Amyloid-β peptide protects against microbial infection in mouse and worm models of Alzheimer’s disease

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The amyloid-β peptide (Aβ) is a key protein in Alzheimer’s disease (AD) pathology. We previously reported in vitro evidence suggesting that Aβ is an antimicrobial peptide. We present in vivo data showing that Aβ expression protects against fungal and bacterial infections in mouse, nematode, and cell culture models of AD. We show that Aβ oligomerization, a behavior traditionally viewed as intrinsically pathological, may be necessary for the antimicrobial activities of the peptide. Collectively, our data are consistent with a model in which soluble Aβ oligomers first bind to microbial cell wall carbohydrates via a heparin-binding domain. Developing protofibrils inhibited pathogen adhesion to host cells. Propagating β-amyloid fibrils mediate agglutination and eventual entrapment of unattached microbes. Consistent with our model, Salmonella Typhimurium bacterial infection of the brains of transgenic 5XFAD mice resulted in rapid seeding and accelerated β-amyloid deposition, which closely colocalized with the invading bacteria. Our findings raise the intriguing possibility that β-amyloid may play a protective role in innate immunity and infectious or sterile inflammatory stimuli may drive amyloidosis. These data suggest a dual protective/damaging role for Aβ, as has been described for other antimicrobial peptides.

INTRODUCTION

Neurodegeneration in Alzheimer’s disease (AD) is mediated by soluble oligomeric intermediates generated during fibrillation of the amyloid-β protein (Aβ) (1). Overwhelming evidence supports Aβ’s pivotal role in AD. However, despite remarkably high sequence conservation across diverse species (humans share Aβ42 sequences with coelacanths, a 400 million–year–old fish taxon) (2) and extensive data showing broad activity spectra for Aβ, the peptide has traditionally been characterized as a functionless catabolic byproduct. Activities identified for Aβ in vivo are most often described as stochastic pathological behaviors. Oligomerization, in particular, is viewed as a pathogenic pathway, and Aβ oligomers are assumed to be intrinsically abnormal. Scant consideration has been given to possible physiological roles for Aβ.

Members of the evolutionarily ancient family of proteins, collectively known as antimicrobial peptides (AMPs), share many of Aβ’s purportedly abnormal activities, including oligomerization and fibrillation (3, 4). For AMPs, these activities mediate key protective roles in innate immunity. AMPs are the first line of defense against pathogens and act as potent broad-spectrum antibiotics and immunomodulators that target bacteria, mycobacteria, enveloped viruses, fungi, protozoans, and, in some cases, transformed or cancerous host cells (5). AMPs are widely expressed and are abundant in brain and other immunoprivileged tissues where actions of the adaptive immune system are constrained. Although AMPs are normally protective, AMP dysregulation can lead to host cell toxicity, chronic inflammation, and degenerative pathologies (6–8). Particularly germane to Aβ’s role in AD, AMPs are deposited as amyloid in several disorders (3, 4, 9), including senile semisel vesicle amyloid and isolated atrial amyloidosis, two of the most common human amyloidopathies. Consistent with identity as an AMP, we recently reported that synthetic Aβ exhibits potent in vitro antimicrobial activity toward eight common and clinically relevant microbial pathogens (3). Furthermore, whole-brain homogenates from AD patients show Aβ-mediated activity against Candida albicans. More recently, synthetic Aβ has been shown to protect cultured cells from influenza A virus (10) and herpes simplex virus (11). However, the biological relevance of protective in vitro Aβ activities requires validation in vivo. Here, we extend our original findings and show that Aβ expression inhibits infection in a transgenic mouse model of AD (5XFAD), in the nematode Caenorhabditis elegans, and in cultured mammalian cell models. Mice lacking the amyloid precursor protein (APP) that have low Aβ expression also show a trend toward attenuated survival after bacterial infection. Most surprisingly, oligomerization and fibrillation appear to mediate Aβ’s protective activity, and cerebral infection with microbial cells seeds and markedly accelerates β-amyloid deposition in 5XFAD mice and transgenic C. elegans.

RESULTS

Aβ-mediated protection was characterized in mice, C. elegans, and cell culture models of infection. Salmonella enterica serotype Typhimurium (S. Typhimurium) was used as an infecting agent in mouse models. Nematode and cultured cell experiments used pathogenic (hyphal) C. albicans (Candida) and S. Typhimurium.

Aβ protects against meningitis in genetically modified mice

We first used genetically modified mice to test for protective effects of elevated Aβ expression and attenuated resistance with decreased peptide. Four-week-old 5XFAD transgenic mice constitutively express human Aβ in the brain at high levels but lack the β-amyloid deposits and features
of neuroinflammation found in older animals (12). APP knockout (APP-KO) mice lack the precursor protein required for murine Aβ generation (13). One-month-old 5XFAD mice (n = 12), APP-KO mice (n = 15), and wild-type littermates (n = 11 and 15, respectively) received a single intracerebral injection of 65,000 colony-forming units (CFU) of S. Typhimurium. Clinical progression to the moribund state was followed according to established grading criteria for mouse encephalomyelitis (fig. S1A). Survival of Aβ-expressing 5XFAD mice was significantly increased compared to that of nontransgenic littermates (P = 0.009) (Fig. 1A). Consistent with increased resistance to infection, 5XFAD mice also ranked significantly higher in clinical tests grading mouse encephalomyelitis progression (P < 0.0001). 5XFAD mice also showed reduced weight loss (P = 0.0008) and lower cerebral S. Typhimurium loads (P = 0.03) compared to wild-type controls (Fig. 1, B to D). Consistent with immunodeficiency associated with low Aβ, APP-KO mice showed a trend (P = 0.10) toward increased mortality after infection (Fig. 1E). Control injections using heat-killed bacteria did not lead to clinical decline or death in 5XFAD and wild-type mice (Fig. 1F), consistent with mouse mortality being mediated by S. Typhimurium infection. Next, we confirmed high amounts of soluble Aβ and low amounts of insoluble Aβ in 4-week-old 5XFAD mouse brain using formic acid extraction and anti-β-amylloid enzyme-linked immunosorbent assays (ELISAs) (fig. S1B). To confirm that inflammation did not immunologically prime and protect 5XFAD mice against infection, we compared the immune profiles in 1-month-old transgenic and wild-type mouse brains. Consistent with previous reports showing an absence of immune activation (12), there was no significant increase in glial fibrillary acidic protein–positive (GFAP+) astrocytes, Iba1+ microglia, and the amounts of 10 cytokines in 4-week-old 5XFAD mice compared to wild-type littermates (fig. S1, C to E).

**Aβ increases survival of transgenic C. elegans infected with Candida**

To further explore the ability of Aβ to afford protection against infection, we next tested transgenic C. elegans for resistance to Candida. Our nematode infection model uses two previously described C. elegans transgenic strains: GMC101 that expresses the 1–42 residue human Aβ isoform (Aβ42) (14) and CL2122, a control strain that expresses intestinal green fluorescent protein (GFP) (mtl-2:gfp) marker (as does GMC101) but does not express Aβ. Adult GMC101 nematodes ultimately develop age-progressive paralysis and β-amyloid deposition in the body wall muscle. For our experiments, developmentally synchronized L4 larvae were infected 5 days before the onset of paralysis. Aβ expression is driven by the unc-54 promoter (which encodes a myosin heavy chain), active in body wall muscle (14) as well as in other tissues, including muscle cells of the gastrointestinal tract (15). Amyloidogenic peptides under the unc-54 promoter have also been shown to translocate via vesicular transport to the gut of transgenic worms, and Aβ has been proposed as a likely candidate for translocation via this mechanism (16). Immunohistological analysis of adult GMC101 using three different anti-Aβ antibodies confirmed Aβ localization in the body wall muscle and the gut lumen (fig. S2, A and B). Anti-Aβ antibodies did not label negative control strain CL2122 intestine or body wall cells. In addition, excreta from healthy GMC101 but not CL2122 worms were positive for anti-Aβ signal by immunoblot (fig. S2C). Although an origin for gut Aβ remains unclear, strong empirical evidence supports the localization of Aβ peptides in the intestinal lumen of GMC101 nematodes. Thus, transgenic GMC101 nematodes appear to be suitable models for testing Aβ-mediated protective activities against intestinal pathogens.

