Review

Virus sensing receptors in cellular infectivity of influenza A virus

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Abstract
An innate immune response is essential to mobilize protective immunity upon the infection of respiratory epithelial cells with influenza A virus (IAV). The response is classified as early (nonspecific effectors), local systematic (effector cells recruitment) and late (antigen to lymphoid organ transport, naive B and T cells recognition, effector cells clonal expansion and differentiation). Virus particles are detected by the host cells as non-self by various sensors that are present on the cell surface, endosomes and cytosol. These sensors are collectively termed as pattern recognition receptors (PRRs). The PRRs distinguish unique molecular signatures known as pathogen-associated molecular pattern, which are present either on the cell surface or within intracellular compartments. PRRs have been classified into five major groups: C-Type Lectin Receptor (CLR), Toll-like receptor (TLR), Nod-like receptor (NLR), Retinoic acid-inducible gene-I-like receptor (RLR), which play a role in innate immunity to IAV infection, and the pyrin and hematopoietic interferon-inducible nuclear (PYHIN) domain protein. Here, we discuss the role of PRRs in cellular infectivity of IAV and highlight the recent progress.

Key words: IAV; influenza A virus; innate immunity; receptors; PRRs.


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Introduction
Influenza virus of the family Orthomyxoviridae causes the flu, which is an infection of the respiratory tract that has been reported over two millennia ago [1]. Influenza viruses are classified into three genera based on the antigenic differences in their nucleoprotein and matrix protein, of which only the influenza A Virus (IAV) is known to cause pandemics. IAV causes influenza in birds and a few other species of mammals like humans and pigs. Its wide host range allows it to cause pandemics like the 1918 Spanish flu. IAV annually infects 3-5 million people by direct contact with infected humans and fomites. The disease also spreads due to inhalation of the aerosols that are expelled by infected humans when they cough or sneeze. It is characterized by an assortment of direct and indirect pathological effects [2]. The decimation of the infected respiratory cells, damage to the respiratory epithelium, and the immunological responses that cause pneumonia and general discomforts like body ache, coughing, sneezing, high fever, sore throat, rhinorrhea, myalgia, and fatigue are considered as direct effects [3]. While the secondary bacterial infection due to tissue damage and the worsening of other diseases like cardiovascular diseases and diabetes are considered as indirect effects [4]. IAV infection claimed an emphatically enormous number of lives during flu seasons of the last century worldwide [5]. It commonly affects children, adolescents, elderly people, and immunocompromised humans. IAV has several subtypes, labelled as per the two proteins on the surface of the viral envelope. These proteins are haemagglutinin (HA), which causes red blood cells to agglutinate, and neuraminidase (NA), which is an enzyme that breaks the glycosidic bonds of neuraminic acid. Both proteins are essential for the entry and exit of virus particles at the plasma membrane of the target cell [6]. Currently, there are 18 known variations of HA and 11 of NA, theoretically allowing 198 different combinations [7]. Famous combinations include H1N1 (1918 Spanish flu, 1977 Russian Flu, 2009 pandemic flu), H3N2 (1968 Hong Kong flu) and H2N2 (1957 Asian flu) [8].
Influenza A virus gene products and pathogenesis

IAV genome consists of eight segments. The segmented nature allows a range of variabilities in antigens that can be associated with the antigenic shift, which takes place when viral recombination occurs in a cell that may be co-infected with two or more viruses. A good example of this is the 2009 flu virus, which was found to be a “quadruple reassortant” virus [9,10]. IAV produced proteins, their function and their respective RNA segments are illustrated in Table 1.

