Mounting evidence implicates antiretroviral (ARV) drugs as potential contributors to the persistence and evolution of clinical and pathological presentation of HIV-associated neurocognitive disorders in the post-ARV era. Based on their ability to induce endoplasmic reticulum (ER) stress in various cell types, we hypothesized that ARV-mediated ER stress in the central nervous system resulted in chronic dysregulation of the unfolded protein response and altered amyloid precursor protein (APP) processing. We used in vitro and in vivo models to show that HIV protease inhibitor (PI) class ARVs induced neuronal damage and ER stress, leading to PKR-like ER kinase–dependent phosphorylation of the eukaryotic translation initiation factor 2α and enhanced translation of β-site APP cleaving enzyme-1 (BACE1). In addition, PIs induced β-amyloid production, indicative of increased BACE1-mediated APP processing, in rodent neuronal cultures and human APP-expressing Chinese hamster ovary cells. Inhibition of BACE1 activity protected against neuronal damage. Finally, ARVs administered to mice and SIV-infected macaques resulted in neuronal damage and BACE1 up-regulation in the central nervous system. These findings implicate a subset of PIs as potential mediators of neurodegeneration in HIV-associated neurocognitive disorders. (Am J Pathol 2017, 187: 91–109; http://dx.doi.org/10.1016/j.ajpath.2016.09.006)
suggested to contribute to these changes in HAND neuropathology, other risk factors, such as peripheral toxicities of ARVs and potential central effects linked to enhanced amyloidogenesis and aging, remain largely underexplored.

Among the six classes of ARVs currently prescribed, nucleoside reserve transcriptase inhibitors (NRTIs) and HIV protease inhibitors (PIs) have the longest history of use and continue to be prescribed globally. NRTIs inhibit the viral reverse transcriptase enzyme,12 and PIs prevent gag-pol precursor polyprotein cleavage to block viral maturation.13 As a result of combination ARV therapy that incorporates several drugs from different categories with PIs, HIV-infected individuals have experienced a 50% reduction in mortality rates with reduced HIV-associated dementia incidence.14 Despite their numerous benefits, ARVs were also linked to wide-ranging, peripheral metabolic and neural disturbances that could themselves influence HAND progression, including NRTI-associated mitochondrial dysfunction manifesting as peripheral neuropathy and myopathy14,15 and PI-associated dyslipidemia, hypercholesterolemia, and metabolic syndrome.16–22

We previously demonstrated that ARV administration lead to synaptodendritic damage in healthy adult rats and SIV-infected macaques.23 We and others also showed ARV-mediated accumulation of reactive oxygen species and eventual synaptic damage and neuronal death in cortical neurons.23,24 These data support a growing body of evidence for significant ARV-associated CNS neurotoxic effects, as was shown in the peripheral nervous system, and suggest that ARVs may affect long-term neurocognitive outcomes in patients. More importantly, both NRTIs and PIs were shown to induce numerous cellular stress response pathways, including endoplasmic reticulum (ER) and oxidative stress, both of which were demonstrated in HAND and other neurodegenerative diseases, such as Alzheimer disease (AD).25,26 Diverse cellular stressors converge on a ubiquitous intracellular response mechanism termed the unfolded protein response (UPR; alias ER stress response).27,28 In eukaryotes, misfolded protein accumulation and/or other stressors disrupt the interaction between the ER-resident chaperone, binding protein (BiP), and the luminal domain of one or more of the three ER membrane-bound UPR sensors, PKR-like ER kinase (PERK), activating transcription factor 6 (ATF6), and inositol-requiring enzyme 1α, in the ER, permitting their consequent activation29,30; however, UPR activation is promiscuous as it can be activated in response to a wide range of stimuli. Downstream effects of UPR induction include global protein synthesis attenuation via eukaryotic translation initiation factor 2α (eIF2α) phosphorylation, up-regulation of genes containing the ER stress response element, and ER-mediated protein degradation.30

Although transient UPR activation serves a protective function, the chronic nature of cellular stress and stress response activation in the CNS during HIV infection may ultimately overwhelm ER’s buffering capacity, resulting in aberrant UPR activation and eventual cell death.26,31 We and others demonstrated a potential role for ER stress and UPR activation in HAND pathogenesis, as well as in other neurodegenerative diseases.22,26,32,33 In postmortem brain tissue from ARV-treated, HIV+ patients and AD patients, expression levels of BiP, phosphorylated eIF2α (p-eIF2α), ATF6β, and other UPR markers were increased, primarily in neurons.26,30,34 Interestingly, Vassar and colleagues35,36 showed that in vivo UPR activation via glucose deprivation or tunicamycin in the CNS resulted in increased APP and β-amyloid (Aβ) deposition35–37 and that impaired brain energy metabolism lead to chronic PERK activation, sustained eIF2α phosphorylation, and β-site APP cleaving enzyme 1 (BACE1) up-regulation.36 Phosphorylation of eIF2α inhibits global protein translation while up-regulating the translation of a subset of mRNA species that contain multiple upstream open reading frames in their 5’ untranslated regions (5’ UTRs), including BACE1.36,37 BACE1, an aspartyl protease critical for the generation of Aβ from APP, was up-regulated in the CNS of AD patients.38,39 Increased BACE1 expression and activity were correlated with plaque number and cognition status, and BACE1 was implicated as a biomarker for mild cognitive impairment and AD.39 Altogether, these studies suggest that UPR-mediated up-regulation of BACE1 may function as an early event in AD pathogenesis.

In the present study, we determined the mechanisms by which ARVs induced damage and death in primary neuroglial cultures and in vivo models. We also assessed whether altered BACE1 expression and function were contributors. We showed that two PIs, ritonavir and saquinavir, which were previously shown to induce oxidative stress and synaptic loss,23,24 induced neuronal damage and ER stress in neuroglial cultures, leading to UPR activation and PERK-dependent translational BACE1 up-regulation. In addition, we found that PIs enhanced the production of Aβ peptides in neurons and human APP-expressing Chinese hamster ovary (CHO) cells. Moreover, these findings were corroborated with increased expression of BACE1 in macaques and rodents that received chronic ARV treatment in vivo. These studies suggested that PIs might alter APP processing and contribute to the changing neuropathology and persistence of cognitive decline among ARV-treated, HIV+ individuals.