**C. albicans** [American Type Culture Collection (ATCC) 90028] is an Aβ-sensitive...
dimorphic fungus (3) and a well-characterized C. elegans intestinal pathogen that causes distention, penetrative filamentation, and death among wild-type nematodes 2 days after ingestion. Links between fungal brain infections and AD pathology have also recently emerged, including for C. albicans (17) and closely related Candida glabrata (18). We compared survival of control CL2122 (n = 56) and GMC101 (n = 59) nematodes after incubation (2 hours, 25°C) on C. albicans lawns. Consistent with Aβ-mediated protection, GMC101 nematodes infected with C. albicans showed significantly (P < 0.00001) reduced mortality as compared to control CL2122 worms that did not express Aβ (Fig. 2A). Consistent with mouse data, Aβ-expressing nematodes were also protected from the C. elegans intestinal pathogen S. Typhimurium with GMC101 worms showing statistically significant (P = 0.0005) increased survival compared to CL2122 controls after infection with the bacterium (fig. S3A).

The antimicrobial activities of Aβ protects cells in culture
To address the mechanism of protection, we next tested the ability of Aβ to protect cell monolayers from infection using transformed cultured human brain neuroglioma (H4) and Chinese hamster ovary (CHO) cells. H4 lines included stably transformed H4-Aβ40 and H4-Aβ42 cells that selectively secrete the 1–40 residue Aβ isoform (Aβ40) or Aβ42 isoform, respectively (19). Processing of a BRI-Aβ fusion protein expressed by transformed H4 cells led to constitutive high-level expression and secretion of the encoded Aβ protein. For double transfected CHO cells (CHO-CAB), overexpression of APP and the APP-processing protease β-secretase leads to APP cleavage and the generation of multiple Aβ isoforms (20). Non-transformed H4 (H4-N) and CHO (CHO-N) cells were used as control cell lines. C. albicans has been extensively characterized
in cell culture infection models and was used in our experiments as an infectious agent.

We first compared nontransformed and transformed host cells for survival after infection with *C. albicans*. Host cells were prelabeled with bromodeoxyuridine (BrdU). After infection, host cell viability was determined by assaying for anti-BrdU immunofluorescence. Consistent with findings for 5XFAD mice and GMC101 nematodes, survival 28 hours after infection was significantly increased for Aβ- overexpressing H4-Δ4b40 (P = 0.002) and H4-Δ4b42 (P = 0.001) transformed cell lines compared to control H4-N cells, with rank order H4-Δ4b42 > H4-Δ4b40 > H4-N (Fig. 2B). Survival of transformed CHO-CAB cells was also significantly higher (P = 0.004) than that of control CHO-N cell lines. Additional independent assays of host cell viability (fig. S4, A and B) were performed to confirm increased resistance of transformed H4-Δ4b42 cells to *C. albicans* infection. Attenuated *C. albicans* load for H4-Δ4b42 cells was also independently confirmed by comparing wells for yeast CFU (fig. S4C).

Whereas the amount of Aβ in conditioned cell culture medium (fig. S5, A and B) fell within the physiological ranges reported for human cerebrospinal fluid (CSF) (2 to 20 ng/ml) (21), concentrations were two orders of magnitude (log10) lower than the minimal inhibitory concentration (MIC) for fungicidal species in microdilution MIC assays (3). We have previously reported that Aβ’s antimicrobial activities show close parallels with those of LL-37 (3), an archetypal human AMP that remains protective at subfungicidal concentrations (22). Two linked, yet distinct activities mediate LL-37’s protective anti-*Candida* actions at low peptide concentrations (22). The first is disruption of *C. albicans* adhesion to host cells. Host cell attachment is a prerequisite step for infection by many pathogens, including *C. albicans*. The second is agglutination of the resulting unattached yeast cells. Agglutination limits microbial access to host cells and also generates high local AMP concentrations within peptidomicrobe aggregates. Accordingly, we next tested Aβ for adhesion inhibition and agglutination activities using the cell culture infection model. Hyphal *C. albicans* was incubated (2 hours, 37°C) in preconditioned medium with transformed or nontransformed cell cultures prepared in slide chambers. Microscopic examination revealed fewer *C. albicans* attached to transformed Aβ-expressing cells compared to nontransformed monolayers (Fig. 2C and fig. S6A). To confirm these data, we repeated *C. albicans*-cell culture incubation experiments in 96-well microtiter plates, and we assayed the *Candida* load in wells immunochemically using anti-*Candida* antibodies. Data confirmed visual observations with statistically significant attenuation of *C. albicans* adhesion to transformed H4-Δ4b42 (P = 0.001), H4-Δ4b40 (P = 0.001), and CHO-CAB (P = 0.004) cells compared to naive control lines (Fig. 2D). Additionally, after overnight incubation, marked microbial agglutination was observed in wells containing transformed, but not nontransformed, host cells (Fig. 2E and fig. S6B). Images of wells were analyzed for yeast aggregation. *Candida* aggregation was significantly elevated in transformed H4-Δ4b42 (P = 0.00004), H4-Δ4b40 (P = 0.0003), and CHO-CAB (P = 0.002) samples compared to naive controls (Fig. 2F). For H4 cell lines, adhesion inhibition and agglutination activities were consistent with host viability data, with rank orders H4-Δ4b42 > H4-Δ4b40 > H4-N.

We next characterized cell-free conditioned culture medium for Aβ-mediated adhesion inhibition and agglutinating activities. Yeast adhesion and agglutination were assayed in 96-well plates using the methods of Tsai et al. (22). Briefly, synchronized hyphal *C. albicans* were incubated (2 hours, 37°C) with conditioned medium samples in the absence of host cells. After washing, yeast adhering to well surfaces were stained with Calcofluor white, and fluorescence was measured. Well images were analyzed for yeast aggregation after overnight incubation. Immunodepletion with anti-Aβ antibodies significantly attenuated H4-Δ4b42, H4-Δ4b40, and CHO-CAB medium adhesion inhibition (P = 0.009, P = 0.001, and P = 0.004, respectively) and agglutination (P = 0.001, P = 0.0005, and P = 0.004, respectively) activities against *C. albicans* (Fig. 3, A and B). Analysis confirmed that anti-Aβ immunodepletion removed >95% of the Aβ from samples used in experiments to confirm that the anti-*Candida* activities of transformed cell culture medium were specific for Aβ (fig. S5, A and B).

Consistent with yeast data, S. Typhimurium were agglutinated in H4-Δ4b42 conditioned medium (fig. S3B). H4-Δ4b42 cell cultures incubated with S. Typhimurium also have significantly (P = 0.036) lower intracellular infection compared to nontransformed H4-N cells (fig. S3, C and D).

