Viral Factors Important for Efficient Replication of Influenza A Viruses in Cells of the Central Nervous System

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Running title: Viral factors important for replication in CNS cells

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Abstract

Central nervous system (CNS) disease is one of the most common extra-respiratory tract complication of influenza A virus infections. Remarkably, zoonotic H5N1 virus infections are more frequently associated with CNS disease than seasonal or pandemic viruses. Little is known about the interaction between influenza A viruses and cells of the CNS, and therefore it is currently unknown which viral factors are important for efficient replication. Here, we determined the replication kinetics of a seasonal, pandemic, zoonotic, and lab-adapted influenza A virus in human neuron-like (SK-N-SH), astrocyte-like (U87-MG) cells and primary mouse cortex neurons. In general, highly pathogenic avian influenza (HPAI) H5N1 virus replicated most efficiently in all cells which was associated with efficient attachment and infection. Seasonal H3N2 virus and to a lesser extent pandemic H1N1 replicated trypsin-dependent in SK-N-SH but not in U87-MG cells. In the absence of trypsin, only HPAI H5N1 and WSN viruses replicated. Removal of the multi-basic cleavage site (MBCS) from HPAI H5N1 virus attenuated, but did not abrogate replication. Taken together, we showed that the MBCS and to a lesser extent ability to attach and are important determinants for efficient replication of HPAI H5N1 virus in cells of the CNS. This suggests that both an alternative HA cleavage mechanism and preference for α-2,3 linked sialic acids allowing efficient attachment, contribute to the ability of influenza A viruses to replicate efficiently in cells of the CNS. This study further improves our knowledge on potential viral factors important for the neurotropic potential of influenza A viruses.

Importance

Central nervous system (CNS) disease is one of the most common extra-respiratory tract complications of influenza A virus infections and frequency and severity differ between seasonal, pandemic and zoonotic viruses. However, little is known about the interaction of these viruses with cells of CNS. Differences among seasonal, pandemic and zoonotic viruses in replication efficacy and cleavability in CNS cells partially explain the higher frequency and severity of zoonotic viruses. Identifying important viral factors and detailed knowledge of the interaction between influenza virus and CNS cells are important to prevent and treat this potentially lethal CNS disease.
Introduction

One of the most common extra-respiratory complications of influenza virus infection is central nervous system (CNS) disease (1, 2). Clinically, CNS disease can range from mild febrile seizures to severe or even fatal meningo-encephalitis (2, 3). Although most studies on influenza virus associated CNS disease have focuses on influenza A viruses, viruses of type B are also able to cause CNS disease. This is however, less frequently observed (2, 4). Influenza A viruses, hereinafter referred to as influenza virus, have been linked to CNS disease since the 1918 H1N1 pandemic (5, 6) and CNS disease has been observed during all subsequent pandemics (7-12) as well as during seasonal epidemics with sporadic detection of influenza virus in the CNS or cerebral spinal fluid (CSF) of humans (13-15). Zoonotic influenza viruses only occasionally infect humans, but when they do, they are frequently associated with severe and systemic disease (1). Highly pathogenic avian influenza (HPAI) H5N1 and H7N9 viruses, two recent zoonotic influenza viruses, are both associated with CNS disease (16-19). The HPAI H5N1 virus is possibly the most neurotropic influenza virus known and has frequently been associated with CNS disease in humans and in other naturally (20-23), and experimentally infected (24-27) mammalian species.

