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Sialic acid receptors: focus on their role in influenza infection

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Abstract: Receptor-binding specificity is a major determinant of host range, pathogenesis, and interspecies transmission in influenza virus. Recognition of different conformations of sialic acid by the influenza virus hemagglutinin has driven influenza intra- and interspecies transmission for nearly 100 years since the first human pandemic was recorded in 1918. Key amino acid residues near or at the receptor-binding site of the hemagglutinin of different influenza A subtypes correlate with the ability of the virus to infect and replicate in the upper or lower respiratory tract of the host and transmit efficiently. A global concern exists for those highly pathogenic avian viruses that have emerged with the potential to cause a novel pandemic in humans. Studying the receptor-binding affinity to sialic acid receptors as a marker of pandemic risk could provide reliable measures to aid in timely pandemic planning and surveillance worldwide.

Keywords: influenza A, sialic acid, receptor binding, host range

Introduction

The cell entry of influenza virus is governed by receptor biology. Sialic acids (SAs) of cell surface glycoproteins and glycolipids are the receptors for the influenza virus, recognized by the viral glycoprotein hemagglutinin (HA; Figure 1). The site on HA at which the cellular receptors are bound is at the distal end of the molecule.^{1,2} The receptor-binding site (RBS) is recognized as a pocket of highly conserved residues in all subtypes of influenza viruses; mutations in the RBS are required for adaptation of the animal viruses HA to the human host.³

Two major linkages between SA and the penultimate galactose residues of carbohydrate side chains are found in nature, Neu5Ac $\alpha(2,3)$ -Gal and Neu5Ac $\alpha(2,6)$ -Gal.⁴ Most avian and equine influenza viruses preferentially bind to SA joined to the sugar chain through an $\alpha 2-3$ linkage, and viruses isolated from birds possess HAs with high affinity for this type of sugar, whereas human influenza A viruses (IAV) preferentially use $\alpha 2-6$ linked SA as a cellular receptor.⁵

The affinity of the influenza virus HA for respective SA species is one of the factors that determine whether animal viruses can cross the host barrier. This event is facilitated when animal species such as swine or quail function as “mixing vessels”, because they express both types of SA receptors.^{6,7}

Investigating the SA receptor-binding properties of IAV

Influenza virus entry to susceptible cells is governed by the interaction between the HA surface antigen of the virus and the cell receptor SA. The recognition of different

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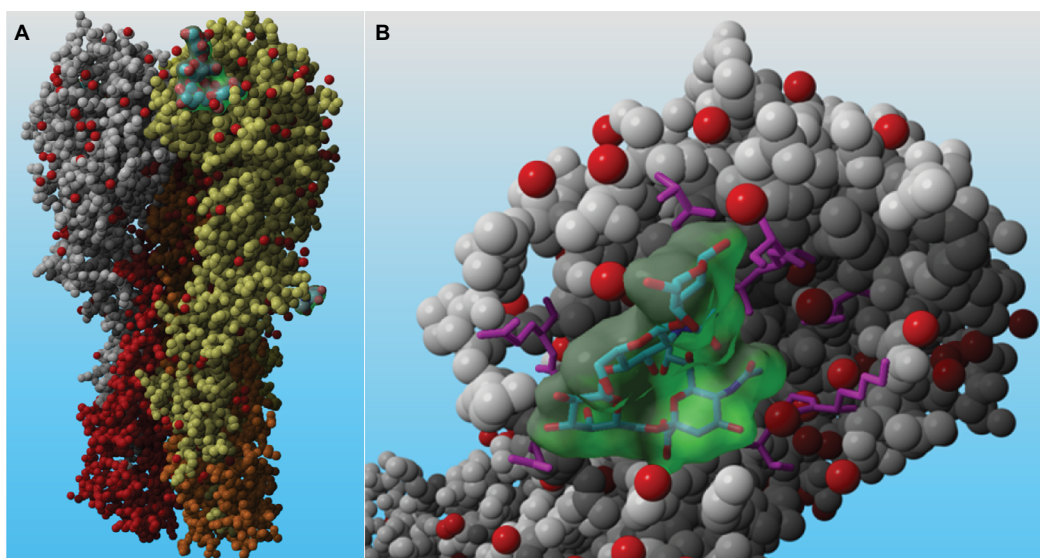


Figure 1 Influenza virus HA molecule from A/California/04/2009 (PDB: 3ube).

