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## The Brazilian Zika virus strain causes birth defects in experimental models

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### Author Contributions

F.R.G. derived the NPCs, neurons and neurospheres, performed the viral infections and cell analyses and analyzed the data. I.R.F. performed the viral infections of cells, processed and analyzed infected brain organoids, prepared manuscript figures and analyzed the data. F.B.R. derived the NPCs, performed immunostainings and analyses, prepared manuscript figures and analyzed the data. B.C.F. revised the manuscript and with S.R. generated the organoid cultures from iPSCs and assisted with the IF staining and experimental design. J.L.M.D. performed macroscopic and microscopic staining and analyses of the mice. K.O.P.G. performed the TEM experiments, RNA extraction and quantification and histopathological analyses. C.B. and N.S. performed RNA extraction and quantification and prepared figures. G.C.P. performed cell culture, analyzed the data and revised the manuscript. C.M.P., I.C., C.L.F., W.N.B. and C.R. performed cell death q PCR array and flow cytometry, D.G.A performed flow cytometry staining protocols and analyzed the data. C.M.P, I.C. and D.G.A. infected and observed the pregnant mice daily. C.M.P., C.L.F., I.C. and C.R. processed the mouse tissues for virus quantification and further analyses. D.P.F., A.T.G. and C.A.B. performed the CT scans and analyzed and plotted the data. C.T.B and E.A.M performed virus amplification, titration and gene expression analysis. A.A.S provided MR766 and YF-17 isolates and serological reagents. P.M.A.Z. designed the experiments and revised the manuscript. J.P.S.P., A.R.M. and P.C.B.B.B. designed the experiments, analyzed the data and wrote the manuscript.

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## Summary

Zika virus (ZIKV) is an arbovirus belonging to the genus *Flavivirus* (Family *Flaviviridae*) and was first described in 1947 in Uganda following blood analyses of sentinel *Rhesus* monkeys<sup>1</sup>. Until the 20<sup>th</sup> century, the African and Asian lineages of the virus did not cause meaningful infections in humans. However, in 2007, vectored by *Aedes aegypti* mosquitoes, ZIKV caused the first noteworthy epidemic on the island of Yap in Micronesia<sup>2</sup>. Patients experienced fever, skin rash, arthralgia and conjunctivitis<sup>2</sup>. From 2013 to 2015, the Asian lineage of the virus caused further massive outbreaks in New Caledonia and French Polynesia. In 2013, ZIKV reached Brazil, later spreading to other countries in South and Central America<sup>3</sup>. In Brazil, the virus has been linked to congenital malformations, including microcephaly and other severe neurological diseases, such as Guillain-Barré syndrome<sup>4,5</sup>. Despite clinical evidence, direct experimental proof showing that the Brazilian ZIKV (ZIKV<sup>BR</sup>) strain causes birth defects remains missing<sup>6</sup>. Here we demonstrate that the ZIKV<sup>BR</sup> infects fetuses, causing intra-uterine growth restriction (IUGR), including signs of microcephaly in mice. Moreover, the virus infects human cortical progenitor cells, leading to an increase in cell death. Finally, we observed that the infection of human brain organoids resulted in a reduction of proliferative zones and disrupted cortical layers. These results indicate that ZIKV<sup>BR</sup> crosses the placenta and causes microcephaly by targeting cortical progenitor cells, inducing cell death by apoptosis and autophagy, impairing neurodevelopment. Our data reinforce the growing body of evidence linking the ZIKV<sup>BR</sup> outbreak to the alarming number of cases of congenital brain malformations. Our model can be used to determine the efficiency of therapeutic approaches to counteracting the harmful impact of ZIKV<sup>BR</sup> in human neurodevelopment.

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The recent increase in microcephaly cases in Brazil has been associated with the outbreak of Zika virus (ZIKV)<sup>7</sup>, originated from an Asian-lineage strain that can be spread by *Aedes aegypti* mosquitoes<sup>8</sup>. The Brazilian ZIKV (ZIKV<sup>BR</sup>) has been detected in the placenta and amniotic fluid of two women with microcephalic fetuses<sup>9–11</sup> and in the blood of microcephalic newborns<sup>10,12</sup>, suggesting that the virus can cross the placental membrane. The virus has also been identified in the brains and retinas of microcephalic fetuses<sup>11–13</sup>. However, there is no direct evidence of the mechanism by which ZIKV<sup>BR</sup> causes brain malformations. A previous study revealed that the African ZIKV (ZIKV<sup>AF</sup>, strain MR-766) has the ability to infect human skin cells<sup>14</sup>. Neurons and astrocytes in the mouse brain could also be infected, inducing hippocampal degeneration and necrosis of pyriform cells seven days post-infection (p.i.)<sup>15</sup>. More recently, ZIKV<sup>AF</sup> was also shown to infect human pluripotent stem cell (hPSC)-derived neural progenitor cells (NPCs) *in vitro*, which induced apoptotic cell death<sup>16</sup>. These studies were performed using the MR-766 ZIKV<sup>AF</sup> strain isolated in Uganda in 1947, which shares 87–90% sequence similarity with the Polynesian and Brazilian isolates<sup>3,17</sup>. Nevertheless, because severe congenital malformations were not reported for African isolates, there is a need to study the association of ZIKV with

microcephaly and birth defects with isolates from affected localities, such as the ZIKV<sup>BR</sup> strain. Therefore, there is an urgent need to develop model systems to determine the relationship between infection with the ZIKV<sup>BR</sup> strain and birth defects.

We used the ZIKV<sup>BR</sup> isolated from a febrile case in the state of Paraíba, northeast of Brazil in 2015 (see Methods). To evaluate the causal relationship between ZIKV<sup>BR</sup> and birth defects, including brain malformation during development, we first used a murine experimental model in which SJL and C57BL/6 pregnant mice were infected with ZIKV<sup>BR</sup> evaluating newborns immediately after birth (Extended Data Fig. 1a). Notably, similar to ZIKV<sup>BR</sup>-infected human newborns<sup>18,19</sup>, pups born from the SJL ZIKV<sup>BR</sup>-infected pregnant females displayed clear evidence of whole-body growth delay or intra-uterine growth restriction (IUGR)<sup>20</sup> compared to pups born from the mock-infected controls (Fig. 1a, b). Using a qPCR assay, we confirmed the presence of ZIKV<sup>BR</sup> genomic RNA in several tissues of newborn animals observing significantly more viral RNA in the brain, confirming the neurotropic nature of the virus (Fig. 1c).

Microcephaly is perhaps the most dramatic birth defects reported in ZIKV<sup>BR</sup> infected newborns<sup>4,18,19</sup>. Mouse models often fail to reproduce the severely reduced brain size and pathological alterations found in human patients<sup>21,22</sup>, likely due to significant differences in gestation time and brain development between the two species. Nevertheless, upon close inspection of the ZIKV<sup>BR</sup>-infected mice brains, we noticed cortical malformations in the surviving animals, with reduced cell number and cortical layer thickness, signs associated to microcephaly in humans (Fig. 1d–f). At a cellular level, the neurons in the cortex, thalamus and hypothalamus displayed a “vacuolar nuclei” appearance. This morphology was characterized by central emptiness and marginalized chromatin pattern with nuclear debris, suggesting ongoing cellular death (Fig. 1d and Extended Data Fig. 2). In addition, we also noticed apparent ocular abnormalities, reminding of what was also observed in human patients<sup>23</sup> (Fig. 1g). Thus, SJL-infected pups presented congenital malformations compatible with ZIKV<sup>BR</sup>-infected human newborns. While the impact of ZIKV<sup>BR</sup> infection in the SJL mice was striking, no major body alterations were detected in pups from the infected C57BL/6 animals (Extended Data Fig. 1a–c). To exclude potential minor alterations in the C57BL/6 mice, computed tomography (CT) scans were performed to quantify the skull/body volumes. No significant changes in the pups born from the ZIKV<sup>BR</sup>-infected C57BL/6 mothers were observed compared to the controls (Extended Data Fig. 1d, e). A diagnostic qPCR assay of the pups from the ZIKV<sup>BR</sup>-infected animals was negative, suggesting that the virus did not cross the placenta in the C57BL/6 mouse strain (Extended Data Fig. 1f). To elucidate the type of cell death induced by the ZIKV<sup>BR</sup> in the SJL pup’s brain, we used a qPCR array to distinguish different molecular pathways involved. Our data clearly indicate that ZIKV<sup>BR</sup> infection misregulates genes intimately linked to autophagy and apoptosis, such as upregulation of *BMF*, *IRGM1*, *BCL2*, *HTT*, *CASP6* and *ABL1*. Conversely, *GADD45a*, *TNFRSF11B*, *FASL*, *ATG12*, *BCL2L11* and *DFFA* were highly suppressed (Fig. 1h and Extended Data Fig. 1g, h).