Human infection by influenza virus is achieved through oral or nasal cavities where the viral NA protein helps the virus permeate the mucus that covers the respiratory epithelium by cleaving the sialic acids that are found in the respiratory tract mucous. If not for this NA protein, the virion would be trapped in the mucous, preventing it from reaching the host cell [11,12]. After successfully penetrating the mucous layer, the virus attaches to and invades the respiratory epithelial cells. The attachment is mediated by the HA protein, which is the primary IAV antigen to which the host’s immune system responds by inducing a neutralizing antibody response. The HA protein is a trimeric glycoprotein on the virus surface, with an inherent specificity for α-2,6-linked sialic acid on host cell surfaces [13,14]. Once replication in these cells becomes productive, the virus spreads to both non-immune and immune cells in the respiratory tract [15].

Years of co-evolution have helped the virus develop strategies to evade the host immune system. Viral matrix protein, M2, is known to play a role in autophagy by blocking auto-phagosome maturation, preventing fusion with the lysosome and significantly affecting host cell apoptosis [16]. The virus is also known to prevent the activation of interferon-inducible dsRNA-dependent protein kinase by binding to dsRNA using the non-structural protein 1 (NS1) [17,18]. The NS1 protein is also known to inhibit NFκB by physically interacting with the C-terminal effector domain of IκB kinase (IKK) [19]. NS1 has been shown to affect the phosphorylation function in both the cytoplasm and nucleus of IKKa/IKKβ. It stops the translocation of NFκB into the nucleus by blocking IKKβ-mediated phosphorylation and degradation of IκBα in the classical pathway of NF-κB activating signalling. It suppresses IKKα-mediated processing of p100 to p52 in the alternative pathway [19]. Another viral protein, PB1-F2, has been shown to have pleiotropic effects [20]. The protein was reported to cause apoptosis in immune cells, thus affecting efficient viral clearance [21]. This is achieved by a PB1-F2 mediated bridging of mitochondrial apoptotic mediators ANT3 and VDAC1, which mediates mitochondrial permeabilization, followed by a cytochrome C release, leading to apoptosis [22]. PB1-F2 increases the chances for the development of secondary bacterial pneumonia, which is a serious life-threatening complication, causing severe lung damage, high inflammatory response, hypertyrosinemia and elevated bacterial loads [20,23,24]. This may be achieved by PB1-F2 interacting with IKKβ, which is known to suppress IFNγ signalling in macrophages by inhibiting STAT1 activation [25]. It has also been shown that PB1-F2 inhibits NFκB binding to the DNA. However, the ability of PB1-F2 to modulate NFκB signalling pathway does not correlate to its ability to interact with IKKβ, suggesting that it may recruit other proteins to interact with IKKβ. Hence, all cytokines expressed under the control of NFκB are potentially susceptible to modulation through PB1-F2 [26].

Table 1. Influenza proteins and their functions.

<table>
<thead>
<tr>
<th>Protein</th>
<th>RNA Segment</th>
<th>Function</th>
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<tbody>
<tr>
<td>RNA polymerase subunit (PB2)</td>
<td>1</td>
<td>RNA-dependent RNA polymerase</td>
</tr>
<tr>
<td>RNA polymerase subunit (PB1) or PB1-F2 protein</td>
<td>2</td>
<td>RNA-dependent RNA polymerase</td>
</tr>
<tr>
<td>RNA polymerase subunit (PA)</td>
<td>3</td>
<td>PA is involved in the assembly of functional viral RNA polymerase complexes from their active intermediates</td>
</tr>
<tr>
<td>Hemagglutinin (HA = HA1 and HA2)</td>
<td>4</td>
<td>HA1 mediates binding to sialic acid of target cells by the virion. HA2 mediates entry of the viral genome into the target cell by acting as the fusion peptide.</td>
</tr>
<tr>
<td>Nucleoprotein (NP)</td>
<td>5</td>
<td>Encapsulates RNA and provides species specificity.</td>
</tr>
<tr>
<td>Neuraminidase (NA)</td>
<td>6</td>
<td>Remove sialic acid from glycoproteins. Prevents trapping in mucus on entry and aggregation on cells during exit. M1 mediates the encapsidation of RNA-nucleoprotein cores into the membrane envelope and assists in virion assembly and egress. M2 facilitates budding of mature virions.</td>
</tr>
<tr>
<td>Two matrix proteins (M = M1 and M2)</td>
<td>7</td>
<td>NS1 suppresses interferon response. NEP mediates export of vRNP complexes from nucleus.</td>
</tr>
<tr>
<td>Non-structural proteins (NS1 and NEP)</td>
<td>8</td>
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Viruses exhibit certain pathogen-associated molecular patterns (PAMPs), like double-stranded RNA, that are usually not found in the hosts [27]. These PAMPs are then recognized by certain receptors known as pattern recognition receptors (PRRs) [28]. These PRRs are subdivided according to their ligand specificity, function, localization, and/or evolutionary relationships. The main receptor to recognize the single-stranded RNA and transcriptional intermediates in the host cell is the retinoic acid-inducible gene-I protein (RIG-I). Toll-like receptors (TLRs) are another group of PRR that detect pathogens near the cells and in endosomes and lysosomes. Other PRRs like nucleotide oligomerization and binding domain (NOD)-like receptor, have been known to activate during an IAV infection [29]. These PRRs trigger a rapid, nonspecific innate immune response, which mostly culminates in the elimination of the virus from the host. In this article, we will focus mainly on the role of virus-sensing receptors in cellular infectivity of IAV.