Notes: Each monomer in the (A) trimer is indicated in color; the front view shows the HA₁ (green or gray) and HA₂ subunits (red or orange). A closer view of the (B) globular head and the interaction with ligand LSTc; in purple are shown some of the amino acid residues at the RBS that directly interact with the sialic acid-like receptor. Images were generated with Yasara software version 17.1.28.

Abbreviations: HA, hemagglutinin; LSTc, sialylneolacto-*N*-tetraose c; RBS, receptor-binding site; PDB, Protein Data Bank.

types of SA linkages by human and animal viruses is a major barrier for host range restriction. Lessons learned from the old pandemics showed that viruses from 1918, 1957, and 1968 all acquired mutations at the HA RBS during the first months of the pandemic that allowed them to transmit between humans.⁸

A simple way to assess the receptor specificity of influenza viruses can be through the agglutination of erythrocytes from different animal species. Human viruses agglutinate erythrocytes from chicken, ducks, guinea pigs, and sheep, but not those from horses or cows; however, avian and equine viruses agglutinate all these erythrocytes.⁹

“Hemagglutination assays” (Figure 2A) have been widely used in routine diagnostics since they were first described by Hirst in 1941. The property of the influenza virus HA to agglutinate red blood cells (RBCs) from different species reflects diversity in SA species. However, virus evolution and host adaptation affect the recognition of SA species, and the HA assay can be poorly specific and even unreliable for certain strains.

For many years, the interaction of the influenza virus HA with the SA receptor was modeled with trisaccharide or pentasaccharide molecules (e.g., Neu5Ac- α 2,3/ α -2,6Gal β 1-4Glc), where the SA is at the end of the oligosaccharide chain (Figure 2B).¹⁰ The interaction between HAs from human viruses and the receptor analogs varies with subtype. H1 HAs were bound strongly to sialylneolacto-*N*-tetraose c and 6'-sialyl-*N*-acetylglucosamine than to 6'-sialyllactose trisaccharide or 3'-sialyllactose trisaccharide, suggesting an interaction with

the asialo portions of the long analogs, whereas H3 HAs were bound equally strong to 6'-sialyllactose trisaccharide or 6'-sialyl-*N*-acetylglucosamine.^{11,12} Otherwise, the glycans preferred by the avian HA could vary depending on the avian species from which the virus was isolated.

The “glycan array” technology has become a powerful technology to study the binding profile of human and animal glycoproteins (Figure 2C). Numerous plant lectins, glycan-specific antibodies, and microbial toxins used in research have been analyzed for their specificity on glycan arrays. Glycan microarrays are also capable of detecting carbohydrate-specific antibodies in human sera, or the binding of bacteria and eukaryotic cells.¹³ In recent years, massive improvements in glycobiology knowledge and the availability of new technology and resources have shown that influenza viruses recognize different forms of glycopolymers with terminal SA.^{14,15}

Glycan arrays present many different receptor mimics and may contain a more biologically relevant molecule. The extent to which influenza HA proteins discriminate between these different carbohydrates is perhaps surprising. The glycan array analysis is a novel and suitable technique to analyze subtle differences in carbohydrate receptor specificities of known and pandemic human influenza strains as the novel H1N1pdm09 subtype.¹⁶ The glycan array format has been used to compare the binding profiles between the pH1N1 virus and the human seasonal H1N1, H3N2, and the avian origin H5N1, H7N7, or the H9N2.¹⁵