In order to infect, replicate in, and spread throughout the CNS, influenza viruses first have to be able to enter the CNS. Entry of influenza viruses into CNS can occur via for example the olfactory (24, 26-28), trigeminal (6, 27, 29, 30) vagal (29-31), sympathetic nerve (27, 31) and possibly other cranial nerves. The primary targets of influenza viruses are, however, epithelial cells of the respiratory tract (32), which differ from cells of the CNS. Influenza virus infection starts with attachment of the virus to sialic acids (SA) present on host cells (33). Human and avian influenza viruses attach preferentially to α-2,6 and α-2,3-linked SA, respectively, present in the upper and lower respiratory tract of humans, respectively (33). In cells of the CNS, little is known about SA distribution on the different cells at different anatomical locations. One comparative study using lectin immunohistochemistry suggested that in humans, both α-2,6 and α-2,3 SA are present on neurons and glial cells in many different regions, including cerebral cortex, hippocampus, brainstem, and cerebellum (34). In the mouse brain, however, SA distribution is less widespread, and regions
with and without detectable SA are infected with influenza viruses (34). In another study, it was found that in human cortex tissue, some neurons only express α-2,3 SA, oligodendrocytes mainly express α-2,6 SA, while astrocytes appear to express both receptors (35). Moreover, both α-2,3 and α-2,6 SA receptors have been found to be present on human neuroblastoma SK-N-SH and SH-SY5Y and human glioblastoma T98G cell lines (36, 37). Given these differential results as well as the fact that SA usage depends on more than α-2,3 and α-2,6 SA linkage, e.g. α-2,8 SA linkage (38, 39) or even SA-independent entry of the virus (40), more studies should reveal which viruses are able to attach to cells in the CNS.

In order for progeny viruses to infect new cells, cleavage of the immature surface protein hemagglutinin (HA) into the biologically activated and infectious form is required (41). Influenza viruses that contain a mono-basic cleavage site can be cleaved by trypsin-like serine proteases such as human airway trypsin-like protease (HAT), transmembrane serine protease 2 (TMPRSS2), TMPRSS4 or matriptase present in the human respiratory tract (42-44). In the human CNS, expression of HAT in the cerebellum (45) and matriptase mRNA in the frontal and temporal cortex, hippocampus, and cerebellum have been reported (46). Viruses that contain a multi-basic cleavage site (MBCS), such as the HPAI H5N1 virus, can be cleaved by ubiquitously expressed subtilisin-like proteases like furin and PC5/6 (41, 47). This MBCS is an important factor contributing to the ability to spread systemically, including the CNS. Although extra-respiratory spread of HPAI H5N1 virus depends on the presence of the MBCS in ferrets, insertion of a MBCS into a seasonal H3N2 virus did not result in efficient systemic replication in ferrets, suggesting that more factors are necessary (24, 48). Other viruses that are associated with CNS invasion in mice or ferrets are 1918 H1N1 and A/WSN/33 viruses (6, 49, 50). These viruses do not possess a MBCS, but use a different protease-mediated mechanism for HA cleavage, allowing trypsin-independent replication. Taken together, virus receptor specificity, receptor availability on host cells, protease distribution and availability, and HA cleavage mechanism all seem to play an important role in influenza virus infection, cell tropism as well as replication efficiency in the respiratory tract and beyond.
To date, not much is known about the replication efficiency of different influenza viruses, especially seasonal viruses, in cells of the CNS. Thus far, evidence from both *in vivo* and *in vitro* studies suggest that HPAI H5N1 viruses are able to infect and replicate in neurons and astrocytes (17, 20-27, 36, 51-54), but a direct comparison of replication efficiency in cells of the CNS between seasonal, zoonotic, and pandemic influenza viruses is currently lacking. Similarly, insights into the role of attachment, protease availability and presence of a MBCS on replication efficiency in cells of the CNS for these viruses is lacking. Therefore, we here determined the virus attachment, infectivity, and replication kinetics of a seasonal H3N2, 2009 pandemic H1N1 (pH1N1), HPAI H5N1, and WSN viruses in human neuroblastoma (SK-N-SH), human astrocytoma (U87-MG), primary mouse cortex neurons (pmCortex), and MDCK cells. Subsequently, we established the importance of the MBCS for the replication efficiency of HPAI H5N1 virus in cells of the CNS.

**Materials & Methods**

**Cells**

Human neuroblastoma (neuron-like, SK-N-SH) and human glioblastoma (astrocyte-like, U87-MG) cells were purchased from Sigma-Aldrich and maintained in EMEM (Lonza, Breda, the Netherlands) supplemented with 10% fetal bovine serum (FBS, Sigma-Aldrich, St Louis, Missouri, USA), 100 IU/ml penicillin (Lonza, Basel, Switzerland), 100 μg/ml streptomycin (Lonza), 2 mM glutamine (Lonza), 1.5 mg/ml sodium bicarbonate (Cambrex, Wiesbaden, Germany), sodium pyruvate (Thermo Fisher Scientific, Waltham, Massachusetts, USA) and 1X (0.1 mM) non-essential amino acids (MP Biomedicals Europe, Illkirch, France). As a control cell line, we have included Madin–Darby Canine kidney (MDCK) cells since these cells are extensively used for influenza virus propagation. MDCK cells were maintained in EMEM supplemented with 10% FBS, 100 IU/ml penicillin, 100 μg/ml streptomycin, 2 mM glutamine, 1.5 mg/ml sodium bicarbonate, 1 mM, 10 mM Hepes (Cambrex) and 1X (0.1 mM) non-essential amino acids.