Next, we evaluated the impact of ZIKV<sup>BR</sup> infection in human neural cells derived from hPSCs to establish a correlation between ZIKV and impairment of neurogenesis (Extended Data Fig. 3a). We generated human cortical NPCs and neurons from healthy donors hPSCs.

First, we determined the expression levels of the *Tyro3*, *AXL* and *MertK* (TAM) receptors tyrosine kinases in NPCs and neurons. This is an important family of receptors used for cell invasion by the Dengue virus and ZIKV, and *AXL* has been recently proposed as a candidate receptor for ZIKV infection during neurogenesis<sup>14,24</sup>. Mock-infected NPCs expressed higher levels of *AXL* when compared to mock-infected neurons (Fig. 2a). However, no significant changes in expression levels were observed upon ZIKV infection in NPCs (Fig. 2b). We then investigated the impact of ZIKV<sup>BR</sup> and ZIKV<sup>AF</sup> infection in NPCs and neurons. After infection using viral multiplicity of infection (MOI) of 10, ZIKV<sup>BR</sup> particles were detected inside the NPCs and neurons at several stages of viral assembly using transmission electron microscopy (TEM) (Fig. 2c). Immunostaining performed on NPCs and neurons at both MOI of 10 and MOI of 1 revealed production of viral protein aggregates (Fig. 2d and Extended Data Fig. 3b, c). With an MOI of 10, the amount of ZIKV<sup>BR</sup> particles in the NPC and neuron culture supernatant increased over time, suggesting the efficient production of infectious viral particles (Fig. 2e, f). With an MOI of 1, NPCs but not neurons, continued to produce ZIKV<sup>BR</sup> RNA in the culture supernatant (Extended Data Fig. 3d, e). After 96 hours p.i. we observed a significant cell death in NPC cultures using fluorescence-activated cytometry (FAC). We quantified the cell death overtime in NPCs cultures and detected an increase in the number of apoptotic/necrotic cells both in the ZIKV<sup>BR</sup>- and ZIKV<sup>AF</sup>-treated cultures compared to the mock-infected cultures at MOI of 10 (Fig. 2g), but not at MOI of 1 during the same time-frame (Extended Data Fig. 3f). No difference was observed between the two ZIKV strains concerning cell death in neurons at MOI of 10 and MOI of 1 (Extended Data Fig. 3g, h).

Next, we challenged two tri-dimensional neural cell culture systems, neurospheres and cerebral organoids, with ZIKV<sup>BR</sup> and ZIKV<sup>AF</sup>. We generated neurospheres by growing human NPCs in suspension. While the mock-infected control neurospheres continued to grow over time, the ZIKV<sup>BR</sup>-infected neurospheres (MOI of 10) displayed evident morphological abnormalities with signs of cell death (Fig. 2h). The sizes of the neurospheres infected with ZIKV<sup>BR</sup> were significantly smaller than the mock-control and ZIKV<sup>AF</sup>-infected at 96 hours p.i. (Fig. 2i). A less dramatic effect is observed at MOI of 1, where both the ZIKV<sup>BR</sup> and ZIKV<sup>AF</sup> infection reduced the size of the neurospheres compared to mock-infected controls (Extended Data Fig. 4a, b). These observations were paired with increased ZIKV<sup>BR</sup> replication in these cultures at both MOI of 10 and 1 (Fig. 2j and Extended Data Fig. 4c). These results suggest that ZIKV<sup>BR</sup> induces cell death in human NPCs, impairing the growth and morphogenesis of healthy neurospheres (Extended Data Fig. 4d–f).

The majority of the described cases of ZIKV<sup>BR</sup>-infected newborns (95%) had malformations of cortical development<sup>4</sup>. Thus, we also used brain organoids generated from hPSCs and human embryonic stem cells, to evaluate the impact of ZIKV<sup>BR</sup> on human cortical development. In the following experiments, alongside the ZIKV<sup>AF</sup> and mock infection, we added the Yellow Fever virus (YFV), a slow replicating attenuated live-vaccine Flavivirus that has a low risk of causing neuropathy. Cerebral organoids are tri-dimensional, self-organized, stem cell-derived models that recapitulate the first trimester of human neurodevelopment, including the molecular and cellular architecture reminiscent of the fetal cortex<sup>25</sup>. Organoids show some degree of lamination and resembled the human neocortex in terms of the spatial relationships of the progenitor populations, defined here as proliferative

ventricular zone (VZ), post-migratory neurons in cortical plate (CP) and a marginal zone (MZ) (Fig. 3a–c). We infected organoids with the ZIKV<sup>BR</sup>, ZIKV<sup>AF</sup> and YFV using a MOI of 0.1 and compared to mock-infected organoids at 24 and 96 hours p.i. We quantified the percentage of different subtypes of cortical neurons, TBR1-positive or CTIP2-positive cells (deep-layer V/VI) in the CP, finding a significant reduction in their number and respective CP thickness in the ZIKV<sup>BR</sup>-infected organoids compared to the others. A significant reduction in TBR1-positive cells was observed in the ZIKV<sup>BR</sup> at 96 hours p.i., while CTIP2-positive cells were significantly reduced in both ZIKV<sup>BR</sup> and ZIKV<sup>AF</sup> at the same time-point (Fig. 3d, e and Extended Data Fig. 5a–f). Consistent with the reduced population of cortical neurons, we observed a significant decrease in Pax6-positive cells (dorsal forebrain progenitor cells) following ZIKV infection (Fig. 3f and Extended Data Fig. 5d). Dividing cells in the VZ, detected by the population of Ki67 and Sox2-positive cells, were only significantly reduced in the ZIKV<sup>BR</sup>-infected organoids (Fig. 3g–h and Extended Data Fig. 5d). As observed in our other *in vitro* models, the number of apoptotic cells (cleavage Caspase 3- and TUNEL-positive cells) was increased in organoids infected with ZIKV<sup>BR</sup>, possibly explaining the decrease in the NPC population (Fig. 3i and Extended Data Fig. 5g–h). The ZIKV<sup>AF</sup> was derived from a zoonotic agent associated with primates in Africa, whereas the ZIKV<sup>BR</sup> is an isolate from a lineage adapted to human-to-human transmission for the past 70 years. As an entry point to establishing the potential mechanistic adaptive differences between ZIKV<sup>BR</sup> and ZIKV<sup>AF</sup> towards human cells, we also generated brain organoids from non-human primate (chimpanzee) pluripotent stem cells. We infected these chimpanzee's cerebral organoids (MOI of 0.1) and measured the impact on cortical neurons at 24 and 96 hours p.i. The ZIKV<sup>BR</sup> failed to reduce the percentage of either TBR1- or CTIP2-positive cells in non-human primates (Extended Data Fig. 5i, j). Consistently, the kinetics of infection was different between the two ZIKV isolates. While the ZIKV<sup>BR</sup> did not replicate in the chimpanzee organoids, the ZIKV<sup>AF</sup> seemed well adapted to these cells (Extended Data Fig. 5k).

To evaluate the causal relationship between ZIKV congenital infection and birth defects, we used a murine experimental model, in which pregnant SJL and C57BL/6 mice were infected with ZIKV<sup>BR</sup>. Notably, the SJL strain was susceptible to viral infection of fetal tissues, causing severe IUGR that resembled the affected Brazilian newborns, including signs of microcephaly, such as cortical malformations<sup>4</sup>. We also showed that the ZIKV<sup>BR</sup> induced apoptosis and autophagy in the mouse neural tissue. This is in accordance with the literature, as it has been previously demonstrated that ZIKV induces and localizes in autophagic phagosomes<sup>14</sup>. To our knowledge, this is the first report showing a gene expression profile that correlates to cell death in the brains of microcephaly newborn ZIKV<sup>BR</sup>-infected mice, corroborating a causal relationship. It is unclear why the virus could not cross the placenta of C57BL/6 mice, but this result may be due to the robust anti-viral immune response of this mouse strain, which secretes significant levels of type I/II interferon, known to confer resistance to ZIKV<sup>14,26</sup>. These data suggest that genetic differences could explain in part why some ZIKV-infected pregnant women give birth to newborns without detectable congenital brain malformations<sup>27</sup>. Nonetheless, our murine model is a valuable tool for future pre-clinical studies, such as vaccine development. The presence of major cortical histological abnormalities in the pups with IUGR prompted us to use an hPSC model to

study the impact of ZIKV on neurodevelopment. ZIKV infects cells at different stages of brain maturation leading to alterations in the cortical layer organization. While this manuscript was under review, two other papers revealed the impact of previously established ZIKV strains on human organoids, confirming our observations with the ZIKV<sup>BR</sup> 28,29. Finally, our data using a non-human primate organoids suggested that the ZIKV<sup>BR</sup> might have experienced adaptive changes in human cells. In fact, it has been demonstrated that the Asian lineage of ZIKV is undergoing codon usage adaptation towards biases observed in highly expressed human genes<sup>30</sup>. Our findings support the hypothesis that microcephaly is a distinctive feature of recent ZIKV Asian-lineage virus, which originated in the Pacific and is now spreading in South and Central America.

