



Avian Influenza Viruses, Inflammation, and CD8⁺ T Cell Immunity

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Avian influenza viruses (AIVs) circulate naturally in wild aquatic birds, infect domestic poultry, and are capable of causing sporadic bird-to-human transmissions. AIVs capable of infecting humans include a highly pathogenic AIV H5N1, first detected in humans in 1997, and a low pathogenic AIV H7N9, reported in humans in 2013. Both H5N1 and H7N9 cause severe influenza disease in humans, manifested by acute respiratory distress syndrome, multi-organ failure, and high mortality rates of 60% and 35%, respectively. Ongoing circulation of H5N1 and H7N9 viruses in wild birds and poultry, and their ability to infect humans emphasizes their epidemic and pandemic potential and poses a public health threat. It is, thus, imperative to understand the host immune responses to the AIVs so we can control severe influenza disease caused by H5N1 or H7N9 and rationally design new immunotherapies and vaccines. This review summarizes our current knowledge on AIV epidemiology, disease symptoms, inflammatory processes underlying the AIV infection in humans, and recent studies on universal pre-existing CD8⁺ T cell immunity to AIVs. Immune responses driving the host recovery from AIV infection in patients hospitalized with severe influenza disease are also discussed.

Keywords: avian influenza virus, CD8⁺ T cells, inflammation, influenza, human, influenza disease

AVIAN INFLUENZA VIRUSES AND THEIR EPIDEMIOLOGY

Avian influenza viruses (AIVs, which belong to type A) circulate in birds as well as avian viral reassortants capable of infecting humans. Although AIV infections of wild birds are mainly asymptomatic, wild birds carry AIVs in their intestines and, thus, shed the virus via feces, saliva, and nasal secretions. The first avian infection was recorded in 1878 in Italy, named a “fowl plague” and subsequently isolated from chickens in 1902. Aquatic birds are considered a natural reservoir of AIVs (1) and, to date, all the AIV subtypes were isolated from aquatic birds, with the exception of H17N10 and H18N11 found in bats (2). Based on the pathogenicity in chickens, AIVs are classified into low pathogenic avian influenza viruses (LPAIVs) and highly pathogenic avian influenza viruses (HPAIVs). LPAIVs, commonly found in wild birds, cause only mild disease in domestic birds, while HPAIVs cause severe influenza symptoms, affect respiratory tract as well as internal organs and lead to a high mortality rate in poultry. So far, influenza H5 and H7 subtypes, originating from non-pathogenic precursor viruses, are the only HPAIVs (3). Primary outbreaks of HPAIV in poultry (turkey and chicken) have been reported 17 times between 1959 and 2000 and 8 times since 1990. They resulted in devastating losses of birds as millions of animals had to be culled to control those AIV outbreaks (4).

Out of all the AIVs, only H5N1, H7N2 (5), H7N3 (6, 7), H7N7 (8, 9), H9N2 (10), H7N9 (11), H5N6 (12), H6N1 (13), H10N7 (14, 15), and H10N8 (16) are known to cause human infections (**Table 1**). Generally, human infections with H7N2, H7N3, H7N7, and H6N1 result in mild clinical symptoms. However, some AIVs, including H5N1 and more recently, H7N9 strains, cause severe disease and high mortality rates in humans. This is mainly due to the absence of pre-existing AIV-specific antibodies (17–20), although both anti-H5 and H7-antibodies can be detected during the course of infection (21, 22). The highly pathogenic H5N1 virus attracted widespread attention in 1997, when a H5N1 outbreak in Hong Kong resulted in 18 human infections and 6 deaths. Subsequent outbreaks of the HPAIV H5N1 subtype occurred in China, Indonesia, Malaysia, Vietnam, and Thailand poultry industries in the early 2004, resulting in 13 human cases with 9 deaths in Vietnam, and 27 cases with 20 deaths in Thailand (23). Since then, multiple reassortant H5 HPAIV strains have spread across Asia and into Europe, the Middle East and Africa, giving rise to multiple subtypes (H5N2, H5N5, H5N6, and H5N8). Humans in close contact with AIV-infected poultry, and on rare occasion with other AIV-infected humans, can become