**Viruses**

Five viruses were included in this study, a seasonal H3N2 virus (A/Netherlands/213/2003), pH1N1 virus (A/Netherlands/602/2009), and zoonotic...
HPAI H5N1 virus (A/Indonesia/5/2005) all isolated from humans. Neurotropic WSN virus (A/WSN/33), and H5N1 virus lacking a MBCS (H5N1\(^{\text{∆MBCS}}\)) were generated using reverse genetics as described before (55) and passed once on 293T cells and once on MDCK cells. Experiments involving HPAI H5N1 and H5N1\(^{\text{∆MBCS}}\) viruses were performed under biosafety level 3 conditions.

**Isolation and Culture of Primary Mouse Cortex Neurons**

Animals were housed and experiments were conducted in strict compliance with European guidelines (EU directive on animal testing 86/609/EEC) and Dutch legislation (Experiments on Animals Act, 1997). Primary mouse cortex tissue was isolated from E17-E19 C57/BL6 mouse embryos (Charles River Laboratories, Wilmington, Massachusetts, USA). The cultures were pooled cortices of several mouse embryo's originating from one mother. In brief, the cortex was dissected in ice-cold Hank's balanced salt solution (HBSS, Life Technologies) supplemented with 20 \(\mu\)g/mL gentamicin (Life Technologies) under the guidance of a stereomicroscope (Nikon). Next, tissues were cut into ~1mm\(^3\) using a scalpel and digested using medium consisting of HBSS supplemented with 10 U/ml papain (Sigma), 2.5 U/ml DNAse I (Roche), and 4 mM MgCl\(_2\) (Sigma Aldrich) at 33°C for 15 minutes. After incubation, cells were washed once in 1 ml of 10% FBS (Life Technologies) in HBSS to stop the digestion. A second “mechanical digestion” was performed by carefully pipetting up and down in digestion buffer (without papain). After washing in HBSS twice, cells were counted using Moxi Go cell counter (Orflo, Ketchum, Idaho, USA) and seeded on laminin (500 \(\mu\)g/ml, Sigma) coated #1.5H 96 well glass-bottomed plates (Cellvis, Sunnyvale, California, USA) at a density of 1.0 x 10\(^4\) cells/well. For the first 2 hours, cells were cultured in culture medium containing 10% FBS. After 2 hours, medium was replaced with fresh culture medium, without FCS. The culture medium contains primary neuron growth medium (PNBM; Lonza), GS-21 supplement (Tebu-Bio, Le-Perray-en-Yvelines, France), 5 \(\mu\)g/ml gentamicin (Thermo Fisher), and 2 mM glutamax (Life Technologies). Half of the medium was changed once a week and cells were cultured for 7-10 days before use.

**Replication kinetics**

Cells were infected at a multiplicity of infection (MOI) of 0.1. Virus dilutions were prepared in the cell specific culture medium without serum (infection medium, see...
virus titrations). After one hour of virus absorption, cells were washed once with fresh infection medium and cultured in infection medium in the presence or absence of TPCK treated trypsin (see virus titration). At time points 1, 6, 24, and 48 hours post infection (hpi), 100 μl supernatant was collected and stored at -80°C for subsequent virus titration. All experiments were performed three times (biological replicates), and each experiment performed in duplicate (technical replicates) from which the average was used for statistical analysis.

