Comparing Influenza Virus Biology for Understanding Influenza D Virus

Raegan M. Skelton and Victor C. Huber*

Division of Basic Biomedical Sciences, University of South Dakota, Vermillion, SD 57069, USA; raegan.skelton@coyotes.usd.edu
* Correspondence: victor.huber@usd.edu

Abstract: The newest type of influenza virus, influenza D virus (IDV), was isolated in 2011. IDV circulates in several animal species worldwide, causing mild respiratory illness in its natural hosts. Importantly, IDV does not cause clinical disease in humans and does not spread easily from person to person. Here, we review what is known about the host–pathogen interactions that may limit IDV illness. We focus on early immune interactions between the virus and infected host cells in our summary of what is known about IDV pathogenesis. This work establishes a foundation for future research into IDV infection and immunity in mammalian hosts.

Keywords: influenza D virus; innate immunity; interferon; virus pathogenesis

1. Introduction

Within the Orthomyxoviridae family, the four genera of influenza viruses are Alphainfluenzavirus (influenza A virus, IAV), Betainfluenzavirus (influenza B virus, IBV), Gammainfluenzavirus (influenza C virus, ICV), and Deltainfluenzavirus (influenza D virus, IDV) [2,11]. Within the human host, influenza-like illness has been observed during infection with IAV, IBV, and ICV.
Symptoms of influenza infection can develop suddenly, and mild infections remain confined mostly to the upper respiratory tract with a sore throat, runny nose, and cough as well as fever, headache, muscle pain, and fatigue [2,9,12]. More severe infections, typically associated with either IAV or IBV, can spread to the lower respiratory tract and other areas in the body, leading to more serious complications such as pneumonia, secondary bacterial infections, myocarditis, encephalitis, and even death [2,13]. Although symptoms usually last for three to eight days, a persistent cough and fatigue can remain for at least two weeks [10,12]. Seasonal influenza continues to have a significant impact in the United States, with the Centers for Disease Control and Prevention (CDC) estimating that the 2018–2019 influenza season (prior to the start of the COVID-19 pandemic) had 36 million illnesses, nearly 500,000 hospitalizations, and over 34,000 influenza-associated deaths [9]. Influenza has a far-reaching impact and is also a significant global health concern. Worldwide, there are an estimated 1 billion infections, 54.5 million hospitalizations, and 200,000 influenza-associated deaths every year due to influenza-like illnesses [14,15]. According to the World Health Organization (WHO), about 10% of the world’s population is affected by influenza annually [15].

3. Host Range of Influenza Viruses

Depending on the genus, the infection cycle of influenza viruses can take place in a wide variety of host species (Figure 1). IAVs have the largest host range and are considered to be “species jumpers” for their ability to rapidly move between host species [16].

![Figure 1. Natural host range of each influenza virus type. The major species of hosts that each influenza virus type naturally infects is shown, with some overlap existing across influenza virus types. Of note, pigs are the only species known to be infected by all four influenza virus types, and influenza D virus demonstrates the second widest host range behind influenza A virus [16]. Figure created with BioRender.com was adapted from ref. [16]: Kuchipudi and Nissly, 2018.](image-url)
In nature, the IAV host range includes birds, which serve as their natural reservoir, as well as humans, pigs, rodents, cats, horses, and even whales [2,16]. Seasonal and epidemic influenza virus infections can be caused by IAV strains, and all previous influenza pandemics have been due to IAVs [2,17]. In contrast, IBV and ICV infect a narrow host range. IBVs, which can also cause seasonal and epidemic outbreaks of influenza, are mostly restricted to the human host, but they can also infect pigs and seals [2,16]. Humans and pigs are the only known hosts for ICV, and most humans acquire immunity against ICV by early childhood while exhibiting either mild symptoms or asymptomatic illness [16,18,19].