infected. Globally, the total number of H5N1 cases reached 842 with 447 deaths. Thus, the fatality rate is >50% for H5N1 infections. Although, several studies addressed the calculations of more realistic rates of H5N1 infections, taking into account mild asymptomatic cases and low transmissibility, these are still estimated to be >30% (24). Countries that had the highest number of documented cases of H5N1 include Egypt, Indonesia, Vietnam, Cambodia, and China, where the total numbers ranged from 52 to 344 (WHO/GIP; data obtained on the 23 June 2015). The overall prevalence of H5 viruses increased in China since 2010 (25), with H5-type HPAIV, H5N6 and H5N8 being prevalent among China's domestic waterfowl. Avian H5N6 influenza virus is a low virulent strain in poultry that has caused several outbreaks in chickens in China, Vietnam, and Laos during 2014 and 2015. Two cases of H5N6 were reported recently in humans in China and the clinical manifestations were severe (12, 26).

H7 subtype AIV has caused numerous outbreaks in poultry in Europe and the Americas. In 2003, H7N7 emerged in poultry in the Netherlands, and then spread to Belgium and Germany (43, 44), leading to the culling of over 30 million birds and resulting in >80 human cases, with one fatality (45). In 2013 in Italy,

TABLE 1 | Summary of human infections caused by the avian influenza viruses.

Influenza strain	Year/location	Cases/fatalities	Clinical manifestation	Reference
H5N1	1997/Hong Kong	18/6	Flu-like symptoms, pneumonia	(27)
	2003/Hong Kong	2/1	Pneumonia	(28)
	2003/China	1/1	Pneumonia	(29)
	2003–2015/Vietnam, Thailand, Cambodia, Indonesia, China, Azerbaijan, Djibouti, Egypt, Iraq, Nigeria, Turkey, Laos, Myanmar, Pakistan, Bangladesh, Canada	844/449	Flu-like symptoms, pneumonia	WHO/GIP, data in HQ as of 13 November 2015
H5N6	2014/China	2/1	Severe respiratory symptoms	(12, 26)
H6N1	2013/Taiwan	1/0	Flu-like symptoms	(13)
H7N2	2002/USA	1/0	Flu-like symptoms	(30)
	2003/USA	1/0	Respiratory tract infection	(30)
	2007/UK	1/0	Conjunctivitis	(31)
H7N3	2004/Canada	2/0	Conjunctivitis	(32)
	2006/UK	1/0	Conjunctivitis	(33)
	2012/Mexico	2/0	Conjunctivitis	(34)
H7N7	1980/USA	3/0	Conjunctivitis	(35)
	1995/UK	1/0	Conjunctivitis	(36)
	2003/Netherlands	89/1	Conjunctivitis, pneumonia, flu-like symptoms	(37)
H7N9	2013–2015/China	665/229	Range from mild to severe respiratory symptoms	WHO, CDC
H9N2	1998/China	5/0	Flu-like symptoms	(38)
	1999/Hong Kong	2/0	Flu-like symptoms	(39)
	2003/Hong Kong	1/0	Flu-like symptoms	(40)
	2007/Hong Kong	1/0	Flu-like symptoms	(30)
	2009/Hong Kong	2/0	Flu-like symptoms	(41)
	2011/Bangladesh	1/0	Flu-like symptoms	(42)
H10N7	2004/Egypt	2/0	Flu-like symptoms	(14)
	2010/Australia	2/0	Conjunctivitis, respiratory tract symptoms	(15)
H10N8	2013/China	3/2	Pneumonia	(16)