**Virus sensing receptors**

Five major groups of PRR have currently been identified, namely retinoic acid-inducible gene-I (RIG-I)-like receptors (RLRs), C-type lectin receptors (CLRs), Toll-like receptors (TLRs), NOD-like receptors (NLRs) and the PYHIN domain containing family, including AIM2 [30]. The first four group are discussed below.

**RIG-I-like receptors (RLRs)**

RLRs consist of three members that have been identified in humans and mice, the retinoic acid-inducible gene-I (RIG-I), melanoma differentiation-associated gene 5 (MDA5), and laboratory of genetics and physiology 2 (LGP2) [31]. RLRs are exclusively cytosolic PRRs. They play an important role as RNA-sensing PRRs belonging to the RNA helicase family [32]. Studies have demonstrated that RIG-I enters antiviral stress granules where viral RNA and interferon-stimulated gene (ISG) products like serine/threonine-specific protein kinase co-localize [33]. RIG-I recognizes virus-associated RNA species, like 5′-triphosphate single-stranded RNA and its transcriptional intermediates, with its helicase domain [34]. It is expressed in a variety of cells like macrophages, fibroblasts, myeloid dendritic cells, and epithelial cells [15,35]. In addition to the RNA helicase domain, RIG-I contains two N-terminal caspase activation and recruitment domains (CARDs) [36]. On recognition of viral RNA, RIG-I exposes its CARDs, which are controlled by dephosphorylation or ubiquitination by E3 ligases. On activation, RIG-I interacts with the adaptor molecule mitochondrial antiviral signalling (MAVS) on the outer membrane of the mitochondria (Figure 1). This promoter leads to phosphorylation of interferon regulatory factor IRF3 and activates nuclear factor kappa-light-chain-enhancer of activated B cells NF-κB; thus, helps in inducing type I interferons, pro-inflammatory cytokines, and chemokines [37]. RIG-I has also been proposed as a molecular adjuvant for the flu vaccine as it was shown to reduce the antigen requirement by 10- to 100-fold [38]. Studies have also established that the individual deletion of either RIG-I or MAVS coding genes did not change survival or innate responses in vivo after IAV infection in mice, but caused an increased risk of infection [39,40].