**Virus Titrations**

The 50% tissue culture infectious dose (TCID\textsubscript{50}) in cell supernatant was determined by end point titration on MDCK cells, as described before (56). Briefly, 10-fold serial dilutions of cell supernatants were prepared in infection medium. Infection medium consisted of EMEM, 100 IU/ml penicillin, 100 μg/ml streptomycin, 2 mM glutamine, 1.5 mg/ml sodium bicarbonate, 10 mM Hepes, 1X (0.1 mM) non-essential amino acids, and 1 μg/μl L-1-Tosylamide-2-phenylethyl chloromethyl ketone (TPCK) treated trypsin (Sigma Aldrich). Before inoculation, MDCK cells were washed twice with PBS to remove remaining FBS. One hundred μl of the diluted supernatant was used to inoculate a confluent monolayer of MDCK cells in 96-well plates. After 1 hour at 37°C the cells were washed once with infection medium and 200 μl new infection medium was added to each well. Three days after infection, supernatants of infected cell cultures were tested for agglutinating activity using turkey erythrocytes as an indicator of virus replication. The titers of infectivity were calculated from three replicates according to the method of Spearman–Karber (57). An initial 1:10 dilution of supernatant resulted in a detection limit of 10\textsuperscript{1.5} TCID\textsubscript{50}/ml.

**Percentage of infection**

After 8 hpi (with a MOI of 3) or after 8 and 24 hpi (with a MOI of 0.1), in the absence of trypsin, cells were collected, fixed, and permeabilized using BD Cytofix/Cytoperm solution (BD Biosciences, Franklin Lakes, New Jersey, USA) according to the manufacturer’s instructions. Cells were incubated with 2% normal goat serum (NGS, Dako, Denmark) in PBS, for 10 min on ice. Next, influenza A virus was detected using a monoclonal antibody against influenza A virus nucleoprotein (clone HB-65; ATCC, 1 μg/ml) or mouse IgG2a isotype control (MAB003; Dako, 1 μg/ml) in BD Perm/Wash containing 2% NGS and incubated for 1 hour on ice and in the dark.
Cells were washed twice, and incubated with goat anti-mouse IgG2a conjugated with Alexa Fluor 488 (Life Technologies, Inc., The Netherlands, 8 µg/ml) for 1 hour in the dark and on ice. After incubation, cells were washed twice and resuspended in FACS buffer. Cells were measured and data collected using a BD FACSCanto II (BD Biosciences, USA). Data were analyzed using FlowJo 10 software (Ashland, OR, USA). All experiments were performed three times (biological replicates), and each experiment included duplicate (technical replicate) measurements from which the average was calculated and used for further analysis.

**Virus attachment**

For influenza virus histochemistry, viruses were grown, inactivated, and labelled as described previously (32). As a control, uninfected MDCK cells, and cell debris, were harvested and processed similarly. Subsequently, in a 12-well plate, 2 x 10^5 cells were seeded and one day later, the near confluent monolayer of MDCK, SK-N-SH, and U87MG cells were harvested, washed in FACS buffer, and incubated with FITC-labelled virus for 1 hour at 4°C. After incubation, the cells were washed twice in FACS buffer and measured using a BD FACSCanto II (BD Biosciences, USA). Data were analyzed using FlowJo 10 software (Ashland, OR, USA). All experiments were performed three times (biological replicates), and each experiment performed in duplicate (technical replicates) from which the average was used for statistical analysis.

**PCR proteases**

Since MDCK cells are of canine origin, we have included primary human nasal (HN) cells (MucilAir™, pool of 14 donors, Epithelix, Geneve, Switzerland), and primary human bronchial/tracheal epithelial cells (HTBE, Lonza, catalog #CC-2540, lot#97366, donor #97366: male, Caucasian, 57 years, healthy) as control cell types for the expression of human HAT, human TMPRSS2, and human TMPRSS4. Total RNA was isolated from SK-N-SH, U87-MG, HN, and HTBE cells using the High Pure RNA isolation kit (Roche, Basel, Switzerland) according to the manufacturer’s protocol. cDNA synthesis was performed using Oligo dT primers and Superscript IV (Applied Biosystems, Foster City, California, USA) according to manufacturer’s instructions. For detection of HAT-, TMPRSS2-, and TMPRSS4-specific mRNAs, primers were used from Böttcher-Friebertshäuser E, et al (58). The GAPDH mRNA
was detected using primers GAPDH-FW (5` TGA ACG GGA AGC TCA CTG G 3`) and GAPDH-RV (5` TCC ACC ACC CTG TTG CTG TA 3`) as a control for sample quality. PCR products were resolved on a 1.5% agarose gel stained with SYBR Safe (Thermo Fisher) and imaged using a ChemiDoc MP Imaging System and ImageLab 5.1 (Bio-rad, Hercules, California, USA). To confirm the specificity of the primers, PCR products were extracted from the gel and sequenced using a BigDye Terminator v3.1 Cycle sequencing kit (Applied Biosystems) and a 3130XL genetic analyzer (Applied Biosystems), according to the instructions of the manufacturer.