monitoring of the workers from a recent H7N7 outbreak at a poultry farm revealed three cases of H7N7 human mild infections. H7N3 viruses were reported in poultry in several countries, including Chile and Canada, and a recent 2013 outbreak of HPAIV H7N3 virus in Mexico led to the culling of >20 million birds and two confirmed human mild infections (6). Since March 2013, a novel reassortant A/H7N9 virus crossed host barrier from birds to humans, resulting in 657 human cases with 38% mortality (46). This is striking given that the 1918 Spanish influenza pandemic had a fatality rate of >2.5% and was considered to be extremely severe (47, 48). Although H7N9 virus causes severe human infections with a 98% hospitalization rate, H7N9 is a low pathogenic influenza virus in birds and can be found in live poultry markets across China, but rarely at the poultry farms.

In 2013, H10N8 virus emerged from live poultry markets. It originated from the internal genes of enzootic H9N2 viruses in chickens, and surface genes of H10 and N8 viruses from domestic ducks. It caused three severe infections in humans, leading to two deaths (16). Interestingly, internal genes of both H7N9 and H10N8 originated from H9N2 virus (49), an ongoing reassortant with local chicken H9N2 strains, thought to have a high epidemic or pandemic potential. However, it is not fully understood how these H9N2 chicken viruses facilitated the genesis of the newly emerged H7N9 virus (50). Since 1990s, H9N2 avian influenza viruses became well established in domestic poultry across Asia and Europe (51). Although of low pathogenicity, five human cases of H9N2 infection were reported during 1998 and 2008 [CIDRAP: Avian Influenza (Bird Flu): Implications for Human Disease]. H9N2 AIVs rapidly differentiated giving rise to more than 102 genotypic variants being recognized based on the nomenclature system (52). Notably, H9N2 viruses belonging to genotype 57 (G57) have a selective advantage in chickens, with increased infectivity and ability to escape selective pressure via antigenic shift. G57 AIVs caused the widespread H9N2 outbreaks among chickens during 2010–2013 in China (50). The virus became predominant in H9N2-vaccinated farm chickens, caused widespread outbreaks prior to the emergence of the H7N9 virus, increased reassortment events between H9N2 and other viral subtype, and finally provided all of the internal genes to the novel H7N9 virus. The prevalence and high mutation rate of the H9N2 influenza viruses at the poultry farms, their capacity to provide internal genes to other avian viruses, and adaptation to the human host could provide an important early warning of the emergence of novel reassortants with pandemic potential.

RISK OF INFLUENZA PANDEMICS

Wide distribution of AIVs in poultry, occasional jumping across the host barrier to humans and the continuing evolution raise concerns about AIV's pandemic potential. Nowadays, H9N2 and H5-type AIVs have nearly a global distribution in birds, and H7N9 and H10N8 viruses are widely circulating in live poultry markets in China (25). Occasional human-to-human transmission of H5, H6, H7, H9, and H10 AIV and their continuous reassortment with local chicken AIV strains (especially local H9N2 strains) have a potential of becoming influenza pandemic. H7N9, in particular, appears to be fulfilling the requirements for a potential influenza

pandemic virus: (1) crosses the host barriers and is widespread in live poultry markets; (2) adapts to human host and further reassorts with local H9N2, while being a “low-pathogenicity” virus in chickens; (3) there are no pre-existing neutralizing antibodies (NABs) at the population level. Overall, there is an urgent need to understand the pathogenesis and immunity of AIVs in humans, especially for those strains that can cross the host barrier to humans.