Serial dilution experiments showed that adhesion inhibition and agglutination activities were dose-dependent for both synthetic and cell-derived Aβ (Fig. 3, C and D). However, synthetic Aβ peptide preparations had lower specific activities compared to cell-derived material. Cofactors secreted by cultured cells were unlikely to account for the increased potency of cell-derived Aβ because synthetic peptide incubations were performed in H4b42-depleted conditioned medium (H4-Δ4b42-1D) from H4-Δ4b42 cell cultures. Anti-Aβ antibodies used to clear Aβ42 from H4-Δ4b42 culture medium before addition of synthetic peptides were specific for Aβ and not likely to deplete species acting as cofactors. Oligomerization has been shown to modulate a range of Aβ activities. Moreover, conditioned medium from experimental cell lines has been reported to contain oligomeric Aβ (23), whereas our synthetic peptide preparations were pretreated to remove oligomer species. Synthetic peptide pretreatments included fractionation by preparative size exclusion chromatography to remove species >6 kD. Characterization experiments using analytical size exclusion chromatography confirmed that immediately before experimental inoculation with yeast, cell-derived material contained a polydisperse population of soluble Aβ oligomers of between 8 and 50 kD, whereas synthetic peptides remained overwhelmingly monomeric (fig. SSC).

To test whether oligomerization modulates Aβ’s AMP activity, we generated synthetic Aβ oligomers and compared the antimicrobial activities of Aβ42 monomer, soluble oligomeric ADDLs (amyloid-β-derived diffusible oligomeric ligands) (24), and high-order protofibril (>600 kD) preparations. Compared to monomeric peptide, ADDLs exhibited potentiated, and protofibrils attenuated, adhesion inhibition (Fig. 3E) and agglutination (Fig. 3F) activities. Our data are consistent with a central role for soluble Aβ low-order (2 to 30 monomer units) oligomers in mediating the peptide’s AMP activities. Consistent with such a role, soluble Aβ is overwhelmingly oligomeric in vivo (25), and oligomers are key for the protective activities of a wide range of AMPs (26–29) including LL-37 (26, 30).

**Antimicrobial actions are mediated by the heparin-binding activity of Aβ oligomers**

Binding of AMP peptides to microbial surfaces is a prerequisite step for adhesion inhibition and agglutination activities. LL-37 contains an XBBBXBX heparin-binding motif (where X is a hydrophobic or uncharged residue and B is a basic residue) that mediates inhibition of host cell adhesion and agglutination activities by facilitating attachment of
oligomeric species (26, 30) to microbial cell wall carbohydrates (22). Aβ also contains an XBBXBX heparin-binding motif between residues 12 to 17 (VHHQKL) (31). Competitive inhibition by soluble microbial sugars is a hallmark for AMPs with activities mediated by lectin-like carbohydrate binding (22). Indeed, fungal and bacterial pathogens secrete specialized scavenging exopolysaccharides that target the heparin-binding domains of AMPs as a countermeasure to defenses mounted by hosts. Soluble forms of mannan and glucan, the two most abundant carbohydrates in the yeast cell wall, have been shown to inhibit XBBXBX-mediated binding of LL-37 to Candida (22, 32). We investigated whether the adhesion inhibition and agglutination activities of Aβ were similarly inhibited by soluble mannan and glucan. Live yeast cells were incubated in H4-Aβ40, H4-Aβ42, and CHO-CAB conditioned medium in the presence or absence of mannan or glucan. Consistent with anti-Candida activity mediated by Aβ’s heparin-binding domain, mannan and glucan significantly attenuated adhesion inhibition (P < 0.008) and agglutination (P < 0.003) activities of conditioned medium from Aβ-expressing transformed cells (Fig. 3, G and H).

We further characterized Aβ’s binding to C. albicans and S. Typhimurium using a new binding immunoassay. For this assay, samples were incubated in wells containing immobilized intact hyphal Candida or S. Typhimurium cells, and bound Aβ was detected immunochromatically with an Aβ42-specific antibody. Aβ binding to Candida and S. Typhimurium was concentration-dependent (Fig. 3I and fig. S3E). Consistent with binding mediated by Aβ’s VHHQKL domain, the anti-Aβ signal from H4-Aβ42 medium was significantly attenuated in the presence of glucan (P = 0.008) or mannan (P = 0.004) (Fig. 3J). The anti-Aβ signal in wells was also significantly reduced (P = 0.006) for anti-Aβ-immunodepleted H4-Aβ42 medium (negative control), which was consistent with assay specificity for Aβ42 binding. Consistent with findings for antimicrobial activities, cell-generated Aβ oligomers showed increased binding to immobilized yeast compared to synthetic monomeric peptide (Fig. 3I). Previous studies have shown that Aβ oligomerization greatly increases carbohydrate-binding activity (31). Heparin-binding AMP oligomers also show potentiated carbohydrate binding compared to monomeric species (33). Overall,
our findings are consistent with soluble Aβ oligomers having an enhanced propensity to bind to cell walls, engendering greater adhesion inhibition and agglutination activities compared to monomeric synthetic peptide.

**Aβ fibrillization mediates Candida agglutination**

Binding by Aβ of glycosaminoglycans found in brain tissue induces peptide fibrillation (34). Aβ’s binding of cell wall and glycocalyx carbohydrates at microbial surfaces seemed likely to also generate Aβ fibrils. Although viewed solely as a part of Aβ’s pathophysiology, fibrillation among AMPs is a normal protective behavior that mediates antimicrobial activities, including microbial cell and viral agglutination (35) and bacterial membrane perturbation (3, 4). Most recently, studies have shown that the human AMP α-defensin-6 (HD6) forms fibrils that entangle and trap microbial cells (36). Thus, we next investigated a possible role for Aβ fibrillation in the peptide’s protective AMP activities. Analysis of early-stage (<3 hours after infection) Candida agglutination in H4-Aβ42 medium using transmission electron microscopy (TEM) revealed clumped microbial cells entwined and linked by fibrils propagating from cell surfaces (Fig. 4, A to D). *C. albicans* lack flagella and are not reported to produce extended fibrillar structures. Moreover, the fibrillar structures on the Candida cell surface were labeled by anti-Aβ immunogold nanoparticles (anti-Aβ-Au). Anti-Aβ-Au binding to fibrils was ablated by co-incubation with synthetic Aβ peptide, consistent with Aβ-specific labeling (Fig. 4D). TEM analysis of early-stage *S. Typhimurium* agglutinates in H4-Aβ42 conditioned medium confirmed that bacterial cells were also bound and linked by fibrils (fig. S3F).

Epifluorescence micrographs of Thioflavin S–stained late-stage (>12 hours after infection) H4-Aβ42 yeast aggregates displayed the enhanced fluorescence and red shifts that mark the presence of amyloid fibrils (Fig. 5A). Enhanced fluorescence was not observed for negative control yeast agglutinates (Fig. 5A). Thioflavin S fluorescence within H4-Aβ42 yeast aggregates colocalized with the signal for anti-Aβ immunoreactivity (Fig. 5B). Congo red–stained H4-Aβ42 yeast aggregates also showed birefringence under polarized light, another marker for β-amyloid (fig. S7). Scanning electron microscopy (SEM) micrographs of yeast aggregates from H4-Aβ42 medium revealed an irregular material adhering to cell surfaces not present in Candida pellets prepared by centrifugation in Aβ–free medium (Fig. 5C). Analysis of the Candida cell surface by TEM revealed the adhering material to be filamentous and immunoreactive to anti-Aβ-Au (Fig. 5D). Co-incubation of soluble synthetic Aβ40 peptide abolished anti-Aβ-Au binding. Collectively, the data are consistent with microbially agglutination and entrapment mediated by Aβ fibrillation in our cell culture infection model.

