-17 by CD4⁺



Figure 5. IL-17 neutralization prevents synaptic dysfunction independently of $A\beta$ and tau pathology or BBB disruption

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(A) Brain sections from AD+IgG and AD+alL-17 immunostained with 6E10 antibody amplification of hippocampal region of each brain section (n = 3). Scale bars, 1 mm.

(B) Western blot (WB) of APP and C-terminal fragments (CTFs) normalized to GAPDH on the hippocampus of WT+IgG, AD+IgG, and AD+alL-17 (n = 5 mice)

(C) WB of tau (N-ter and C-ter) and tau phosphorylation at S396, S199, S404, and S212/T214 (AT100) epitopes, as well as dephosphorylated tau (tau-1) in the hippocampus of 9-month-old Tau22 mice, as positive control, WT+IgG, AD+IgG, and AD+aIL-17 animals (n = 5 mice).

(D) Quantitative analysis of Evans blue leakage into the brain (μ g/mL) normalized to tissue weight (n = 3-7 mice)

(E) Input/output (I/O) curves corresponding to the fEPSP slope evoked by different stimulation intensities (0.8-3.0 mA) of WT+IgG, AD+IgG, and AD+alL-17; n = 3-6 mice; F-test.

(F) Time course (left panels) and magnitude (right panels) of LTP induced by theta-burst stimulation (TBS) in hippocampal slices from WT+IgG, AD+IgG, and AD+alL-17 animals (n = 3-5 mice). Kruskal-Wallis test followed by Dunn's multiple comparisons test is shown.

Data are mean \pm SD. *p < 0.05.

T cells was also reported in the PD population (Sommer et al., 2018). Although scarce, there is increasing evidence of the involvement of Th17 cells in patients with AD (Saresella et al., 2011; Oberstein et al., 2018) as well as in rodent models (Zhang et al., 2013; Cristiano et al., 2019). On the other hand, in mice subjected to EAE, a multiple sclerosis model, IL-17-producing cells-such as $\gamma\delta$ T cells, Th17 cells, and ILC3-infiltrate the meninges and the brain and, in concert action, amplify a detrimental immune response in CNS (Sutton et al., 2009; Hatfield and Brown, 2015; reviewed in Waisman et al., 2015). We unveil that $\gamma\delta$ T cells, followed by CD3⁺CD4⁻ $\gamma\delta^{-}$ T cells, inferred to be CD8⁺ T cells, are the major sources of IL-17 increasing in AD context. In AD, the detrimental role of CD8⁺ T cells is mostly correlated with Tau pathology-but not amyloid pathology-in both humans (Merlini et al., 2018) and rodent models (Laurent et al., 2017). However, tissue-resident CD8⁺ T cells were described to populate the human healthy brain, contributing to immune surveillance (Smolders et al., 2018) and to expand in the cerebrospinal fluid (CSF) of patients with AD (Gate et al., 2020). Altogether, this suggests a potential proinflammatory role of these cells in early stages of AD.

 $\gamma \delta$ T cells are reported to be impaired in elderly subjects (Colonna-Romano et al., 2004), and upon aging, there is an increased differentiation of T cell into Th17 pathway (Lee et al., 2011; Li et al., 2017). Upon aging, the main risk factor for AD, there is an increase of inflammatory cytokines, such as IL-17, as well as cytokines driving IL-17 secretion, namely IL-1 and IL-23 (Chen et al., 2014; Lim et al., 2014; Barrientos et al., 2015). Of note, we observed an increased expression of the receptor for IL-1 in meningeal yo T cells, when compared to their $\alpha\beta$ T cell counterparts, suggesting a potential contribution of this cytokine in the accumulation of $\gamma \delta 17$ T cells in AD mice. In aged mice, IL-17-producing $\gamma\delta$ T cells were shown to dominate the $\gamma\delta$ T cell pool on lymph node, mainly due to the selective increase of V γ 6⁺ $\gamma\delta$ 17 T cells (Chen et al., 2019). Altogether, these data are consistent with increased circulation of Th17 cells (Saresella et al., 2011; Oberstein et al., 2018) and IL-17 levels in the serum of patients with AD (Chen et al., 2014; Hamdan et al., 2014). In addition, it has been recently reported that, upon aging and in AD models, there is an impairment in brain drainage due to the disruption of the lymphatic vessels, where immune cells reside (Da Mesquita et al., 2018; Ahn et al., 2019). This phenomenon can lead to a breakdown in the homeostasis and circulation of meningeal immune population and may underlie the elevated pool of IL-17⁺ cells observed in our study.

