Caspase-1 inhibition alleviates cognitive impairment and neuropathology in an Alzheimer’s disease mouse model

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Alzheimer’s disease (AD) is an intractable progressive neurodegenerative disease characterized by cognitive decline and dementia. An inflammatory neurodegenerative pathway, involving Caspase-1 activation, is associated with human age-dependent cognitive impairment and several classical AD brain pathologies. Here, we show that the nontoxic and blood–brain barrier permeable small molecule Caspase-1 inhibitor VX-765 dose-dependently reverses episodic and spatial memory impairment, and hyperactivity in the J20 mouse model of AD. Cessation of VX-765 results in the reappearance of memory deficits in the mice after 1 month and recommencement of treatment re-establishes normal cognition. VX-765 prevents progressive amyloid beta peptide deposition, reverses brain inflammation, and normalizes synaptophysin protein levels in mouse hippocampus. Consistent with these findings, Caspase-1 null J20 mice are protected from episodic and spatial memory deficits, neuroinflammation and Aβ accumulation. These results provide in vivo proof of concept for Caspase-1 inhibition against AD cognitive deficits and pathologies.
Currently, no treatment significantly delays, or reverses AD-associated cognitive decline or pathologies. Targeting earlier degenerative events may be essential to stem the scourge of AD. Preclinical involvement of inflammation in AD, genetic association of immune-related genes with AD, and increased risk of AD in systemic and CNS conditions that increase inflammation support the hypothesis that inflammation is an integral part of AD.3 In addition, increasing support for the role of inflammation in AD is provided from AD mouse models.

AD mouse models on complement 37, colony stimulating factor 1 receptor 34, inflammasome Nod-like receptor protein 3 (Nlrp3) 38, Caspase-1 (Casp1) 39, scavenger receptor class B type 1 40, poly (ADP-ribose) polymerase 1 41, or complement 1q9 42 null background, show improvements in cognitive behaviour, synaptic pathology, and often amyloid beta peptide (Aβ) pathology. Inhibition of inflammasome activator P2X7R decreases Aβ9, and silencing of inflammasome Nod-like receptor protein 1 (Nlrp1) improves cognition in AD models 43.

We identified a human neurodegenerative pathway, mediated by neuronal Nlrp1 inflammasome activation of Casp1, which then activates Caspase-6 (Casp6), in stressed CNS human primary neuron cultures and AD brains.11 Active Casp6 is abundant in neuritic plaques, neurofibrillary tangles, and neurophil threads of sporadic and familial AD brains 12,13, and Nlrp1 immunopositive neurons are increased 20–25-fold in AD14. Entorhinal cortex and hippocampal CA1 Casp6 activation is associated with age-dependent cognitive impairment in humans 15,16 and in transgenic mice in the absence of other AD pathologies 16. Active Casp6 is associated with axonal degeneration 17–19, cleaves several cytoskeleton or cytoskeletal-associated proteins including Tau, α-Tubulin, Drebrin, Spinophilin, Actinin-1 and -4 synaptic proteins 20, increases Aβ production 21–23 and impairs valosin-containing protein p97 proteasome-mediated misfolded protein degradation 24. Consequently, Nlrp1, Casp1, and Casp6 represent feasible therapeutic targets against age-dependent cognitive deficits and AD.

While there are currently no suitable inhibitors for Nlrp1 and Casp6, VX-765, a pro-drug that is rapidly metabolized in vivo to VRT-043198, is a potent bioavailable, blood–brain barrier permeable, and nontoxic Casp1 small molecule inhibitor.25,26 VX-765 is reported to be selective for Casp1 against Casp3–10, Casp14, granzyme B, trypsin and cathepsin B26,27. VX-765 inhibits IL-1β release and IL-18 in activated peripheral blood mononucleated cells and in lipopolysaccharide-induced inflammation, oxalozone-induced dermatitis, and rheumatoid arthritis mouse models 27. VX-765 blocks HIV-induced pyroptosis of CD4 T cells 28 but lacks significant antiapoptotic activity against hypoxia or Fas-induced apoptosis, eliminating the risk of potential inhibition of physiological apoptotic events in vivo.27 VX-765 is safe for humans by oral administration as tested in a phase 2b human clinical trial against epilepsy 29. Therefore, VX-765 represents a safe drug that could rapidly be tested against human age-dependent cognitive impairment and AD.

Here, VX-765 was given to the J20 APPSw/Ind transgenic mouse model 30. J20 mice show episodic memory deficits by 3–4 months, spatial memory and learning deficits by 6–7 months 31, increased total hippocampal Aβ42 levels at 2 months of age, Aβ plaque formation by 7 months 30, and synaptophysin loss by 2–6 months 30. Microglial inflammation and neuronal loss preclude plaque deposition 32. The early manifestation of these symptoms and pathologies allows rapid analyses of the VX-765 effects.

Results
VX-765 rescues cognitive deficits in symptomatic J20 mice.
VX-765 and VRT-043198 potently and specifically inhibited human Casp1 (IC50 3.68 nM and 9.91 nM) relative to human Casp2–10 (see Supplementary Fig. 1a, b). Similarly, mouse recombinant Casp1 (IC50 52.1 nM and 18 nM) was strongly inhibited compared to inflammatory mouse Casp11 (see Supplementary Fig. 1c, d). Although variable between individual mice, VX-765 crossed the blood–brain barrier of WT and J20 mice, was metabolized into VRT-043198, and reached physiologically active concentrations in both the hippocampus and cortex (see Supplementary Fig. 1e).