**β-Amyloid mediates pathogen entrapment in GMC101 nematodes and 5XFAD mice**

We also investigated infection-associated Aβ fibrillation in our nematode and mouse infection models. Consistent with Aβ targeting and binding to yeast cells in our cell culture model, *Candida* in the gut of recently infected (2 hours after ingestion) GMC101 nematodes were labeled by anti–Aβ-Au nanoparticles (Fig. 6A). Yeast cells in the gut of the control CL2122 nematode were not labeled by anti–Aβ-Au (fig. S8A). Aβ fibrillation in GMC101 worms is normally confined to the body wall muscle. However, compared to infection-free nematodes,
GMC101 worms with late-stage Candida infection showed enhanced Thioflavin S fluorescence in nonmuscle tissue, including the gastrointestinal tract (Fig. 6B). High-resolution micrographs of yeast cells in the gastrointestinal tract of GMC101 nematodes revealed clumped Candida embedded in the material that showed enhanced fluorescence after Thioflavin S staining (Fig. 6C) and was labeled by anti-Aβ antibodies (Fig. 6D). Consistent with Aβ-specific labeling, anti-Aβ signal (fig. S2B) and enhanced Thioflavin S fluorescence (fig. S8B) were absent from uninfected or Candida-infected negative control CL2122 nematodes that did not express Aβ. Findings for C. albicans–infected GMC101 nematodes were consistent with the agglutinating and entrapment roles of Aβ fibrils observed in our cell culture infection models. Thus, Aβ fibrillation on the surface of yeast cells infecting the gut of GMC101 nematodes may mediate the resistance to infection observed for these worms.

Four-week-old 5XFAD mouse brain is normally negative for β-amyloid deposits (12). However, Thioflavin S and anti-Aβ staining of 5XFAD mouse brain revealed widespread β-amyloid deposition 48 hours after infection with S. Typhimurium (Fig. 7, A and B). Moreover, anti-Salmonella and β-amyloid signal colocalized in the 5XFAD mouse brain, suggesting that bacterial cells may have induced Aβ fibrillation. TEM analysis also revealed that bacterial cells were embedded in fibrous material labeled by anti-Aβ-Au nanoparticles in 5XFAD but not wild-type mouse brain sections (fig. S8). A video of Z-section projections rotating through 360° shows that bacteria are not confined to the surface of Aβ accretions but are embedded within the β-amyloid deposits (video S1). Consistent with fibrillation driven by proliferation of S. Typhimurium cells, β-amyloid deposits were absent from sham-infected 1-month-old 5XFAD control mice injected with heat-killed bacteria. Thioflavin S staining and anti-β-amyloid antibodies did not label mouse brain from negative control nontransgenic littermates (Fig. 7A).

**DISCUSSION**

Our findings are consistent with a potential protective role for Aβ in vivo as an AMP. Expression of Aβ was associated with increased host survival in cell culture, nematode, and mouse infection models (Figs. 1 and 2). Low Aβ expression was associated with higher mortality after infection of APP-KO mice. Our data are consistent with a protective role for Aβ in innate immunity that uses a classic AMP mechanism characterized by reduced microbial adhesion to host cells and agglutination and entrapment of microbes by Aβ fibrils. Moreover, well-characterized Aβ activities mediate the peptide’s antimicrobial actions. However, these same properties, oligomerization, fibrillation, and carbohydrate binding, are also linked to Aβ’s pathophysiology. Whereas a protective/damaging duality is a new proposition for Aβ’s activities, this is not the case for classical AMPs. For example, LL-37 offers a germane model for the potential pathological consequences of normally protective AMP actions. LL-37 is essential for normal immune function, and low expression leads to lethal infections (37). However, at elevated concentrations, LL-37 is cytotoxic to host cells, particularly smooth muscle cells (38). The cytotoxic and pro-inflammatory activities of LL-37 are implicated in the pathogenesis of several major late-life diseases, including rheumatoid arthritis, lupus erythematosus, and atherosclerosis (39). Thus, a normally protective Aβ activity spectrum that, when dysregulated, also leads to AD pathology is consistent with the actions of classical human AMPs.

Adhesion blocking and agglutination activities are distinct from AMP microbicidal activities, which typically require micromolar concentrations of peptide and involve different mechanisms (22). The adhesion inhibition and agglutination activities that we observed in vitro for cell-derived Aβ (Fig. 3) fall within physiological concentration ranges reported for normal human CSF (1 to 5 ng/ml). Consistent
were infected with transgenic GMC101 nematode gut. Fig. 6. Intestinal infection with Candida induces Aβ fibrillization in transgenic GMC101 nematode gut. Aβ42-expressing GMC101 C. elegans were infected with C. albicans (Candida) and probed for anti-Aβ immunoreactivity and β-amyloid markers using TEM and confocal fluorescence microscopy (CFM). (A) Micrograph shows positive labeling of yeast cell surface in GMC101 worm gut by immunogold nanoparticles coated with anti-Aβ antibodies (α-Aβ-Au) after Candida ingestion. (B to D) Visible (VIS) and fluorescence signals from freeze-fracture nematode sections with advanced Candida infections. (B) Comparison of uninfected and infected worms. (C and D) Thioflavin S and anti-Aβ staining for gut yeast aggregates. Signals include anti-Candida immunoreactivity (α-Candida), Thioflavin S and enhanced fluorescence (ThS), anti-Aβ immunoreactivity (α-Aβ), and superimposed (Overlay) signals. Yellow denotes signal colocalization. Uninfected and infected CL2122 nematode controls were negative for anti-Aβ immunoreactivity and enhanced Thioflavin S fluorescence (figs. S2 and S8). Micrographs are representative of data from three or more replicate experiments and multiple discrete image fields (table S1B).

with a normal in vivo protective role, the highest cerebral concentrations of Aβ are in the leptomeninges (10 to 50 ng/ml) (40), the brain’s first line of defense against infection and a tissue enriched for LL-37 and other innate immune proteins (41). The high specific activity observed for cell-derived material is consistent with our previous finding that Aβ in human brain extracts is a potent anti-Candida agent (3). Classical AMP expression can be either constitutive or inducible (5). In our transgenic mouse, nematode, and cell culture models, constitutive expression of Aβ is maintained artificially. Hence, our models are not suitable for testing whether infection normally results in Aβ up-regulation. However, data from other investigators suggest that Aβ may be an inducible AMP. Host cell exposure to herpes simplex virus-1 (42), HIV-1 (42), spirochetes (43), or Chlamydia (44, 45) increases Aβ expression.

In in vitro assays, cell-derived and synthetic Aβ oligomers were more potent against Candida than were monomeric forms (45). The specific activities of synthetic ADDLs, although higher than nonoligomerized peptide, remain lower than cell-derived Aβ species. Peptide posttranslational modifications may enhance the AMP activity of cell-derived Aβ oligomers. However, oligomer conformation is also likely to play a key role. Neurotoxicity has been shown to be highly dependent on the arrangement of Aβ peptides within oligomeric assemblies. Oligomer morphology may also modulate Aβ’s protective antimicrobial activities. Protocols for preparing ADDLs and other synthetic Aβ assemblies are optimized for oligomer populations with neurotoxic, not antimicrobial, activities. Future protocols optimized for enhanced AMP activities may generate soluble synthetic Aβ oligomers with potencies that approach that of cell-derived material.

Aβ pathophysiology is thought to arise from an abnormal propensity to generate soluble oligomers. However, oligomerization is not a pathogenic behavior for AMPs, and it plays a key role in normal protective activities across this diverse group of proteins, including microbe agglutination and entrapment (35), the targeting (26, 30) and disruption of microbial cell membranes (4, 46), resistance to bacterial proteases (26, 27, 46), and expanding of the molecular diversity and protective functions of AMP families without commensurate genome expansion (28, 29). Our data and the widespread involvement of oligomerization in the protective actions of AMPs suggest that the brain’s pool of soluble Aβ may normally include physiologically functional oligomeric species that mediate protective antimicrobial activities. The intrinsic polymorphic stoichiometry of Aβ oligomers may also play a protective physiological role. As has been shown with classical AMPs, diverse polymorphic oligomer pools target a broader spectrum of pathogens and are more resistant to AMP-targeting microbial proteases than are homogeneous peptide populations.