To clarify whether IL-17 could contribute to the initial phase of memory impairment, we neutralized IL-17 before the onset of cognitive deficits, from 3.5-month-old to 5-month-old AD mice. Anti-IL-17 treatment was sufficient to prevent the short-term cognitive deficits observed in the onset of disease in AD models. However, the cognitive deficits in the MWM could not be prevented by the treatment. These go in line with the absence of



role of IL-17 in MWM at steady state, suggesting that other ADrelated mechanisms independent of IL-17 signaling might be in place. For instance, MWM was shown to be dependent on the presence of IL-4⁺ CD4 T cells (Derecki et al., 2010).

The role of IL-17 in AD has been so far described in the context of established Aβ pathology (Zhang et al., 2013; Cristiano et al., 2019). Here, we unveil an early role for IL-17 in AD that precedes the formation Aβ plaques, tau pathology, and BBB disruption, key features of AD. This further supports the involvement of IL-17 in initial stages of the disease. We also show that prolonged treatment with anti-IL-17 prevents the appearance of shortterm cognitive deficits in later stages of disease (7 to 8 months old), suggesting that IL-17 is a triggering event in AD. We hypothesize that an accumulation of meningeal IL-17 may initiate the inflammatory cascade in the brain as well as leukocyte recruitment. Because the BBB is still intact at 5 to 6 months old, cerebral infiltration may occur through the choroid plexus (Bouzerar et al., 2013). IL-17 is a strong chemoattract of neutrophils, which are described to accumulate at the onset of cognitive deficits in the same AD model promoting AD-related pathology (Zenaro et al., 2015). Consistently, we observed a positive correlation between the absolute numbers of $\gamma\delta$ T cells and neutrophils in the meninges, pointing at a potential cross-talk between these two populations upon disease onset, as previously reported in the context of infection (Nakasone et al., 2007; Shibata et al., 2007; Hamada et al., 2008). Neutrophils may act as perpetuators of neuronal damage following initiation of neuroinflammation downstream of IL-17 elevations. It is plausible that neutralization of IL-17 might constrain the recruitment of these and other populations into the CNS. This was previously shown in the pathogenesis of cerebral ischemia upon yo17 T cell infiltration (Shichita et al., 2009). Anti-IL-17 treatment after stroke blocked neutrophil invasion into the brain (Gelderblom et al., 2012). In addition, $\gamma \delta 17$ T cells are reported to amplify Th17 responses (Sutton et al., 2009; Pikor et al., 2015). This could explain why the deleterious role of Th17 cells is only described when AD pathology is fully established.

IL-17 can act directly on interneurons (Chen et al., 2017) and cortical glutamatergic neurons (Alves de Lima et al., 2020), as well as indirectly on glial cells, amplifying neuronal responses through the promotion of brain-derived neurotrophic factor (BDNF) production in the steady state (Ribeiro et al., 2019). Of note, the characterization of BDNF in AD has been largely covered (Tapia-Arancibia et al., 2008), namely in the 3xTg-AD model where BDNF signaling pathway has been shown to be impaired (Corona et al., 2010). This highlights the complexity of BDNF regulation upon neurodegeneration, implying additional and different mechanisms than the one involved at steady state.