Materials and Methods

Chemicals and Reagents

The following antibodies used in this study were purchased from the indicated vendors or provided by the indicated colleagues: Abcam (Cambridge, MA): synaptophysin (ab8049), a disintegrin and metalloprotease 10 (ADAM10) (ab1997), and microtubule-associated protein (MAP2) (5392); BD Biosciences (San Jose, CA): BIP (610978); Cell Signaling Technology (Danvers, MA): BACE1 (56065S), phospho-eIF2α (9721), total eIF2α (9722), lamin A/C (2032), and presinilin-1 (5643); Covance (Princeton, NJ):...
APP (6E10); Santa Cruz Biotechnology (Dallas, TX); glyceraldehyde-3-phosphate dehydrogenase (sc-32233) and actin (A2066). The mouse monoclonal antibody against BACE1 (3d5) was a generous gift from Dr. Robert Vassar (Feinberg School of Medicine, Northwestern University, Chicago, IL). The antibody against activating transcription factor 4 (ATF4) was generously provided by Dr. Ron Wek (School of Medicine, Indiana University, Indianapolis). The rabbit monoclonal antibody against PERK was a generous gift from J.A.D. The following chemical reagents used in the study were purchased from the indicated vendors: Citifluor, Ltd (London, UK): citifluor AF1; BioRad (Hercules, CA): bradford protein assay dye, polyvinylidene difluoride membrane, and prestained broad-range molecular weight ladder; Jackson ImmunoResearch Labs (West Grove, PA): all dye-conjugated secondary antibodies; Life Technologies (Carlsbad, CA): Dulbecco’s modified Eagle’s medium, neurobasal media, and B27 supplement; Millipore (Temecula, CA): 20S proteasome activity assay; Nanotherics (Newcastle under Lyme, UK): neuromag transfection reagent; Sigma (St. Louis, MO): DAPI, fast green FCF, protease inhibitor cocktail, 4-hydroxytamoxifen (H7904), anisomycin (A9789), and actinomycin D (A1410); Peptides International (Louisville, KY): poly-L-lysine; New England Biolabs (Ipswich, MA): tyramide amplification system; Scytek Labs (Logan, UT): normal antibody diluent; Thermo Fisher Scientific (Waltham, MA): SuperSignal West Dura extended-duration substrate and all horseradish peroxidase—conjugated secondary antibodies. Antiretroviral reagents were kindly provided by the AIDS Research and Reference Reagent Program, Division of AIDS, National Institute of Allergy and Infectious Diseases, NIH (Bethesda, MD).

Ethics Statement

All pigtailed macaques (Macaca nemestrina) were housed and cared for in an indoor ABSL2 facility at the fully Association for Assessment and Accreditation of Laboratory Animal Care International—accredited Johns Hopkins University Research Animal Resources facility in accordance with standards of the Animal Welfare Regulations, the eighth edition of the Guide for the Care and Use of Laboratory Animals, and the Weatherall Report. All animal studies were approved by the Johns Hopkins University Institutional Animal Care and Use Committee (approval number PR12M310). Animal welfare was optimized through social housing and provision of both edible and manipulanda environmental enrichment by a dedicated behaviorist. Water was supplied ad libitum, and animals were provided commercial monkey chow twice daily and fresh produce at least three times per week. Animals were monitored daily by animal care and veterinary staff and sedated with ketamine anesthesia for all sampling by skilled veterinary personnel. Euthanasia was performed under deep pentobarbital anesthesia by skilled veterinary personnel, and tissues were perfused to decrease contamination by blood and peripheral virus. The negative control animals were mock-inoculated, and samples were obtained at the same time points as for the inoculated, untreated macaques.

SIV/Pigtail Macaque Model of Lentiviral CNS Disease

Pigtailed macaques (M. nemestrina) were inoculated with SIV/DeltaB670 and SIV/17E-Fr, as described previously.11 Twelve days after infection, animals were divided into placebo and ARV-treated groups until necropsy (range, day 161 to 175). ARV treatment was composed of the NRTI tenofovir (Gilead, Oceanside, CA) at 30 mg/kg s.c. given once daily, saquinavir (Roche, Basel, Switzerland) and atazanavir (Bristol-Myers Squibb, New York, NY) at 205 and 270 mg/kg, respectively, given twice daily via oral administration, and an integrase inhibitor, L-870812 (Merck, Kenilworth, NJ), at 10 mg/kg orally twice daily.12 Tenofovir dosing was based on previous studies, whereas saquinavir and atazanavir dosing was based on pharmacokinetic analysis performed in pigtail macaques to reflect the area under the curve parameters similar to those detected in humans treated with atazanavir and saquinavir.13 L-870812 dosing was based on previous studies conducted in rhesus macaques.42

Rodent Model of ARV-Induced Neurotoxicity

All surgical procedures were performed with the approval of the Institutional Animal Care and Use Committee of the University of Pennsylvania (Philadelphia; approval number 803156). Adult male C57BL/6 mice were catheterized via the jugular vein, as described previously.15 Before surgery, mice were anesthetized with 80 mg/kg ketamine and 12 mg/kg xylazine. An indwelling silastic catheter was placed into the right jugular vein, sutured in place, and connected to a backmount platform (Strategic Applications, Inc., Lake Villa, IL) that was s.c. secured on the back of the animals with sutures. Catheters were flushed daily with 0.1 mL of the antibiotic clavulanic acid and ticarcillin (Timentin; bioWORLD, Dublin, OH) dissolved in heparinized saline (0.93 mg/mL). Treatment regimens were as follows: vehicle [dimethyl sulfoxide (DMSO)], ritonavir (20 mg/kg), or saquinavir (25 mg/kg), administered twice daily for 14 days by continuous i.v. injection. These drug doses were based on previously published pharmacokinetic studies in rodents.46–48 At the time of euthanasia, catheter patency was verified by response to pre-euthanasia sedation (80 mg/kg ketamine and 12 mg/kg xylazine). Euthanasia was performed by carbon dioxide administration, followed by cervical dislocation and declamping to confirm death. Brains were removed, frontal cortex and hippocampus were dissected on ice, and tissue samples were stored at −80°C until immunoblot analyses were performed (vehicle, n = 6; ritonavir, n = 7; saquinavir, n = 7).
Preparation of Primary Neuroglial Cultures

Primary rat or mouse cortical neuroglial cultures were isolated from brains of embryonic day 16 Sprague-Dawley rat pups or embryonic day 16.5 to 17.5 floxed PERK (PERKloxP/loxP) mice, with modifications of previously described protocols. Briefly, to generate neuroglial cultures, cortical cell suspensions were isolated from rat or mouse embryos and were plated on poly-L-lysine-coated tissue culture dishes. Cultures were maintained in neurobasal media supplemented with B27 at 37°C with 5% CO2, as described previously. Unless otherwise indicated, all experiments were performed at 21 days in vitro (DIV) in cultures containing approximately 90% neurons and 10% astrocytes/glia at a density of 500,000 cells/mL. Pure neuronal cultures were prepared by treating neuroglial cultures with 10 μmol/L cytosine arabinoside 48 hours after plating and were maintained in neurobasal media with B27 at 37°C with 5% CO2.