In contrast with the other three influenza virus types, IDV has not yet been described clinically in humans. This virus naturally infects cattle, pigs, horses, and other ruminants around the world such as camels, goats, and sheep [5,6,20,21]. As research focuses on this new influenza virus type, the potential host range of IDV continues to increase. In the research setting, mammalian infection by IDV has been demonstrated in mice [22,23], guinea pigs [24], and ferrets [6], and in vitro studies show that IDV can infect a range of cells while replicating at different temperatures [5,6,25]. In the absence of known clinical infection, seropositivity against IDV remains our only evidence that IDV can infect humans. Studies using archived sera have detected antibodies against IDV at least as far back as 2005, pre-dating the isolation of this virus [8]. In fact, a published report showed that greater than 90% of those who regularly work with cattle have antibodies against IDV [7], suggesting that IDV is capable of inducing adaptive immunity in humans that have high exposure rates [5]. In the absence of clinical cases of IDV, it is assumed that these infections are associated with either mild symptoms or asymptomatic illness. Thus, at this time neither the full host range nor the breadth of disease caused by IDV are completely known, and additional research is needed to fully appreciate the global reach of this newly identified influenza virus type.

4. Influenza Virus Proteins and Their Function

All four types of influenza virus are enveloped, take on a spherical (∼100 nm) or filamentous (300 nm+) shape, and contain a single-stranded, segmented RNA genome [4,11,26]. While IAV and IBV use eight RNA gene segments to express anywhere from 10–17 proteins, ICV and IDV utilize seven RNA gene segments that express nine proteins [4,27,28]. Influenza viruses are packaged with everything needed to infect a host cell and make more virus, with replication cycles highly conserved across all four types of influenza virus [4]. The first step in the infection cycle is binding and entry, when influenza viruses use surface proteins to bind sialic acids or sialic acid derivatives located on host cell receptors [4,29,30]. For IAV and IBV, the surface protein that targets host cells is hemagglutinin (HA), while hemagglutinin esterase fusion (HEF) is used by ICV and IDV. HA and HEF binding targets can be found within the respiratory tract of mammalian hosts, as well as in the intestinal tract of avian hosts [2]. The interaction between the virus and the host cell triggers receptor-mediated endocytosis of the virus into the host cell and initiation of the infection cycle [4,30].

Upon entry into a host cell, ion channels along the surface of the virus function similarly across influenza virus types (M2 for IAV, BM2 for IBV, CM2 for ICV, and DM2 for IDV), allowing for acidification of the endosome to facilitate fusion of viral and endosomal membranes. Membrane fusion allows for release of viral RNA gene segments, packaged as viral ribonucleoprotein complexes (vRNPs), into the host cell cytoplasm. Each vRNP (Figure 2) consists of an RNA gene segment neatly wrapped with several copies of influenza nucleoprotein (NP) and bound by the viral RNA-dependent RNA polymerase (RDRP) complex, which consists of PB2, PB1, and PA for IAV/IBV or PB2, PB1, and P3 for ICV/IDV [4,30]. The vRNPs are transported into the nucleus where the RDRP complex bound to each RNA segment initiates replication and transcription of viral RNA (vRNA) [30]. The vRNA that enter the nucleus as negative-sense templates are transcribed by the RDRP complex into “host” positive-sense mRNA that are translated into viral proteins within the cytoplasm using the host’s ribosomal machinery [4,30]. Newly formed vRNPs are transported out of
the nucleus with help from the influenza nuclear export protein (NEP), expressed by all four influenza virus types, while appropriate viral are transported to the cell surface. These viral proteins work together to package the newly created proteins and genetic material into budding virions [30].

Each RNA gene segment packaged in an influenza virus encodes at least one protein that will help facilitate viral replication, with some gene segments encoding accessory proteins that perform additional functions within infected cells. The names and functions of the major protein products encoded by each influenza gene segment are summarized in Table 1. Accessory proteins of influenza A viruses are reviewed in more detail elsewhere [27,31].