Mechanistically, synaptic dysfunction is described to underlie the cognitive deficits observed in 3xTg-AD mice (Oddo et al., 2003b). Here, we demonstrate that anti-IL-17 treatment prevented synaptic deficits observed in basal transmission and long-term potentiation at the onset of short-term cognitive deficits of AD mice. These data suggest that exacerbated levels of IL-17 are detrimental for synaptic signaling, mirroring the findings for other cytokines, such as IL-1 β and tumor necrosis factor alpha (TNF- α), involved in both homeostatic synaptic plasticity and pathology (Avital et al., 2003; Stellwagen and Malenka,

Cell Reports

2006; Ren et al., 2011; Pribiag and Stellwagen, 2013; Prieto et al., 2015). Of note, exacerbated levels of IL-17 were shown to promote gliosis in several different settings (Sarma et al., 2009; Zimmermann et al., 2013; You et al., 2017). On the other hand, we have recently shown that IL-17 promotes synaptic plasticity and short-term memory in the physiological context, by modulating AMPA/NMDA ratio of glutamatergic synapses (Ribeiro et al., 2019). Adding to these findings, this study suggests that IL-17 can mediate opposite effects, seemingly depending on its concentration in the microenvironment, and implies a dual role for IL-17, supporting the idea that IL-17 action follows a bell-shaped curve with a threshold above which IL-17 promotes a deleterious neuroinflammation. Thus, a relevant open guestion remains as "how much of IL-17 is too much?". In an attempt to address this issue, we have titrated the supplementation of IL-17 in hippocampal slices of WT mice, using electrophysiology as a readout, and observed that a preincubation of 30 ng/mL of IL-17 was sufficient to significantly impair LTP, as opposed to a dose 10 ng/mL that had no effect. Of note, this concentration range is in accordance with our previous observation that 10 ng/mL was able to partially rescue LTP in IL-17 knockout (KO) mice (Ribeiro et al., 2019). Consistently, for proof of concept, we could phenocopy the mild cognitive impairment observed in AD mice by injecting a high dose of IL-17 (300 ng/mL) in the brain ventricle of WT mice.

In conclusion, we have described a pathogenic role of IL-17 in promoting synaptic dysfunction and cognitive function underlying the onset of AD. Given the pro-cognitive role of IL-17 in healthy meninges (Ribeiro et al., 2019), our data suggest that IL-17 levels are finely tuned to foster cognition in the steady state, but their pathophysiological dysregulation promotes neurodegeneration. These findings shed light in the early events driving AD pathogenesis, opening perspectives for early biomarkers of disease and combinatorial treatments where the core pathologies (A β plaques and tau pathology) and underlying inflammation are collectively targeted to halt the disease.

STAR*METHODS

Detailed methods are provided in the online version of this paper and include the following:

- KEY RESOURCES TABLE
- RESOURCE AVAILABILITY
 - Lead contact
 - Materials availability
 - Data and code availability
- EXPERIMENTAL MODEL AND SUBJECT DETAILS
 O Ethics statement
 - Mice
- METHOD DETAILS
- Behavioral Tests
 - O OF

 - EPM
 - Y-Maze
 - O MWM
 - Surgical procedures
 - Flow cytometry





- Immunohistochemistry
- Evans Blue Quantification
- ELISA Measurements
- Western Blots
- Electrophysiological fEPSPs recordings
- QUANTIFICATION AND STATISTICAL ANALYSIS

SUPPLEMENTAL INFORMATION

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AUTHOR CONTRIBUTIONS

H.C.B. designed and performed the experiments and analyzed the data, except when otherwise stated, and wrote the manuscript. M.R., J.E.C., S.C.-P., J. Darrigues, J. Dunot, H.M., and A.A.d.A. assisted in some experiments of flow cytometry, electrophysiology, ICV injection, and behavioral tests. R.G. assisted in RNA extraction and RT-PCR experiments, and S.C.-P. assisted in experiments of immune-cell isolation. Histology and western blot experiments were performed by H.C.B. at the University of Lille, INSERM – Lille France, in a PhD exchange period with assistance and collaboration of D.B., E.F., V.G.-M., and K.C. P.A.P. and B.S.-S. assisted in the experimental design and provided key research tools; L.V.L. and J.C.R. coordinated the study, performed some experiments, supervised the research, and wrote the manuscript.

DECLARATION OF INTERESTS

The authors declare no competing interest.