Five-month-old mice were behaviourally and longitudinally assessed before treatment (baseline), after 3 injections per week with 50 mg kg⁻¹ VX-765 (Treatment 1; T1), after an additional 2 weeks of injections (Treatment 2; T2), after 4 weeks without treatment (Washout; WO), and after 3 more injections per week (Treatment 3; T3) before sacrificing the mice at 8 months of age (Fig. 1a). At baseline, J20 and littermate WT mice showed normal motivation behaviour determined by % time moving (see Supplementary Fig. 2a) and did not exhibit thigmotaxis indicative of anxiety (see Supplementary Fig. 2b), but J20 mice showed a strong deficit in the novel object recognition (NOR) episodic (retention) memory discrimination index (Fig. 1b). J20 NOR deficits were reversed and reached near-normal levels after VX-765 T1 and T2, reappeared after WO, and renormalized after T3. Results were consistent across individual mice (see Supplementary Fig. 2c). J20 mouse hyperactivity, measured by distance travelled in the open field task, was attenuated by VX-765 after T2, reappeared after WO, and again was significantly reduced after T3 (Fig. 1c and see Supplementary Fig. 2d). At T2, J20 mice showed learning acquisition deficits in the Barnes maze spatial memory training phase evidenced by increased primary errors to identify the target correctly compared to WT mice. This deficit was not significantly attenuated by VX-765 (Fig. 1d). Primary latency to find the target did not differ between the three groups (see Supplementary Fig. 2e). During the probe test, primary latency, primary errors (Fig. 1e) and the ability to find the target (Fig. 1f) were clearly impaired in vehicle-injected J20 compared to WT mice. VX-765 eliminated these spatial memory deficits in J20 mice. After WO, vehicle-injected J20 showed learning deficits early on, but all groups performed equally well by the end of training (Fig. 1g and see Supplementary Fig. 2f). The difference in WO primary latency compared to T2 is likely due to a learning effect by repetition or a reduction of stress to the apparatus. VX-765-injected, but not vehicle-injected, J20 mice were normal in primary latency and errors during the probe, suggesting spatial memory retention even after 1 month without drug (Fig. 1h). VX-765-injected mice also performed better than vehicle-injected J20 mice in their ability to find the target (Fig. 1i). J20 mice performance in the Y maze working memory task was consistently low but not always statistically different from WT or treated J20 mice (see Supplementary Fig. 2g).

VX-765 was administered at 25 and 10 mg kg⁻¹ to assess the dose-response of J20 to VX-765 (Fig. 2). All groups showed normal motivation (see Supplementary Fig. 3a) and thigmotaxis behaviour (see Supplementary Fig. 3b). NOR discrimination index was normalized with 25 mg kg⁻¹ dose at T1 and T2, deficits reappeared after WO, and returned to normal at T3, similar to the results with the 50 mg kg⁻¹ dose (Fig. 2a and see Supplementary Fig. 3c). Mice treated with 10 mg kg⁻¹ showed normalized NOR behaviour only after T2, the effect was lost with WO, and reappeared after T3. J20 hyperactivity showed a nonsignificant improvement at T2 with 25, but not 10, mg kg⁻¹ (see Supplementary Fig 3d, e). In the T2 Barnes maze, learning acquisition did not change after VX-765 treatment by the end of the training phase (Fig. 2b and see Supplementary Fig. 3f), and J20 remained impaired in the probe primary latency at both the 25 and 10 mg kg⁻¹ dose (Fig. 2c). However, primary errors were
decreased with 25 mg kg$^{-1}$ and both doses showed that the J20 mice had improved ability to recognize the target during the probe (Fig. 2d). After WO, primary latency was shorter initially during training (see Supplementary Fig. 3g), but no differences in learning were observed among the three groups by the end of the training (Fig. 2e). No differences were observed in primary latency and errors, with or without drug during the probe (Fig. 2f). However, mice treated with 25 mg kg$^{-1}$, but not 10 mg kg$^{-1}$, retained their ability to recognize the target (Fig. 2g).

To confirm that VX-765's effect on re-establishing normal cognition in J20 mice was due to Casp1 inhibition, J20 mice were generated with a Casp1 null background (J20/Casp1$^{-/-}$) and
Two-way repeated-measures ANOVA, Dunnett 

\( F(3,24) = 20.42, p < 0.0001 \)

Vehicle-treated (black circles), and 50 mg kg\(^{-1}\) (blue triangles), 25 mg kg\(^{-1}\) (pink triangles), or 10 mg kg\(^{-1}\) (purple hexagon) VX-765-treated J20 mice. Each mouse tested is represented by one symbol. Data represent mean and s.e.m.

**Fig. 2** VX-765 dose-dependently restores cognitive function in J20 mice. **a** NOR discrimination index at T1 (\( F(3,24) = 20.42, p < 0.0001 \)), T2 (\( F(3,23) = 12.83, p < 0.0001 \)), WO, and T3 (\( F(3,23) = 8.828, p = 0.0004 \)). Vehicle-treated (black circles), and 50 mg kg\(^{-1}\) (blue triangles), 25 mg kg\(^{-1}\) (pink triangles), or 10 mg kg\(^{-1}\) (purple hexagon) VX-765-treated J20 mice. Each mouse tested is represented by one symbol. Data represent mean and s.e.m.

Increased Iba1-positive microglia in the hippocampus and cortex of 8-month-old vehicle-treated J20 mice, compared to vehicle-treated WT mice, were normalized in VX-765-treated J20 mice, indicating that VX-765 reversed inflammation (Fig. 4a).

This was confirmed quantitatively as the number of Iba1-immunopositive microglia measured from the pyramidal cell layer to the stratum lacunosum moleculare (SLM) layer of the hippocampal CA1 region, or from the cortical retrosplenial area to the S1, returned to normal after VX-765 treatment (Fig. 4b). Quantitation of microglial activation levels based on morphological measurements (see Supplementary Fig. 5a) showed decreased type I resting microglia and increased type II and III activated microglia in vehicle-treated J20 hippocampus and cortex compared to vehicle-treated WT and VX-765-treated J20 mice (Fig. 4b). Iba1-activated microglia in vehicle-treated J20 compared to baseline or VX-765-treated mice (see Supplementary Fig. 5b) or other inflammatory brain protein levels, although TNF-\( \alpha \), KC-GRO, and IFN-\( \gamma \) were slightly elevated in vehicle-treated J20 and normalized after VX-765 treatment (see Supplementary Fig. 5c).
Analyses of activated microglia were conducted in the J20/WT, WT/Casp1−/− and J20/Casp1−/− to confirm that VX-765 effects occurred through inhibition of Casp1. The microglia in J20/Casp1−/− were as activated as those of J20/WT (Fig. 4a and see Supplementary Fig. 5d) but as observed in VX-765-treated J20 (Fig. 4b), the number of Iba1-positive microglia increased in J20/WT compared to WT and were normalized in J20/Casp1−/− (Fig. 4c). However, in contrast to VX-765 treatments, the absence of Casp1 did not modify the distribution of the subtypes of microglia; fewer resting microglia at the expense of increased activated microglia was observed in J20/Casp1−/− mimicking levels of the J20/WT. IL-1β levels were unchanged in J20/Casp1−/− compared to J20/WT and surprisingly these were not different from WT mice. These results suggest that cremophor vehicle injections in the VX-765 study may have an inflammatory effect that exacerbates APP-mediated inflammation in J20.