Human primary cortical neurons were obtained from the Comprehensive NeuroAIDS Center, Temple University (Philadelphia, PA). Cultures were prepared from fetal brain tissue (gestational age, 16 to 18 weeks) performed in full compliance with the NIH and Temple University ethical guidelines, as described previously. A total of 50% of the culture media was changed every 4 days, and cultures were maintained at 37°C with 5% CO2 for 21 DIV.

Previously characterized PERKloxP/loxP mice were crossed with tamoxifen-inducible cre (Cre-ERT2) heterozygous mice, and the resulting PERKloxP/loxP; Cre-ERT2 mice were kindly provided by Dr. Alan Diehl. PERK deletion was achieved by treating 10 DIV mouse neuroglial cultures with various concentrations of 4-hydroxytamoxifen. Antiretroviral drug treatments were performed 4 days after 4-hydroxytamoxifen treatment to allow for maximal PERK excision. Sixteen hours after antiretroviral drug treatments, PERK−/− neuroglial cultures were harvested for immunoblotting and genotyping using routine methods.

Drug Treatments

Twenty-one DIV rat cortical neuroglial cultures or human fetal neurons were treated with indicated log-fold concentrations of zidovudine (AZT; 25 mmol/L stock in sterile water), or saquinavir or ritonavir (both drugs prepared in DMSO at 25 mmol/L stock concentrations). The final concentration of DMSO was 0.04% for all treatments. All antiretroviral drugs were added to neuroglial cultures 16 hours before harvest. Anisomycin and actinomycin D were prepared in DMSO at 50 and 5 mmol/L stock concentrations, respectively. Twenty-one DIV rat cortical neuroglial cultures were pretreated with either 10 or 50 μmol/L anisomycin or 5 μmol/L actinomycin D for 12 hours, followed by treatment with 10 μmol/L ritonavir for 16 hours.

Immunoblotting

Whole cell extracts of primary rat cortical cultures were prepared with ice-cold whole cell lysis buffer containing 50 mmol/L Tris, pH 7.5, 120 mmol/L NaCl, 0.5% NP-40, and 0.4 mmol/L Na3VO4, supplemented with protease inhibitors. Lysates were centrifuged at 14,000 × g at 4°C for 10 minutes. Protein concentrations were determined using the Bradford method, and equal amounts of protein (15 to 20 μg) were loaded into each well of 4% to 12% Bis-Tris gradient gels or 3% to 8% Tris-acetate gradient gels (for confirmation of PERK overexpression). After transfer at 25 V at room temperature for 1 hour, polyvinylidene difluoride membranes were blocked in tris-buffered saline containing Tween 20 and 5% bovine serum albumin for at least 30 minutes at room temperature. Membranes were then incubated in primary antibodies overnight at 4°C, washed with tris-buffered saline and Tween 20, and incubated with appropriate horseradish peroxidase–conjugated secondary antibodies for 30 minutes at room temperature. Membranes were developed using SuperSignal West Dura extended duration substrate, and densitometric analyses were performed using NIH ImageJ software program version 1.44 (NIH, Bethesda, MD). Equal loading and transfer were confirmed by fast green staining of membranes. Band densities of interest were normalized to the indicated loading controls or fast green–stained membranes. Immunoblots shown are representative of at least three independent biological replicates.

RNA Isolation and Quantitative RT-PCR

Total RNA was isolated from primary rat neuroglial cultures using the RNeasy Mini Kit (Qiagen, Valencia, CA). To assess the expression of spliced XBP-1 and BACE1 genes, 1 μg total RNA from each sample was processed by SuperScript II First-Strand Synthesis system (Invitrogen, Carlsbad, CA). Approximately 100 ng cDNA was amplified via quantitative RT-PCR using Custom TaqMan Gene Expression Assays from Applied Biosystems (Foster City, CA). Relative expression of mRNA levels from samples was quantified using the comparative CT method, and data were normalized to unspliced XBP-1 and TBP for spliced XBP-1 and BACE1 genes, respectively. All samples were run in triplicate, and experiments were repeated in at least three biological replicates.

Human/Rat Aβ ELISAs

Cell media from 21 DIV primary rat neuroglial cultures and CHO cells stably expressing human wild-type APP were diluted 25-fold and assessed with a human/rat Aβ12- or Aβ40-specific sandwich enzyme-linked immunosorbent assay (ELISA) (Wako Chemicals, Richmond, VA), according to the manufacturer’s protocols. For experiments including CHO cells, cultures were pretreated with 10 nmol/L, a dps protocols. For experiments including CHO cells, cultures were pretreated with 10 nmol/L.
100 nmol/L, or 1 μmol/L BACE inhibitor IV (Calbiochem, Billerica, MA) for 12 hours, followed by treatment with 10 μmol/L ritonavir or vehicle (DMSO) for 16 hours, at which time media were collected. For rat neuroglial cultures, cells were treated with combinations of antiretroviral drugs, followed by media collection at 4 and 16 hours.

Human Soluble APP ELISAs

Cell culture media from CHO cells stably expressing human wild-type APP were pretreated with 10 nmol/L, 100 nmol/L, or 1 μmol/L BACE inhibitor IV for 12 hours, followed by treatment with 10 μmol/L ritonavir or DMSO vehicle for 16 hours, at which time media were collected. Conditioned media were then diluted 25-fold, and human soluble APPα (sAPPα) and sAPPβ levels were measured with human ELISA assays (IBL International, Toronto, ON, Canada), according to the manufacturer’s recommendations.

20S Proteasome Activity Assay

Whole cell lysates prepared from 21 DIV rat cortical neuroglial cultures treated with ritonavir, saquinavir, or lactacystin for 16 hours, as described above, were used to measure 20S proteasome activity (assay from Millipore), according to the manufacturer’s recommendations. A total of 10 μg protein per sample was used.

Immunofluorescence Staining of Neuroglial Cultures

To assess the neuroprotective effect of BACE1 inhibition, neuroglial cultures grown on coverslips were pretreated with 1 μmol/L BACE inhibitor IV for 30 minutes, followed by 12 hours treatment with 10 μmol/L ritonavir. Propidium iodide (15 μmol/L) was added to neuroglial cultures 15 minutes before the conclusion of treatments. Cultures were washed twice with phosphate-buffered saline (PBS) and fixed for 30 minutes with 4% paraformaldehyde/4% sucrose in PBS, followed by permeabilization in 0.2% bovine serum albumin plus 0.1% Triton X-100 in PBS for 1 hour at room temperature. Coverslips were then washed twice with PBS and incubated with anti-MAP2 antibody (1:1000) in normal antibody diluent overnight at 4°C. After washing in PBS–Tween 20, coverslips were incubated with a fluorescein isothiocyanate–conjugated goat anti-mouse IgG secondary antibody (1:200) for 30 minutes at room temperature. Coverslips were mounted onto slides, and the number of MAP2-positive cells was determined by blind counting of all fields (15 at 200×; by P.J.G., N.S.K., and G.V.D.W.) in three adjacent vertical columns through the center of each coverslip. Images were captured using a Nikon Eclipse E400 fluorescent microscope (Nikon Corp., Tokyo, Japan) equipped with an Olympus DP70 digital camera (Olympus Corp., Tokyo, Japan). Propidium iodide–positive puncta were hand-counted using NIH ImageJ software program version 1.44 and were plotted as fold changes compared to untreated controls. Averages were expressed as means ± SEM.