Table 1. Influenza virus genes and proteins. Gene segments are listed for influenza A virus (IAV), influenza B virus (IBV), influenza C virus (ICV), and influenza D virus (IDV). The protein product(s) of each segment and their associated functions are summarized for each influenza species [4,27,31].

| Gene Segment | IAV | IBV | Viral Function |
|--------------|-----|-----|---------------|
| 1            | PB2 | PB2 | RNA-dependent RNA polymerase (RDRP) component |
| 2            | PB1 | PB1 | RDRP component |
|              | PB1-F2 |     | Inflammation, apoptosis, regulation of host immune responses |
|              | PB1-N40 |     | Regulates PB1 expression and activity |
| 3            | PA  | PA  | RDRP component |
|              | PA-X |     | Enhances viral gene expression, facilitates host mRNA degradation, regulation of cell-mediated host responses |
|              | PA-N155 |     | Functions unknown, likely involved with viral replication |
|              | PA-N182 |     | Functions unknown, likely involved with viral replication |
### Table 1. Cont.

| Gene Segment | ICV | IDV | Viral Function                                                                 |
|--------------|-----|-----|--------------------------------------------------------------------------------|
| 1            | PB2 | PB2 | RNA-dependent RNA polymerase (RDRP) component                                  |
| 2            | PB1 | PB1 | RDRP component                                                                |
| 3            | P3  | P3  | RDRP component                                                                |
| 4            | HEF | HEF | Host receptor binding, membrane fusion; esterase; assists release of new virions |
| 5            | NP  | NP  | Packages viral RNA in vRNPs with RDRP components                               |
| 6            | M1  | M1  | Facilitates packing of vRNPs into new virions                                  |
|              | CM2 | DM2 | Ion channel; assists in release of vRNPs into host cytoplasm                   |
| 7            | NS1 | NS1 | Host immune response antagonism                                                |
|              | NS2/NEP | NS2/NEP | Nuclear export protein for newly synthesized vRNPs                            |

1 Accessory protein; 2 vRNPs: viral ribonucleoproteins.

All four influenza virus types use the M1 protein to package nascent virions with vRNPs and the virions then bud from the host cell with everything they need to infect new cells [4,30]. At budding, virus-binding residues on the host cell surface are cleaved through either neuraminidase (NA) activity for IAV and IBV or the esterase function of the HEF protein for ICV and IDV [4,30]. One of the unique features of ICV and IDV is that the single HEF protein performs both the binding and residue-cleaving functions of the IAV and IBV HA and NA proteins [4]. Ultimately, this allows ICV and IDV to utilize only seven RNA segments rather than the eight RNA segments required for IAV and IBV. Interestingly, the mild illness that ICV causes may be partly due to the fact that folding of the ICV HEF protein in infected host cells is more efficient at the lower temperature of 33 °C that is found in the upper respiratory tract. This temperature sensitivity may limit progression of ICV into the warmer lower respiratory tract [4]. It is interesting that, unlike the HEF protein of ICV, the IDV HEF protein demonstrates surprising temperature and acid stability [32]. In fact, the IDV HEF is the most stable surface protein expressed across all influenza virus types and it may contribute to the ability of IDVs to infect and spread between host species [4]. Thus, the HEF protein presents a unique feature of IDV that sets it apart from the other three types of influenza viruses, and its potential etiological significance should continue to be explored.

### 5. Innate Immunity against Influenza

In the human host, IAVs continue to cause significant morbidity and mortality each year, while IDVs have not been associated with clinical illness. Long before antibodies that are highly specific for a given pathogen can be produced, many early signals of the innate immune response serve as the first line of defense against infection [33–35]. Activation of this highly specialized attack system with multiple levels of protection, communication, and functionality involves two critical antiviral pathways that are activated within hours of a viral infection: the interferon regulatory factor (IRF) pathway and the nuclear factor kappa-light-chain-enhancer of activated B cells (NFκB) pathway [33,34]. The IRF and NFκB pathways induce cytokine expression from host cells, and these cytokines act as key com-
municators of the immune response [34–36]. The most important agents produced during viral infections are the cytokines known as interferons (IFNs), which induce hundreds of IFN-stimulated genes (ISGs) and effectively initiate the antiviral state within infected tissues. Through induction of ISGs and additional cytokines, IFNs elicit specific functions including: (1) telling neighboring infected cells to undergo apoptosis [37,38], (2) altering receptor expression on the surface of infected cells to increase recognition and destruction by surrounding immune cells [37,38], (3) signaling neighboring uninfected cells to destroy RNA and reduce protein synthesis to limit spread of a viral infection [36,38], and (4) increasing the recruitment and activation of innate immune response cells to more effectively eliminate the viral infection [37,39,40]. Activation and secretion of IFN by infected cells can be visualized in Figure 3, using infected lung epithelial cells as an example.