CLINICAL FEATURES OF SEVERE AVIAN INFLUENZA DISEASE

During seasonal influenza epidemics, elderly individuals (65+ years of age) showed higher rates of hospitalization (53). Severe disease from H5N1 infection was observed across different age groups, although higher hospitalization was found in several countries, including Vietnam and Indonesia in children and young adults (54). Conversely, during the first wave of H7N9 infections, more severe disease was observed in males aged >59 years old (55). These men shopped at the live bird markets in China and, therefore, were more likely exposed to H7N9 viruses. Furthermore, during aging both the innate and adaptive immune systems undergo immunosenescence, and results in decreased responsiveness to influenza vaccination and infection (56, 57). Some of the clinical and pathological features of severe A/H7N9 disease induced by AIVs infections include multiple organ failure, leukopenia, lymphopenia, viral dissemination in extra-pulmonary sites, and prolonged viral shedding of H7N9 viral RNA in feces and urine (11, 58–62). Indeed, high viral loads were also associated with death in H5N1-infected patients (63). During the recent H7N9 epidemic, decreased viral load was associated with recovery and outcome (59). Fatalities from avian influenza often result from pneumonia and acute respiratory syndrome (ARS) (64). Most hospitalized patients present with viral pneumonia, and the prevalence of bacterial co-infections is limited (51, 65). However, many hospitalized patients undergo broad-spectrum antibiotic treatment (20, 64), which possibly confound the detection of bacterial co-infections. This is in contrast to the 1918 pandemic during which secondary bacterial pneumonia was the leading cause of mortality (62, 66). Additionally, co-morbidities and underlying medical conditions, including type I/II diabetes and hypertension, were highly prevalent in the hospitalized cases of H7N9 influenza (22, 64). Interestingly, mild influenza disease was observed in young children with documented cases of H5N1 and H7N9 infections (67). Clinically, these children presented with fever, but no ARS-associated symptoms, such as pulmonary edema, were observed (67). However, the study had a small sample size ($n = 2 \leq 4$), and the inflammatory milieu was not addressed. These observations warrant more analyses on the age-associated changes in the human immune system.

High fatality rates in AIV cases raise the question of how the avian viruses interact with the host immune system to cause such severe disease. Indeed, high viral titers and exuberant inflammatory responses contribute to fatal and severe avian influenza clinical disease. The mechanisms driving this in humans are likely to involve both intrinsic viral pathogenicity factors and dysregulation of inflammatory responses. The following sections

of this review focus on the host immune responses to AIVs in humans.

INFLAMMATION DURING AVIAN INFLUENZA INFECTION

Inflammation acts as a double-edged sword during viral infections. While inflammation can promote the recruitment of immune cells, uncontrolled and exacerbated inflammation, as observed in many AIV human cases, is associated with systemic edema and extensive tissue damage. It is well established that human infection with avian viruses, including H7 and H5 subtypes, can lead to “cytokine storms.” Analyses of serum and plasma from hospitalized patients revealed that hypercytokinemia of pro-inflammatory mediators (MCP-1, MIF, IL-6, IL-8, IP-10) and angiogenic/growth factors (HGF, SCF) is a common factor shared by H5N1 and H7N9 infections (63, 68, 69), and can be linked to fatal disease outcomes (68). Furthermore, analyses of bronchoalveolar lavage samples from hospitalized H7N9-infected patients showed elevated levels (by 100- to 1000-fold) of pro-inflammatory modulators, such as IL-6, IL-8, MIP-1 β , or IL-1 β (69). Thus, plasma/serum cytokine analyses give an indication of spillover into the periphery of patients with severe infection, although the concentrations are considerably lower. Additionally, airway epithelial cells infected with H5N1 viruses contribute to this inflammatory milieu (70, 71).