Statistical analysis
Statistical analyses were performed using GraphPad Prism 6.0h software (La Jolla, CA, USA) for Mac. Each specific test is indicated in the figure legend. P values of ≤0.05 were considered significant. All data are presented as mean ± SD of at least three independent experiments with ns, P > 0.05; *, P ≤ 0.05; **, P ≤ 0.01 *** P ≤ 0.001; ****, P ≤ 0.0001.

Results
HPAI H5N1 virus replicates more efficiently in CNS cells than H3N2 and pH1N1 viruses.
The replication kinetics of pH1N1, H3N2, HPAI H5N1, and WSN viruses were determined in SK-N-SH, U87-MG, pmCortex, and MDCK cells in the presence of trypsin. All viruses replicated efficiently in MDCK cells, where HPAI H5N1 virus and H3N2 virus replicated to higher titers than pH1N1 virus and WSN virus (figure 1d). In both SK-N-SH and U87-MG cells, HPAI H5N1 virus replicated to a significantly higher titer (~7 log_{10}TCID_{50}/mL) compared to all other viruses (figure 1a, 1b). In U87-MG and pmCortex cells, only HPAI H5N1 and WSN viruses were able to replicate (figure 1b, 1c). In addition to HPAI H5N1 virus, SK-N-SH cells supported replication of pH1N1, H3N2, and WSN viruses, reaching a virus titer of ~5.7, ~3.9, ~3.3, respectively (figure 1a). Overall, our results show that HPAI H5N1 and WSN viruses replicated in all cells investigated and that H3N2 and pH1N1 viruses replicated less efficiently in SK-N-SH cells not at all in U87-MG cells and pmCortex cells.
HPAI H5N1 and WSN viruses infected cells more efficiently than H3N2 and pH1N1 viruses.

To determine whether efficient replication was associated with the ability of the virus to enter and infect host cells, we determined the percentage of infection 8 hpi (MOI 3) in SK-N-SH, U87-MG, and MDCK cells, measured by flow cytometry (figure 2). In MDCK cells, pH1N1, HPAI H5N1, and WSN viruses infected significantly more cells than H3N2 virus. In SK-N-SH and U87-MG cells, HPAI H5N1 and WSN viruses infected significantly more cells than H3N2 or pH1N1 viruses.

pH1N1 and H3N2 viruses attach less efficiently to SK-N-SH and U87-MG cells than HPAI H5N1 and WSN viruses

To determine whether there were differences in attachment between the viruses and whether this was associated with the infection percentages, we performed a virus attachment assay. The attachment efficiency was scored as follows: inefficient attachment (0-5%), low attachment (6-25%), intermediate attachment (26-75%), and efficient attachment (>76%). HPAI H5N1 virus attached efficiently (>90%) to all cell lines investigated (figure 3). WSN virus attached efficiently (>95%) to SK-N-SH and U87-MG cells and intermediately (70%) to MDCK cells (figure 3). Pandemic H1N1 virus attached with intermediate efficiency to SK-N-SH (37%) and MDCK cells (30%), (figure 3a, 3b, 3d). Seasonal H3N2 virus attached with low efficiency to SK-N-SH (12%) and intermediate efficiency to MDCK cells (39%), respectively (figure 3a, 3b, 3d). Neither pH1N1 nor H3N2 viruses attached to U87-MG cells (<2%, figure 3a, 3c). Overall, these results show that pH1N1 and H3N2 viruses attach less efficiently to SK-N-SH and U87-MG cells than H5N1 and WSN viruses.