Pulse-Chase and Immunoprecipitation

Primary rat neuroglial cultures were grown in 60-mm dishes for 21 DIV in neurobasal media supplemented with B27. Cultures were pulse-labeled with 2 mL methionine/cysteine-free Dulbecco’s modified Eagle’s medium containing 0.4 mCi 35S-labeled methionine and cysteine for 30 minutes. Cultures were washed twice with sterile Dulbecco’s PBS and chased with neurobasal media plus B27 with or without 10 μmol/L ritonavir. At 0, 6, 12, 18, and 24 hours after labeling, cultures were harvested with ice-cold lysis buffer (50 mmol/L Tris, pH 7.5, 120 mmol/L NaCl, 0.5% NP-40, and 0.4 mmol/L Na3VO4, supplemented with protease inhibitors). Lysates were immunoprecipitated with Dynabeads Protein A (Thermo Fisher Scientific) and 2 μg anti-BACE1 (Cell Signaling Technology), run on a 4% to 12% Bis-Tris gradient gel for separation, and transferred onto polyvinylidene difluoride membranes for approximately 7 hours. Membranes were dried and exposed via autoradiography.

Subcellular Fractionation

To evaluate ATF4 nuclear translocation in response to antiretroviral drug treatments, 21 DIV rat neuroglial cultures were treated with 10 μmol/L ritonavir or vehicle (DMSO) for 16 hours, and cytoplasmic and nuclear fractions were prepared. Briefly, cultures were washed twice in ice-cold PBS and lysed for 10 minutes in cytoplasmic lysis buffer (10 mmol/L HEPES, pH 7.9, 10 mmol/L KCL, 10 mmol/L EDTA, 1 mmol/L dithiothreitol, and 0.4% NP-40, supplemented with protease inhibitors). Extracts were centrifuged at 4°C for 3 minutes at 15,000 × g, and cytoplasmic supernatants were stored at −80°C. The remaining nuclear pellets were resuspended in nuclear lysis buffer (20 mmol/L HEPES, pH 7.9, 400 mmol/L NaCl, 1 mmol/L EDTA, 10% glycerol, and 1 mmol/L dithiothreitol, with protease inhibitors) and incubated at 4°C, with end-to-end rotation, for 2 hours. The nuclear supernatants were collected after centrifugation at 15,000 × g for 5 minutes at 4°C. Protein concentrations were determined using the Bradford method, and lysates were run on 4% to 12% Bis-Tris gradient gels. Membranes containing nuclear fractions were probed with rabbit anti-ATF4 antibody.

Transfection Using Magnetic Nanoparticles

Pure neuronal cultures at DIV 10, prepared as described above, were transfected with plasmid DNA via magnetofection using the Magnefect-Nano II system (Nanotherics). Briefly, 0.5, 1, or 2 μg of the following plasmid DNAs were complexed with 2.4 μL of Neuromag transfection reagent (Nanotherics) for 15 to 20 minutes in serum-free neurobasal...
media: pcDNA3.1Zeo+/BACE1 5′ UTR, pcDNA3.1Zeo+/BACE1-Δ5′ UTR (both generous gifts from Robert Vassar), pLenti CMV/TO Puro destination vector (backbone), pcDNA3.1Zeo+/wild-type mouse PERK, and ppcDNA3.1Zeo+/PERK-K618A. Complexes were added drop-wise to 24-well plates of pure neuronal cultures containing 500,000 cells per well, and plates were incubated for 30 minutes at 37°C with 5% CO2 over a magnet array oscillating at 3 Hz with 0.2-mm amplitude of displacement. At the end of magnetofection, plates were removed from the magnet array and were processed after 48 to 72 hours for either immunoblotting or cell staining, following routine methods.

Statistical Analysis

All data were analyzed with Prism software version 5.0 (GraphPad Software, San Diego, CA). All quantifications were expressed as means ± SEM. Paired results were analyzed by Mann-Whitney U test or unpaired t-test, and data with multiple categories were analyzed by one-way analysis of variance plus Newman-Keuls or Dunnett post hoc test, as indicated. P < 0.05 was considered significant for all statistical analyses performed.

Results

ARVs Induce Axonal Damage and UPR in Vivo

Several clinical studies have reported an inverse relationship between ARV regimens that have high CNS penetration effectiveness scores and neurocognitive outcomes. However, it has been difficult to separate CNS toxicity induced by HIV from that of ARV. We previously showed that ARVs administered to SIV-infected pigtail macaques lead to decreases in markers of synaptodendritic integrity, including synaptophysin, in the hippocampus and calmodulin kinase II in the frontal cortex, compared to SIV-infected, placebo-treated, or uninfected controls. In agreement with our previous studies in rats, animals displayed no overt signs of behavioral alterations or distress during the course of treatment. However, immunoblotting analysis revealed a statistically significant increase in BACE1 expression in the hippocampus in ritonavir- and saquinavir-treated mice compared with the vehicle-treated controls, complementing our findings in SIV+/ART macaques (Figure 2, A and B). ARV-treated mice also displayed significant increases in p-eIF2α expression, whereas saquinavir-treated mice had elevated levels of BiP in the hippocampus compared to ritonavir- and vehicle-treated animals. These data further support a role for ARV-associated UPR activation and BACE1 up-regulation in vivo independent of virus, we administered ritonavir or saquinavir i.v. to healthy adult mice for 2 weeks. Ritonavir and saquinavir doses were based on published pharmacokinetic ARV studies in rodents. In agreement with our previous studies in rats, animals displayed no overt signs of behavioral alterations or distress during the course of treatment. However, immunoblotting analysis revealed a statistically significant increase in BACE1 expression in the hippocampus in ritonavir- and saquinavir-treated mice compared with the vehicle-treated controls, complementing our findings in SIV+/ART macaques (Figure 2, A and B). ARV-treated mice also displayed significant increases in p-eIF2α expression, whereas saquinavir-treated mice had elevated levels of BiP in the hippocampus compared to ritonavir- and vehicle-treated animals. These data further support a role for ARV-associated UPR activation and BACE1 up-regulation in the CNS.