Increased GFAP immunopositive astrogliosis in the hippocampus and cortex of J20 mice was normalized to WT levels with VX-765 treatment and absence of Casp1 (Fig. 4d–f). Overall, these results indicate that VX-765 treatment reversed both microglial and astroglial activation in J20 brains.

**VX-765 prevents Aβ accumulation in J20 mice brains.** To determine if APP or Aβ levels were altered by VX-765 treatment or the absence of Casp1 in J20 mice, western blots against full length and C-terminal fragments (CTF) of APP, immunohistochemistry with an anti-Aβ1–40 antiserum, and ELISA measurement of Aβ1–40, 42 were conducted. Thioflavin S staining of histological brain sections confirmed the amyloid status of Aβ immunoreactivity. Thioflavin S and immunopositive Aβ levels were substantially reduced in VX-765-treated compared to vehicle-injected J20 hippocampus and cortex (Fig. 5a, b and see Supplementary Fig. 6a). However, Aβ deposits did not completely disappear and were comparable to those observed in baseline J20 brains (Fig. 5a). These remaining Aβ deposits were mainly localized in the SLm and dentate gyrus regions of the hippocampus (see Supplementary Fig. 6a). Levels of immunopositive Aβ plaques were similarly reduced in the J20/Casp1−/− hippocampus and cortex (Fig. 5a, c and see Supplementary Fig. 6b). In addition, a much lower level of Aβ was observed in untreated J20 (J20/WT; Fig. 5c) compared to vehicle-injected J20 (Fig. 5b), suggesting an exacerbation of Aβ deposition with the cremophor vehicle injections.

**Fig. 3 Casp1 KO restores cognitive function in J20 mice.** a–e Genotypes: J20+/−/Casp1+/−/ WT/WT (grey squares), J20+/−/Casp1+/−/ WT/Het (pink diamonds), J20+/−/Casp1+/−/ WT/KO (blue symbol), J20+/−/Casp1+/−/ J20/WT (black circles), J20+/−/Casp1+/−/ J20/Het (purple triangles), J20−/−/Casp1+/−/ J20/KO (blue triangles). Each mouse tested is represented by one symbol. Data represent mean and s.e.m. a NOR discrimination index (F(5,263) = 16.22, p < 0.0001) and b distance travelled during open field task (F(5,67) = 3.717, p = 0.005). c–e Barnes maze: c # of primary errors during learning acquisition (Genotype, F(5,263) = 6.469, p < 0.0001; Training day, F(3,263) = 29.44, p < 0.0001), d probe primary latency and errors (Primary errors, F(5,66) = 4.8, p = 0.008) and e target preference. a, b, d ANOVA and e two-way ANOVA, Dunnett’s post-hoc versus WT/WT. *p < 0.05, **p < 0.01, ****p < 0.0001.
RIPA-soluble $\alpha$-soluble $\beta_42$/total $\beta_38+40+42$ ratios were reduced to baseline J20 levels in the hippocampus and less obviously in the cortex of VX-765-treated mice (Fig. 5d), whereas total $\beta$ levels did not change (Fig. 5e). Formic acid-soluble $\alpha$-soluble $\beta_42$/total $\beta_38+40+42$ ratios were reduced in the hippocampus, but not in the cortex (Fig. 5f), whereas total $\beta$ levels were more variable than RIPA $\beta$ levels but tended towards a reduction in both hippocampus and cortex (Fig. 5g). VX-765 treatment had no significant effect on $\beta_38$ or $\beta_40$ subtypes in hippocampus and cortex (see Supplementary Fig. 6c, d). The reduction in $\beta_42$ levels with VX-765 treatment was not due to decreased APP mRNA (see Supplementary Fig. 7a) or protein levels measured with the...
human-specific 6E10 antibody (Fig. 5, m) or the mouse and human anti-CTF antibody (see Supplementary Fig. 7c, d).

RIPA-soluble Δβ42/total Δβ38+40+42 ratios were reduced in the hippocampus and cortex in J20/Casp1+/− and J20/Casp1−/− mice, with no changes in total Δβ levels although there was a trend towards an increase in the hippocampus (Fig. 5h, i). These reduced ratios were due to increased Δβ38 and Δβ42 rather than decreased Δβ42 levels in J20/Casp1−/− (see Supplementary Fig. 6e). J20/Casp1+/− and J20/Casp1−/− mice had similar formic acid-soluble Δβ38/total Δβ38+40+42 ratios, total Δβ levels (Fig. 5j, k), and Δβ subtypes in the hippocampus or cortex (see Supplementary Fig. 6f). The absence of Casp1 in J20 mice had no effect on APP protein levels (Fig. 5n, o and see Supplementary Fig. 7e, f).

To determine if Δβ clearance might have increased with VX-765 treatment or the absence of Casp1, we measured the levels of Δβ-degrading nephrilysin (Nep) and insulin degrading (IDE) enzymes. Increased degradation of Δβ was unlikely since IDE or Nep protein (see Supplementary Fig. 8a, b) and mRNA levels (see Supplementary Fig. 8c, d) were unchanged with VX-765 treatment or the absence of Casp1 (see Supplementary Fig. 8e, f). Together, these results indicate that VX-765 treatment and the absence of Casp1 stopped the progressive deposition of Δβ in J20 brains.