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STAR*METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Brilliant Violet 510 anti-mouse CD45 (Clone 30-F11),	Biolegend	Cat# 103138, RRID AB_2563061
FITC anti-mouse CD45 (Clone 30-F11),	Biolegend	Cat# 103107, RRID AB_312972
Brilliant Violet 711 anti-mouse CD3 (Clone 145-2C11),	Biolegend	Cat# 100349, RRID AB_2565841
APC anti-mouse TCRd (Clone GL3),	Invitrogen	Cat# 17-5711-82, RRID AB_842756
eFluor450 anti-mouse TCRd (Clone GL3),	Invitrogen	Cat# 48-5711-82, RRID AB_2574071
Brilliant Violet 605 anti-mouse CD4 (Clone GK1.5)	Biolegend	Cat# 100451, RRID AB_2564591
PerCP/Cyanine5.5 anti-mouse CD8a (Clone 53-6.7)	Biolegend	Cat# 100734, RRID AB_2075238
PE/Cyanine7 anti-mouse NK1.1 (Clone S17016D)	Biolegend	Cat# 156514, RRID AB_2888852
Brilliant Violet 421 anti-mouse CCR6 (Clone 29-2L17)	Biolegend	Cat# 129818, RRID AB_11219003
PE/Cyanine7 anti-mouse F4/80 (Clone BM8),	Biolegend	Cat# 12311, RRID AB_893478
Brilliant Violet 605 anti-mouse CD44 (Clone IM7)	Biolegend	Cat# 103047, RRID AB_2562451
PE anti-mouse CD69 (Clone H1.2F3)	Biolegend	Cat# 104508, RRID AB_313111
Brilliant Violet 711 anti-mouse CD11b (Clone M1/70)	Biolegend	Cat# 101242, RRID AB_2563310
FITC anti-mouse CD62L (Clone MEL-14)	Biolegend	Cat# 104406, RRID AB_313093
PE/Cyanine7 anti-mouse TCRVg4 (Clone UC3-10A6),	Biolegend	Cat# 137707, RRID AB_10899574
APC anti-mouse TCRVg1 (Clone 2.11),	Biolegend	Cat# 141107, RRID AB_10897806
Brilliant Violet 711 anti-mouse Ly6G (Clone 1A8)	Biolegend	Cat# 127643, RRID AB_2565971
PerCP/Cyanine5.5 anti-mouse Ly6C (Clone HK1.4)	Biolegend	Cat# 128012, RRID AB_1659241
APC anti-mouse IL-1R (Clone JAMA-147)	Biolegend	Cat# 113509, RRID AB_2264757
PE anti-mouse/human Ki-67 (Clone 11F6),	Biolegend	Cat# 151210, RRID AB_2716008
FITC anti-mouse IL-17A (Clone TC11-18H10.1)	Biolegend	Cat# 506908, RRID AB_536010
PE anti-mouse IFN-g (Clone XMG1.2),	Biolegend	Cat# 505808, AB_315402
Brilliant Violet 421 anti-mouse TNF-a (Clone MP6-XT22)	Biolegend	Cat# 506328, RRID AB_2562902
APC anti-mouse RORgt (Clone AFKJS-9)	Invitrogen	Cat# 17-6988-82, RRID AB_10609207
PE/Cyanine7 anti-mouse Tbet (Clone 4B10),	Invitrogen	Cat# 25-5825-82, RRID AB_11042699
inVivoMAb anti-mouse IL-17A (Clone 17F3)	BioXCell	Cat #BE0173, RRID: AB_10950102
InVivoMAb mouse IgG1 isotype control (MOPC-21)	BioXCell	Cat #BE0083, RRID: AB_1107784
anti-Aβ (6E10)	Covance	Cat #SIG-39340, RRID AB_2564652
anti-APP (C17)	This paper	N/A
anti- C-terminal fragments (CTF)	This paper	N/A
anti-Tau N-Ter (M19G)	This paper	N/A
anti-Tau C-ter	This paper	N/A
anti-tau phosphorylation at S199	This paper	N/A
anti-phospho-Tau (Ser202, Thr205) (AT8)	Millipore	Cat # MN1020, RRID: AB_223647
anti-phospho-Tau (Ser396) Polyclonal Antibody	ThermoFisher Scientific	Cat #44-752G, RRID: AB_2533745
anti-phospho-Tau (Ser262) Polyclonal Antibody	ThermoFisher Scientific	Cat#44-750G, RRID: AB_2533743
anti-phospho-Tau (Ser404) Polyclonal Antibody	ThermoFisher Scientific	Cat#44-758G, RRID: AB_2533746
anti-phospho-Tau (Thr212, Ser214) (AT100)	ThermoFisher Scientific	Cat# MN1060, RRID: AB_223652
anti-Tau-1 (PC1C6)	Millipore	Cat #MAB3420, RRID: AB_94855
Peroxidase labeled goat anti-rabbit	Vector Laboratories	PI-1000
Peroxidase labeled horse anti-mouse	Vector Laboratories	PI-2000
Iba1 polyclonal Antibody	Wako	019-197441
GFAP polyclonal Antibody	Dako	Z0334