ARVs Activate the UPR and Up-Regulate BACE1 Expression

We and others have shown that ARVs are toxic to primary neurons. Although it is suggested that such toxicity is mediated by mitochondrial DNA damage and/or pro-oxidant properties of these drugs, additional mechanisms may also contribute to ARV-mediated neuronal damage and death. Given the established role of PIs inducing ER stress in numerous cell types, including macrophages and hepatocytes, we hypothesized that ARV drugs may be activating the UPR in neurons. Therefore, we examined canonical UPR markers in primary rat neuroglial cultures in
response to increasing concentrations of AZT, saquinavir, or ritonavir (Figure 3A). Drug dose ranges used in our in vitro studies were based on reported plasma and cerebrospinal fluid levels of ARVs.63,64 At 16 hours after treatment with ritonavir or saquinavir, neuroglial cultures displayed a fivefold increase in BiP expression and more than a fourfold increase in p-eIF2α (Figure 3, B and C). AZT treatment (25 μmol/L) resulted in a threefold increase in BiP expression but did not significantly alter p-eIF2α. As an added parameter to ensure the presence of ARV-mediated UPR activation, we treated primary neuroglial cultures with 10 μmol/L ritonavir, 1 μmol/L saquinavir, 25 μmol/L AZT, or 1 μmol/L thapsigargin (positive control) for 16 hours and assessed the levels of spliced X box binding protein (XBP)-1 relative to unspliced XBP-1 by TaqMan quantitative RT-PCR (Figure 3G). We observed nearly a twofold increase in spliced XBP-1 mRNA in ritonavir- and saquinavir-treated cultures, but no change in AZT-treated neurons, compared to vehicle controls.

Recent studies have shown that induction of ER stress in neurons could increase protein expression of the APP-cleaving enzyme, BACE1.36,65 Therefore, we sought to determine whether ARV-treated neuroglial cultures, which showed robust activation of ER stress, displayed changes in BACE1 expression. Immunoblotting analysis revealed that 16-hour treatment with AZT, saquinavir, or ritonavir resulted in fourfold to fivefold increases in BACE1 expression (Figure 3D). To rule out the possibility of species-specific effects, we also treated 21 DIV human fetal neurons with increasing doses of AZT, saquinavir, and ritonavir for 16 hours and found similar, albeit statistically insignificant, changes in both BACE1 and BiP expression in saquinavir- and ritonavir-treated cultures (Figure 3, H–K).

In addition to BACE1, APP may be cleaved by two additional secretases, γ-secretase and α-secretase, the latter of which cleaves APP to preclude Aβ production. To examine whether ARVs affect the expression of α- and/or γ-secretases, we also immunoblotted for presenilin-1 C-terminal fragment (γ-secretase) and mature ADAM metalloproteinase domain 10 (α-secretase) (Figure 3A). We found no significant changes in the expression of either of these enzymes regardless of drug treatment or dose (Figure 3, E and F). Taken together, these data indicate that antiretroviral drugs, and PIs in particular, induce UPR and BACE1 in neurons.
Ritonavir Increases BACE1 Processing of APP in Primary Neurons

Based on our observation that ARVs induce BACE1 expression in neuroglial cultures, we sought to determine whether this translated into functional changes in BACE1’s enzymatic activity by assessing APP processing. To that end, we treated CHO cells that are stably expressing human APP with 10 μmol/L ritonavir, as a representative PI, for 16 hours and measured the relative amount of Aβ species in the culture media using a commercially available human/rat Aβ40- or Aβ42-specific sandwich ELISA. Although ritonavir had no effect on secreted Aβ40 levels, Aβ42 levels were increased by nearly twofold (Figure 4A). In addition, pretreatment with a cell-permeable BACE1 inhibitor blocked ritonavir-mediated increases in Aβ42 in a dose-dependent manner. These findings differ from previous reports that changes in BACE1 activity affect the generation of all Aβ isoforms equally.66 To further understand how ritonavir may be affecting human APP processing, CHO culture media were measured using a commercially available human/rat Aβ40- or Aβ42-specific sandwich ELISA. Although ritonavir had no effect on secreted Aβ40 levels, Aβ42 levels were increased by nearly twofold (Figure 4A). In addition, pretreatment with a cell-permeable BACE1 inhibitor blocked ritonavir-mediated increases in Aβ42 in a dose-dependent manner. These findings differ from previous reports that changes in BACE1 activity affect the generation of all Aβ isoforms equally.66

Figure 2 ARV drugs induce the UPR and BACE1 in the CNS independent from immunodeficiency virus. Adult male C57BL/6 mice catheterized through the jugular vein were administered one the following: dimethyl sulfoxide (DMSO), 20 mg/kg ritonavir, or 25 mg/kg saquinavir twice daily for 14 days by continuous i.v. injection. A: Hippocampal brain lysates were immunoblotted, and actin was used as a loading control. B: Densitometric analysis revealed significant Increases in BACE1 and p-eIF2α levels in both ritonavir- and saquinavir-treated mice versus vehicle controls, whereas BiP was elevated only in saquinavir-treated mice. Data represent means ± SEM (B). n = 6 (DMSO-treated mice); n = 7 (ritonavir- and saquinavir-treated mice). *P < 0.05, ***P < 0.001 (one-way analysis of variance, Newman-Keuls post hoc).

BACE1 Inhibition Prevents MAP2 Loss in Ritonavir-Treated Neuroglial Cultures

Several lines of evidence suggest that BACE1 may act as a stress response protein in the CNS.36,67,68 Therefore, we sought to determine whether BACE1 has a protective or deleterious effect on neuronal survival after chronic exposure to ritonavir. Primary rat cortical cultures were pretreated with vehicle (DMSO) or 1 μmol/L BACE1 inhibitor for 30 minutes, followed by 10 μmol/L ritonavir for 48 hours. Cultures were labeled with propidium iodide to assess cellular viability and immunostained for MAP2 and synapsin. BACE1 inhibition had no effect on MAP2+ cell count, synapsin intensity, or propidium iodide uptake compared with untreated and vehicle controls (Figure 5A). Forty-eight hour ritonavir treatment resulted in a threefold decrease in MAP2+ cell count and nearly a threefold increase in propidium iodide uptake, but no change in synaptic integrity relative to MAP2 area in 10 DIV neuroglial cultures (Figure 5, A–D). More important, pretreatment with BACE1 inhibitor rescued MAP2+ cell loss in ritonavir-treated
neurons and decreased propidium iodide uptake (Figure 5D). Together, these results suggest BACE1 may contribute to PI-mediated neuronal damage and death.