HPAI H5N1 and WSN viruses replicate in the absence of trypsin, but H3N2 and pH1N1 viruses not

To test whether efficient replication is dependent on the presence of trypsin, we determined the replication kinetics in the absence of trypsin. Replication of HPAI H5N1 and WSN viruses was not affected by the absence of trypsin (figure 4). In the presence of trypsin, H3N2 and pH1N1 viruses replicated efficiently in SK-N-SH cells, but not in the absence of trypsin (figure 4a). To further understand this finding, we analyzed the presence of specific host cell proteases known to cleave the HA protein.
of H3N2 and pH1N1 viruses (43). We found that neither SK-N-SH nor U87-MG cells expressed HAT, TMPRSS2, nor TMPRSS4 mRNA, whereas these transcripts were present in human nasal cell (HN) cultures and human bronchial/tracheal epithelial (HBTE) cultures (figure 5). These results show that HPAI H5N1 and WSN viruses replicate independently of trypsin, and that pH1N1 and H3N2 viruses are dependent on trypsin for replication in SK-N-SH cells.

The MBCS of HPAI H5N1 virus is important, but not solely responsible for replication in SK-N-SH cells

To determine whether efficient replication of HPAI H5N1 virus in cell culture solely depends on the presence of the MBCS, we generated a H5N1 virus without MBCS. The replication kinetics of HPAI H5N1 WT virus was not affected by the presence or absence of trypsin in all cell lines (figure 6a). However, H5N1ΔMBCS virus, without trypsin, replicated to a lower titer on each cell line investigated. This phenotype was restored by the addition of trypsin to the culture medium, allowing the virus to replicate to wildtype levels (figure 6a). The reduced replication efficiency of H5N1ΔMBCS virus was not explained by the ability of the virus to infect cells, since this was not affected (figure 6b). In order to determine if there were multiple rounds of infection, and that virus detected in the supernatant was not solely the result of primary infected cells, we investigated the percentage of infection at 8 and 24 hpi with a MOI of 0.1 -without trypsin-, measured by flow cytometry. We found that HPAI H5N1 WT virus efficiently replicated in MDCK, SK-N-SH, and U87-MG cells as indicated by the increase of infection percentages (figure 6c). In contrast, a significant increase for H5N1ΔMBCS virus only was observed in MDCK and U87-MG cells. In SK-N-SH cells, no increased infection percentage was observed. These results reveal that the MBCS is important but not solely responsible for efficient replication in MDCK and U87-MG cells in the absence of trypsin.

Discussion

Here we show that HPAI H5N1 virus replicates more efficiently in human and mouse neuronal cells than seasonal H3N2 and 2009 pandemic H1N1 viruses. Both the ability to attach efficiently and the presence of a MBCS of HPAI H5N1 virus contributed to efficient replication in cells of the CNS, indicative that these are viral
factors that contribute to the neurotropic potential of influenza viruses. This fits with the facts that HPAI H5N1 virus is more frequently associated with CNS disease in humans than seasonal and pandemic viruses (1) and that this virus is also more often detected in tissues of the CNS in experimentally inoculated laboratory animals than seasonal and pandemic viruses (17, 20-27, 51, 59-62).

The ability of HPAI H5N1 virus and WSN to replicate efficiently in cells of the CNS seems to be associated with the ability to attach to and infect host cells efficiently. Especially HPAI H5N1 virus replicated efficiently in SK-N-SH and U87-MG cells, and attached to high percentages of cells with high intensity and infected these cells efficiently. WSN attached to, and infected high percentages of neuronal cells which resulted in multiple round infection in all cells, although to lower titers on SK-N-SH and U87-MG cells than observed for HPAI H5N1 virus. The latter could be due to the fact that WSN is extensively passage in suckling mouse brains, thereby adapting to mouse neuronal cells, and not human neuronal cells. Seasonal H3N2 viruses replicated in SK-N-SH cells, even in the absence of efficient attachment, or infection. Whether this is due to low affinity binding of H3N2 virus, which cannot be detected by our assays is unknown, but it does suggest that efficient attachment, as observed for H5N1 virus and WSN on neuronal cells, is not the only viral factor involved in replication of influenza viruses in cells of the CNS. The lack of replication of H3N2 and pH1N1 virus in U87-MG cells could be explained by both inefficient attachment and infection.