## Methods

### Viral culture and amplification

A lyophilized ZIKV isolate from a clinical case in Brazil (ZIKV<sup>BR</sup>), gently provided by the Evandro Chagas Institute in Belém, Pará, was reconstituted in 0.5 mL of sterile DEPC water. The African-lineage MR-766 (ZIKV<sup>AF</sup>), a reference strain isolated in Uganda in 1947 and the Yellow Fever Vaccine strain (YFV-17D)<sup>31</sup>, both used here as controls, were gently provided by the Institute Pasteur in Dakar, Senegal. *Aedes albopictus* mosquito cells (C6/36 cells) were previously prepared to culture the three viruses. C6/36 cell culture was maintained using Leibovitz's L-15 medium supplemented with 10% fetal bovine serum (FBS) (Gibco<sup>TM</sup>), 1% non-essential amino acids (Gibco<sup>TM</sup>), 1% sodium pyruvate (Gibco<sup>TM</sup>), 1% penicillin/streptomycin (Gibco<sup>TM</sup>), 0.05% of Amphotericin B (Gibco<sup>TM</sup>) and kept at 27°C in the absence of CO<sub>2</sub>. After reaching an approximately 70% confluent monolayer, 50µL of each viral samples were inoculated into C6/36 with an hour of adsorption, with gentle shaking every 10 minutes to allow the homogeneous adsorption of the viruses. At the end of the adsorption period, 5mL of the culture media were added, plus 2% FBS, 1% non-essential amino acids and 1% sodium pyruvate. The cultures were then incubated under the same adsorption conditions. In the first subculture (T1), the infected cells were less confluent compared to the control cells but had few noticeable morphological changes. On the fourth day after infection, the second subculture (T2) was made blindly by transferring 500µL of the T1 supernatant, followed by the third subculture, which was collected on the eighth day after infection (T3). Pronounced cytopathic effects were perceived beginning at T2. The supernatants were harvested, titrated and T3 was used for the experimental inoculation.

### Virus titration

Titration (in PFU/mL) of each C6/36 subculture was obtained by plaque assay to determine the amount of infectious viral particles (PFU). The virus titration was performed in porcine kidney epithelial (PS) cells and in L15 medium with 5% FBS. Briefly, the virus titration was done using 200 µL of L15 medium (5% FBS, 1% penicillin-streptomycin, and 1% glutamine) in a 24-well plate. Then, a serial dilution of each virus stock from ZIKV<sup>BR</sup>, ZIKV<sup>AF</sup> and YFV-17D in L15 medium was performed, from 10<sup>-1</sup> to 10<sup>-11</sup>. Then, 200 µl of each dilution was added in each 24-well plate. After this, 1×10<sup>6</sup> PS cells were seeded in the each 24-well plate for at least 3 hours at 37°C to allow virus adsorption and PS cells adherence. Later, each well was overlaid with complete carboxymethyl cellulose (CMC)

medium (0.6% in L15 supplemented with 3% FBS). After 5 days of incubation at 37°C, the plaque visualization was made using blue black staining solution. The most appropriate viral dilution was estimated to determine the amount of infected cells visible (PFU per mL). For ZIKV<sup>BR</sup>, the first C6/36 subculture had a titer of  $6 \times 10^8$ . The following subcultures had, respectively, titers of  $7.5 \times 10^6$  (T2) and  $4 \times 10^{12}$  (T3). All the subculture aliquots were stored in cryovials and maintained in liquid nitrogen or were distributed to the ZIKV São Paulo-task force.

### ***In vivo* infection**

Pregnant mice, 6–8 weeks of age, C57BL/6 or SJL (JAX), were infected intra-venously with 200 µL of ZIKV<sup>BR</sup>-infected C6/36 cell supernatant containing  $10^3$ ,  $4 \times 10^{10}$  or  $1 \times 10^{12}$  PFUs/mL of virus on day 10–13 of gestation. The animals were observed daily. All the experiments were performed with the approval of the Institute of Biomedical Sciences Ethics Committee protocol number 05/2016.

### **Real-time PCR**

RNA was extracted from each sample (cells, supernatant of cell culture or mouse tissue) using the QIAamp UltraSens Virus Kit (Qiagen) or TRIzol® reagent (Invitrogen). All RNA pellets were resuspended in 30 µL of RNase-free distilled water, quantified using a NanoDrop spectrophotometer (NanoDrop Technologies) and stored at –80°C. The set of primers/probes specific for ZIKV were synthesized by Sigma Life Science, with 5- FAM as the reporter dye for the probe. The set of primers/probes ZIKV 835, ZIKV 911c and ZIKV 860-FAM were previously described<sup>2</sup>. The real-time reaction was performed with 10 µL of each sample and 10 µL of the AgPath-IDTM One-Step RT-PCR reagents (Applied Biosystems). The amplification was done in an Applied Biosystems 7500 real-time PCR system, and involved activation at 45°C for 15 min, 95°C for 15 min followed by 40 amplification cycles of 95°C for 15 s, 60°C for 15 s, and 72°C for 30 s. The real-time data were analyzed using SDS software from Applied Biosystems. For the detection and quantification of viral RNA, the real time PCR of each sample was compared with threshold cycle (CT) value with a ZIKV plasmid standard curve, which was obtained carrying out serial dilutions of a clone of the envelope gene of an isolate from the 2007 Yap Island outbreak (provided by the Institute Pasteur in Dakar, Senegal). For the detection and quantification of YFV RNA, a YFV-specific real time assay was applied. The fold changes of gene expression were calculated in comparison to the values for the YFV controls<sup>31</sup>. The positive PCR control CT value was used to normalize gene expression and determine fold changes along the 96 hours p.i. The RPLA27 gene (60S ribosomal protein L27) was used as endogenous control for the PCR reactions<sup>30</sup>. For the TAM receptors detection, NPCs or neurons from two donors were infected or not with ZIKV<sup>BR</sup>, at 10 MOI for 48 hours and were submitted to standard TRIzol (Invitrogen®) protocol for RNA extraction. Total RNA was quantified using a NanoDrop spectrophotometer and 2 µg was used for further cDNA synthesis using the superscript III reverse-transcription polymerase (Invitrogen®) according to manufacturer's protocol. qPCR was performed using Taqman® probes (Supplementary Table 1) and the QuantStudio® 12K Flex Real-Time PCR System (Applied Biosystems). For normalization β-actin was used as housekeeping gene.

### Cell death pathway analysis

One microgram of total RNA from brains of 4 pups, 2 pooled mock and 2 pooled ZIKV<sup>BR</sup> infected from SJL mothers were submitted to gene expression analysis for cell death target genes using the RT<sup>2</sup> Profiler™ PCR Array Mouse Cell Death PathwayFinder (Cat# PAMM-212ZA-Qiagen®) according to manufacturer's protocols. qPCR was performed in the QuantStudio® 12K Flex Real-Time PCR System (Applied Biosystems). To evaluate gene expression, we established a fold change threshold of at least 2-fold up- or downregulation compared to mock infected samples. Statistical analysis was performed using the RT2 profiler RT-PCR array data analysis software V 3.5.

### NPCs, neurons, neurospheres and organoids

We used 3 human and 2 chimpanzee iPSC clones that were previously characterized in the Beltrão-Braga and in the Muotri lab<sup>32-35</sup> plus H9 human embryonic stem cells (hESC) for all the experiments using pluripotent stem cells. All the cell lines tested negative for mycoplasma contamination. Briefly, high passages of iPSC/hESC colonies on feeder-free plates were maintained for 5 days with mTSeR media (Stem Cell Technologies). On the fifth day, the medium was changed to N2 media (DMEM/F12 medium supplemented with 1X N2 supplement (Invitrogen) and the dual SMAD inhibitors, 1µM dorsomorphin (Tocris) and 1 µM SB431542 (Stemgent), for 48 hours. Further, the colonies were detached from the plate and cultured in suspension as Embryoid Bodies (EBs) for 5 days at 90 rpm in N2 media with the dual SMAD inhibitors. The EBs were plated on matrigel-coated plates with NBF media composed of the following: DMEM/F12 media supplemented with 0.5X N2, 0.5X B27 supplement (Gibco™), 20 ng/mL of FGF2 and 1% penicillin/streptomycin. The emerged rosettes containing the NPCs were manually picked, dissociated and plated in a double-coated plate with poly-ornithine (10 µg/mL, Sigma-Aldrich) and Laminin (2.5µg/mL, Gibco™). The NPC population was expanded using NBF media. The neuronal differentiation induction protocol consisted of treating the confluent NPC plate with 10 µM ROCK inhibitor for 48 hours (Y-27632, Calbiochem) in the absence of FGF in the media, with regular media changes every 3 or 4 days. Neurons were considered completely differentiated and ready to experiments after 28 days. To produce neurospheres, NPC were scrapped from the plates and submitted to continuous shaking for 5–7 days at 90 rpm in NBF media. Cerebral organoids were generated as previously described<sup>25, 36</sup>. All experiments were performed with the approval of the Institute of Biomedical Sciences Ethics Committee protocol number 1001.