Ritonavir Up-Regulates BACE1 Expression in a Translation-Dependent Manner

Vassar and colleagues first reported that chronic activation of the UPR results in translational up-regulation of BACE1. To determine whether ARVs increase BACE1 expression by altering mRNA levels, we first treated 21 DIV rat neuroglial cultures with ritonavir, saquinavir, AZT, or thapsigargin for 16 hours and harvested mRNA for quantitative RT-PCR. We observed no changes in BACE1 mRNA in any of the ARV-treated cultures compared to controls; however, there was a significant decrease in BACE1 mRNA in thapsigargin-treated neuroglial cultures (Figure 6A). To confirm that BACE1 regulation is occurring via a post-transcriptional mechanism, we pretreated neuroglial cultures with a translation inhibitor, anisomycin, or a transcription inhibitor, actinomycin D, for 12 hours, followed by 16 hours treatment with ritonavir, and harvested protein lysates for analysis by immunoblotting for BACE1. Inhibition of transcription with 5 μmol/L actinomycin D did not affect

**Figure 3** ARV drugs activate the UPR and up-regulate BACE1 expression in primary neurons. **A:** Twenty-one DIV rat neuroglial cultures treated with ARVs for 16 hours were immunoblotted. Actin was used as a loading control for all targets, except p-eIF2α, which was normalized to total eIF2α. Representative blots shown. B-F: Quantification of band intensities was normalized to actin and expressed as fold changes relative to vehicle. BIP, p-eIF2α, and BACE1 protein levels were significantly elevated compared to controls. A disintegrin and metalloprotease 10 (ADAM10) and presenilin-1 C-terminal fragment (PS-1 CTF) did not show any significant changes. **G:** Rat neurons were treated with 0.04% dimethyl sulfoxide (DMSO; Vehicle), 10 μmol/L ritonavir (Rit), 1 μmol/L saquinavir (Saq), 25 μmol/L zidovudine (AZT), or 1 μmol/L thapsigargin for 16 hours. Total mRNA was isolated, and expression levels of spliced X box binding protein (XBP)-1 mRNA were measured via TaqMan using the comparative Ct (ΔΔCt) method relative to unspliced XBP1. Quantitative RT-PCR assays were performed in triplicate. Both ritonavir and saquinavir induced significant increases in spliced XBP1. **H:** Twenty-one DIV human fetal neurons were treated as in **A,** and lysates were immunoblotted. One representative blot is shown. **I-K:** Quantification of band intensities was normalized to actin and expressed as fold changes relative to vehicle. All values represent means ± SEM (B–G, and I–K). n = 5 (B–F, vehicle: 0.04% DMSO); n = 4 (G); n = 3 (H). ∗P < 0.05, ∗∗P < 0.01, and ∗∗∗P < 0.001 (one-way analysis of variance, Dunnett post hoc). UT, untreated.
up-regulation of BACE1 protein levels by ritonavir; however, translation inhibition by 10 or 50 μmol/L anisomycin abrogated ritonavir-mediated BACE1 protein up-regulation (Figure 6, B and C), indicating that PIs are exerting their effects on neuronal BACE1 expression through either a translational or a post-translational mechanism.

ATF4 is a canonical target of translational control by p-eIF2α. Similar to BACE1, ATF4 mRNA contains multiple upstream open reading frames in its 5′ UTR, which act to suppress translation under basal conditions, but undergo scan-through after phosphorylation of eIF2α, leading to selective translational up-regulation during conditions of ER stress.36,65,69 ATF4 subsequently translocates to the nucleus to up-regulate the expression of genes involved in the UPR.

To confirm that translational control of BACE1 is occurring through its 5′ UTR, we transfected rat cortical neuroglial cultures with vectors containing full-length human BACE1 with the 5′ UTR (þ 5′ UTR) or without the 5′ UTR (− 5′ UTR). Deletion of the 5′ UTR resulted in a significant increase in overall BACE1 expression in transfected cells, as described previously36,70; however, overexpression of the -5′ UTR/BACE1 construct prevented ritonavir-mediated up-regulation of BACE1 in neurons (Figure 6, F and G). We also assessed whether a PI-mediated increase in BACE1 occurred via proteasome inhibition, based on previous studies on dividing cells.19,71

As seen in Figure 6H, compared to cultures treated with the proteasome inhibitor lactacystin, neither ritonavir nor saquinavir led to proteasome inhibition in neuroglial cultures after 16 hours, providing further evidence for the translational control of BACE1 in neurons exposed to these PIs.
To rule out the involvement of a post-translational mechanism in BACE1 up-regulation by ritonavir, we performed a pulse-chase experiment using 35S-metabolic radiolabeling to determine the half-life of BACE1 in ritonavir-treated neuroglial cultures compared to vehicle controls. After confirming immunoprecipitation of BACE1 from neurons (Figure 7A), we pulse-labeled 21 DIV neuroglial cultures for 30 minutes and then chased with neurobasal media with or without 10 μmol/L ritonavir for up to 24 hours (Figure 7B). Compared to controls, we observed no apparent difference in BACE1 half-life in ritonavir-treated neuroglial cultures over the course of 24 hours (Figure 7C). In both treatment conditions, we observed that BACE1 half-life was approximately 12 hours, similar to what has been reported previously. Collectively, these data strongly suggest that ritonavir-mediated up-regulation of BACE1 in neurons occurs through a 5' UTR-dependent translational mechanism.

PERK Is Necessary for ARV-Mediated BACE1 Up-Regulation in Neurons

Several reports have proposed a role for p-eIF2α in post-transcriptional regulation of BACE1 under conditions of oxidative and ER stress. However, eIF2α can be phosphorylated by one of four kinases (PERK, PKR, heme-regulated inhibitor kinase, and general control nonderepressible 2), depending on the type of stress impinging on cells. Given that PERK is the central kinase responsible for eIF2α phosphorylation under conditions of ER stress, we assessed whether PERK was involved in the stress response signaling in neuroglial cultures treated with ARVs. To that end, we generated neuroglial cultures from brains of tamoxifen-inducible cre recombinase conditional PERK knockout embryonic mice. At DIV 10, neuroglial cultures were treated with 100 nmol/L or 1 μmol/L 4-hydroxytamoxifen to excise the PERK gene. The estimated efficiency of PERK excision in neurons was approximately 50% to 60% after 96 hours of 4-hydroxytamoxifen treatment (Figure 8A). Wild-type and PERK-excised neurons were treated with 10 μmol/L ritonavir or 1 μmol/L saquinavir for 16 hours. As shown in Figure 8, B and C, PERK-excised neuroglial cultures treated with ritonavir or saquinavir displayed reduced p-eIF2α and BACE1 compared to wild-type cultures treated with the same drugs. These findings indicate that ARV-mediated BACE1 up-regulation in neuroglial cultures is dependent on UPR signaling through PERK.
Discussion

The unanticipated persistence of neurocognitive deficits in HIV+ patients is a growing public health concern as the average age of patients in the United States is expected to be >50 years by 2015. Mounting clinical and experimental evidence suggests ARV-mediated neurotoxicity; however, potential mechanisms remain elusive. Previously, we have reported ARV-mediated reactive oxygen species production in primary neurons, leading to activation of the endogenous antioxidant response. Our findings in this study provide a mechanistic link between ARV-mediated UPR in neurons and increased BACE1 in the CNS of mice and SIV-infected, ART-treated macaques.