(Continued on next page)



Continued		
REAGENT or RESOURCE	SOURCE	IDENTIFIER
Chemicals, peptides, and recombinant proteins		
Complete EDTA free protease inhibitor cocktail	Merck Life Sciences	Cat#11873580001
Mouse IL-17A Recombinant Protein,	Invitrogen	Cat #34-8171-82,
Evans Blue	Sigma	E2129, CAS 314-13-6
Cresyl violet acetate	Sigma	C5042, CAS 10510-54-0.
Critical commercial assays		
LIVE/DEAD Fixable Near-IR Dead Cell Stain Kit	ThermoFisher	Cat #L10119
DC protein assay reagent kit,	Biorad	Cat#5000116
DAB (used for A β IHC)	Sigma	D9015
ABC kit (for Aβ IHC)	Vector Laboratories	PK6100
Amyloid beta 40 Human ELISA Kit	ThermoFisher Scientific	Cat # KHB3481
Amyloid beta 42 Human ELISA Kit	ThermoFisher Scientific	Cat # KHB3441
4%-12% NuPage Novex gels	Invitrogen	Cat # NP0323BOX
chemiluminescence kits ECL [™]	Amersham Bioscience	RPN2106
Experimental models: organisms/strains		
3xTg-AD C7BL/6-129SvJ mice	The Jackson Laboratory	Stock No: 34830-JAX,
WT C7BL/6-129SvJ mice	The Jackson Laboratory	Stock No: 101045,
Software and algorithms		
SMART video-tracking	Panlab	RRID SCR_002852
FlowJo	Tree Star	RRID SCR_008520
GraphPad Prism 8.0	GraphPad	RRID SCR_002798
ImageJ software	Scion Software	RRID:SCR_003070

RESOURCE AVAILABILITY

Lead contact

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact Julie C. Ribot (jribot@medicina.ulisboa.pt).

Materials availability

There are restrictions to the availability of home-made antibodies due to our need to maintain the stock. We are glad to share reagents with compensation by requestor for its processing and shipping.

Data and code availability

- All data reported in this paper will be shared by the lead contact upon request.
 - This study did not generate any code.
 - Any additional information required to reanalyze the data reported in this paper is available from the lead contact upon request.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Ethics statement

All handling, surgical, and post-operative care procedures were approved by Instituto de Medicina Molecular Internal Committee (ORBEA) and the Portuguese Animal Ethics Committee (DGAV), in accordance with the European Community guidelines (Directive 2010/63/EU) and the Portuguese law on animal care (DL 113/2013). All efforts were made to minimize the number of animals used in the study.

Mice

C7BL/6-129SvJ mice bearing three mutations (3xTg-AD) associated with familial AD (amyloid precursor protein [APPswe], presenilin-1 [PSEN1] and microtubule-associated protein tau [MAPT]) were purchased from the Mutant Mouse Research and Resource Center at The Jackson Laboratory. Mice were bred and housed at the Instituto de Medicina Molecular animal facility under conventional conditions. Male and female animals were tested at an early stage (2-3 months old (mo), i.e., when no cognitive deficits are observed),





at the onset of disease (5-6 mo, when the cognitive deficits initiate), and at later stages (8-9 mo, when pathology and memory deficits are well established) (Oddo et al., 2003a; Billings et al., 2005, 2007; Giménez-Llort et al., 2007). WT mice from the C7BL/6-129SvJ background were used as controls.