VX-765 does not alter Casp8 or Casp9 processing in J20 mice. Casp5, Casp8, Casp9 and Casp10, which were inhibited in the nanomolar range in vitro by VX-765 and VRT-043198, were considered to determine whether VX-765 could affect other caspases in J20 mice (see Supplementary Fig. 9). Since Casp5 and Casp10 are not expressed in mouse brain, they were not examined. Pro-Casp9 levels were significantly reduced in the hippocampus, but not in cortex, of vehicle-treated J20 compared to WT mice, but normalized after VX-765 treatment (see Supplementary Fig. 9a). This could indicate that pro-Casp9 has been converted to active subunits, and that activation was inhibited by VX-765. Yet, activated Casp9 subunits could not be detected in the mouse brain. Since Casp3 is an excellent substrate of Casp9, we verified if Casp3 was processed into its active subunits in the hippocampal tissues. No active Casp3 subunit was observed in vehicle- or VX-765-treated J20 hippocampus, and there was also no effect of VX-765 on pro-Casp3 levels. Cleaved Casp8 20 kDa active subunit could not be detected in the hippocampus in either WT or J20 mice. A faint band corresponding to 43 kDa cleaved Casp8 active subunit (p43) was detected in the cortex, but its levels remained unaffected after VX-765 treatment (see Supplementary Fig. 9a).

Because processed Casp8 and Casp9 levels in the adult mouse brain were low, VX-765’s potential effects on Casp8 and Casp9 were further analysed in the spleen, which expresses higher levels of these enzymes and is known to naturally contain abundant active Casp8 (see Supplementary Fig. 9b, c). There were no differences in active Casp8 subunits, pro-Casp9 levels, or Casp8 or Casp9 substrate pro-Casp3 and active Casp3 20 kDa subunit, among the three groups (see Supplementary Fig. 9c). Together, these results suggest that VX-765 did not inhibit Casp8 and Casp9 in J20 mice.

VX-765 normalizes synaptophysin in J20 mice brains. At 5 months of age, J20 hippocampus showed decreased immunopositive synaptophysin levels (Fig. 6a). Synaptophysin levels remained low in 8-month-old vehicle-treated J20 hippocampus but were significantly increased and returned to normal levels in VX-765-treated mice hippocampus (Fig. 6a, b). There was no significant difference in the cortex. Measures of 84 different synaptic gene mRNA levels in vehicle-treated J20 hippocampi revealed significantly increased mRNA levels in three genes involved in synaptic function: Camk2a, Grin2b, Kif17 (Fig. 6c). Levels were decreased with VX-765 treatment. Additionally, TNF-α mRNA was increased in J20 and VX-765-treated J20 hippocampi (Fig. 6c), whereas TNFα protein levels increased slightly in J20 mice hippocampi and returned to normal with the VX-765 treatment (see Supplementary Fig. 5c). These results indicate a normalizing effect of VX-765 on several synaptic components that may account for the reversal to normal cognition.

VX-765 prevents axonal degeneration in human neurons. To determine if VX-765 can protect neurons against neurodegeneration, VX-765 was assessed in human CNS human primary neuron cultures (HPN). Treatment of HPN with 25, 50, 100, or 200 µM VX-765 for 72 h was not toxic (Fig. 7a). As previously described19, EGFP was homogeneously distributed within the cell body and neurites of HPN, whereas coexpression with APPWT resulted in EGFP-positive beading indicative of neurodegeneration (Fig. 7b). HPN were pretreated with VX-765 for 1 h before APP transfection and treatment continued thereafter. VX-765 treatment at 25 and 50 µM prevented neurotropic beading in APPWT-transfected neurons at 48 and 72 h post transfection, similar to the Casp1 Z-YVAD-fmk peptide inhibitor (Fig. 7c). Similarly, VX-765 protected against, albeit less strongly, serum-deprivation-induced neuritic beading. To assess if neuritic beading was reversible, VX-765 was administered 48 h after APPWT transfection or serum deprivation. VX-765 treatment did not reverse but prevented further neuritic beading in APPWT-transfected neurons. In contrast, Z-YVAD-fmk was unable to reverse or prevent neuritic beading at any time point. The 50 µM VX-765 reduced secreted and cellular Δβ42/Δβ38+40+42 levels, whereas 25 µM reduced secreted, but not cellular, Δβ42/total Δβ levels
IL-1β and TNF-α levels were significantly increased after serum deprivation but reduced after VX-765 (Fig. 7e). These results indicate that VX-765 can prevent human neuronal degeneration.

Discussion

These results strongly indicate that VX-765 is a promising drug against AD. Our idea of using Casp1 as a therapeutic target is based on the identification of the Nlrp1-Casp1-Casp6 neuronal...
Fig. 5 VX-765 prevents progressive 
Aβ accumulation in J20 mice. a Aβ micrographs of hippocampal SLM and S1 cortex (scale bar = 50 μm). b Quantitative analysis comparing Aβ immunostaining density between J20 + vehicle (black circles) and J20 + VX-765 (blue triangles) mice in the CA1 hippocampal pyramidal cell layer to the SLM (p = 0.0029) and cortex (p = 0.0314, unpaired t test). c Quantitative analysis comparing Aβ immunostaining density between J20/−/−/Casp1−/−/+ J20/WT (black triangles) and J20/−/−/Casp1−/−/− J20/KO (blue triangles) mice in the CA1 hippocampal cell layer (p = 0.0040) and cortex (0.0276, unpaired t test). d–g RIPA- (d, e) and formic acid- (f, g) soluble Aβ levels in the hippocampus and cortex of 5-month baseline J20/−/− (black diamonds), J20 + vehicle (black circles), and J20 + VX-765 (blue triangles) mice. d, f Aβ1-42/ Aβ1-40 + Aβ1-38 levels (RIPA soluble, hippo: F(2,17) = 27.85, p < 0.0001) and e, g total Aβ ANOVA, Dunnett’s post-hoc versus J20 + vehicle. ***p < 0.001, ****p < 0.0001. h–k J20/−/−/Casp1−/−/+ J20/WT (black circles), J20/−/−/Casp1−/−/− J20/KO (blue triangles) RIPA- (h, i) and formic acid- (j, k) soluble Aβ levels in the hippocampus and cortex of J20/WT, J20/Casp1+/− (J20/He) and J20/Casp1−/− J20/KO mice. h, j Aβ1-42/ Aβ1-40 + Aβ1-38 levels (RIPA-soluble, hippo: F(2,26) = 16.03, p < 0.0001; RIPA-soluble, cortex: F(2,26) = 6.602, p < 0.0048) and i, k total Aβ ANOVA, Dunnett’s post-hoc versus J20/WT, ***p < 0.001, ****p < 0.0001, ***p < 0.001. l–o Human APP protein levels (6E10 immunostaining) and quantification in the hippocampus and cortex of l, m J20 + vehicle (black circles) and J20 + VX-765 mice (blue triangles), and n–o J20/WT (black circles), J20/He (purple triangles), and J20/KO (blue triangles) mice.