We herein show, for the first time, that ARVs, PIs in particular, activate the UPR in primary rat neuroglial cultures as well as human fetal neurons. We also observed increased nuclear ATF4, indicating functional changes in the translational profile of neurons as a result. The specific mechanism by which PIs and NRTIs activate the UPR in neurons, albeit unclear, is likely multifactorial. In other cell types, PIs induce UPR via proteasome inhibition and...
AZT, saquinavir, and ritonavir increased BACE1 expression in both rodent and human neurons after 16 hours, in parallel to increased nuclear ATF4. This increase in BACE1 was at least partially mediated by PERK; however, potential contribution of other eIF2α kinases could be ruled out. Interestingly, PERK excision decreased p-eIF2α and BACE1 levels, albeit had no effect on BiP or APP, indicating divergent mechanisms at play in ARV-treated neurons. Although BiP is transcriptionally regulated during ER stress, the mechanism of APP up-regulation remains unclear and may be unrelated to UPR activation.

More important, neither presenilin-1 C-terminal fragment, the γ-secretase subunit, nor ADAM metalloproteinase domain 10 showed any changes in expression after ARV treatment. Notably, mature γ-secretase is tightly regulated at multiple levels; thus, further studies will be necessary to conclusively determine whether ARVs alter the frequency and/or efficiency of γ-secretase cleavage via regulation of other essential subunits or cellular modulators of γ-secretase, including the γ-secretase activating protein. However, our results strongly suggest ARV-mediated specific BACE1 up-regulation is because of its unique mRNA structure. Unlike presenilin-1 and ADAM10, BACE1 mRNA contains three upstream open reading frames in its long, GC-rich, 5′ UTR that suppress translation under normal cellular conditions and allow for more efficient BACE1 mRNA translation because of scan-through p-eIF2α to bypass inhibitory upstream open reading frames; this is an evolutionarily conserved mechanism among many species to control translation of stress-related genes. However, it remains unclear what protective, if any, role BACE1 might have in neurons undergoing UPR.

One potentially detrimental consequence of elevated BACE1 expression is increased APP processing to generate Aβ peptides. Giunta et al. was the first to show ARV-mediated Aβ production in Swedish mutant APP-overexpressing N2a cells. Our findings of rönitavir-mediated Aβ42 increases in human APP-expressing CHO cells did not extend to Aβ40 production, despite the detected ARV-mediated Aβ40 increases in rat neuroglial culture media. This divergence may result from potential differences in APP processing in CHO cells versus neurons, as selective Aβ40 degradation was shown in these cells. Alternatively, APP overexpression may precipitate ER stress and UPR. Nonetheless, PI-induced induction of BACE1 accompanies increased BACE1 activity, which may alter APP processing in PI-exposed neurons.

In a previous study, Lan et al. showed PIs, including ritonavir and saquinavir, modestly inhibited Aβ degradation in human macrophages, and decreased Aβ40 and BACE1 in human cortical neurons, whereas oral dosing with other PIs, nelfinavir or lopinavir/ritonavir, did not affect Aβ40 and BACE1 degradation. Alternatively, APP overexpression may precipitate ER stress and UPR. Nonetheless, PI-induced induction of BACE1 accompanies increased BACE1 activity, which may alter APP processing in PI-exposed neurons.
BACE1 up-regulation is PERK dependent. A–C: Floxed PERK mouse cortical neuronal cultures were generated by isolating neurons from brains of tamoxifen-inducible cre recombinase conditional PERK knockout mice at embryonic day 16.5 to 17.5, and treatment with 0.1 or 1 μmol/L 4-hydroxytamoxifen (4-OHT) at DIV 10 to excise PERK. At DIV 14, neuronal cultures were used for experiments (estimated PERK half-life is 13 hours). A: Efficiency of PERK excision was estimated to be approximately 50% to 60% via densitometric analysis. Wild-type and PERK-excised neuronal cultures were treated with 10 μmol/L ritonavir (Rit; B) or 1 μmol/L saquinavir (Saq; C) for 16 hours and lysates were immunoblotted. Representative images are shown. Fold changes are indicated below the blots, after normalization to actin. Compared to wild-type cultures, ARV-treated and PERK-excised neuroglial cultures showed repressed under basal conditions but is selectively up-regulated during ER and oxidative stress; thus, we hypothesized a protective role for BACE1 in cellular stress adaptation and/or survival. In fact, BACE1-mediated Aβ was protective against neuronal hyperactivity/excitotoxicity in vitro. However, our findings suggest that BACE1 may contribute to neuronal damage during ER stress over 48 hours, and raise questions regarding BACE1’s physiological functions in neurons, which remain largely elusive. BACE1 has been implicated in myelination via cleavage of neuregulin-1 and in maintenance of synaptic functioning. BACE1 is also up-regulated in neurons in response to a host of inflammatory mediators, including interferon-γ, tumor necrosis factor-α, and Aβ, and contains transcription factor binding sites for specificity protein 1 (Sp1), cAMP response element binding protein (CREB), and NF-κB, among others. These aspects of BACE1 need to be further investigated in future in vivo studies.

Based on our findings and previous reports, neuronal BACE1 up-regulation may be deleterious to cell survival during stress conditions. However, the cellular context of the specific stressor may be the critical determinant. As shown by Baleriola et al., Aβ42 can induce p-eIF2α accumulation and local translation of a subset of mRNAs, including ATF4, when applied to distal rat hippocampal neuronal axons. ATF4, typically considered neuroprotective in cellular adaptation to ER stress, was shown to propagate a neurodegenerative signaling cascade in Aβ42-treated neurons, leading to cell death. Thus, stress-induced genes, such as ATF4 and BACE1, that are translationally up-regulated in response to elevated p-eIF2α detected in HAND and AD CNS may be playing dual roles in neuronal viability, depending on stressor chronicity and localization. In addition, viral infection and inflammation are well-known UPR inducers. Furthermore, BACE1 expression is largely limited to neurons, whereas p-eIF2α is ubiquitously; changes in p-eIF2α might be masked in the cellular extracts containing all cell types. Overall, these factors may account for our in vivo findings showing p-eIF2α elevation in SIV+/placebo macaques, but not in ART-treated animals. These caveats should be examined in careful time-course experiments. Although our data suggest PI-mediated BACE1 up-regulation with concomitant functional activation, the implications of increased Aβ, an intriguing question beyond the scope of the current study, will be investigated in future studies.