Similar findings were observed in macaques, ferrets, and mice challenged with HPAIV H5N1. Extensive tissue damage and hypercytokinemia are consistently present in H5N1-inoculated animals, although HPAIV-infected mice have greater inflammatory responses in the brain (72–75). Indeed, the clinical observations of hospitalized patients and animal studies to date showed that high levels of pro-inflammatory responses were strongly associated with severe disease from both H5N1 and H7N9 infections. Interestingly, anti-inflammatory molecules, including IL-10, were detected in serum of H5N1-infected patients (63). Furthermore, limited production of anti-viral type I IFN, which in part serves to limit viral replication, was observed in respiratory epithelial cells infected with H5 or H7 subtype *in vitro* (76, 77). This indicates that there may be an attempt by the immune system to modulate inflammation, although this could be too little too late. Murine studies demonstrated that blocking pro-inflammatory cytokines, such as TNF, IL-6, and CCL2, did not improve survival of mice challenged with HPAIV (78), suggesting that multiple mechanisms, in addition to inflammation, are likely to exacerbate pathology. Mechanistically, blocking pro-inflammatory mediators in humans has not yet been performed for AIVs. Further studies examining the use of immunomodulating cytokines and chemokines are warranted if alternative therapies are to be utilized for the treatment of severe influenza disease.

THE INNATE IMMUNE RESPONSES DURING AVIAN INFLUENZA INFECTION

The innate immune response, composed of both cellular and soluble mediators, forms the first line of defense during viral

infections. Indeed, the protective role of innate immunity is clearly demonstrated during HPAI infection of birds whereby, ducks exhibit asymptomatic or mild disease, while chickens become ill. Following H5N1 infection, primary lung cells isolated from chickens showed highly elevated immune and pro-inflammatory responses compared to duck cells, which appeared to be mediated via inhibition of STAT-3 (79). Another study also found that RIG-I is absent in chickens, providing a plausible explanation for their increased susceptibility to influenza viruses as compared with ducks (80).

In humans, immunohistological analyses of lung tissues in five fatal cases of H5N1 showed an influx of neutrophils into the alveolar spaces, while TNF and IL-6 were detected in monocytes/macrophages *in situ* (81). The role of monocyte-derived macrophages (MDMs) and alveolar macrophages in protection versus pathology in avian influenza remains controversial. Robust pro-inflammatory responses, including TNF and IP-10 were detected in culture supernatants in H5N1-infected human MDMs *in vitro* (82–84). This was dependent on the viral isolate and independent of viral replicative capacity in MDMs (82–84). Conversely, alveolar macrophages isolated from human lungs showed limited cytokine secretion after HPAIV infection (85). The latter study demonstrated that the permissiveness to infection was lower in alveolar macrophages than in MDMs (85). These data indicate differential roles of tissue-resident macrophages versus circulating and differentiated monocytes during avian influenza infections. Although, H7N9 viruses isolated from humans can replicate more efficiently in cultured primary alveolar macrophages, they induce lower levels of cytokine production, including TNF and CCL5 as compared to H5N1 viruses (86).

Indeed, during the steady state, tissue macrophages have been shown to provide homeostatic functions, including resolution of inflammation via removal of apoptotic debris and cells (87). Conversely, MDMs have an inflammatory effector profile, suggesting that they have the potential to contribute to inflammation-induced pathology (87, 88). Together, these studies highlight the importance of understanding how microenvironments drive differential levels of inflammation during avian influenza infection.

Natural killer (NK) cells, which comprise up to 10% of the lymphocyte compartment, contribute to immune homeostasis and viral immunity in both cancer and chronic viral infections. Their activation is regulated by the interaction of combinations of activating and inhibitory receptors with self and altered-self ligands, e.g., killer Ig-like receptors, CD16 (Fc γ receptor), NKp46 (89–91). Expanded numbers of NK cells were observed in H7N9-infected hospitalized patients with prolonged hospital stays. Furthermore, in these patients robust IFN γ responses were observed following *ex vivo* H7N9 virus stimulations, indicating preserved effector functionality (22). Additionally, antibody-dependent cell cytotoxic (ADCC) antibody isotypes could be detected in H7N9- and H5N1-infected patients (92). Recent studies have also shown that NK cells from healthy influenza-seropositive donors, can be activated via ADCC, where NK cells robustly upregulate IFN γ production and the surrogate marker of degranulation, CD107a (89). In the absence

of neutralizing H5-antibodies, these responses were also cross-reactive with H5N1 hemagglutinin (84). Together, these studies demonstrate a role of NK cells during avian influenza virus infection and their potential to kill AIV-infected cells and aid in the viral control.