Fig. 6 VX-765 reverses loss of synaptophysin in J20 mice. a Synaptophysin immunopositive micrographs. Scale bar = 200 μm for hippocampus and 50 μm for cortex. b Synaptophysin immunopositive density in WT + vehicle (grey squares, n = 4), J20 + vehicle (black circles, n = 4), and J20 + VX-765 (blue triangles, n = 4) hippocampus (F(2,9) = 7.974, p = 0.0102, ANOVA, Tukey’s post-hoc, **p < 0.01) and cortex. c Significantly altered (Kruskall–Wallis) synaptic protein mRNA levels in vehicle or VX-765-treated J20 mice hippocampi (n = 3 per group)

degeneration pathway in CNS primary human neurons, the association of this pathway with memory and cognitive impairment in aged humans and mice1–16,36,37, and the genetic association between NLRP1 and CASP1 with AD progression or AD38,39. VX-765 effect is exquisitely rapid against episodic and spatial memory impairment, which are normalized after only 1 (3 injections) or 3 (9 injections) weeks of treatment, respectively. Reversal of cognitive impairment is accompanied by normalized microglial and astroglial reactivity and synaptophysin immunohistochemical staining, a normalization of gene expression of four different synaptic components, and the prevention of progressive amyloid pathology in the brain. Target engagement for VX-765 against Casp1 was confirmed in the J20 Casp1 null mice, which also exhibited normal episodic and spatial memory, and reduced inflammation and Aβ accumulation. To our knowledge, this drug provides the most rapid response observed with a preclinical treatment in the J20 AD mouse model40–42.

Casp1 represents a feasible therapeutic target against age-dependent cognitive impairment and AD. Rare naturally occurring human CASP1 variants associated with very low activity, in vitro and in vivo, suggest that inhibition of Casp1 activity is compatible with normal human life.44,45. In these CASP1 variants, proinflammatory signalling still occurs through NFKB.46. Casp1 null mice are fertile and healthy indicating that Casp1 is not essential for proper development.47. The implication of microglia in AD progression has been nicely demonstrated in the APP/PS1 AD transgenic mouse model where Aβ-mediated activation of microglial Nlrp3 inflammasome results in cognitive deficits and
AD-like pathologies\(^5\). LTP and dendritic spine density are restored, and increased Aβ levels are reduced, in APP/PS1/Casp1\(^{-/-}\) mice\(^5\). While the deficit in AD immunity is largely attributed to CNS resident microglia, astrocytes and perivascular myeloid immune cells, we have identified abnormal activation of the Nlrp1 inflammasome in AD neurons, resulting in Casp1 activation, which then can activate Casp6-mediated neurodegeneration and II-β-mediated glial inflammation\(^1\). By targeting Casp1 in both neurons and glia, VX-765 provides a double protection against these coexisting degenerative AD features.

The reduction of Aβ levels in VX-765-treated mice was unexpected. After only 12 treatments over a 3-month span, RIPA- and formic acid-soluble, immunostained, and thioflavin S-positive Aβ levels were considerably lower than in vehicle-treated J20 mice and remained similar to levels in pretreated 5-month-old J20 brains. Similarly, the absence of Casp1 in the J20/Casp1\(^{-/-}\) decreased Aβ deposits in the brain. These results indicate that VX-765 and the absence of Casp1 stop the progressive accumulation and deposition of Aβ. Furthermore, it became apparent that cremophor vehicle injections increased Aβ accumulation in the J20 mouse of the VX-765 study compared to the Casp1 null study. It is tempting to speculate that either the cremophor or the injections mimic increased inflammation in the aged brain.

Since VX-765 inhibits inflammatory Casp1, the results indicate that in the J20 mouse model, inflammation may be driving Aβ accumulation and contrast with the more popular view that Aβ drives inflammation (see Supplementary Fig. 10). These results are consistent with recent evidence indicating that microglia elimination prevents amyloid deposition\(^14\), and increased peripheral inflammatory markers in midlife humans are associated with brain atrophy and decreased cognition in ageing\(^43\). Whereas the J20 is a familial AD mouse model, the presence of the Nlrp1-Casp1-Casp6 pathway in mild cognitively impaired and in the early and late stages of sporadic human AD suggests that the treatment will also work in sporadic AD. VX-765’s
selectivity for Casp1,25,26, nontoxicity to human neurons and tolerability in phase 2b clinical trials against epilepsy22, blood–brain barrier permeability, and prevention of neurodegeneration in cultured human CNS neurons suggest that it is a most promising novel therapy against both sporadic and familial forms of AD.

It is likely that an efficient treatment against AD will require very early treatment to prevent neuronal degeneration. Similar to antihypertensive or anticholesterol medications that protect against stroke or heart infarcts, we believe that the brain requires preventative early treatments before serious damage occurs. With this in mind, we chose to treat J2O mice only 1 month after the onset of cognitive decline. The outcome was unexpectedly positive. These results suggest that an efficient therapeutic intervention with VX-765 is likely to be more successful in mild cognitively impaired individuals or at the very early onset of clinically diagnosed AD.

Methods

Study design. VX-765 study: Five-month-old vehicle-injected WT (WT + vehicle; n = 18) and J2O mice (J2O + vehicle; n = 14), and VX-765-injected J2O mice (J2O + VX-765; n = 13) mice were longitudinally assessed at five different time points: baseline before treatment (baseline untreated J2O were grouped together; n = 19), after three IP injections per week of VX-765 or vehicle (Treatment 1), after six additional injections over 2 weeks (Treatment 2), after 4-week washout period (WO), and after an additional three injections in 1 week (Treatment 3) (Fig. 1a). All animals tested were included in the analyses unless (a) the animal died during the experiment, or (b) the animal did not behaviourally respond. A total of 25 litters, spread across four different cohorts and tested at different times, were used. Three of these cohorts underwent NOC and open field testing, while cohort 4 was used for Barnes maze. After animals were baseline tested to confirm that J2O animals were behaviourally deficient, J2O animals were randomly assigned to either vehicle or VX-765 treatment independent of behavioural performance. Of all animals used, four J2O mice showed no behavioural deficits at the beginning of the experiment and were not used. Two J2O + vehicle mice did not move at all during the experiment and their behavioural data excluded; however, these animals were kept and were still used for post-mortem analysis.