The lectin activity of Aβ oligomers is thought to promote brain amyloidosis (34). Studies to date have focused on accelerated Aβ fibrilization induced by binding of endogenous brain proteoglycans and glycosaminoglycans. However, our findings suggest that Aβ oligomers also bind to microbial carbohydrates with high affinity (Fig. 3, G to J). Carbohydrate-binding activity among AMPs is widespread and normally protective, playing a key role in helping peptides to recognize and bind to microbial pathogens (22). Heparin-binding AMPs have high affinities for the unique microbial carbohydrates found in cell walls but also bind to host glycosaminoglycans (47). Consistent with our findings for Aβ, binding of classical AMPs to microbial carbohydrates can lead to rapid peptide fibrillization and amyloid-mediated antimicrobial activities (48). Dysregulated carbohydrate binding by
Aβ may play a role in AD amyloidogenesis. However, a normal role as an AMP would suggest that polymeric microbial cell surface carbohydrates may be the normal in vivo target for the heparin-binding activity of oligomeric Aβ species. Long recognized as a key defensive strategy among lower organisms, AMP-mediated microbial agglutination is also emerging as an important part of human immunity (49). AMP fibrillization appears to play a central role in this important protective activity (35). Most recently, in vivo fibrillization of HD6 has been shown to mediate not only agglutination but also microbial entrapment within an amyloid fibril network (36). Our findings suggest that fibrillization is also involved in Aβ-mediated agglutination and leads to the entrapment of microbial cells by Aβ fibrils. On the basis of our findings, we propose a three-stage model for the protective activity of Aβ in vivo. Our model parallels the agglutination and entrapment actions of amyloidogenic HD6 (36). First, the VHHQKL heparin-binding domain of Aβ mediates targeting and binding of soluble oligomeric species to cell wall carbohydrates (Fig. S10A). Bound oligomers then provide a nidus and anchor for Aβ fibril propagation. Second, growing protofibrils interfere with microbial adhesion to host cells (Fig. S10B). Third, Aβ fibrils link, agglutinate, and then entrap the unattached microbial cells in a protease-resistant network of β-amyloid (Fig. S10C). Consistent with our model for the antimicrobial activities of Aβ, classical human AMPs have also been shown to generate amyloid fibrils on microbial surfaces that agglutinate pathogens and inhibit infection (35).

Consistent with our AMP model for Aβ, APP-KO mice show a trend for reduced pathogen resistance (Fig. 1E). However, the increase in infection-driven mortality among APP-KO mice was less marked than the increase in survival observed in the 5XFAD mouse model (Fig. 1A). For AMP-deficient models, immune impairment is often moderate because redundant activities among related members of AMP families can partially offset the loss of protection associated with low expression of individual AMP species (50). The well-studied human AMP LL-37 that serves as our model for Aβ’s AMP activity (3) is a member of the cathelicidin protein family. In humans, serious immunodeficiency associated with low LL-37 expression typically leads to fatal infections in childhood if untreated (37). However, mice lacking the murine LL-37 precursor protein (mCRAMP) show only a modest increase in mortality (5.1%) due to bacterial meningitis (51). Conversely, survival with infection among transgenic mice overexpressing human LL-37 is increased several-fold (52). APP-KO mice generate at least two Aβ homologs from amyloid precursor-like protein 1 (APLP1) and 2 (APLP2), which may help to mitigate loss of Aβ-mediated protection (53). Consistent with this model, APP, APLP1, and APLP2 and their nonamyloidogenic processing products show extensive functional redundancy (54), likely because of the gene duplication origin for this protein family. APP-KO mice also have an important additional limitation as models for the loss of Aβ-mediated protection. APP itself may be involved in central nervous system (CNS) immunity (55). It remains unclear how loss of activities normally mediated by full-length APP can be excluded as the source of attenuated infection resistance in APP-KO mice.

Genetically modified mice that lack proteases [BACE1 (β-site APP cleaving enzyme 1) and BACE2] for generating the Aβ family of peptides provide an alternative Aβ-null model. Consistent with our data, knockout BACE-KO mice that lack BACE have been reported to have marked immunodeficiency. Whereas neonatal mortality is below 2% under sterile conditions, in less stringently antiseptic environments, up to half of pups born to BACE-KO mice die from infections within the first 2 weeks of life (56). Benchmark tests for adaptive immunity have failed to identify defects in the response of BACE-KO mice to immune challenges. Findings for BACE-KO mice appear consistent with an innate immune deficiency and a possible normal protective role for Aβ. However, as with APP-KO mice, it is unclear how to demonstrate that the immunodeficiency in BACE-KO mice is specific for a loss of members of the Aβ family of peptides. Additional data are required to conclusively link the etiology of BACE-KO mouse immunodeficiency to low Aβ.
Our findings for Aβ and β-amyloid may have corollaries for amyloidopathies beyond AD. Protein fibrillization may be important not only for Aβ’s AMP activities but also for the normal actions of other amyloidosis-causing proteins. An association between amyloidosis and chronic bacterial infections has been recognized for almost a century (57), but the potential protective activities of host-generated amyloid have only recently emerged (4, 35, 58). At least six amyloidosis-associated peptides show antimicrobial activities, including amylin (59), atrial natriuretic factor (9), prion protein (60), cystatin C (61), lysozyme (5), and superoxide dismutase (62). Conversely, host AMPs have been identified that generate protective amyloids localized to infection sites (4). AA-type amyloidosis involves both systemic deposition of the acute-phase opsonin AMP serum amyloid A and has an infection-driven etiology (63). It remains to be determined whether serum amyloid A or other amyloidosis-causing AMPs also engage in nonpathogenic fibrillation pathways that help to protect against infection. However, should this prove to be the case, Aβ may be the first member of a new class of AMPs in which amyloid-generating activities protect against local infections but can also lead to widespread pathological amyloidosis.

If confirmed, our model carries important implications for understanding the pathogenesis of amyloidosis in AD. Excessive β-amyloid deposition may arise not from an intrinsically abnormal propensity of Aβ to aggregate but instead may be mediated by dysregulation of the brain’s innate immune system, for example, the consequence of an immune response mounted to microbial or sterile inflammatory stimuli. Our new model is congruent with the amyloid hypothesis and the importance of Aβ and β-amyloid in the neurodegenerative cascade of AD. However, our model would shift the modality of Aβ’s pathophysiology from abnormal stochastic behavior toward dysregulated antimicrobial activities.

Our study used genetically modified cell and animal models to generate data consistent with a normal physiological role for Aβ as an AMP. However, it remains unclear from these data how important a role Aβ plays in normal infection resistance. To address this question, additional data will be needed from wild-type animals modeling common physiological routes of infection. Further investigation will also be needed to clarify the extent to which the normal antimicrobial activities of Aβ identified in our study affect AD pathology.

It is important to emphasize that although infection of 5XFAD mice with S. Typhimurium seeded and accelerated β-amyloid deposition, the presence of a CNS infection is not implicit in our proposed AD amyloidosis model. Our work has identified what we believe is the normal role of Aβ. What drives widespread β-amyloid deposition in AD remains unclear. Among sterile inflammatory diseases, dysregulated innate immune responses rather than infections are emerging as drivers of pathology. Notably, two of the three confirmed AMP amyloidopathies are not linked to obvious infections (4, 9, 64). However, a large body of data accrued over nearly a century suggests that genuine infection may also play a role in AD etiology (65). Moreover, although a causal link to amyloidosis remains to be conclusively demonstrated, recent epidemiological findings have given increased prominence to the “infection hypothesis,” including studies linking brain fungal infection to AD (17, 18) and data showing that risk for the disease increases with infectious burden (66). Our findings do not constitute direct evidence of a role for infection in AD etiology. However, they do suggest a possible mechanism for pathogen-driven β-amyloid amyloidosis. Our data also suggest the possibility that a range of microbial organisms may be able to induce β-amyloid deposition, a possible reason for why a single pathogen species has not yet been identified that is overwhelmingly associated with AD. Future studies systematically characterizing microbial pathogens (viral, bacterial, and fungal) in the brains of AD patients, for example, by RNASeq, will be necessary to further interrogate whether specific clinical pathogens seed β-amyloid as part of the brain’s innate immune system. In any case, whether infectious or sterile inflammatory stimuli drive AD pathology, the pathways that regulate innate immunity in the brain may offer significant new targets for therapeutic intervention.