Efficient replication of influenza viruses in cells of the CNS seems to depend in part on the presence of a MBCS or alternative HA cleaving proteases. Studies in ferrets, mice, macaques, and chickens show that introduction or removal of a MBCS has different outcomes based on the virus backbone and host species (24, 48, 63-65). In vitro, in the absence of trypsin, only HPAI H5N1 and WSN viruses were able to replicate indicating that pH1N1 and H3N2 viruses are not able to circumvent the need for trypsin-like protease for HA cleavage. Removal of the MBCS from HPAI H5N1 virus resulted in attenuated replication in SK-N-SH and U87-MG cells, in the absence of trypsin, but not in pmCortex cells. Viruses without a MBCS that are associated with replication in cells of the CNS, in vivo, such as the 1918 H1N1 and WSN viruses, have an alternative HA cleavage mechanism (6, 66). The WSN virus
lacks a conserved glycosylation site in the neuraminidase making the virus trypsin-independent (67) by using the serine-protease plasmin, which is present in many organ systems besides the respiratory tract (68). The 1918 H1N1 virus grows trypsin-independent, neuraminidase-dependent in MDCK cells and polarized Calu-3 cells, but not in huh-H7 cells (69, 70). Our observation that neither HAT, TMPRSS2 nor TMPRSS4 are found in SK-N-SH and U87-MG cells, supports the hypothesis that for efficient replication in cells of the CNS, influenza viruses require alternative HA cleavage as shown for the HPAI H5N1, WSN, and 1918 H1N1 viruses. However, it must be noted that there is limited knowledge on expression and accessibility of proteases in tissues other than the respiratory tract.

Previous studies on the replication kinetics of influenza viruses in cells of the CNS observed some differences, which can in part be explained by the use of different cells, virus isolates and experimental approach (36, 54). Replication of HPAI H5N1 viruses in differentiated astrocytic cells lines resulted in efficient replication, similar to our observations. However, in SH-SY5Y cells, a sub-clone cell line derived from SK-N-SH cells, two HPAI H5N1 viruses did not replicate efficiently. This discrepancy could be due to the relatively high MOI used in this study compared to the low MOI we used and which resulted in efficient replication (36). Furthermore, two pandemic H1N1 viruses did not replicate in the neuronal or astrocytic cell lines, which fits with our observation, in the absence of trypsin (54).

Pandemic H1N1 and seasonal H3N2 viruses are occasionally detected in the CNS or CSF in humans, and from experimentally inoculated ferrets and mice even though pH1N1 and H3N2 viruses did only replicate in SK-N-SH cells in the presence of trypsin (11-15, 71-74). However, these viruses are rarely isolated in high titers or by immunohistochemistry in the CNS of humans or experimentally inoculated ferrets and mice, indicative that these viruses might be able to enter the CNS, but that replication is inefficient. This could be attributed to the limited attachment and infection and lack of an alternative HA cleavage mechanism allowing efficient replication. Previously we have shown that even in the absence of active virus replication, pro-inflammatory cytokines, such as IL-6, IL-8 and TNFα are induced in the CNS of pH1N1 experimentally inoculated ferrets (75). Future studies should reveal how both efficient and inefficient replication in neuronal cells can trigger local
pro-inflammatory responses, for which HPAI H5N1 and H5N1∆MBCS viruses might be a good model.

Taken together, our study has shown that the presence of a MBCS and to a lesser extent ability to attach are important determinants for replication of HPAI H5N1 virus in cells of the CNS. This suggests that, at least for replication within the CNS, neurotropic influenza viruses contain an alternative HA cleavage mechanism and prefer α-2,3 linked sialic acids.

Acknowledgments

The authors thank Mathilde Richard for providing the human nasal cells and Laurine Rijsbergen for providing human bronchial/tracheal epithelial cells.

Financial support

This work was supported by a fellowship from the Netherlands Organization for Scientific Research (contract 91614115 and 91718308), the Erasmus MC Foundation and an ESCMID research grant.

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Figure 1. In the presence of trypsin, HPAI H5N1 virus replicates most efficiently in SK-N-SH, U87-MG, pmCortex, and MDCK cells. Growth kinetics of pH1N1, H3N2, HPAI H5N1, and WSN viruses in SK-N-SH, U87-MG, pmCortex, and MDCK cells (MOI 0.1) in the presence of trypsin. Data are presented as mean ± SD of at least three independent experiments. Two-way ANOVA with Dunnett’s multiple comparisons test against HPAI H5N1 virus, with *, P ≤ 0.05; **, P ≤ 0.01 ***, P ≤ 0.001; ****, P ≤ 0.0001.