Other innate cellular components include the innate-like T cells, $\gamma\delta$ T cell, NKT, mucosal-associated invariant T cells, which express re-arranged T cell receptors, and also the innatelymphoid cells, which lack re-arranged receptors. Collectively, these innate-like cells produce inflammatory and immune-modulating cytokines (e.g., IFN γ , TNF, IL-22, IL-17, IL-33), which are found in the periphery and can be enriched in peripheral tissues (93–96). Although there is a paucity of data regarding their role in AIV infection, the immunoregulatory roles of ILCs and NKTs were demonstrated in murine models of influenza. Both ILCs and NKTs have been shown to aid in the protection of the lung epithelium during IAV infection via the production of IL-33 and IL-22, respectively (97, 98). An example of cytolytic potential during avian influenza virus infection comes from human V γ 9 δ 2 T cells, where the killing of H5N1-infected MDMs were demonstrated when expanded prior with a TCR-cognate phosphoantigen (99). Together, innate lymphocytes may be utilized as alternative biological therapies for severe avian influenza.

CD8⁺ T CELL IMMUNITY AGAINST AVIAN INFLUENZA VIRUSES

Influenza virus-specific CD8⁺ cytotoxic T lymphocytes (CTLs) provide protection against severe influenza disease, accelerate viral clearance, and promote rapid host recovery (22, 100–102). CTL-based heterosubtypic immunity against seasonal influenza viruses has been extensively studied in animal models (103, 104). In murine models, primary influenza-specific CTLs emerge around day 7, peak around day 10, and then contract via apoptosis. This is in response to the influenza virus replication, which peaks at days 3–5 and gets resolved around days 8–10 (105–107). A few studies were performed to investigate how CD8⁺ T cell responses are induced during HPAIV (e.g., H5N1) infection in mice and whether virus-specific CD8⁺ T cells play a protective or immunopathologic role during a primary HPAIV infection. Hatta et al. showed that different virulent H5N1 viruses had different replication kinetics in mice and that rapid replication could outpace the CD8⁺ T cell response (108). Some studies reported that H7N9 replication in mice peaked at day 5 (109) and cleared around day 8 (110), while others reported that during the primary infection of B6 mice, virus peaked on day 6, decreased by days 8–10, and cleared by day 12 (111). In any case, this appears to differ from human H7N9 disease, where H7N9 viral RNA could still be detected in nasal swabs, blood, urine, or feces at days 8–14 following disease onset. Since the virus replication kinetics and the pathogenicity of HPAIV in mice differ to those found in humans, it is of key importance to understand the influenza viral replication and CTL responses in humans.

Published reports provide evidence that prominent pre-existing influenza-specific CD8⁺ or CD4⁺ memory T cell pools

are associated with more rapid recovery from experimentally mild H3N2 or naturally occurring H1N1-2009 influenza virus infection, as manifested by milder disease symptoms, shorter illness time, diminished viral shedding, and less transmission (100, 101, 112, 113). However, the exact kinetics and the role of CD8⁺ or CD4⁺ T cell responses in the recovery from infection, especially severe AIV infection, is not well understood. Recently, Sridhar et al. showed that the pre-existing IFN- γ -producing memory CD8⁺ T cells directed at the conserved epitopes inversely correlated with disease illness and symptom scores (101). Wilkinson et al. reported that influenza-specific T cell response increased largely at day 7 when virus was completely undetectable in nasal samples, while serum antibodies were present (112). Pre-existing memory CD4⁺ T cells, capable of killing virus-infected cells responded to influenza internal proteins and inversely correlated with less virus shedding and lower disease scores. The above-mentioned studies used the household cohort approach during the 2009/2010 pH1N1 infection season (101) and the human influenza challenge with experimentally low dose of seasonal H3N2 and H1N1 infections (112), with viral clearance around days 6–8 after influenza infection (114). This might, however, differ to the viral replication and the resultant cellular immune responses following natural severe AIV infection.