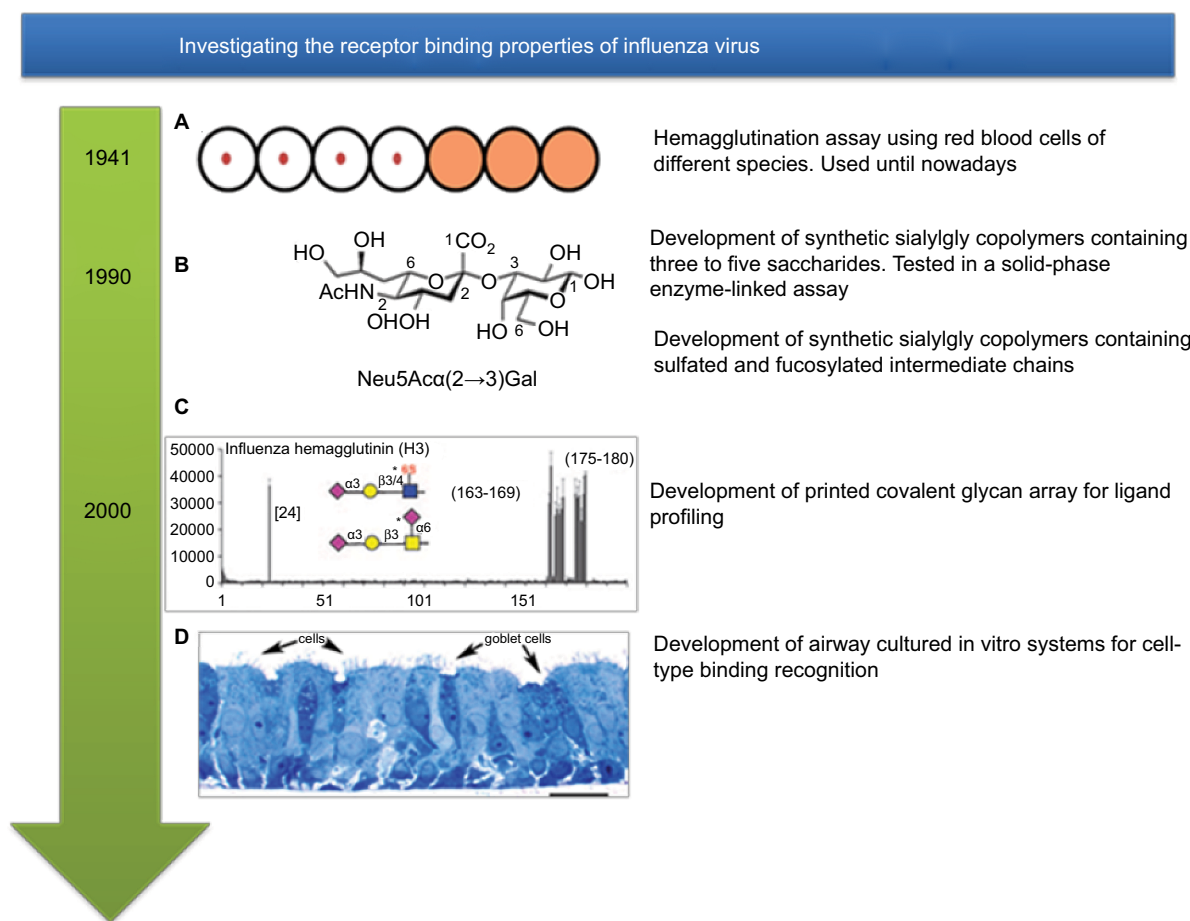


Figure 2 Schematic representation of the methods and techniques used throughout influenza virus history to identify the receptor type recognition.

“Cultured human airway cells and excised human trachea sections” have been used for studying the preferred cell type of infection by the influenza virus (Figure 2D). Human influenza virus infects the upper respiratory tract with major involvement of the pharynx and trachea. Human viruses need to access their target cells as well as to overcome the presence of mucins, which are produced by the goblet cells and are abundantly expressed in trachea.¹⁷ Several groups have shown that cultures of fully differentiated human airway epithelia or human tracheobronchial epithelia are readily infected with influenza viruses, although the cell tropism differs between viruses of avian and human origin in these systems.^{17–19} Accordingly, lectin staining of the cultures showed that some, but not all, ciliated cells expressed SA α 2,3, whereas non-ciliated cells expressed only SA α 2,6. Antigen detection confirmed that only ciliated cells were infected with avian influenza viruses (AIV), whereas non-ciliated cells were infected with human viruses. Although avian viruses replicated, a restricted spread to adjacent cells was observed, because not all ciliated cells expressed the

avian type of receptor, suggesting restrictions to adaptation by a reduced cellular tropism.

Receptor-binding profile of influenza H1N1: from Spanish flu to H1N1pdm09

The IAV H1N1 subtype has co-circulated in humans together with the H3 subtype since 1977, and both have adapted for human replication and transmission during decades of being human pathogens.^{20,21}

For the H1 HA, it is clear that mutations in amino acids 190 and 225 (hereafter considered in H3 numbering) were essential for the adaptation of avian origin H1N1 viruses to replicate and transmit either in swine or in humans. Receptor-binding analysis of the HA variants demonstrated that the 1918 virus with the combination D190 and G225 was bound to both SA α 2,3 and SA α 2,6 molecules, whereas the variant with D190 and D225 behaved like typical human viruses and showed preference for SA α 2,6. A mutated 1918 virus with the combination E190 and G225 (avian-like genotype)

showed preference for the avian type of receptor SA α 2,3.^{15,22} Complementary studies using the preferred animal ferret model suggested that amino acid changes at conserved residues 190 and 225 affected not only the receptor-binding phenotype of the 1918 H1N1 virus, but also most importantly the efficiency of transmission.²³