Previous studies investigating amyloid pathology in brain specimens from ARV-mediated HIV+ subjects have
HIV PIs Up-Regulate BACE1

yielded conflicting results. Green et al were the first to show increased APP accumulation, primarily intraneuronal, and amyloid-associated pathology in ARV-treated, HIV+ individuals compared to ARV-naïve, HIV+ patients; these findings were confirmed by Xu and Ikezu, who observed intraneuronal Aβ immunoreactivity and Aβ plaques in ARV-treated, HIV encephalitic brains but only intraneuronal Aβ in ARV-untreated, HAD brains, with no evidence of extracellular deposition. One major caveat in these studies is the pathological effects associated with variables such as comorbidities, drug abuse, and potential ARV-mediated adverse effects. Thus, our studies in pigtailed macaque and rodent models of chronic ARV drug exposure were designed to tease out these confounders. In addition to increased hippocampal synaptopendritic damage we previously observed in ARV-treated SIV+ macaques, we herein report elevated neuronal APP and BACE1 in the same cohort. Elevated APP has been previously reported to be associated with increased neuronal injury; however, numerous reports indicate that this might be a neuroprotective response. In addition, roles for APP in synaptogenesis suggest that elevated APP may be an attempt at neuronal repair. Thus, our observation that altered APP levels among our treatment groups may be difficult to interpret. In addition, several caveats must be considered. First, the time to euthanasia was different between SIV+/placebo and SIV+/ART groups, as these cohorts were originally designed to address a distinct question regarding use of ARVs. Second, SIV+/ART macaques harbored persistent CNS viral DNA, although it was not clear if it was replication competent. Finally, this retrospective study did not include an SIV−/ART control group to more effectively account for ARV effects in the absence of virus. Future studies with SIV+/ART animals will be invaluable for isolating viral contribution to CNS damage from that of ARVs. More important, ARV concentrations in that model closely reflect dose ranges measured in human plasma and cerebrospinal fluid.

Similar to our observations in SIV+/ART macaques, adult mice in our rodent model of chronic ARV exposure displayed increased hippocampal BACE1 expression, indicating virus-independent, ARV-mediated CNS BACE1 up-regulation. However, unlike ART-treated macaques, ARV-treated mice had increased p-eIF2α and BiP expression, likely attributable to the use of different ARVs or species-specific differences in ARV effects in the hippocampus. These results build on our previous studies showing decreased MAP2 and synaptophysin in ARV-treated rat hippocampus, and taken together with our macaque data, suggest that ARVs, independent from potential viral effects, induce neuronal damage and UPR activation in the CNS and may contribute to the neurodegenerative processes associated with HAND.

Although CNS effects of ARVs are still poorly understood, clinical studies have begun to evaluate the effect of ARV regimens with high CNS penetration on HIV+ patients. Although numerous studies have shown ARVs’ benefit in improving cognition, others have found a significant association between ARV regimens with high CNS penetration effectiveness scores and worse neurocognitive performance. Inherent and significant confounders in these studies are the variation in brain-barrier permeability among different ARVs, CNS viral load persistence, and the resulting difficulty in distinguishing neurotoxicity of HIV from that of ARVs. ARV drugs vary in their physicochemical properties, particularly regarding their blood-brain barrier permeability; however, extensive evidence demonstrates increased CNS entry of small molecules and serum proteins after HIV-mediated disruption of blood-brain barrier endothelial cells and tight junctions. Thus, higher-than-anticipated concentrations of ARV drugs may be occurring in HIV+ patients with a compromised blood-brain barrier. Notably, pharmacokinetic analysis of ARVs in guinea pigs revealed significantly higher uptake of [3H]-ritonavir into brain parenchyma compared to cerebrospinal fluid, suggesting that clinical estimates of CNS penetration for PIs may be lower than actual CNS drug concentrations. In addition to uncertainties in ARV CNS penetration with currently prescribed regimens, efforts to enhance ARV delivery to the CNS via nanotherapeutics necessitate a careful examination of their potential neurotoxic effects.

Although ARV concentrations used herein were significantly higher than peak CNS levels measured in patients, their effects on neuroglial cultures are likely to occur over long-term exposures in vivo. We sought to model their chronic, low-dose effects in a more acute time frame with higher doses. Our results indicated that PIs, at nanomolar concentrations, robustly activated the UPR in primary cortical neurons, leading to PERK-dependent increases in p-eIF2α and BACE1 translational up-regulation. ARVs, either alone or in combination, also increased Aβ production in multiple cell types, which was dose-dependently blocked by a cell-permeable BACE1 inhibitor. Taken together with our previous reports of elevated BiP, p-eIF2α, and ATF6 in HAND tissue, these data strongly suggest a role for UPR in CNS HIV pathogenesis. These results bear striking similarities to observations made in AD, Parkinson disease, amyotrophic lateral sclerosis, and prion diseases in which UPR markers, including phospho-PERK, are elevated in multiple brain regions. Although transient eIF2α phosphorylation is protective during ER stress, sustained global protein synthesis suppression in the CNS may ultimately lead to impaired long-term potentiation and synaptic failure. However, promising therapeutics targeted at the UPR have already been reported. Several studies showed genetic and pharmacological suppression of PERK activity decreased p-eIF2α expression, improved synaptic plasticity, and rescued memory deficits in an AD mouse model, and alleviated prion-mediated pathology and cognitive deficits in prion protein-inoculated mice. In addition, BACE1 inhibitors, which are currently being aggressively pursued in...
climatic trials, may have therapeutic benefits in patients experiencing HAND.\(^8\) Future investigations will be necessary to determine the role of BACE1 in UPR. We also propose to investigate ARVs' potential effects on BACE1 redistribution to the trans-Golgi network and early endoplasmic compartments as a mechanism of increased APP processing, similar to what was shown by Tan et al.\(^13\) Such approaches, in a precision medicine setting, will aid the clinicians in determining the most efficient ART regimens with minimal adverse effects that can be reinforced with the most appropriate adjunctive therapeutic approaches, such as BACE1 inhibitors.

Acknowledgments


References


patterns of HIV p24 immunohistochemistry in different brain areas of AIDS patients. AIDS 2001, 15:571–575


100. Wang H, Sun J, Goldstein H: Human immunodeficiency virus type 1 infection increases the in vivo capacity of peripheral monocytes to cross the blood-brain barrier into the brain and the in vivo sensitivity of the blood-brain barrier to disruption by lipopolysaccharide. J Virol 2008, 82:7591—7600