The kinetics of CD8⁺ T cell responses in human influenza infection has been addressed by Rimmelzwaan's laboratory. Hillaire et al. reported that virus-specific CD8⁺ T cells peaked within 1 week after influenza virus infection and then contracted rapidly (115), consistent with the observations in mice (116). However, Ennis et al. found that virus-specific CD8⁺ T cells expanded around days 6–14 post-infection, and then declined around day 21 after infection (117), whereas Wagar et al. showed that total IFN γ ⁺CD8⁺ T cells peak at around 3 weeks after the disease onset in case of pH1N1 infection. Interestingly, the CD8⁺ T cell number moderately decreased around day 78 and reached the baseline level at day 700 post-disease onset (118), consistent with Ennis' report. As mentioned previously, the new AIV H7N9 could replicate actively between days 8 and 14 (59), which differs greatly from the mild infection in humans reported for pandemic H1N1 (101) and experimental H3N2 infection (112). This raises the following questions: (1) whether the kinetics of CD8⁺/CD4⁺ responses during AIV infection are similar to those observed during a mild influenza disease; (2) whether CD8⁺ T cell responses are mainly protective or could also be immunopathogenic; (3) whether the expanded CD8⁺ T cell responses are derived predominantly from the memory pools or possibly recruited from naïve precursors; and (4) what is the relationship between CD8⁺ T cells, CD4⁺ T cells, and humoral immunity.

The reports that looked at human CD8⁺ T cell responses elicited to HPAIV are scarce and focused mainly on pre-existing and cross-protective immunity (Table 2). Early reports indicated that memory CD8⁺ (and CD4⁺) T cells from healthy individuals with no documented history of prior exposure to avian influenza could recognize epitopes from avian-derived viruses, including 1997 Hong Kong H5N1 (119). The cross-reactive nature of influenza-specific CD8⁺ T cells was further

TABLE 2 | Relevant publications assessing human T cell immunity toward avian H5N1 and H7N9 influenza viruses.

Reference	Avian influenza subtype	T cells	Responses toward avian viruses
Jameson et al. (119)	H5N1	CTLs reactive to H1N1, H2N2, and H3N2, obtained from healthy individuals from USA	Cross-reactivity with H5N1
Lee et al. (122)	H5N1	CTLs reactive to H3N2 obtained from healthy individuals from Vietnam and UK	Cross-reactivity with H5N1
Goy et al. (120)	H5N1	CTLs reactive to H1N1 and H3N2 obtained from healthy individuals from Australia	Cross-reactivity with H5N1
Krejtz et al. (121)	H5N1	<i>In vitro</i> expanded CTLs toward H3N2 obtained from healthy HLA-typed donors from the Netherlands	Cross-reactivity with H5N1-infected BLCL
van de Sandt et al. (123)	H7N9	CTLs expanded <i>in vitro</i> toward H1N1 (seasonal and pandemic) or H3N2 (seasonal) from healthy HLA-typed donors from the Netherlands	CTL cross-reactivity with H7N9-peptide loaded BLCLs
Quinones-Parra et al. (124)	H7N9	<i>In vitro</i> peptide-expanded PBMCs obtained from healthy HLA-typed individuals from Australia	CTL cross-reactivity with H7N9-derived immunogenic peptides is restricted by certain HLA haplotypes and varies across ethnicities
Chen et al. (20)	H7N9	PBMCs obtained from H7N9-infected patients in China	High numbers of peripheral blood T lymphocytes (CD4 ⁺ and CD8 ⁺) correlate with better clinical outcomes
Wang et al. (22)	H7N9	<i>Ex vivo</i> longitudinal analyses of PBMCs obtained from hospitalized H7N9-infected patients in Shanghai, China	Rapid recovery from H7N9 infection is associated with early CD8 ⁺ T cell responses