Table 1 Casp1 validation study: To validate Casp1-specific effects of VX-765, J2O mice were generated on a Casp1−/− background. Seventy-three mice were used (n = 12 –13 per genotype, six different genotypes), all of which were bred through in vitro fertilization (IVF) from 35 donor females. Breeding details are described in the Animal Studies section below. All mice were behaviourally assessed between 7 and 8 months of age, and no animals were excluded from this study.

All mice were sacrificed and processed at 8 months of age. Behavioural scoring was blinded to mouse genotype and treatment, and all animals were randomly assigned for either biochemical or histological analysis and blindly analysed.

VX-765, VRT-043198 IC50 assays on recombinant caspases. Recombinant caspases were commercially obtained (Biovision, Milpitas, CA, USA; K233-10-25) and used according to the manufacturer’s protocol. Caspase activity was measured using a fluorescence-based assay using the following substrates (Enzo Life Sciences, NY, USA): N-Ac-Tyr-Val-Ala-Asp-7-amino-4-trifluoromethylcoumarin (Ac-YVAD-AMC) for human and mouse Casp1, N-Ac-Val-Ala-Asp-7-amino-4-trifluoromethylcoumarin (Ac-VDVAD-AMC) for human Casp2, N-Ac-Glu-Ala-Val-Asp-7-amino-4-trifluoromethylcoumarin (Ac-AEVED-AMC) for human Casp3 and Casp7, Ac-Tryp-Glu-His-Asp-7-amino-4-tri-fluoromethylcoumarin (Ac-WEHD-AMC) for human Casp4 and Casp5, and mouse Casp5, Ac-Glu-Ala-Asp-7-amino-4-trifluoromethylcoumarin (Ac-GEVLF-AMC) for human Casp6, Ac-Ile-Glu-Thr-Asp-7-amino-4-trifluoromethylcoumarin (Ac-IETD-AMC) for human Casp8 and Casp10, and N-Ac-Val-Glu-His-7-amino-4-trifluoromethylcoumarin (Ac-LEHD-AMC) for human Casp9. VX-765 (100 µM–100 µM) and VRT-043198 (100 µM–1 µM) (DSK Biopharma, Morrisville, NC) were used with 0.25 units of caspase and 10 µM substrate in a buffer containing 50 mM Tris-Cl (pH 7.2), 50 mM NaCl, 0.1% 3-(3-cholamidopropyl)dimethylammonio)-1-propanesulfonate (CHAPS), 10 mM ethylenediaminetetraacetic acid (EDTA), 5% glycerol, and 10 mM dithiothreitol. Enzyme activity was measured in a Synergy H1 plate reader (Bio-Tek, Winooski, VT, USA) at 340 nm excitation, 405 nm emission wavelengths every 30 s for 30 min at 37°C. Cleavage rates were calculated from the linear phase of the assay. Fluorescence units were converted to the molar concentration of released 7-amino-4-trifluoromethylcoumarin (AMC) based on a 0–250 µM free AMC standard curve. VX-765 IC50 values were determined by plotting the percentage of the DMSO control inhibitory concentrations. Data were fitted to a dose–response curve equation containing a 1-slope with top and bottom constraints of 1 and 0, respectively.

Animals. All animal procedures followed the Canadian Council on Animal Care guidelines and were approved by the McGill University Animal Care committee. The C57 transgenic line (Lex Stock No. 006292, Jackson Laboratories, Bar Harbor, ME, USA) expresses the Swedish (670/671KMNL) and Indiana (717V–42) human APP mutations under the PDGF-β promoter. Male J2O mice were used as breeders and their sperm was used for IVF colony expansion.

The 120/Casp1 cohort was generated following the animal breeding standard protocols of the Institut de recherche en immunologie et en cancérologie (IRIC) at the Université de Montréal (Montreal, QC, Canada). Casp1 null mice were obtained from Jackson Laboratories (B6.N.129S2-Casp1tm1Byj)), IVF was carried out using fresh sperm from 120/Casp1+/− males and the eggs of WT/Casp1−/− donor females. Fertilized eggs were implanted in 20 C57Bl/6 females, generating pups with six possible genotypes: WT/Casp1−/− (WT/WT), WT/Casp1+/− (WT/WT), WT/Casp1+/− (WT/Het), J2O/Casp1+/− (J2O/WT), J2O/Casp1+/− (J2O/Het), J2O/Casp1−/− (J2O/KO). Both males and females were used, aged at the IRIC, and then transferred to our laboratories prior to the start of experiments.

Genotypes were determined by tail biopsy and RT-PCR. Mice were house-moused (2–4 animals per cage) in standard macromolecules under a 12-h reverse light/dark cycle and controlled environmental conditions. Food and water were available ad libitum.

Blood–brain barrier permeability analysis. Five-month-old J2O and WT mice were anesthetised with isoflurane and the right carotid artery was separated. A small incision was made along the carotid wall and a micro-catheter was inserted into the entrance of the internal carotid artery. The catheter was secured in place using suture thread. VX-765 (50 mg kg−1) was infused at 50 µl min−1 (90–120 µl total volume). The micro-catheter was left an additional 5 min to allow diffusion, after which it was collected by syringe. The mice were transcardially perfused with ice-cold saline and the hippocampus and cortex were dissected out. Samples were sent to the Biopharmacy platform (Université de Montréal) for liquid chromatography and tandem mass spectrometry analysis.

Behavioural analysis. Open field: locomotor activity was measured using an HVS2100 automated tracking system (HVS Image, Hampton, UK). Mice were placed in the open field chamber (40 × 40 cm Plexiglas box, no ceiling, and white floor) and allowed to explore for 5 min.

Novel object recognition (NOR): mice were pre-exposed to two identical objects in the open field chamber for 5 min. Two hours after pre-exposure, mice were placed back into the chamber and exposed to one familiar object and one novel object for 5 min. Mice were exposed to a specific object only once across different test sessions, and novel object placement was counterbalanced within each treatment group. Percentage touches of the novel object during the test phase and discrimination index (% touches novel – % touches familiar/total touches) were assessed.