Influenza H1N1 viruses were introduced to humans and swine around the same time and evolved independently in each host. Analyzing 154 full-length swine H1 HA sequences (classical swine, human-like, and avian-like) shows the predominance at the RBS of the D190 G225 phenotype, with only 25% of the strains presenting the D190 D225 sequence. In contrast, the analysis of 451 full-length human H1 HA sequences shows the predominance of the D190 D225 phenotype, with some strains having the D190 G225 sequence, corresponding to those viruses from the 40s and 50s right at the end of the H1N1 era, or from the 70s when the H1N1 virus was reintroduced to humans (Table 1). Thus, it is generally the case that D225 paired with D190 is required for efficient human-to-human transmission. Indeed, the ferret transmission studies indicated that viruses with this genotype showed 100% transmission, whereas viruses with G225 (Table 1) only transmitted to one in three exposed animals.^{15,22,23}

Differences in receptor usage by avian and human viruses indicate that AIV do not transmit between humans; therefore, this raised the hypothesis of an intermediate host, likely the pig, in the generation of true pandemic viruses.²⁴ This hypothesis is supported by the presence of both SA α 2,3 and SA α 2,6 linkages in pig trachea.²⁵ Viruses of swine origin infecting humans had a dual pattern of binding for the SA α 2,3 and SA α 2,6 glycans.¹⁶ These viruses differ between them by a

residue in position N225 or G. In contrast, seasonal viruses of the H1 subtype showed two differences at residues D225 and 227A→E and affinity only for glycans with SA α 2,6 linkages when analyzed using the glycan array.

The HA gene of the 2009 pandemic influenza H1N1 virus has its origin from the classical swine lineage. After emergence in the human population, classical swine viruses isolated from American pigs between 1945 and 2009 from the α , β , and γ antigenic groups, as well as swine viruses isolated in the same year as H1N1pdm09 were analyzed in search of those amino acid changes at the RBS required for the novel 2009 virus to transmit to humans. Swine viruses showed selective binding to SA α 2,6 sialosides besides the term of circulation and the host from which they were isolated (humans or swine), suggesting that no adaptation steps were required for the transmission of the novel 2009 virus from pigs to humans.^{26,27}

The novel H1N1 virus contained the amino acid sequence D190 D225 at the RBS of the HA. The early pdmH1N1 isolates almost exclusively recognized α 2,6-linked SAs. Strikingly, some human isolates of the pandemic influenza H1N1 virus possess the sequence G225, such as H1 viruses of birds; data from other influenza viruses, such as the highly pathogenic avian H5N1, showed that viruses with the avian-like phenotype at the RBS of the HA may target SA α 2,3 that is present in the human lung (non-ciliated bronchiolar cells), resulting in a severe lower respiratory tract infection in the human host.^{28,29}

Severe cases of H1N1 2009 virus were characterized by viral pneumonitis, indicating a more prominent role of lower respiratory tract infection for this virus than seen

Table 1 Residues at the RBS of the HA protein of H1N1 viruses isolated from humans, avian, or swine

Virus subtype	Amino acid						SA receptor	
	190	222	225	226	227	228	SA α 2,3	SA α 2,6
Subtype H1 (1918)								
Avian	E	K	G	Q	A	G	+++	–
Human	D	K	D	Q	A	G	–	+++
Variant	D	L	G	Q	A	G	++	++
Human seasonal H1N1	D	K	D/G	Q	E	G	+/-	+++
Pandemic H1N1/09	D	K	D	Q	E	G	+++	+++
Human/swine H1N1	D	K	D	Q	A	G	+	+++
Swine H1N1 ^a	D	K	G/D	Q	A/E	G	–	+++
Subtype H7N9				L			+++	–
Subtype H5 (1997–2009)								
Avian	E		G	Q	S	G	+++	–
Human	E		G	Q	S	G	+++	–

Notes: Mutations at the residues 190 and 225 are considered key to the receptor specificity by H1 viruses, for some of them also showed the SA receptor affinity. Avian H5 amino acid sequence at the same residues showed exclusive preference for the SA α 2,3 receptor. For H7N9 AIV, the presence of a human-like residue Q226L does not change the receptor affinity for the avian-like phenotype. ^aViruses included in the analysis of 154 sequences correspond to classical swine, avian-like, and human-like.