### *In vitro* infection

NPCs, neurons, neurospheres and organoids were infected with ZIKV<sup>BR</sup>, ZIKV<sup>AF</sup>, YFV and mock (culture supernatant from uninfected C6/36 cells). NPCs were seeded in plates in 24 well plates and after 24 hours viral samples were diluted to the desired MOI (0.1; 1 or 10) and added to the cells. For viral adsorption, cells in monolayer were incubated for 1 hour at 4°C with gentle agitation every 10 min. Next, the inoculum was removed and cells were washed once with PBS (USB Corporation). Culture medium was added to each well, and cells were incubated at 37°C and 5% CO<sub>2</sub> for the duration of the experiment. For neurospheres, NPCs were kept in constant shaking. For neuronal infection, NPCs were



previously differentiated for 28 days and then neurons were infected with the desired MOI. For organoids, the number of cells available for infection was estimated to be  $2.5 \times 10^4$  cells, as calculated by dividing the average surface area of a typical organoid by the average area of a typical cell (*i.e.*, a fibroblast). This calculus was used to stipulate the desired 0.1 MOI. For mock controls, the same volume of supernatant was added to each experiment, and the same procedures were followed.

### Immunofluorescence

Cells were fixed using paraformaldehyde, 4% in PBS, for 15 minutes at room temperature. After washing, the cells were permeabilized with 0.1% Triton X-100 (Promega) diluted in PBS for 15 minutes. After blocking with 2% of BSA (Sigma-Aldrich) for 4 hours, primary antibodies directed against the following were added: anti-ZIKV (polyclonal mouse, Institute Pasteur in Dakar, 1:80), anti-Flavivirus D1-4G2-4-15 (polyclonal mouse, Millipore, 1:100), 1:50, anti-MAP2 (chicken, Abcam ab5392, 1:200), anti-cleaved caspase-3 (rabbit, Cell Signaling #9661, 1:400), anti-Sox2 (mouse, Abcam ab97959), anti-GFAP (rabbit, Abcam, 1:500) and anti-Mushashi1 (rabbit, Abcam, 1:1000) (Supplementary Table 2). The cells were incubated overnight at 4°C. Secondary antibodies were added for a one-hour incubation at room temperature, being the following ones: anti-mouse Alexa Fluor 488, anti-chicken Alexa Fluor 647, anti-rat Alexa Fluor 555 and anti-rabbit Alexa Fluor 555 (Invitrogen). The nuclei were stained with DAPI (Invitrogen, 1:10,000) diluted in a PBS 1x solution for 5 minutes and mounted with DPX (Sigma). Images were acquired with Nikon Eclipse 80i. Analysis of data was performed using software NIS Elements 3.22 (Tokyo, Japan).

### Cerebral organoids analyses

Human and Chimpanzee organoids were infected with MOI of 0.1 and analyzed after 24 and 96 hours p.i. Organoids were cryosectioned at 20  $\mu$ m. Immunofluorescence was performed after blocking sections in a solution with 0.1% Triton and 3% BSA (Gemini) for 1 hour at room temperature. The primary antibodies were diluted in a solution with 0.1% Triton and 3% BSA, and the sections were incubated with following antibodies: anti-ZIKV, anti-flavivirus, anti-MAP2, anti-cleaved-caspase-3 and anti-Sox2, all mentioned above, and anti-PAX6 (rabbit, Covance PRB-278P, 1:100), anti-TBR1 (rabbit, Abcam ab31940, 1:300), CTIP2 (rat, Abcam ab18465, 1:100) and Ki67 (rabbit, Abcam ab15580, 1:100). The sections were blocked with 0.1% Triton (Sigma-Aldrich) and 3% BSA for 30 min at room temperature and the secondary antibodies previously diluted, the same mentioned above, were added. The nuclei were stained with DAPI, as mentioned above and slides were mounted with DPX (Sigma-Aldrich).

### Transmission Electron Microscopy (TEM)

Cell pellet was fixed using a 3% glutaraldehyde solution (Merck) at 4°C for 2 hours, rinsed in 3 changes of PBS for 1 hour, and incubated for 16 hours at 4°C. The next day, post-fixation was performed with 1% of osmium tetroxide for 30 minutes at room temperature. Dehydration was carried out gradually with a series of ethanol concentrations: 70%, 95% and 100%. Sample was taken through two changes of propylene oxide and placed at a 1:1 ratio with embedding medium for 1 hour in a rotary mixer followed by 100% embedding

medium at room temperature for 24 hours. Fresh embedding medium was placed overnight at 37°C and polymerized in oven to 24 hours. Ultrathin sections were cut and stained with uranyl acetate and lead citrate. The cells were visualized with a transmission electron microscope (JEOL, JEM 1011, Peabody, MA, USA). All experimental analyses were performed blinded to the treatment.

### Flow Cytometric Analysis (FAC)

The cells were infected under a MOI of 10 and 1, prepared using supernatants from infected C6/36, and equal volume of mock. Cellular infection occurred at 4°C for 1 hour with cell homogenization every 10 minutes. After that, the cells were washed once and then maintained at 37°C in CO<sub>2</sub> incubators with medium, as described before. After 24, 48, 72 and 96 hours p.i. the cells were harvested and then submitted to a staining protocol for annexin V and propidium iodide (PI) (BD Biosciences®). The cells were washed twice with PBS and were harvested with 200 µL of trypsin 0.25 % (LGC®) for 10 minutes at 37°C. Next, the cell suspensions were washed in DMEM with 10% of FBS and centrifuged for 5 minutes at 450 g and 4°C. The cells were then resuspended in 20 µL of annexin V binding buffer in 96-well round bottom plates and with 1 µL of FITC-annexin V + 1 µL of PI and then incubated at room temperature for 15 minutes protected from light. After incubation period, the samples were added to 80 µL of binding buffer and acquired in the BD FACS Accuri C6 (BD Biosciences®) flow cytometer.

### Computed Tomography (CT)

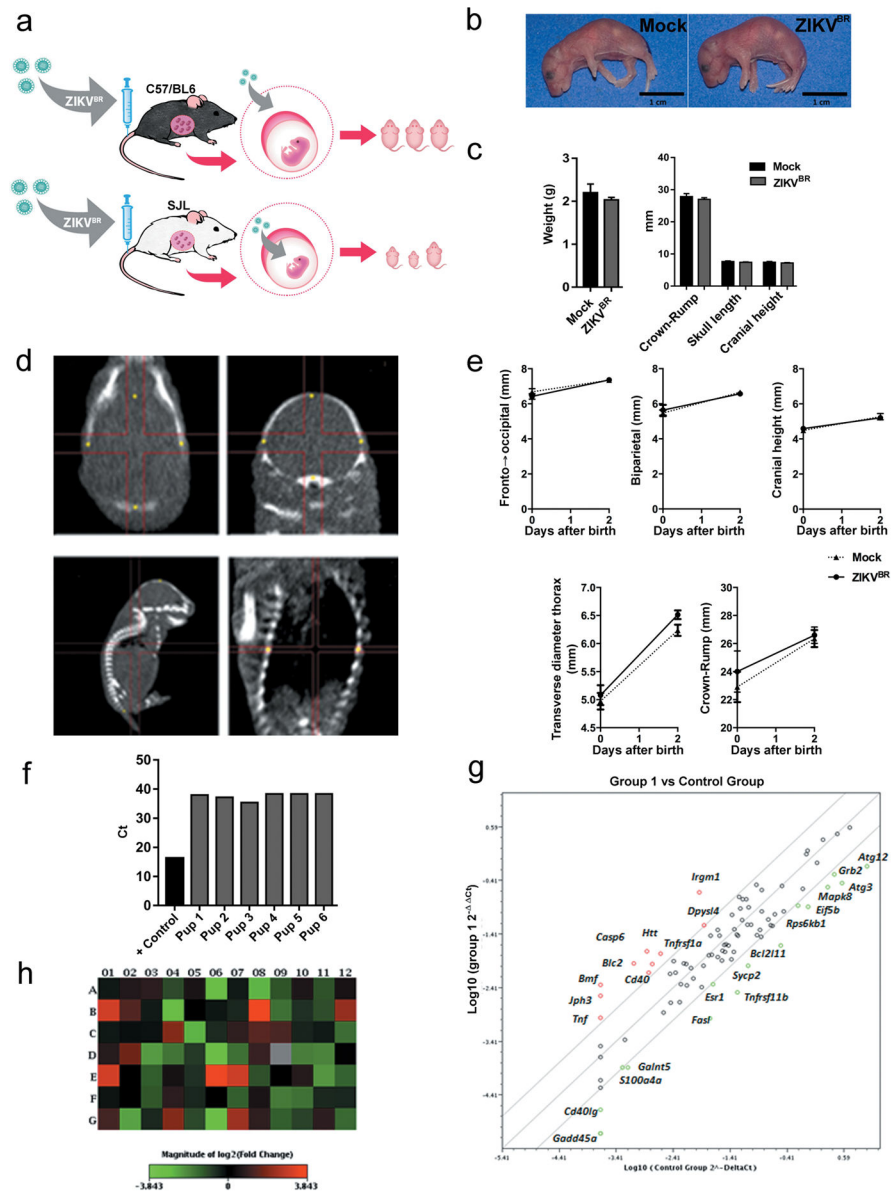
Mice were properly anesthetized with isoflurane and immobilized on their right side on the bed with a piece of gauze and positioned with the whole body in the field of view (FOV). CT images were acquired using small animal imaging equipment (Triumph™ Trimodality Gamma Medica Ideas) with 45 kVp, 0.4 mA and 2.13 minutes of X-Ray exposition (512 projections over 360 degrees). The images were reconstructed using the FBP (Filtered Back Projection) algorithm, a matrix of 512 X 512, smoothing filter and a pixel size of 92–117 µm (according to the animal size). Experienced evaluators, who were blinded to the animal group assignments, performed images analyses using the AMIDE 1.0.4, General Public License software. Fiducial marks were added to measure the distance between points considering, visually, the larger axis of the brain in the sagittal plane for posterior-anterior and superior-inferior distances and the coronal plane for the lateral right-left. For measuring the whole body length, the distance between the superior point of the brain to the first vertebra of the tail was used. The thorax measure was made from the 3<sup>rd</sup> rib (right-left) in the coronal plane to the spinal cord level. The results are presented in mm.