Barnes maze: the Barnes maze platform (91 cm diameter, elevated 90 cm from the floor) consisted of 20 holes (each 5 cm in diameter). All holes were blocked except for one target hole that led to a recessed escape box. Spatial cues, bright light, and white noise were used to motivate the mice to find the escape during each session. For the adaptation phase, each mouse explored the platform for 60 s. Any mouse that did not find the escape box was guided to it and remained there for 90 s. After the adaptation phase, each group followed the same protocol with the goal to train each mouse to find the target and enter the escape box within 180 s. Mice remained in the box for an additional 60 s. Four trials per day, approximately 15 min apart, were performed for 4 consecutive days. In the probe test (day 5), each mouse performed one 90 s trial. The target was still located in the same position, but the escape box was blocked. The latency and errors to reach the target, plus the number of mouse pokes to each of the holes (target preference) were measured.

The Y-Maze was composed of a symmetrical Y-shaped maze with three arms 120° from each other, designated A, B, and C. Animals were placed at the distal end of arm A and allowed to explore the maze for 5 min. Total arm entries and spontaneous alternation, defined as consecutive entries in three different arms, were recorded. Percentage alternation (% alternations/total number arm entries – 2) × 100 was determined.

Tissue processing. For immunohistochemistry, mice (n = 4–6 per group) were anesthetised with isoflurane and transcardially perfused with ice-cold saline followed by ice-cold 4% paraformaldehyde (Sigma-Aldrich, St Louis, MO, USA) in 0.1 M phosphate-buffered saline. Brains were fixed in 10% neutral-buffered formalin (NBF) for 48 h. Tissue was cryoprotected in 30% DMSO containing 10% NBF for 48 h, and transferred to 70% ethanol. Brains were paraffin embedded and sectioned at 4 µm. For western blot and ELISA (n = 4–8 per group), anesthetised mice were cervical dislocated, the hippocampus and cortex dissected out and immediately frozen on dry ice. Proteins were extracted in 5x volume/weight with radiolabels solution (2x HPLC buffer with 10 µg µl−1 protease inhibitors), pH 7.4, 1% NP-40, 150 mM NaCl, 0.25% Na-deoxycholate, 1 mM EDTA, 1 mM Na3VO4, 1 mM NaF, 1 mM PMSF, 10 µl µl−1 of Proteases Inhibitors Cocktail (Sigma-Aldrich) on
ice with a tissue homogeniser (OMNI International, Kennesaw, GA, USA). Samples were centrifuged (20,000 g, 4 °C) for 20 min, the supernatant was removed, and the protein was quantified using the BCA method. The RIPA-insoluble pellet was further extracted in 70% formic acid in dH2O, evaporated, and resolubilized in 200 mM Tris-HCl, pH 7.5.

**Immunohistochemical staining.** Epitopes were demasked in either citrate (10 mM Tris-Na Citrate, pH 6.0) or EDTA (10 mM Tris Base, 1 mM EDTA, 0.05% Tween-20, pH 9.0) antigen retrieval buffer, and immunostained using the DAKO Autostainer Plus automated slide processor and EnVision Flex system (Dako, Burlington, ON, Canada). Following deparaffinization and rehydration, sections were peroxidase treated, blocked in serum-free protein block, and immunostained with the following antibodies diluted in 2% serum-free protein block: DAB. Sections were treated with the kit-supplied appropriate mouse or rabbit HRP-conjugated secondary antibody and visualized with DAB. Sections were hematoxylin counterstained, dehydrated, and coverslipped with Permount (Fisher Scientific). Sections were digitally scanned with MIRAX SCAN for analysis (Zeiss, Don Mills, ON, Canada). After deparaffinization and rehydration, slides were stained with filtered 1% aqueous Thioflavin-S (Sigma-Aldrich).

**Quantitative immunohistochemical analysis.** Iba1-positive cells in the cortex and anterior portion of the CA1 region of the hippocampus were quantified using a modified version of the areal counting frame. Briefly, every fifth slide was quantified (resulting in four sections per brain). Multiple >80 images were taken with the MIRAX SCAN within the area of interest and the number of Iba1-positive cells were manually counted. The numerical density (number of cells mm⁻²) in the anterior CA1 region and cortex was estimated. A minimum number of 150 hits per area were needed to make reliable estimates. Additional sections were counted if we were unable to reach our minimum number. Quantification was performed blinded for treatment and genotype. Image software (NIH, Bethesda, MD, USA) was used to measure Aβ accumulation, GFAP and synaptophysin immunopositive staining in the CA1 region and cortex. Four sections per brain were analysed, and multiple images were taken within each section to cover the CA1 region and cortex. Thresholds were equally adjusted across all brains to highlight the stained area, and particles were analysed to determine total area of staining. Results are expressed as total area stained per total area analysed (μm² mm⁻²). Representative images have only been cropped to conform to size constraints and have not undergone any post-processing.

**Western blotting.** Mouse protein samples (25–100 μg) were prepared in Laemmli (5% 2-β-mercaptoethanol, 1 mM Na₂VO₄, 1 mM NaF, 1 mM PMSF, 10 μM 1-β-methyl 2-proteasome Inhibitors Cocktail (Sigma-Aldrich)), boiled for 5 min, and separated by polyacrylamide gel electrophoresis (PAGE). Samples were probed with the following primary antibodies, diluted in either 5% nonfat dry milk in Tris-buffered saline (TBS) or 5% BSA blocking solution: B-actin (1:1000), Aβ (1-16) (6E10) (BioLegend 803001, San Diego, CA, USA), β1-40 (F25276, laboratory developed), and 1:1000 mouse antisympathophysin (Sigma-Aldrich S5768). Sections were treated with the kit-supplied appropriate mouse or rabbit HRP-conjugated secondary antibody and visualized with DAB. Sections were hematoxylin counterstained, dehydrated, and coverslipped with Permount (Fisher Scientific). Sections were digitally scanned with MIRAX SCAN for analysis (Zeiss, Don Mills, ON, Canada). After deparaffinization and rehydration, slides were stained with filtered 1% aqueous Thioflavin-S (Sigma-Aldrich).