**C-Type Lectin Receptors (CLRs)**

Dendritic cell-specific intercellular adhesion molecule-3-grabbing nonintegrin (DC-SIGN) is a C-type lectin, mainly present at the surface of dendritic cells (DC) [41]. Studies have shown that the lectin-mediated recognition of mannose-rich glycans on IAV...
facilitated the infection of the cells by endocytosis in a sialic acid-independent manner [42]. DC-SIGN also acts as an alternative receptor for IAV in addition to sialic acid and the extent of glycosylation of HA mediates the efficiency of DC-SIGN recognition [43]. Besides, DC-SIGN blocking antibodies helps in reducing cellular infectivity [44]. DC-SIGN has a human homologous protein in mouse which is known as SIGN-R1 and is expressed in macrophages of spleen and lymph nodes. This SIGN-R1 is also reported to interact with IAV strain A/PR/8/34(PR8) by DCs of lymph node both in vitro and in vivo, thus making this receptor essential for humoral immunity [45]. Additional CLR, such as MGL and MMR, also interact with IAV and are salicylic acid (SA)-independent but correlate with the different HA glycosylation levels of a few IAV strains [46]. Figure 2 sheds light on some of these interactions.

**Toll-Like Receptors (TLRs)**

Among the PRRs, the Toll receptor in drosophila was first shown to be necessary for protection against fungal infection [47]. Subsequently, the ten mammalian homologues of the TLRs were described for humans [48]. TLRs are important key sensors belonging to PAMPs. TLR3, TLR7, TLR8, and TLR9 recognize nucleic acid of intracellular or endosomal origin and TLR1, TLR2, TLR4 and TLR6 which are present on cell surface detect the envelope glycoproteins and lipoproteins. These intracellular TLRs appear to be sensors of foreign nucleic acids and trigger an antiviral innate immune response by producing type 1 interferon and inflammatory cytokines [49]. TLRs, in cooperation with other PRRs such as NLRs and RLRs, induce innate immunity against IAV [50]. Figure 3 shows that the binding of TLRs activates the transcription factor, interferon regulatory factor (IRF3), followed by activation of type 1 interferon.

**Figure 2. CLR signalling overview.**

**Figure 3. TLR signalling overview.**

Macrophage galactose-type lectin (MGL), macrophage-mannose receptor (MMR) and dendritic cell specific intercellular adhesion molecule 3 grabbing non integrin (DC-SIGN) augment activation of NF-kB with P50 and P65 subunits. The activation of NF-kB plays a critical role in the induction of innate immune and inflammatory responses as it leads to subsequent secretion of proinflammatory cytokines.
replication of IAV, dsRNA is not generated due to the action of RNA helicase DDX39B. Consequently, it is important to identify the ligand for TLR3 within the influenza-infected cells [15]. Intranasal treatment of mice with poly I:C, a TLR3 agonist, provided a high level of protection against lethal challenge with a highly pathogenic avian influenza virus (H5N1) and showed enhancement of inflammatory reaction and CD8+ T cell response in viral clearance. This suggests that TLR3-triggered innate response inhibits viral spread [53,54].

**TLR7 and TLR8**

Both TLR7 and TLR8 recognize the viral genome of various viruses such as influenza viruses, flaviviruses, coxsackie B virus, HIV-1, and vesicular stomatitis virus. TLR8 is mostly expressed in monocytes, macrophages, and myeloid DCs whereas TLR7 is primarily expressed in DCs and, to some extent, in B cells, monocytes, and macrophages. TLR8 and, to a lesser extent, TLR7 recognizes the GU-rich and AU-rich ssRNA sequences of influenza viruses [55]. Once the replication begins, the viral RNA is released into endosomes where TLR7 and TLR8 reside. Interaction of ssRNA and TLR7/TLR8 triggers the recruitment of a protein involved in signaling within immune cells called myeloid differentiation primary response 88 protein (MyD88), which leads to the activation of NF-κB and other transcription factors and the production of proinflammatory cytokines and chemokines [56].