### Histologic processing

Tissue histology was performed using a dehydrating protocol with two alcohol baths of 95% (the first for 1 hour and 15 minutes and the second for 30 minutes), three absolute alcohol baths (1 hour for the first followed by 3 hours for the second and 2 hours for the third), followed by clarification with three baths of xylene (the first for 30 minutes and the next two for 1 hour each). Finally, the material was added to three paraffin baths (the first for 30 minutes and the second two for one hour each). Then the material was immersed in paraffin and cut with a microtome into a thickness of 5 mm. The deparaffinization protocol consisted

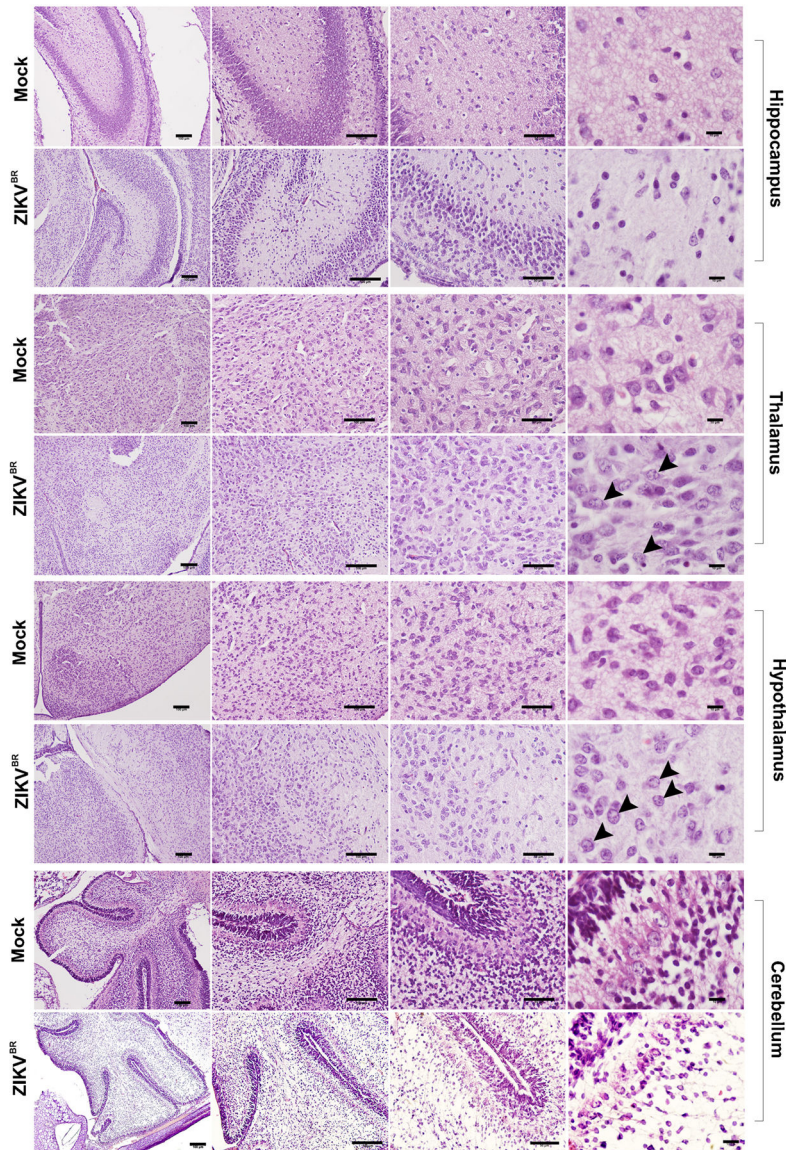
of three xylol baths heated in an oven for 30 minutes each, two baths of absolute alcohol for 2 minutes each, two baths of 95% alcohol lasting 2 minutes each, an alcohol in water bath (85%) for 2 minutes each and the last bath in 70% alcohol for 2 minutes. The hematoxylin-eosin staining protocol began with 2 quick baths in running water, followed by a 2-minute bath in distilled water, a 2-minute bath in hematoxylin, a 5-minute bath in running water, a 1-minute in Eosin in a fast flowing water bath, two baths of 95% ethanol for 2 minutes each, two baths of absolute ethanol for 2 minutes each, ending with three baths in xylene for two minutes each. Slides were mounted using Permount (Sigma-Aldrich) and analyzed on multiple coronal slices in glass slides using light microscopy (Olympus BX40, ZEISS KS400) on a genotype-blinded fashion.