**MATERIALS AND METHODS**

**Study design**

Protective activities associated with Aβ expression were investigated in murine, nematode, and cell culture models of infection. Transgenic mice, nematode, and cell culture models were used that constitutively express human Aβ at high levels. Experiments also included a null-Aβ mouse model. Modulation of infection resistance with peptide expression is considered a hallmark for identity as an AMP. Initial experiments tested for Aβ-mediated increase (high-expression models) or decrease (null-Aβ mice) in survival after infection. End points were death for cultured cells and nematodes and moribundity for mice in accordance with Institutional Animal Care and Use Committee guidelines. Experiments were conducted blind as to cell, nematode, and mouse genotypes. The mechanism of protection afforded by high Aβ expression was then characterized in our cell culture monolayer infection model. We have previously shown parallels between Aβ activities and LL-37, a highly characterized human AMP. LL-37 was used as a model to elucidate the mechanisms for Aβ targeting, adhesion inhibition, and agglutination activities against microbial cells. Finally, nematode and mice models were tested to confirm in animals the potential protective microbial entrapment role of Aβ fibrillization revealed by cell culture experiments. Figure legends include details of replicate experiments used to generate data sets.

**Monomeric and oligomeric synthetic peptide preparation**

Synthetic Aβ1–40 (Aβ40), Aβ1–42 (Aβ42), scrambled Aβ42 (scAβ42), and LL-37 peptides were prepared and purified by J. L. Elliott at Yale University (New Haven, CT) using solid-phase peptide synthesis. Bulk powdered Aβ peptides were initially dissolved and incubated (18 hours) at room temperature (RT) in 30% trifluoroethanol (1 mg/ml) before lyophilization and storage (−20°C) under nitrogen. Before experimentation, dried peptide films were solubilized in 10 mM NaOH. For preparation of monomeric Aβ stocks, peptide solutions were diluted into phosphate-buffered saline (PBS) and fractionated by size exclusion chromatography, and peak monomer fractions (3 to 6 kDa) were pooled. Monomer stocks were stored on ice at 100 μM and used within 2 hours of preparation. Synthetic Aβ42 oligomer preparations (ADDLs and prototribilis) were generated from NaOH peptide stocks using established protocols (67). Peptide concentrations in stock solutions were determined by bicinchoninic acid protein assay and confirmed in experimental serial dilutions by densitometry analysis of anti-Aβ immunoblots.

**Candida inoculants and lawns**

Freezer stocks of C. albicans strain 90028 were obtained from the ATCC. C. albicans stocks were maintained on yeast extract peptone media.
dextrose (YPD) agar at 4°C with subculture to fresh plates every 2 weeks.
C. elegans pathogenicity plates were prepared by streaking (10 μl) sterile 35-mm tissue culture plates (BD Falcon) with yeast grown overnight (30°C) in YPD broth. Plates were incubated at 25°C for 2 hours to generate C. albicans lawn.
Synchronized hyphal yeast for cell culture experiments were prepared by single-colony transfer of C. albicans stock to 5 ml of minimal sugar medium (Formedium) and 48-hour static incubation at RT (68). After pelleting (1750 relative centrifugal force for 2 min) and PBS washing, starved yeast were resuspended in RPMI 1640 medium (HyClone) and concentration was adjusted to 2.5 × 10^6 cells/ml. Stock yeast in RPMI were diluted 10-fold into unconditioned culture medium immediately before inoculation of host cell slide or culture plate wells. Yeast concentration in inoculates was determined using a Bio-Rad TC20 automated cell counter and confirmed by counting CFU after serial dilution and streaking on agar.

S. Typhimurium inoculants
S. enterica serotype Typhimurium SL1344 stocks were provided by B. Cherayil (Mucosal Immunology Department, Massachusetts General Hospital, Boston, MA). Colonies were maintained on agar and subcultured to fresh plates every 3 weeks. Inoculant stocks were prepared by single S. Typhimurium colony to transfer to Luria-Bertani agar with streptomycin (100 μg/ml) and incubation overnight in a shaker incubator (225 rpm at 37°C). After PBS washing, pelleted (10,000g × 2 min) bacteria were resuspended in inoculation medium and diluted to required concentration. Bacterial concentrations in stocks were determined by comparing inoculum turbidity to McFarland turbidity standards and confirmed by streaking on agar and counting CFU.

For mouse experiments, S. Typhimurium inoculants were pathologized before infection by incubation in Luria broth with streptomycin (100 μg/ml) overnight at 37°C. Pathogenicity plates for C. elegans were prepared by streaking inoculate (10 μl) onto Pseudomonas aeruginosa and S. enterica–killing assay plates and overnight incubation at 37°C. For host cell monolayers, inoculant was added directly to culture medium.

Immunodepletion
Protein G Plus Agarose slurry (Pierce) was pelleted, washed, and incubated for 2 hours at RT with 4G8 (epitope: Aβ17–24) monoclonal antibody (mAb) (Covance) or control mouse IgG in PBS. After washing, beads were incubated with medium samples for 2 hours at RT under conditions equivalent to 10 μg of antibody per milliliter of medium. Beads were pelleted, and soluble fractions were removed, filtered (0.2 μm), and assayed to confirm Aβ depletion.

Aβ binding ELISA
The wells of 96-well plates were coated with live yeast by overnight incubation (37°C) with synchronized C. albicans (50 to 250 CFU per well) in RPMI medium (200 μl per well). Wells were washed to remove unattached yeast, and adhering C. albicans cells were then killed and covalently fixed in place by incubation (15 min at RT) with 4% paraformaldehyde. Wells were blocked (2 hours at RT) with 2% bovine serum albumin (BSA) in PBS before incubation with experimental samples. Bound Aβ in wells was detected immunochemically by incubation (overnight at 4°C) with α-Aβ42-HRP (horseradish peroxidase) (Covance) diluted 1:1000 in blocking buffer and development with 100 μl of chemiluminescence reagent (Pierce). Wells were washed (five times) with PBS between incubations.

Mouse infection model
Female 5XFAD (12) APP/PS1 doubly transgenic mice co-overexpress and co-inherit FAD mutant forms of human APP (the Swedish mutation: K670N/M671L; the Florida mutation: I716V; the London mutation: V717I) and PS1 (M146L/L286V) transgenes under transcriptional control of the neuron-specific mouse Thy-1 promoter (Tg6799 line). 5XFAD lines (B6/SJL genetic background) were purchased from The Jackson Laboratory and maintained by crossing heterozygous transgenic mice with B6/SJL F1 breeders. All 5XFAD transgenic mice were heterozygotes with respect to the transgene. Animal experiments were conducted in accordance with institutional and National Institutes of Health guidelines.

One-month-old mice received a single injection of 65,000 CFU (0.18 to 0.20 μl) of S. Typhimurium suspension at anterior/posterior, −1.6; medial/lateral, +1.5; dorsal/ventral, −1.6/−1.1/−0.7 using a 5-μl Hamilton syringe with a 30-gauge needle attached to a digital stereotaxic apparatus and an infusion pump at a rate of 0.15 μl/min. After infusion was completed, the needle remained in place for 10 min before slow withdrawal. Mice were given food and water on the cage floor starting 24 hours after the injection. Control sham infections used S. Typhimurium heat-killed before injection.