**PCR and real-time PCR.** HAP transgenic PCR amplification was performed using Tqy DNA polymerase (New England Biolabs, Whitby, ON, Canada) following the manufacturer’s protocol (bAAPP 5′-AACACAGAAAACAAGGT-3′ and 5′-CCCATGGTGGATGAAGCA-3′; 480 bp amplicon) (15), (Hyoxanthine-guanine phosphoribosyltransferase, HPRT) 5′-GTAATGATCAGTAACGGGAGC-3′ and 5′-CCA GACAGTGTTGACTACCAA-3′ (177 bp amplicon) (Carcino genesis). Products were visualized on RedSafe (FroggBio, North York, ON, Canada) stained 2% agarose gels. Real-time PCR experiments were performed with SYBR Green Taq Mastermix (Quanta Biosciences, Gaithersburg, MD, USA) on an Applied Biosystems 7500 Fast Real-Time PCR system machine (Applied Biosystems, Foster City, CA, USA). Gene amplification was performed with the following primers: (18s) 5′- GTACCGGTGTAACCGG-3′ and 5′-CCATTGATCGTGGTACGG-3′; (β-tubulin) 5′-ACTAAGCTTGTTGAAG-3′ and 5′-GATCTCGTATGGGAAATT-3′; (Nefrin) 5′-TCTGTGAACGACGTCCGACCG-3′ and 5′-CTCCACCAAC CATTCTCC-3′. Results are expressed as fold-induction values normalized to mean HPRT and 18s reference genes using the Pfaff’s method.

**Mouse synaptic plasticity RT²-Profiler PCR array.** The mouse synaptic plasticity RT²-Profiler PCR array (PAMM-126Z, Qiagen) profiles the expression of 84 genes involved in synaptic alterations during learning and memory and five housekeeping genes (β-actin; β-2-microglobulin; glyceraldehyde 3-phosphate dehydrogenase, GAPDH; β-glucuronidase, Gus; Heat shock protein 90 alpha (cytosolic), Hsp90ab1) with real-time PCR. Total RNA (465 ng for each sample) was reverse transcribed (which included a genomic DNA elimination step) following the manufacturer’s protocol (First Strand cDNA Synthesis Kit, Qiagen). The real-time PCR reaction was performed in a real-time cycler ABI 7500 (Fast Block) (Applied Biosystems). Real-time PCR experiments were performed with the RT² SYBR Green PCR master mix (Qiagen). Cycling conditions were as follows: 2 min at 50 °C, 10 min at 95 °C. 40 cycles, 15 s each at 95 °C, and 1 min at 60 °C. Threshold and baseline were set manually according to the manufacturer’s instructions. Data were normalized to the geometric mean of the five housekeeping genes and analysed by the comparative Ct method (2⁻ΔΔCt).

**Neuronal cell culture and axonal degeneration assay.** Twelve- to sixteen-week-old fetal cortical tissue were obtained from the Birth Defects Research Laboratory (University of Washington, Seattle, WA, USA) in accordance with a approved protocol by the McGill institutional review board. HPN were cultured from brains that were dissected with ethyl trypsin, treated with trypsin, and then resuspended through 130, 70, and 30 μm nylon mesh. Dissected cells were centrifuged for 10 min at 300 x g and resuspended in minimal essential medium (MEM) with Earle’s salts, and supplemented with 5% NCS (decapitated BCS, 1× Penicillin-Streptomycin, 1 mM sodium pyruvate, and 2 mM -glutamine (Invitrogen, Carlsbad, CA, USA). Cells were cultured in supplemented MEM supplemented with 5% FBS on 35 mm well tissue culture plates (Sigma-Aldrich). MEM medium was changed every 2 days and astrocyte proliferation was limited with fluorodeoxyuridine (FdU) antimotic treatment (1 mM, Sigma) for the first 4 days. Neuronal cultures grown were 7-10 days before conducting experiments. Four different neuron preparations at different times were tested. One of the neuron preparations was not stable and the culture, including the Serum + control that did not receive any drug, died in the middle of the experiment. Therefore, three different neuron preparations from three genetically different donors were used.

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**NEURODEGENERATION ASSAY.** VX-765 toxicity was assessed with 3-(4,5-dimethylthiazol-2-yl)-2,5- diphenyltetrazolium bromide (MTT). Neurons were treated with 25–200 μM VX-765 or 0.1% DMSO (highest DMSO concentration used) for 72 h. Neurons were then incubated with 0.5 μg ml⁻¹ MTT (Sigma-Aldrich) for 4 h at 37 °C (5% CO₂). The formazan crystals were dissolved in 100 μl DMSO for 30 min. Absorbance was measured at 560 and 670 nm using the Synergy H4 plate reader.

**Axonal degeneration was measured by APβ40 and 42 FdU incorporation or serum deprivation stress. VX-765 was administered either as a pretreatment strategy (1 h prior to stressor) or as a posttreatment stressor strategy (48 h after stressor). Neurons were transfected by Gene gun (BioRad, Mississauga, ON, Canada) with gold beads coated with pBudCE41-eGFP for fluorescence visualization. Neurons that were APβ40/eGFP stressed were transfected with pBudCE41-eGFP/APβ40. Neurons that were APβ40/eGFP stressed were transfected with pBudCE41-eGFP/APβ40. Neurons that were APβ40/eGFP stressed were transfected with pBudCE41-eGFP/APβ40. Neurons that were APβ40/eGFP stressed were transfected with pBudCE41-eGFP/APβ40. Neurons that were APβ40/eGFP stressed were transfected with pBudCE41-eGFP/APβ40. Neurons that were APβ40/eGFP stressed were transfected with pBudCE41-eGFP/APβ40. Neurons that were APβ40/eGFP stressed were transfected with pBudCE41-eGFP/APβ40.
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Author contributions
J.F.: animal behaviour and analyses, immunohistochemistry analyses, ELISA, cell culture experiments, blood–brain barrier permeability assays, designed experimental paradigm, made figures, wrote methods, results and revised manuscript. B.F.: molecular biology, immunostaining, prepared figures, wrote methods and results, revised the manuscript. A.N.: western blot analyses, prepared figures, wrote methods and results, revised the manuscript. J.L.: performed experiments for IC50 on recombinant enzymes, wrote methods and results. Revised the manuscript. C.L.: performed blood–brain barrier permeability experiments, revised the manuscript. A.C.L.: Conception of the experimental idea and design, supervision of data collection and analysis, wrote manuscript.

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