**TLR4**

TLR4 is usually expressed by myeloid cells and weakly expressed on the surface of epithelial cells. Although the ligand for TLR4 is not known in the case of IAV, the DDX21-TRIF-S100A9-TLR4-MyD88 signalling network is triggered by the DAMP molecule, S100A9, regulating inflammation during infection [57]. This molecule is released in IAV-infected lungs and helps in inducing exaggerating proinflammatory response, cell-death, and virus pathogenesis following lethal infection [54]. Additional studies showed that the treatment of mice with the TLR4 antagonist, Eritoran, was found to protect from lethal IAV infection by alleviating lung pathology, clinical symptoms, cytokine, and oxidized phospholipid expression, as well as by controlling viral loads [58].

**Nucleotide-binding oligomerization domain-like receptors (NLRs)**

NLRs are intracellular sensors of PAMPs that form a multiprotein inflammasome complex consisting of NLRP, the adaptor ASC, and procaspase. They also help in triggering several signalling pathways including MAPK, NF-κB, and MAVS-IRF3, which help in the production of IL-6, TNF-α, pro-IL-1β/IL-18, and also type I interferons [59]. NACHT, LRR and PYD domains-containing protein 3 (NLRP3) are known to be activated by membrane damage, inflammation or stress such as infection with pathogens [60]. These receptors are known to cooperate with TLRs to regulate inflammatory and apoptotic responses. Studies have shown that NLRP3, induces IAV in two steps, namely enhancing the transcription of genes encoding pro-IL-1β and pro-IL-18 and NLRP3 (signal 1) and activating NLRP3 inflammasome (signal 2) as seen in Figure 4 [61]. The ssRNA of the virus, the proton flux by viral M2 in the trans-Golgi network and the aggregation of PB1-F2 in the lysosome of lipopolysaccharide primed macrophages are known activators of NLRP3 inflammasome [58,62,63]. The NLRP3 inflammasomes complex was found essential for reducing the viral load in the later stages of infection, suggesting that NLRP3-inflammasome induced inflammatory response and mediates the protective immunity to IAV infection, possibly via adaptive immune responses [64]. NLR2 (or NOD2) is believed to recognize the viral genomic ssRNA which assists the recruitment of MAVS adaptor protein in activating IRF3-mediated type I interferon production in DCs in response to IAV. NOD2 and NLRX1 are also found to negatively regulate the NLRP3 and RIG-induced inflammatory response to lethal doses of IAV, respectively, and to control the
imunopathology of IAV [61,64]. These findings suggest that NLRs execute a differential role in response to IAV infection.

Conclusions
Our knowledge of innate immune response and the possibly damaging role of innate immunity in the pathogenesis of influenza remain preliminary. The past ten years have expanded our understanding of innate sensing and response to IAV infection. The innate immunity plays an important role upon infection with IAV in rapid control, particularly natural killer cells (NK) which possess potent antiviral activity [65]. This also helps in initiation of adaptive immunity, particularly the production of HA and NA specific antibodies [66].

The role of PRRs in the recognition and activation of signalling pathways, which ultimately results in a pro-inflammatory innate immune response has been discussed in various studies [30,48,58]. PRRs recognize unique molecular signatures which are non-self and possess evolutionarily conserved structures and trigger innate immune response, including the production of pro-inflammatory cytokines and type 1 interferons. Identification of immunological receptors has helped in the development of the concept of pattern recognition and thus has significantly changed the perception of the innate immune system. PRRs play an important part in cellular injury and in recognizing immunological stress. Perception of various endogenous danger signals has put PRRs in a central position in the regulation of immunity and other aspects of cellular physiology and is regarded as sensors of both microbe-associated molecular pattern and danger associated molecular pattern. In this review, we examined available data on the inflammatory response of PRRs which coordinates proinflammatory cytokines (TNF, IL-1, IL-6), regulates cell death, helps in modifying vascular permeability, and induces the production of acute phase proteins as useful markers of inflammation.

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Authors’ contributions
SA, IS, and MNAA performed literature review and prepared the manuscript. SMA, AAAQ and MNAA edited and provided comments. All authors contributed to the design and preparation of this scientific review.

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