**Figure 3.** Key players in the innate immune response. Human lung epithelial cells are shown as an example for infected host cells. The innate immune response detects a viral infection at multiple points in the infection cycle through pathogen recognition receptors. These receptors activate the IRF and NFκB pathways, leading to transcription factors that will enter the nucleus and induce the expression of antiviral interferons (IFNs). These IFNs will be secreted from the infected cell and signal through IFN receptors to induce expression of IFN-stimulated genes (ISGs), establishing an antiviral state within the host to limit virus replication and spread [39,41]. (Author dissertation. Labels “nucleus” and “cytoplasm” added from original Dissertation Figure 1.4, ProQuest Document ID 2572559761).

*Influenza Evasion of Host Innate Immunity*

Due to the effectiveness of innate immunity through signals such as IFN, influenza viruses have evolved to express several proteins that counteract these host responses. Understanding host–pathogen interactions against IDV is critical for understanding why pathogenesis differs between IDV and other types of influenza virus. Protein components of the RNA-dependent RNA polymerase of both IAV and IBV, as well as the M1 protein of IAV, have all demonstrated antagonism of IFN response [4,11,42–46]. The IAV accessory proteins PB1-F2 and PA-X each stem from an alternate reading frame or single frameshift of the polymerase protein components of IAVs and have also been linked to anti-immune functions. PB1-F2 is a virulence factor for IAVs [47–51], demonstrating increased cell
death [52], reduced viral clearance [51], and enhancement of viral gene expression [48,53,54] in infected hosts. Regarding PA-X, this protein not only decreases type I IFN expression by preventing innate immune receptor signaling [55], it also facilitates degradation of host mRNAs without affecting vRNAs [56–58]. The most notable and well-studied innate immune antagonist expressed by influenza viruses is the nonstructural 1 (NS1) protein, a multifunctional NS gene product that utilizes specific domains to antagonize numerous components of the host innate immune response [59]. For example, the NS1 protein of influenza A and B viruses lead to global effects within infected cells that block host gene expression, particularly at the level of the host IFN response while simultaneously promoting expression of viral genes, as reviewed in [27,60].

While accessory proteins such as PB1-F2 and PA-X have yet to be described for IDV, the literature has identified two IDV proteins with anti-immune function: matrix 1 (M1) and nonstructural 1 (NS1). As with other influenza virus types, the M1 protein of IDV is encoded on the matrix (M) gene. Recently, the M1 protein of IDV was found to negatively regulate type I IFN signaling using a similar mechanism to the M1 protein of IAV [46]. So far, the NS1 protein of IDV has only proved similar to its IAV and IBV analogs through its ability to interfere with innate immunity in vitro, while research suggests this immune response antagonism is lost in vivo [41,61]. Nogales, et al. (2019) demonstrated for the first time that IAVs encoding the NS1 protein of IDV could antagonize IFN-β responses in human lung epithelial cells in vitro similarly to wildtype IAV [61]. However, the IDV NS1 protein-encoding IAVs did not demonstrate the same virulence or viral replication in vivo as wildtype IAV infection. As the NS1 protein of IAV is well-known for being multifunctional (capable of interacting with many different factors of host signaling pathways within infected cells), this work suggests functional differences exist between the IAV NS1 protein and NS1 proteins of other influenza virus types such as IDV [61]. Because IDV is the only type of influenza virus that is not known to cause clinical disease in the human host, it is of particular interest to understand host–pathogen interactions that determine virus infectivity as compared to other types of influenza virus. Additional research exploring the possible anti-immune functions of other IDV genes, accessory proteins of IDV, and the relationship between the NS1 protein of IDV and host immune response pathways will allow us to better understand what may be driving IDV’s lack of virulence in human hosts.