METHOD DETAILS

Behavioral Tests

Cognitive performance, as well as locomotion and anxious behavior were evaluated according to Ribeiro et al. (2019). Mice were handled for 5 days before behavioral tests, which were performed in the following sequence: open-field (OF), elevated plus maze (EPM), Y-Maze and Morris water maze (MWM). Mazes were cleaned with a 30% ethanol solution between each trial. Animals were randomized for the behavioral testing. All behavioral tests were performed during the light phase between 8 a.m. and 6 p.m, under dim light, in a sound attenuated room. Mice movements were recorded and analyzed using the video-tracking software – SMART®.

OF

The mice were placed in the center of a square apparatus, surrounded by vertical walls ($66 \text{ cm} \times 66 \text{ cm} - \text{open-field}$ arena. They freely explored the maze for 5 min. The total distance traveled was determined. At the end of the 5 min test, mice were removed from the open-field arena and placed into its home cage.

EPM

The maze is shaped like a plus sign and consists of two "open" (no walls, $5 \text{ cm} \times 29 \text{ cm}$) and two "closed" 122 ($5 \text{ cm} \times 29 \text{ cm} \times 15 \text{ cm}$) arms, arranged perpendicularly, and elevated 50 cm above the floor. Each animal was placed on the center of the equipment, facing an open arm, and given 5 minutes to explore the maze. The total number of transitions between the open arms and the total arms were used as anxiety and locomotor parameters as previously done in our Institute (Coelho et al., 2014).

Y-Maze

The Y-maze is a two-trial recognition test. Mice are placed in a Y-shaped maze with 3 arms (each with 35 cm length x 10 cm width x 20 cm height), angled at 120° ; on the first trial (learning trial), the animal explored the maze for 10 min with only two arms opened (start and other arm); after returning to his home cage for 1 h, the same animal was re-exposed to the maze for 5 min (test trial) with the novel arm available. The time spent exploring each arm was quantified. Discrimination ratio is calculated dividing time in the N or O arm, by the sum of the time in both arms (N+O).

MWM

MWM was performed as described (Ribeiro et al., 2019) during five consecutive days and consisted of a four day acquisition phase and a one day probe test. During the acquisition phase each mouse was given four swimming trials per day (30-min intertrial interval). A trial consisted of allowing the mouse to explore and reach for the hidden platform. If the animal reached the platform before 60 s, it was allowed to remain there for 10 s; if the animal failed to find the target before 60 s, it was manually guided to the platform, where it was allowed to remain for 20 s. On the probe test, the platform was removed and animals were allowed to swim freely for 60 s while recording the percentage of time spent on each quadrant.

Surgical procedures

The ALZET Brain Infusion Kit is used for the prolonged administration of the antibody against IL-17 (1.36 mg/animal, during 6 weeks at a constant rate of 0.15 μ I/hr), or mouse IgG1 as control. This dose has been determined based on previous studies (Ribeiro et al., 2019; Brosseron et al., 2014). Mice were anesthetized under 1.5% isoflurane in 100% oxygen in a transparent acrylic chamber. After induction, mice were moved to a stereotaxic frame, maintaining isoflurane anesthesia. A small incision was made in the skin between the scapulae, and a pocket was formed by spreading the connective tissues apart, for the subcutaneous placement of the mini-osmotic pump. The cannula was inserted in the right ventricle, in the following stereotaxic coordinates: 0.5 mm anterior-posterior, 1 mm medial-lateral and 2.5 mm dorsal-ventral to Bregma. The skin incision was closed with sutures. Behavior tests were performed 6 weeks after implanting the cannulas, when treatment is concluded and the mice fully recovered from surgery. For the prolonged treatment, the mini-osmotic pump was removed after the initial 6 weeks of treatment, and replaced with a new pump, filled with either alL-17 or mouse IgG1, to prolong the treatment for another 6 weeks. Behavior was tested immediately after the conclusion of this extended treatment period. Alternatively, C57BL/6 WT mice were administered with IL-17 (0.1 mg/ml, total volume of 3 ul) or control saline solution (PBS, total volume of 3 ul) as previously described (Ribeiro et al., 2019). Briefly, mice were anesthetized under 1.5% isoflurane in 100% oxygen. A single intracerebroventricular injection was performed into the right ventricle of the brain using the stereotaxic coordinates of 0.6 mm posterior, 1.2 mm lateral, and 2.2 mm ventral to bregma. A 10 μ L Hamilton syringe was used for intracerebroventricular injection. Behavioral assessment was performed 24 hours after surgery in the Y-maze.