## Extended Data

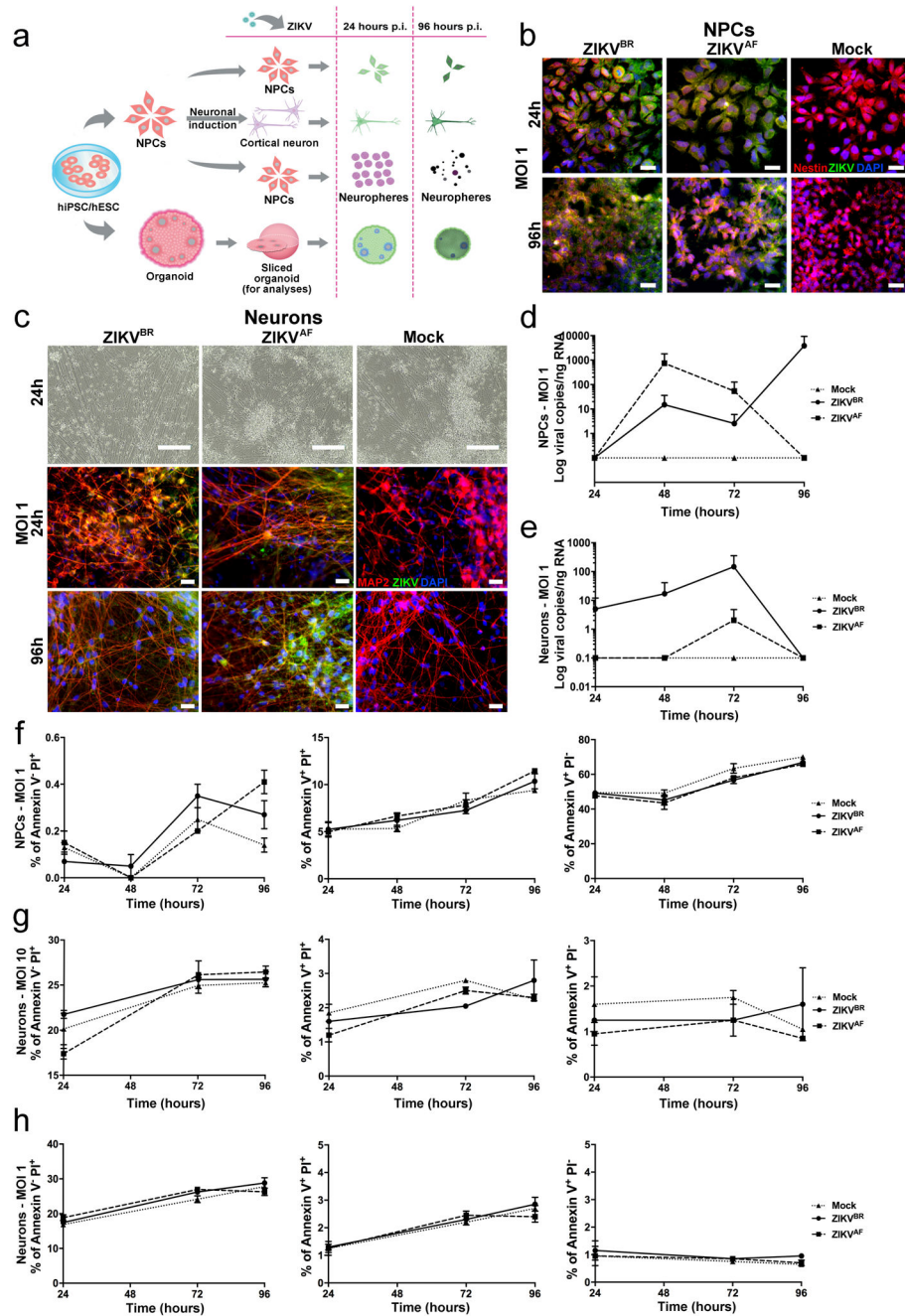


**Extended Data Figure 1. Impact of ZIKV<sup>BR</sup> infection in the C57BL/6 and SJL mice**  
**a**, Scheme for infecting mice and the follow up analyses. Pregnant females at approximately day 10–13 of gestation were challenged with  $4 \times 10^{10}$  PFUs of ZIKV<sup>BR</sup> via an intra-venous route. Their pups were analyzed immediately after birth for signs of malformation. **b**, A representative pup from mock-infected and the ZIKV<sup>BR</sup>-infected C57BL/6 mice. Scale bar = 1 cm. **c**, C57BL/6 pups born with no gross morphological changes or size differences compared to mock controls ( $n = 21$  pups from three separate litters, error bars, s.e.m,  $t$ -test). Scale bar = 1 cm. **d**, **e**, CT analysis confirmed lack of anatomical alterations ( $n = 21$  pups from three separate litters, error bars, s.e.m,  $t$ -test). **f**, ZIKV<sup>BR</sup> RNA was not detected in the brains of six C57BL/6 pups. **g**, Cell death pathway signature revealed by qPCR gene

expression in the brains of the ZIKV<sup>BR</sup>-infected SJL pups ( $n = 2$  technical replicates of pooled RNA from two pups each group; threshold = two-fold). **h**, Heatmap representation of misregulated genes in “g”.



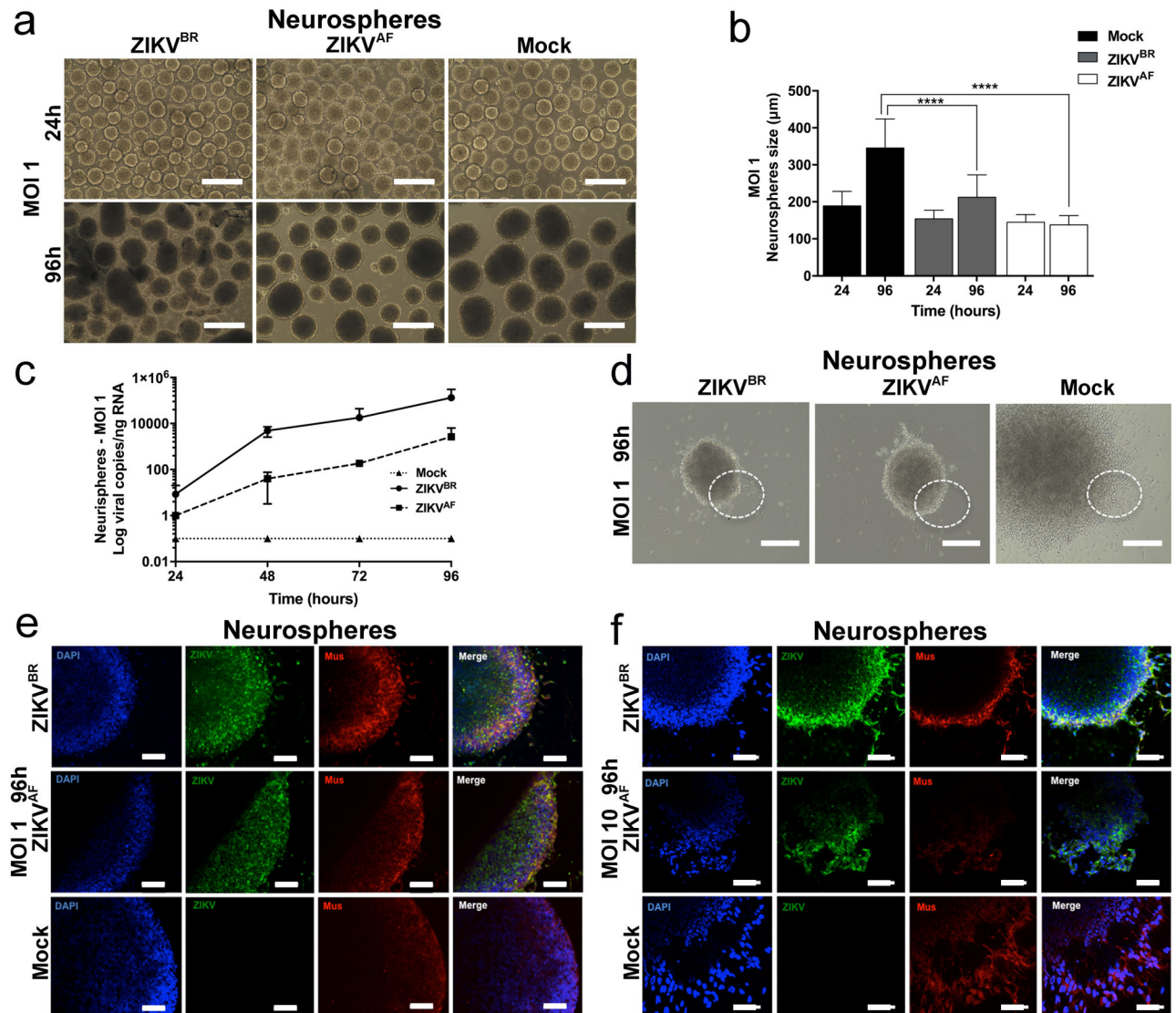
**Extended Data Figure 2. Histopathological analysis of brains from ZIKV<sup>BR</sup>-infected SJL pups**  
Morphological aspect of hippocampus, thalamus, hypothalamus and cerebellum from brains of pups born from mothers infected with the ZIKV<sup>BR</sup>. Arrowheads indicate intranuclear vacuoles and “empty” nuclei aspect with chromatin margination observed in thalamus and hypothalamus. Scale bar from left to right = 100 µm, 100 µm, 50 µm and 10 µm.



### Extended Data Figure 3. Impact of ZIKV infection in human NPCs and neurons

**a**, Scheme of the *in vitro* experiments using hPSCs. The cells were differentiated into NPCs, neurons, neurospheres and cerebral organoids to test the impact of ZIKV<sup>BR</sup> over time. **b**, Infection of NPCs with the ZIKV<sup>BR</sup> and ZIKV<sup>AF</sup> (MOI = 1) at 24 and 96 hours p.i. Scale bar = 25  $\mu$ m. **c**, Aspects of iPSC-derived human neurons after ZIKV infection (MOI = 1) at 24 and 96 hours p.i. Scale bar = 200  $\mu$ m (Bright field) and scale bar = 25  $\mu$ m (immunofluorescence). **d**, Viral replication dynamics in human NPCs over time (MOI = 1) ( $n = 2$  technical replicates from two different donors; error bars, s.e.m). **e**, Viral replication