Clinical scores were recorded every 8 hours according to modified grading criteria for mouse encephalomylitis (69, 70). Clinical criteria are summarized in fig. S1A. Clinical progression was followed to morbidity, and then mice were sacrificed. Scores were recorded for each mouse and expressed as means ± SEM.

Mouse tissue preparation and sectioning
For immunofluorescence, mice were deeply anesthetized with a mixture of ketamine and xylazine and perfused transcardially with 4% paraformaldehyde in cold PBS. Brains were postfixed overnight and then transferred into a 30% sucrose solution until sedimented. Coronal sections (40 μm) were cut from an ice-cooled block using a sliding microtome (Leica). Sections were stored at −20°C in cryoprotective buffer containing 28% ethylene glycol, 23% glycol, and 0.05 M phosphate until processing for analysis.

Immunofluorescence labeling of mouse sections
Immunofluorescence labeling was performed as previously described (12). Primary antibodies include rabbit anti-GFAP (1:500, Dako) for astrocytes, rabbit anti-ionized calcium-binding adaptor molecule 1 (Iba1, 1:500, Wako) for microglia, and anti-Salmonella polyclonal rabbit for S. Typhimurium. Bound primary antibodies were detected with rabbit anti-Alexa Fluor 594 (Invitrogen, 8889S). Cell nuclei in sections were stained with TO-PRO-3 iodide (1:500, Life Technologies).

Immunoo- and Thioflavin S–costained mouse sections
Immunofluorescence labeling was performed as described previously (71). Briefly, sections were incubated with primary anti-Salmonella polyclonal rabbit IgG (1:1000) (PA1-7244, Thermo Fisher Scientific), followed by secondary anti-rabbit Alexa Fluor 594 (1:500) (Invitrogen, 8889S) antibodies. For Aβ staining, sections were incubated with mouse mAbs 3D6 (Ell Lilly) (mouse brain sections) or 4G8 (nematode sections). Bound anti-Aβ antibodies were detected by incubation with anti-mouse Alexa Fluor 488 (1:500) (Life Technologies, A11001)

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antibodies (Invitrogen). After immunostaining, free-floating sections were incubated (8 min) with 0.002% Thioflavin S in tris-buffered saline (TBS), rinsed twice for 1 min in 50% methanol washed for 5 min in TBS, and mounted with ProLong Gold antifade reagent (Life Technologies). Stained sections were analyzed by CFM (Leica TCS SL, Leica Microsystems).

C. elegans model
Two previously described transgenic C. elegans strains were used in experiments. GMC101, dvlIs100 [pCL354(unc-54a:DA-AB1-42) + pCL26 (mtl-2: GFP)] nematodes express human Aβ42 in body wall muscle and GFP in intestinal cells (14). Control C. elegans CL2122 dvlIs15(mtl-2: GFP) nematodes express GFP but not the Aβ42 peptide (14). Worms were synchronized before experimental treatments according to established protocols (72). Briefly, unhatched eggs were released by treating gravid worms with bleach. After an overnight incubation, arrested L1 larvae were added to Escherichia coli OP5050 lawns and incubated at 20°C to generate synchronized L4 larvae (48 hours) or adult (60 hours) nematodes.

For infection experiments, 100 to 150 synchronized L4 stage worms were incubated (2 hours at 25°C) on C. albicans lawns, washed with M9 buffer to remove surface C. albicans, and transferred to six-well culture plates containing 1.5 ml per well of incubation medium (79% M9 buffer, 20% brain-heart infusion, cholesterol (10 μg/ml) in ethanol, and kanamycin (90 μg/ml)]. Nematodes were incubated at 25°C and monitored daily for the distinctive distention and penetrative filamentation that characterize Candida-induced mortality.

Nematode freeze-fracture and immunostaining
Worms (L4) were transferred dropwise to poly-lysine–coated slides and covered with a coverslip. Gentle pressure was applied to the coverslip before the slide assembly was placed on a metal block and flash-frozen using liquid nitrogen. The coverslip was flicked off, and fractured samples were fixed by 5-min incubations with absolute alcohol, followed by acetone. Dried samples were ringed with petroleum jelly and covered with a second coverslip. Slide staining was performed in a wet chamber. For immunostaining, slides were blocked for 15 min with blocking buffer [10% Tween and powdered milk (0.2 g/ml) in PBS]. Wells were incubated (2 hours at RT) with rabbit polyclonal anti-A. albicans antibodies (Invitrogen). After immunostaining, free-floating sections were incubated (8 min) with 0.002% Thioflavin S in tris-buffered saline (TBS), rinsed twice for 1 min in 50% methanol washed for 5 min in TBS, and mounted with ProLong Gold antifade reagent (Life Technologies). Stained sections were analyzed by CFM (Leica TCS SL, Leica Microsystems).

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Host cell monolayer model
Host cell monolayers were prepared from nontransformed and transformed human neuroglioma (H4) or CHO cell lines. Stable transformed H4 cell lines that secrete Aβ40 (H4-Aβ40) or Aβ42 (H4-Aβ42) without overexpression of the APP have been described previously (19). Stable transformed CHO-CAB cells coexpressing human ATCC Swedish mutation and BACE1 were generated by transfecting a pcDNA3.1- BACE1-myc construct into CHO-APP751 cells that overexpress mutant APP751 (K670N/M671L: Swedish mutation) (20).

Nontransformed H4-N and CHO-N cell lines were maintained in complete medium containing Dulbecco’s modified Eagle’s medium (DMEM), 10% fetal bovine serum (FBS), 2 mM l-glutamine, 100 U penicillin, and streptomycin (100 μg/ml). Complete medium for transformed H4-Aβ40 and H4-Aβ42 cells included hygromycin (150 μg/ml) and media for CHO-CAB Zeocin (200 μg/ml) and G418 (200 μg/ml).

To prepare experimental HCMs, trypsinned host cells suspended in antibiotic-free DMEM with 5% FBS and 2 mM l-glutamine were transferred (300,000 and 500,000 cells/ml for H4 and CHO lines, respectively) to the wells of Lab-Tek eight-chamber glass slides (Thermo Scientific) (200 μl per well) or 96-well culture plates (100 μl per well) and incubated for 24 hours. Cell confluence in chamber slides and plate wells was confirmed by microscopic examination. Automated cell counter analysis of well trypsin extracts confirmed that, in control uninfected cell monolayers, nontransformed and transformed cell numbers did not diverge by more than 6% before infection or after the final experimental incubation (fig. S4D).

Nontransformed and transformed culture media were conditioned for 36 hours before inoculation with Candida. HCMs in culture plates were infected by addition of Candida inoculant aliquots (10 μl) containing 2000 or 250,000 CFU, respectively. For host cell survival experiments, Candida were incubated with H4 and CHO cells for 28 and 36 hours, respectively. HCMs were then washed and assayed for host cell survival.

Host cell Brdu labeling
Subconfluent nontransformed and transformed H4 and CHO cells were incubated overnight (10-cm culture dishes) in complete culture medium containing 10 mM BrdU. Confluent BrdU-labeled cell cultures were PBS-washed (three times) to remove free BrdU, then trypsinned, and used for preparation of HCMs in 96-well culture plates. After experimental treatments, plate wells were washed with PBS (three times), then fixed and permeabilized, and assayed according to the manufacturer’s instructions (Cell Proliferation BrdU ELISA, Roche).