Flow cytometry

Flow cytometry was performed according to Ribeiro et al. (2019). Mice were sacrificed with CO_2 and immediately transcardiacally perfused with ice-cold PBS. Meninges were collected and processed as previously described (Derecki et al., 2010). Brains were cut into 2 mm² pieces and incubated for 30 minutes at 37°C with stirring in RPMI 5% fetal bovine serum (FBS) medium supplemented with collagenase D (1.5 mg/ml; Roche) and DNase I (100 μ g/mL, Roche). Supernatants were collected and live cells were isolated on a gradient of Percoll 70% - 30% (GE Healthcare). Spleens and cLNs were homogenized and washed in RPMI medium 10% FBS. Meninges and cLNs were pooled from up to 3 mice, brains and spleens were analyzed individually.

FACS stainings were performed as previously described (Ribeiro et al., 2019) using indicated monoclonal antibodies (mAbs). Dead cells were excluded using LiveDead Fixable Viability Dye (Invitrogen). Samples were acquired using FACSFortessa (BD Biosciences). Data were analyzed using FlowJo software (Tree Star).

Immunohistochemistry

For immunohistochemical studies, females were deeply anaesthetized with pentobarbital sodium (50 mg/kg, intraperitoneally), then transcardially perfused with cold NaCl (0.9%) and with 4% paraformaldehyde in PBS (pH 7.4). Brains were removed, post-fixed for 24 h in 4% paraformaldehyde and cryoprotected in 30% sucrose before being frozen at -40° C in isopentane (methyl-butane) and stored at -80° C. Coronal brains sections (35 µm) were obtained using a Leica cryostat. Free-floating sections were chosen according to the stereological rules, with the first section taken at random and every 12 sections afterward, and were stored in PBS-azide (0.2%) at 4°C.

For A β immunohistochemistry (IHC), sections were pretreated with 80% formic acid for 3 min and were permeabilized with 0.2% Triton X-100/sodium phosphate buffer. Sections were then blocked with 10% "Mouse On Mouse" Kit serum (Vector Laboratories) for 1 h before incubation with mouse biotinylated anti-A β antibody (6E10) at 4°C overnight. After washing in PBS, the sections were incubated with the ABC kit (Vector Laboratories) for 2 h and developed using DAB (Sigma). Images were acquired using Leica ICC50 HD microscope. Quantification of the 6E10 staining intensity was performed using Mercator software (Explora Nova, Mountain View, CA, USA). The number of plaques, the average plaque size and the plaque burden, expressed as percentage of analyzed area, were calculated in the cortex and hippocampus of the 3xTg-AD mice.

Evans Blue Quantification

3xTg-AD and C7BL/6-129SvJ WT mice, previously treated with either alL-17 or IgG1, were injected *i.v.* with 1% Evans Blue dye (Sigma), and perfused with PBS 45-60 min after injection. The brain was then removed, weighted, and incubated at 37°C for 48h in N,N-dimethyl formamide. A standard curve is prepared by serially diluting the Evans Blue dye from 50 to 2,5 µg/ml. The content of dye was determined by spectrophotometer at 620 nm, subtracting absorbance at 740 nm. Evans Blue concentration was normalized to brain weight.