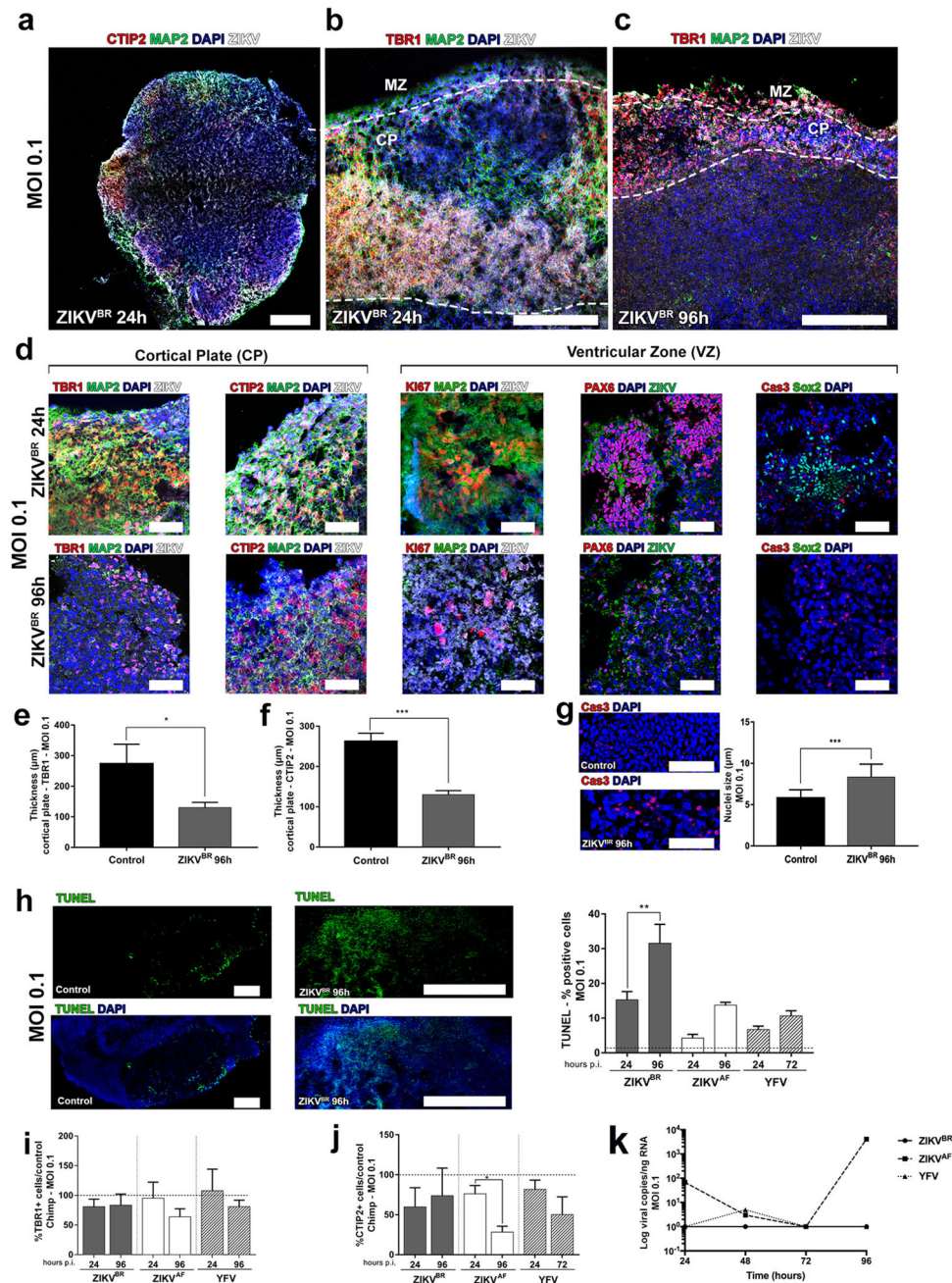
dynamics in human neurons over time (MOI = 1) ( $n = 2$  technical replicates from two different donors; error bars, s.e.m). **f**, Dynamics of NPCs toxicity over time after ZIKV infection (MOI = 1), indicating no significant differences among the different viruses ( $n = 2$  technical replicates from two different donors; error bars, s.e.m). **g, h**, Viral replication dynamics of ZIKV in human neurons over time at MOI = 10 and MOI = 1, respectively ( $n = 2$  technical replicates from two different donors; error bars, s.e.m; one-way ANOVA).



#### Extended Data Figure 4. Impact of ZIKV infection in human neurospheres

**a**, Representative bright-field images of ZIKV infection (MOI = 1) at 24 and 96 hours p.i. Scale bar = 400  $\mu\text{m}$ . **b**, Alterations in neurospheres diameter over time (MOI = 1) ( $n = 22$  neurospheres from two different donors for each time-point in each condition; unpaired  $t$ -test, \*\*\*\* $P < 0.0001$ ). **c**, ZIKV replication dynamics in neurospheres (MOI = 1) ( $n = 3$  technical replicates from two different donors). **d**, Representative bright-field images of ZIKV infection (MOI = 1) at 96 hours p.i. Scale bar = 400  $\mu\text{m}$  and 1000  $\mu\text{m}$  (Mock). The

dotted circle describes the neurospheres borders indicating the immunostained regions in **e** and **f**. **e**, Immunostaining of neurospheres infected with ZIKV at 96 hours p.i. at MOI = 1 (left panel) or MOI = 10 (right panel), revealing a qualitative reduction of proliferative cell migration from Mushashi (Mus)-positive cells. Scale bar = 50  $\mu$ m.



### Extended Data Figure 5. Human and chimp cerebral organoids infected with ZIKV

**a**, Representative image of an entire cross-section of a cerebral human organoid infected with the ZIKV<sup>BR</sup> (MOI = 0.1, 24 hours p.i.). Scale bar = 200  $\mu$ m. **b**, Detail of the surface of a human organoid infected with the ZIKV<sup>BR</sup> at 24 hours p.i. (MOI = 0.1). Marginal zone



(MZ) and cortical plate (CP) delineated by dotted white lines. Scale bar = 200  $\mu\text{m}$ . **c**, Detail of the surface of a human organoid infected with the ZIKV<sup>BR</sup> at 96 hours p.i. (MOI = 0.1). Notice the significant tissue damage and reduction in the CP related to 24 hours p.i. Scale bar = 200  $\mu\text{m}$ . **d**, A representative characterization of CP and ventricular zone (VZ) in human organoid infected with the ZIKV<sup>BR</sup> at 24 and 96 hours p.i. (MOI = 0.1). Scale bar = 50  $\mu\text{m}$ . **e**, Reduction in the cortical thickness measured by the extension of TBR1-positive layer of cells in human organoids at 96 hours p.i. (MOI = 0.1). ( $n = 3$  replicates from three human cell lines; error bars, s.e.m;  $t$ -test,  $*P = 0.0203$ ). **f**, Reduction in the cortical thickness measured by the extension of CTIP2-positive cells layer in human organoids at 96 hours p.i. (MOI = 0.1). ( $n = 3$  replicates from three human cell lines; error bars, s.e.m;  $t$ -test,  $***P = 0.001$ ). **g**, Nuclear size (diameter) of cleaved caspase-3 positive apoptotic cells in human organoids at 96 hours p.i. (MOI = 0.1). ( $n = 10$  organoids/slides from three human cell lines; error bars, s.e.m;  $t$ -test,  $***P = 0.0004$ ). Scale bar = 50  $\mu\text{m}$ . **h**, Percentage of TUNEL-positive cells in relation to controls (dotted line) at 24 and 96 hours p.i. (MOI = 0.1). ( $n = 10$  organoids/slides from three human cell lines; error bars, s.e.m; ANOVA,  $**P = 0.0042$ ). **i**, Percentage of TBR1-positive cells in non-primate organoids (Chimp) in relation to controls (dotted line) at 24 and 96 hours p.i. (MOI = 0.1) ( $n = 3$  organoids from two donors, error bars, s.e.m; ANOVA). **j**, Percentage of CTIP2-positive cells in non-primate organoids (Chimp) in relation to controls (dotted line) at 24 and 96 hours p.i. (MOI = 0.1) ( $n = 3$  organoids from two donors; error bars, s.e.m; ANOVA). **k**, Viral replication dynamics in chimpanzee organoids over time (MOI = 0.1) ( $n = 3$  replicates from two donors; error bars, s.e.m; ANOVA).