6. Importance of Studying Influenza D Virus

IDV has not been shown to cause clinical signs or symptoms of disease within the human host and there is not as much known about how the immune system responds to this novel influenza virus type [5]. So far, researchers have made a handful of interesting observations that warrant further investigation about IDV, but many questions remain. For example, we still do not know the full host range of IDV and do not know the breadth of disease caused by IDV in these hosts. At the viral gene level, it remains unclear how the increased stability of the IDV HEF protein impacts its ability to spread and persist in infected hosts, and how interactions between the IDV NS1 protein and the host influence innate immune responses. Since IAV and IBV remain the most aggressive influenza viruses in the human host, there has been a lot of research into the host–pathogen interactions induced by these viruses. Using this information to begin understanding host–pathogen interactions with IDV will allow us to determine what causes the lack of illness associated with IDV in the human host. This review identifies some of the complex interactions that exist between host and virus, with emphasis on the critical role of innate immunity. While there is still much to learn about IDV, we can use our knowledge of IAV, IBV, and ICV to guide future research efforts.

Author Contributions: Conceptualization, R.M.S. and V.C.H.; writing—original draft preparation, R.M.S.; writing—review and editing, R.M.S. and V.C.H. All authors have read and agreed to the published version of the manuscript.
Viruses 2022, 14, 1036

Funding: Support was provided by an Institutional Development Award (IDeA) from the National Institute of General Medical Sciences (NIGMS) of the NIH (P20 GM103443, V.C.H.). The content is solely the responsibility of the authors and does not necessarily represent the official views of the NIH.

Institutional Review Board Statement: Not applicable.

Informed Consent Statement: Not applicable.

Data Availability Statement: Not applicable.

Conflicts of Interest: The authors declare no conflict of interest. The funder had no role in the design of the study, writing of the manuscript, or decision to publish.