ELISA Measurements

Brain levels of human A β 1-40 and A β 1-42 were measured using ELISA kits (Invitrogen, Carlsbad, CA, USA; IBL-International, Hamburg, Germany) following manufactured' instructions. Briefly, for hippocampal and cortical samples, 20 µg of protein were diluted in Guanidine/Tris buffer (Guanidine HCl 5 M and Tris 50 mM pH 8), sonicated and incubated for 1h at 4°C under agitation. Samples were then diluted in a BSAT-DPBS solution (KCl, KH₂PO₄, NaCl, Na₂HPO₄, BSA 5%, Tween-20 0.03% pH 7.4). The homogenates were centrifuged at 12,000 g for 15 min at 4°C. Supernatants were collected for the analysis of A β ₁₋₄₀ and A β ₁₋₄₂ by colorimetric immunoassays. Absorbance was measured in a TECAN Infinite 200 plate reader. The amounts of A β were expressed as pg/mg of total protein.

Western Blots

For all biochemical experiments, tissue was homogenized in 200 µL Tris buffer (pH 7.4) containing 10% sucrose and protease inhibitors (Complete; Roche Diagnostics GmbH), sonicated, and kept at −80°C until use. Protein amounts were evaluated using the BCA assay (Pierce), subsequently diluted with LDS 2X supplemented with reducing agents (Invitrogen) and then separated on 4%–12% NuPage Novex gels (Invitrogen). Proteins were transferred to nitrocellulose membranes, saturated (5% non-fat dry milk or 5% BSA) in TNT (Tris 15 mM pH 8, NaCl 140 mM, 0.05% Tween) and incubated with primary (APP and C-terminal fragments (CTF), home-made, 1:5000), tau (tau-1 Millipore #MAB3420, 1:10000, N-Ter, C-ter, home-made, 1:10000) and tau phosphorylation at S396 (Invitrogen #44-752G, 1:10000), S199 (home-made, 1:2000), S262 (Invitrogen #44-750G, 1:1000), S404 (Invitrogen #44-758G, 1:10000) and S212/T214 (AT100, Invitrogen #MN1060, 1:1000) overnight and then corresponding secondary antibodies (peroxidase labeled horse anti-rabbit 1/5000 or anti-mouse 1/50,000, Vector Laboratories). Immunoreactivity was visualized using chemiluminescence kits (ECLTM, Amersham Bioscience) and a LAS3000 imaging system (Fujifilm). Results were normalized to GAPDH (Sigma-Aldrich) and quantifications were performed using ImageJ software (Scion Software).

Electrophysiological fEPSPs recordings

Electrophysiological fEPSP recordings were performed according to Ribeiro et al. (2019), as described in the previous section. Briefly, 3xTg-AD females and WT controls were sacrificed by cervical dislocation, the brain was rapidly removed and the hippocampi





dissected free in ice-cold Krebs solution (124 mM NaCl; 3 mM KCl; 1.25mM NaH2PO4; 26 mM NaHCO3; 1 mM MgSO4; 2 mM CaCl2 and 10 mM D-glucose), previously gassed with 95% O2 and 5% CO2, pH 7.4. 400 µM transverse hippocampal slices were obtained with a McIlwain tissue chopper and field excitatory postsynaptic potentials (fEPSPs) were recorded in stratum radiatum of the CA1 area as previously described (Ribeiro et al., 2019). After obtaining a stable 10 minutes baseline, Input/Output (I/O) curves and long-term potentiation (LTP, 10c trains with 4 pulses at 100 Hz separated by 200 ms, induced at 0.5mV/ms) were recorded. Recordings were performed at 32°C, 3 mL/min. For the titration experiments, hippocampal slices were pre-incubated with IL17 (10ng/mL or 30ng/mL; Ebioscience) for 1 hour before LTP induction.

QUANTIFICATION AND STATISTICAL ANALYSIS

Data were analyzed using GraphPad Prism 8.0 (GraphPad, San Diego, CA). The values presented are mean \pm SEM of n independent experiments. To test the significance of the differences between 2 conditions, a Student's t test, Mann-whitney and F-Test were used. In statistical tests between 3 or more conditions, a one-way ANOVA or Kurskal-Wallis Test followed by a Bonferroni's or Dunnett's multiple comparison post hoc test as specified in the figure legends. *P*-values of < 0.05 were considered to be statistically significant. Sample sizes and p values can be found in figure legends.