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

## Acknowledgments

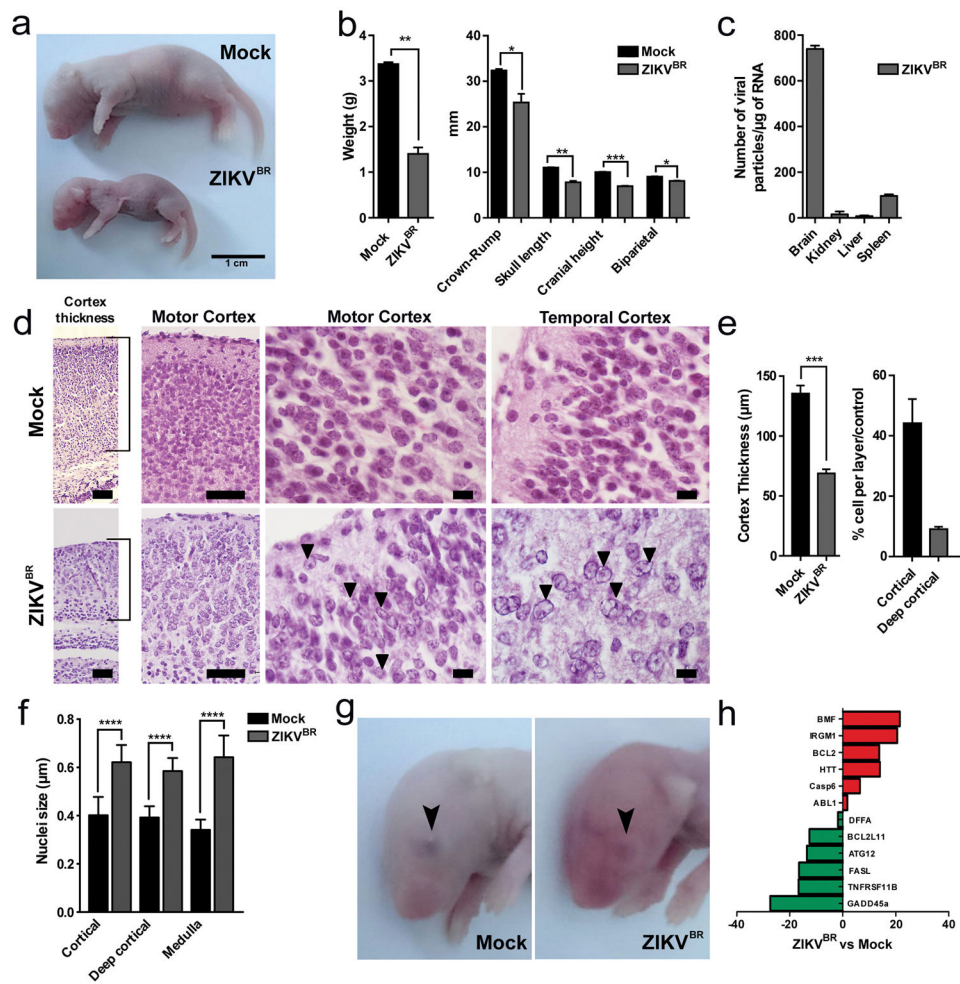
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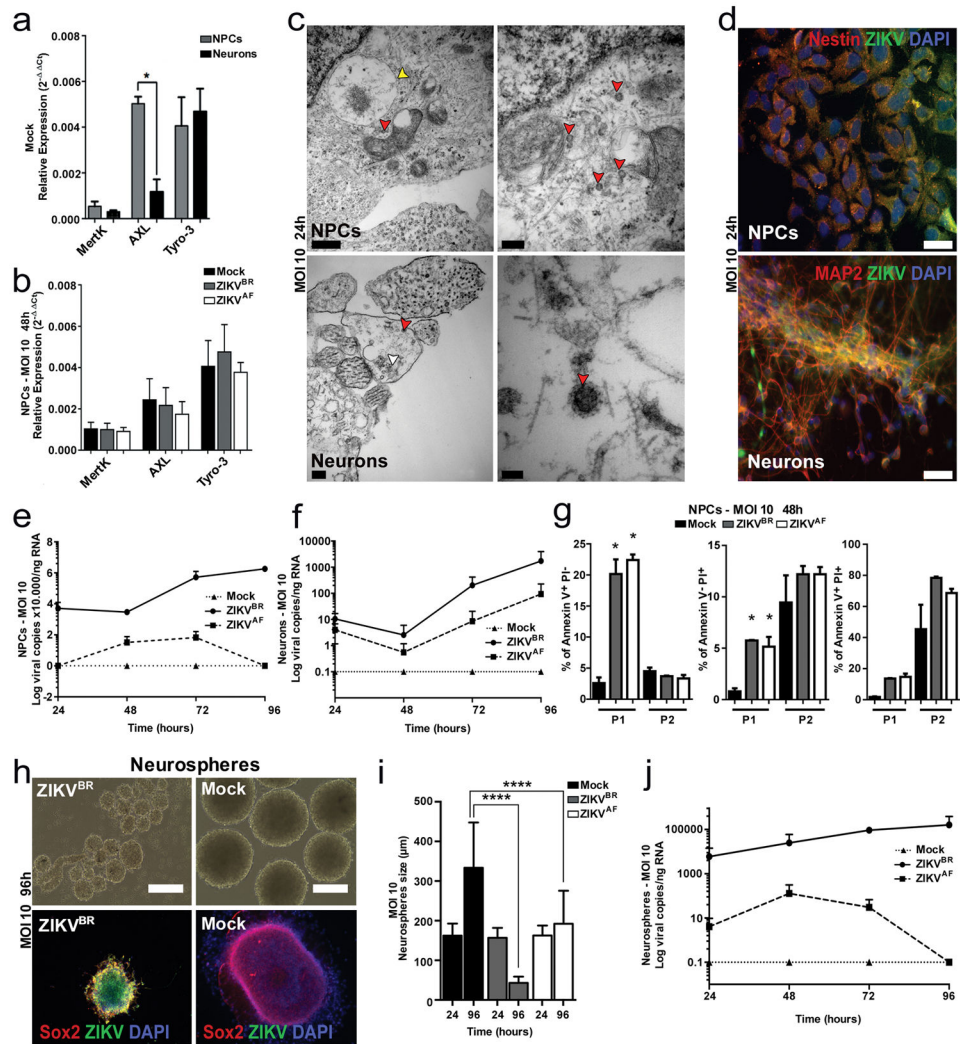
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### Figure 1. ZIKV<sup>BR</sup> infection in SJL mice

**a**, SJL pups born with IUGR. Scale bar = 1 cm. **b**, Total body weight, crown-rump and skull measurements in pups born from infected animals ( $n = 6$  pups, comprising 3 mice from 2 separate litters; error bars, s.e.m;  $t$ -test,  $**P < 0.01$ ). **c**, ZIKV<sup>BR</sup> RNA detected in SJL pup tissues ( $n = 6$  pups, comprising 3 mice from 2 separate litters; error bars, s.e.m;  $t$ -test). **d**, Histopathological aspect of the cortical organization (brackets) in infected brains, including intranuclear vacuoles, and “empty” nuclei aspect with chromatin margination in neurons (arrows). Scale bar = 100 μm (left panels), 50 μm (middle panels) and 10 μm (right panels). **e**, ZIKV<sup>BR</sup>-infected brains displayed a reduced cortical layer thickness ( $n = 6$  pups, comprising 3 mice from 2 separate litters; error bars, s.e.m;  $t$ -test,  $***P < 0.001$ ). Infected brains have fewer cells/layer ( $n = 6$  pups, comprising 3 mice from 2 separate litters; error bars, s.e.m;  $t$ -test,  $**P < 0.1$ ). **f**, ZIKV<sup>BR</sup>-infected cortical neurons have pronounced nuclei (diameter) (cortical  $n = 31$ ; deep cortical  $n = 21$  and medulla  $n = 41$  nuclei; error bars, s.e.m; two-way ANOVA,  $****P < 0.001$ ). **g**, Ocular malformations (arrow) in the ZIKV<sup>BR</sup>-infected pups. **h**, Cell death gene expression signature in the brains of ZIKV<sup>BR</sup>-infected pups ( $n = 2$  mice per group; threshold = two-fold).



**Figure 2. ZIKV infection *in vitro***

**a**, Relative expression of TAM receptors ( $n = 2$  technical replicates from two pooled different donors, error bars, s.e.m.;  $t$ -test; \*\*\* $P < 0,01$ ). **b**, Expression of TAM receptors in NPCs after ZIKV<sup>BR</sup> infection (MOI = 10) at 48 hours p.i. ( $n = 2$  technical replicates from two pooled different donors, error bars, s.e.m.). **c**, TEM detection of ZIKV<sup>BR</sup> viral particles 24 hours p.i. at MOI = 10 (red arrowheads) inside NPCs (top panels) and neurons (bottom panels). Viral factories (yellow arrowhead). Immature viral particles (white arrowheads). Scale bars = 0.5  $\mu\text{m}/40.000\times$  (top left); 200nm/80.000 $\times$  (top right); 0.2  $\mu\text{m}/50.000\times$  (bottom left); 50 nm/3000.000 $\times$  (bottom right). **d**, Immunofluorescence revealed susceptibility to infection in NPC and neurons with the ZIKV<sup>BR</sup> (MOI = 10) at 24 hours p.i. Scale bar = 25  $\mu\text{m}$ . **e**, ZIKV<sup>BR</sup> replication dynamics in NPCs (MOI = 10) ( $n = 2$  technical replicates from RNA of two different donors). **f**, ZIKV<sup>BR</sup> replication dynamics in neurons (MOI = 10) ( $n = 2$  technical replicates from RNA of two different donors). **g**, NPC death measured by FAC with two different cell gating sizes (P1 and P2). Apoptosis (left panel), necrosis (middle panel) and late apoptosis (right panel) (MOI = 10), 48 hours p.i. ( $n = 2$  technical replicates from two different donors; error bars, s.e.m.; two-way ANOVA, \* $P < 0.5$ ). **h**, Representative

images of human neurospheres infected with ZIKV<sup>BR</sup> (MOI = 10; 96 hours p.i.). Scale bar = 200  $\mu\text{m}$ . **i**, Alterations in neurosphere diameter over time (MOI = 10) ( $n = 25$  neurospheres from two different donors for each time point; unpaired  $t$ -test, \*\*\*\* $P < 0.0001$ ). **j**, ZIKV replication dynamics in neurospheres (MOI = 10) ( $n = 2$  technical replicates from two different donors).