References

1. Hutchinson, E.C.; Yamauchi, Y. Understanding Influenza. Methods Mol. Biol. 2018, 1836, 1–21. [CrossRef] [PubMed]
2. Krammer, F.; Smith, G.J.D.; Fouchier, R.A.M.; Peiris, M.; Kedzierska, K.; Doherty, P.C.; Palese, P.; Shaw, M.L.; Treanor, J.; Webster, R.G.; et al. Influenza. Nat. Rev. Dis. Primers 2018, 4, 3. [CrossRef] [PubMed]
3. Ghebrerehewet, S.; MacPherson, P.; Ho, A. Influenza. BMJ 2016, 355, i6258. [CrossRef] [PubMed]
4. Wolff, T.; Veit, M. Influenza B, C and D Viruses (Orthomyxoviridae). In Encyclopedia of Virology, 4th ed.; Elsevier: Cambridge, MA, USA, 2021; Volume 2.
5. Liu, R.; Sheng, Z.; Huang, C.; Wang, D.; Li, F. Influenza D virus. Curr. Opin. Virol. 2020, 44, 154–161. [CrossRef] [PubMed]
6. Hause, B.M.; Ducatez, M.; Collin, E.A.; Ran, Z.; Liu, R.; Sheng, Z.; Armien, A.; Kaplan, B.; Chakravarty, S.; Hoppe, A.D.; et al. Isolation of a novel swine influenza virus from Oklahoma in 2011 which is distantly related to human influenza C viruses. PLoS Pathog. 2013, 9, e1003176. [CrossRef]
7. White, S.K.; Ma, W.; McDaniel, C.J.; Gray, G.C.; Lednicky, J.A. Serologic evidence of exposure to influenza D virus among persons with occupational contact with cattle. J. Clin. Virol. 2016, 81, 31–33. [CrossRef]
8. Trombetta, C.M.; Marchi, S.; Manini, I.; Kistner, O.; Li, F.; Piu, P.; Manenti, A.; Biuso, F.; Sreenivasan, C.; Druce, J.; et al. Influenza D Virus: Serological Evidence in the Italian Population from 2005 to 2017. Viruses 2019, 12, 30. [CrossRef]
9. National Center for Immunization and Respiratory Disease CDC. Estimated Influenza Illnesses, Medical Visits, Hospitalizations, and Deaths in the United States—2018–2019 Influenza Season. Available online: https://www.cdc.gov/flu/about/burden/2018-2019.html (accessed on 22 June 2021).
10. Taubenberger, J.K.; Morens, D.M. The pathology of influenza virus infections. Annu. Rev. Pathol. 2008, 3, 499–522. [CrossRef]
11. Bouvier, N.M.; Palese, P. The Biology of Influenza Viruses. Lancet 2017, 390, 697–708. [CrossRef]
12. Catharine Paules, K.S. Influenza. Vaccine 2008, 26, D49–D53. [CrossRef]
13. Sellers, S.A.; Hagan, R.S.; Hayden, F.G.; Fischer, W.A., 2nd. The hidden burden of influenza: A review of the extra-pulmonary complications of infection. Influenza Other Respir. Viruses 2017, 11, 372–393. [CrossRef] [PubMed]
14. Troeger, C.E.; Blacker, B.F.; Khalil, I.A.; Zimsen, S.R.M.; Albertson, S.B.; Abate, D.; Abdela, J.; Adhikari, T.B.; Aghayan, S.A.; Agrawal, S.; et al. Mortality, morbidity, and hospitalisations due to influenza lower respiratory tract infections. The Global Burden of Disease Study 2017. Lancet Respir. Med. 2019, 7, 69–89. [CrossRef]
15. Clayville, L.R. Influenza update: A review of currently available vaccines. Pharm. Ther. 2011, 36, 659–684.
16. Kuchipudi, S.V.; Nissly, R.H. Novel Flu Viruses in Bats and Cattle: “Pushing the Envelope” of Influenza Infection. Vet. Sci. 2018, 5, 71. [CrossRef] [PubMed]
17. Saunders-Hastings, P.R.; Krewski, D. Reviewing the History of Pandemic Influenza: Understanding Patterns of Emergence and Transmission. Pathogens 2016, 5, 66. [CrossRef]
18. Matsuzaki, Y.; Katsushima, N.; Nagai, Y.; Shoji, M.; Itagaki, T.; Sakamoto, M.; Kitaoka, S.; Mizuta, K.; Nishimura, H. Clinical features of influenza C virus infection in children. J. Infect. Dis. 2006, 193, 1229–1235. [CrossRef] [PubMed]
19. Sederdahl, B.K.; Williams, J.V. Epidemiology and Clinical Characteristics of Influenza C Virus. Viruses 2020, 12, 89. [CrossRef] [PubMed]
20. Lee, J.; Wang, L.; Palinski, R.; Walsh, T.; He, D.; Li, Y.; Wu, R.; Lang, Y.; Sunwoo, S.Y.; Richt, J.A.; et al. Comparison of Pathogenicity and Transmissibility of Influenza B and D Viruses in Pigs. Viruses 2019, 11, 905. [CrossRef] [PubMed]
21. Hause, B.M.; Collin, E.A.; Liu, R.; Huang, B.; Sheng, Z.; Lu, W.; Wang, D.; Nelson, E.A.; Li, F. Characterization of a novel influenza virus in cattle and Swine: Proposal for a new genus in the Orthomyxoviridae family. MBio 2014, 5, e00301-14. [CrossRef]
22. Oliva, J.; Mettier, J.; Sedano, L.; Delverdier, M.; Bourges-Abella, N.; Hause, B.; Loupiais, J.; Parde, I.; Bleuart, C.; Bordignon, P.J.; et al. Murine Model for the Study of Influenza D Virus. J. Virol. 2020, 94, e01662-19. [CrossRef]
23. Skelton, R.M.; Shepardson, K.M.; Hatton, A.; Wilson, P.T.; Sreenivasan, C.; Yu, J.; Wang, D.; Huber, V.C.; Rynda-Apple, A. Contribution of Host Immune Responses Against Influenza D Virus Infection Toward Secondary Bacterial Infection in a Mouse Model. Viruses 2019, 11, 994. [CrossRef] [PubMed]
24. Sreenivasan, C.; Thomas, M.; Sheng, Z.; Hause, B.M.; Collin, E.A.; Knudsen, D.E.; Pillatzki, A.; Nelson, E.; Wang, D.; Kaushik, R.S.; et al. Replication and Transmission of the Novel Bovine Influenza D Virus in a Guinea Pig Model. J. Virol. 2015, 89, 11990–12001. [CrossRef] [PubMed]
51. Zamarin, D.; Ortigoza, M.B.; Palese, P. Influenza A virus PB1-F2 protein contributes to viral pathogenesis in mice. J. Virol. 2006, 80, 7976–7983. [CrossRef]