Figure 2. HPAI H5N1 and WSN viruses infect cells more efficiently than pH1N1 and H3N2 viruses. Percentages of infection in SK-N-SH, U87-MG, and MDCK cells were determined by FACS 8 hpi with either pH1N1, H3N2, WSN, or HPAI H5N1 viruses (MOI of 3). Data are presented as mean ± SD of at least three independent experiments. Statistical analysis was performed using the two-way ANOVA with Tukey’s multiple comparisons test with *, P ≤ 0.05; **, P ≤ 0.01 ***, P ≤ 0.001; ****, P ≤ 0.0001.

Figure 3. HPAI H5N1 and WSN viruses attach more efficiently to SK-N-SH and U87-MG cells than pH1N1 and H3N2 viruses. Virus attachment of pH1N1, H3N2, HPAI H5N1, and WSN viruses (using 100 hemagglutination units [HAU] units) to SK-N-SH, U87-MG, and MDCK cells. A) Percentage of cells to which viruses attached. B-D) Representative histogram of SK-N-SH, U87-MG, and MDCK cells, respectively. Dotted line indicates MDCK cell control. Data panel A are presented as mean ± SD of at least three independent experiments. Two-way ANOVA with Tukey’s multiple comparisons test with *, P ≤ 0.05; **, P ≤ 0.01 ***, P ≤ 0.001; ****, P ≤ 0.0001.

Figure 4. In the absence of trypsin, HPAI H5N1 virus grows most efficiently in SK-N-SH, U87-MG, and MDCK, and pmCortex cells. Growth kinetics of pH1N1, H3N2, HPAI H5N1, and WSN viruses in SK-N-SH, U87-MG, MDCK cells, and pmCortex cells (MOI 0.1) in the absence of trypsin. Data are presented as mean ± SD of at least three independent experiments. Two-way ANOVA with Dunnett’s multiple comparisons test against H5N1 virus, with *, P ≤ 0.05; **, P ≤ 0.01 ***, P ≤ 0.001; ****, P ≤ 0.0001.
Figure 5. HAT, TMPRSS2, and TMPRSS4 mRNAs are not present in SK-N-SH and U87-MG cells. Presence of three known HA cleaving enzymes human airway trypsin (HAT), transmembrane serine protease 2 (TMPRSS2), and TMPRSS4. Positive control cell lines human nasal cells (HN) and human bronchial/tracheal epithelial (HBTE) cells did express HAT, TMPRSS2, and TMPRSS4 mRNA.

Figure 6. H5N1 virus without a MBCS replicates less efficiently in the absence of trypsin. A) Replication kinetics of HPAI H5N1WT and H5N1ΔMBCS viruses in SK-N-SH, U87-MG, and MDCK cells (MOI 0.1) in the presence or absence of trypsin. Statistical analysis was performed using a two-way ANOVA with Tukey’s multiple comparisons test against “H5N1WT + trypsin”. B) Percentages of infection in SK-N-SH, U87-MG, and MDCK cells were determined by FACS at 8 hpi with HPAI H5N1WT and H5N1ΔMBCS viruses (MOI 3). Statistical analysis was performed using the Two-way ANOVA with Bonferroni’s multiple comparisons test. C) Percentages of infection in SK-N-SH, U87-MG, and MDCK cells were determined by FACS at 8 and 24 hpi with HPAI H5N1WT and H5N1ΔMBCS viruses at a MOI of 0.1 in the absence of trypsin. Statistical analysis was performed using the two-way ANOVA with Bonferroni’s multiple comparisons test (8 vs 24 hpi). All data are presented as mean ± SD of at least three independent experiments with *, P ≤ 0.05; **, P ≤ 0.01 ***, P ≤ 0.001; ****, P ≤ 0.0001.
Figure A shows the virus titer (log10 TCID50/ml) of pH1N1, H3N2, H5N1, and WSN in SK-N-SH cells over time (0-48 hours). Figure B depicts the virus titer in U87-MG cells. Figure C illustrates the virus titer in pmCortex cells, while Figure D shows the virus titer in MDCK cells. The graphs indicate the growth of these viruses over time, with significant differences indicated by asterisks (**** for statistical significance).