Imaging C. albicans host cell adherence
Cell monolayers in eight-well chamber slides were infected with synchronized hyphal yeast (10,000 CFU per well) by addition of a 10-μl aliquot of freshly prepared C. albicans inoculate to culture medium (200 μl per well) preconditioned for 36 hours with host cells. Infected slides were incubated for 2 hours, medium was removed by aspiration, and wells were washed with PBS (three times) then fixed by 10-min incubation with 4% paraformaldehyde. Fixative was removed, and wells were washed (three times) before incubation (30 min) with Calcofluor white M2R fungal surface stain (Life Technologies) (73). Wells were water-washed and coverslipped before imaging by fluorescence microscopy (excitation, 360 nm/emission, 460 nm).

Immunohistochemical detection of C. albicans adhering to cell monolayers
Experiments were performed using HCMs prepared in white opaque 96-well culture plates. HCMs were infected with synchronized hyphal yeast (1000 CFU per well) by addition of 10 μl of freshly prepared C. albicans inoculate to wells containing preconditioned (36 hours) culture medium (100 μl per well). Wells were incubated 18 hours with yeast before aspiration of medium, gentle washing with PBS (three times), and fixation by 10-min incubation with 4% paraformaldehyde. Fixative was removed, and wells were washed (three times) before
incubation (1 hour) with blocking buffer [2% albumin in tris-buffered saline–TWEEN 20 (TBST)]. Wells were then incubated (2 hours) with fresh blocking buffer containing a 1:5000 dilution of α-Candida-HRP antibody (Abcam). Wells were washed with TBST (five times) and fluorescent-captured (excitation, 320 nm/emission, 420 nm) after development with QuantaBlu (Pierce), a fluorescent HRP substrate.

**C. albicans** adhesion assay for abiotic surfaces

Experiments used a modified method of Tsai *et al.* (22) to assay *C. albicans* adhesion to polystyrene in conditioned culture medium. Synchronized hyphal yeast (1000 CFU per well) were incubated (37°C) in the wells of clear, flat-bottom polystyrene 96-well microtiter plates containing host cell–conditioned (36 hours) culture medium (200 µl per well). Incubation medium was removed by aspiration, and wells were washed with water. For dye staining, slides were incubated in the buffer was blotted off. Slides were air-dried to fix yeast and then carefully rinsed with water. For immunolabeling experiments, specimens pre-stained with Thioflavin S (5 min) or staining solution and antibody labeling of *C. albicans* aggregates

Aggregated yeast were pelleted (2 min × 500g), washed with PBS (twice), and transferred to glass slides in minimal volume, and excess buffer was blotted off. Slides were air-dried to fix yeast and then carefully rinsed with water. For dye staining, slides were incubated in the dark at RT with 50 µl of Calcofluor white fungal stain solution 6726 (Eng Scientific). After washing, attached hyphae were detected by measuring well fluorescence (excitation, 360 nm/emission, 460 nm).

**C. albicans** aggregation assay

Host cell–conditioned (48 hours) culture medium (200 µl per well) was incubated (overnight at 37°C) with synchronized yeast (200 cells per well) in the wells of clear 96-well microtiter plates. Incubation medium was removed, and yeast pellets were washed twice with PBS. During aspiration, care was taken to minimize disturbance of settled yeast at the well bottom. Settled yeast pellets were resuspended in PBS and transferred to fresh wells. Low-magnification (×4) bright-field well images were captured at maximum condenser aperture. Images were then analyzed for yeast aggregates using ImageJ software (version 1.47) with the following procedure. Captured image files were first converted from 8-bit RGB to 8-bit grayscale and then further transformed to 1-bit black and white images using a conversion threshold of 86%. Well area covered by yeast aggregates was determined from pixel counts of transformed black and white images using the Analyze Particle tool with lower size threshold set to 50 pixels. Isolated black pixel counts of transformed black and white images were captured at maximum condenser aperture. Images were then analyzed for yeast aggregates using ImageJ software (version 1.47) with the following procedure. Captured image files were first converted from 8-bit RGB to 8-bit grayscale and then further transformed to 1-bit black and white images using a conversion threshold of 86%. Well area covered by yeast aggregates was determined from pixel counts of transformed black and white images using the Analyze Particle tool with lower size threshold set to 50 pixels. Isolated black areas of less than 50 pixels (four to six yeast cells) were not included in aggregate totals.

Staining and antibody labeling of *C. albicans* aggregates

Aggregated yeast were pelleted (2 min × 500g), washed with PBS (twice), and transferred to glass slides in minimal volume, and excess buffer was blotted off. Slides were air-dried to fix yeast and then carefully rinsed with water. For dye staining, slides were incubated in the dark at RT with 50 µl of Thioflavin S (5 min) or staining solution and antibody labeling of *C. albicans* aggregates

Aggregated yeast were pelleted (2 min × 500g), washed with PBS (twice), and transferred to glass slides in minimal volume, and excess buffer was blotted off. Slides were air-dried to fix yeast and then carefully rinsed with water. For dye staining, slides were incubated in the dark at RT with 50 µl of Thioflavin S (5 min) or staining solution and then incubated (30 min at RT) with mAb 4G8 diluted 1:1000 in blocking buffer. The grids were washed with PBS (three times) and incubated with goat anti-mouse IgG antibody covalently linked to nanogold particles. After three 5-min PBS washes and four 10-min water washes, specimens were fixed with 1% glutaraldehyde (10 min at RT). Specimens were washed with water, stained with uranyl acetate, and then viewed using a JEM-1011 transmission electron microscope (JEOL Institute).

**Statistical analysis**

Statistical analyses were performed with Prism software (version 6.0c). Arithmetic means were compared using two-tailed *t* tests. Survival curves were compared using log-rank (Mantel-Cox) test and confirmed by Gehan-Breslow-Willcoxon test. *P* values <0.05 were considered statistically significant.

**SUPPLEMENTARY MATERIALS**

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Materials and Methods

Fig. S1. AJ deposition and inflammation in 5XFAD mice before infection and criteria used for assessing clinical performance after infection. Fig. S2. AJ42 localizes to gut and muscle in GMC101 nematodes. Fig. S3. Aj protein protects GMC101 nematodes and CHO-CAB cells against S. Typhimurium. Fig. S4. Confirmation of increased Candida resistance among transformed host cells using three independent assays. Fig. S5. Transformed cell lines generate Ajl oligomers at levels found in the soluble fraction of human brain. Fig. S6. Transformed H4-AJ40 and CHO-CAB host cells resist Candida colonization and agglutinate yeast. Fig. S7. Birefringence of Congo red–stained yeast aggregates from H4-AJ42 medium. Fig. S8. Anti-Aj antibodies do not label CL2122 tissues or yeast. Fig. S9. B-Amyloid colocalizes with S. Typhimurium cells in 5XFAD brain. Fig. S10. Model for antimicrobial activities of soluble Ajl oligomers. Table S1. Figure micrographs are representative of data from multiple repeat experiments and image fields.

Video S1. Z-section projection of 5XFAD mouse brain showing β-amyloid entrapment of S. Typhimurium cells.

**REFERENCES AND NOTES**


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Amyloid-β peptide protects against microbial infection in mouse and worm models of Alzheimer's disease

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Rehabilitation of a β-amyloid bad boy

A protein called Aβ is thought to cause neuronal death in Alzheimer's disease (AD). Aβ forms insoluble aggregates in the brains of patients with AD, which are a hallmark of the disease. A β and its propensity for aggregation are widely viewed as intrinsically abnormal. However, in new work, Kumar et al. show that Aβ is a natural antibiotic that protects the brain from infection. Most surprisingly, A β aggregates trap and imprison bacterial pathogens. It remains unclear whether A β is fighting a real or falsely perceived infection in AD. However, in any case, these findings identify inflammatory pathways as potential new drug targets for treating AD.