52. Chen, W.; Calvo, P.A.; Malide, D.; Gibbs, J.; Schubert, U.; Basta, S.; O’Neill, R.; Schickli, J.; Palese, P.; et al. A novel influenza A virus mitochondrial protein that induces cell death. Nat. Med. 2001, 7, 1306–1312. [CrossRef]

53. Kosik, I.; Krejnísova, I.; Bystricka, M.; Polakova, K.; Russ, G. N-terminal region of the PB1-F2 protein is responsible for increased expression of influenza A viral protein PB1. Acta Virol. 2011, 55, 45–53. [CrossRef]

54. Mazur, I.; Anhlan, D.; Mitzner, D.; Wixler, L.; Schubert, U.; Ludwig, S. The proapoptotic influenza A virus protein PB1-F2 regulates viral polymerase activity by interaction with the PB1 protein. Cell Microbiol. 2008, 10, 1140–1152. [CrossRef]

55. Rigby, R.E.; Wise, H.M.; Smith, N.; Digard, P.; Rehwinkel, J. PA-X antagonises MAVS-dependent accumulation of early type I interferon messenger RNAs during influenza A virus infection. Sci. Rep. 2019, 9, 7216. [CrossRef] [PubMed]

56. Jagger, B.W.; Wise, H.M.; Kash, J.C.; Walters, K.A.; Wills, N.M.; Xiao, Y.L.; Dunfee, R.L.; Schwartzman, L.M.; Ozinsky, A.; Bell, G.L.; et al. An overlapping protein-coding region in influenza A virus segment 3 modulates the host response. Science 2012, 337, 199–204. [CrossRef] [PubMed]

57. Khaperskyy, D.A.; Schmaling, S.; Larkins-Ford, J.; McCormick, C.; Gaglia, M.M. Selective Degradation of Host RNA Polymerase II Transcripts by Influenza A Virus PA-X Host Shutoff Protein. PLoS Pathog. 2016, 12, e1005427. [CrossRef] [PubMed]

58. Khaperskyy, D.A.; McCormick, C. Timing Is Everything: Coordinated Control of Host Shutoff by Influenza A Virus NS1 and PA-X Proteins. J. Virol. 2015, 89, 6528–6531. [CrossRef]

59. Hale, B.G.; Randall, R.E.; Ortin, J.; Jackson, D. The multifunctional NS1 protein of influenza A viruses. J. Gen. Virol. 2008, 89, 2359–2376. [CrossRef] [PubMed]

60. Rosario-Ferreira, N.; Preto, A.J.; Melo, R.; Moreira, L.S.; Brito, R.M.M. The Central Role of Non-Structural Protein 1 (NS1) in Influenza Biology and Infection. Int. J. Mol. Sci. 2020, 21, 1511. [CrossRef]

61. Nogales, A.; Aydillo, T.; Ávila-Pérez, G.; Escalera, A.; Chiem, K.; Cadagan, R.; DeDiego, M.L.; Li, F.; García-Sastre, A.; Martínez-Sobrido, L. Functional Characterization and Direct Comparison of Influenza A, B, C, and D NS1 Proteins in vitro and in vivo. Front. Microbiol. 2019, 10, 2862. [CrossRef]