demonstrated by several groups (120–122), which reported heterosubtypic memory T cell responses against avian H5N1 in healthy individuals without prior exposure to AIV. This heterosubtypic immunity was proposed to provide at least some level of protection toward the new emerging, possibly pandemic, influenza AIV strains. Nevertheless, there are no reports addressing the role of CD8⁺ T cells in individuals infected with primary H5N1 virus in humans. Similarly, it has been reported that a high percentage of CD8⁺ T cells specific for seasonal (H1N1 and H3N2) and pandemic (H1N1) viruses cross-react with the newly emerged H7N9 AIV and suggested that those cross-reactive CD8⁺ T cells can reduce disease severity and virus spread within the population (123). However, the population coverage of these broadly cross-reactive pre-existing CTLs varied greatly across various ethnicities, indicating that especially indigenous populations are at the higher risk of succumbing to H7N9 infection (124). Our recent study showed the importance of CD8⁺ T cell responses against H7N9 AIV in the absence of pre-existing NAb (22). We showed that H7N9-infected patients discharged from the Shanghai Public Health Clinical Centre within 2–3 weeks following admission had early prominent H7N9-specific CD8⁺ T cell responses. On the other hand, individuals with prolonged hospitalization showed late recruitment of CD8⁺/CD4⁺ T cells and antibodies simultaneously (recovery by week 4), augmented by prominent NK cell responses (recovery >30 days). By contrast, those with fatal disease outcomes had minimal influenza-specific immunity and little evidence of T cell activation at the transcriptional level (69).

However, all of the above human studies are based on the assessment of PBMCs within the peripheral blood, which might not totally reflect what happens at the site of infection. Although the previous reports from mice studies (125, 126) showed that

blood CD8⁺ T cell response could be a surrogate for the CD8⁺ T cell response in the lung and lymph nodes, there are still no published studies in humans that investigate whether similar kinetics occur within the blood and lungs during human influenza infection. In addition, it is difficult to know how many lung resident memory T cells are present following the natural influenza virus infection and how protective they are against the heterologous challenge in humans.

CONCLUSION

Taken together, it seems that the pathogenicity and virulence of AIVs in humans are determined by the interactions between the virus and the host immune response. For mild infections, whether caused by seasonal influenza viruses or occasional asymptomatic AIVs, the pre-existing CD8⁺ and/or CD4⁺ T cells can provide a great level of protection (101, 112, 115). In this situation, innate immunity provides a rapid first-line immune defense, slows down the virus, and assists development of adaptive immunity, without any damaging “cytokine storm” effect. This is then followed by adaptive immunity capable of clearing the viral infection and promoting host recovery. However, in case of a severe human influenza infection, caused by either seasonal influenza viruses or HPAIV, due to as yet unknown reasons, infected people are more susceptible to severe outcome. High replication of AIV may either inhibit the innate immunity or drive the overactive innate immune response. This may then interfere with generation of rapid and effective adaptive immunity and, thus, subsequently result in detrimental cytokine storm. Defective innate and/or adaptive immune mechanisms cannot control the infection in a timely fashion, as shown in our H7N9 study (22). Delayed CD8⁺ T cell responses in patients with prolonged hospital stays could have also resulted from the lack of robust cross-strain protective

memory pools, thereby reflected the need to recruit the naïve CD8⁺ T cell precursors into the primary influenza response rather than rapidly recalling the available memory CD8⁺ T cells. Thus, establishment of robust memory pools appears to be of a great importance for the rapid recruitment of adaptive immune responses after HPAIV, so that the innate immunity (and inflammation) does not get to the point that is immunosuppressive for cellular immunity and, thus, cannot control the rapidly replicating virus (22).

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AUTHOR CONTRIBUTIONS

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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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