Abbreviations: AIV, avian influenza viruses; HA, hemagglutinin; RBS, receptor-binding site; SA, sialic acid.

with recently circulating influenza strains. Investigating the binding properties of H1N1pdm09 virus has offered clues on those specific amino acid changes near or at the RBS associated with the dual target of SA α 2,3 and SA α 2,6 and the severity of influenza disease. The first marker of pathogenicity and severe clinical outcome has been associated to a single point mutation D225G at the RBS of the HA, which has been detected in viruses from several countries around the world.^{30–34} Studies using experimental in vivo models analyzing the pathogenic effect of the mutation D225G are controversial;^{34–37} foremost, the amino acid residue 225 has shown polymorphism for 225D/G/N in the HA of the H1N1pdm09 virus.^{38,39} Overall, viruses with 225G phenotype showed changes in the virus receptor-binding profile and the cell/tissue tropism and dual binding for SA α 2,6 and SA α 2,3. On the contrary, viruses with 225N showed preference for SA α 2,6, though they showed reduced ability to infect human tracheobronchial epithelial cells, an effect that was compensated with a supplementary mutation S185N, expanding their cell tropism, increasing the number of infected cells, and likely, in consequence, the severity of the infection and clinical outcome of the patient.^{34,36,40,41}

The evolution of the H1N1pdm09 virus has been followed in detail worldwide; this has been the basis for the selection of vaccine viruses, as well to monitor changes at the antigenic sites, or changes in HA that result in the selection of viruses with better receptor-binding properties. Eight monophyletic groups have been identified based on the sequence variation of the HA1 domain of the pH1N1 virus, actually with group 6B.1 being the most represented with viruses from around the globe.⁴² The most frequent amino acid substitutions are contained in the head domain of the HA protein of pH1N1. Some mutations such as S206T were acquired soon after the beginning of the pandemic; others, such as A137T, S186P, D100N, S188T, A200T, and S146G, occurred during virus evolution and adaptation, defining the monophyletic groups. These amino acid changes have a direct effect on the receptor-binding properties of the pH1N1 viruses and are suggested to have occurred during the evolution and adaptation of the virus as a result of the antigenic drift and the emergence of viral variants that escape from the immune system.^{43–45}

Seasonal H3N2 viruses and the progression for differential SA α 2,6 linkages

Human influenza viruses of the H3N2 subtype emerged in the human population during the 1968 pandemic and remain in circulation today. Early during the pandemic, viruses isolated

from humans showed acquisition of the mutations Q226L and G228S, which made the virus highly transmissible between humans. The effects of these mutations in viruses of the H3 subtype demonstrate how the avian precursor of the 1968 human pandemic (A/duck/Ukraine/1/1963) exclusively recognized SA α 2,3, whereas a humanized H3 virus represented by the A/Victoria/3/75 strain recognized not only SA α 2,6 but also SA α 2,3. More recent H3 influenza viruses represented by A/Panama/2007/1999 showed a complete switch in its affinity for the SA receptor, with exclusive recognition of SA α 2,6 linkages (Figure 3). The transition between an avian H3 virus toward a humanized H3 virus and the effect of mutations 226 and 228 in HA were also observed in human airway epithelium, where avian viruses possessed a restricted infection of ciliated cells in human airway epithelial cell cultures, whereas more recent human H3 viruses primarily infected non-ciliated cells.¹⁸

Human influenza H3N2 viruses represent the best example of adaptability and evolution in humans. In consequence, the H3 virus has presented along those years high antigenic and genetic variability affecting the vaccine efficacy and the need to update the vaccine composition. Major antigenic changes during H3 evolution occurred in the HA in amino acid positions immediately adjacent to the RBS, substantially changing the virus antibody recognition.^{46,47}

Generally, characterization of human H3N2 viruses for vaccine selection is through the agglutination of chicken RBC.¹⁸ However, antigenic and genetic changes over time have caused a progressive change in the affinity and avidity of H3 viruses for SA receptors. Initial observations showed that viruses isolated between 1997 and 2005 lost affinity for SA receptors recognized in chicken RBCs, and that several substitutions at residues E190D or Q226L→I→V or G225D and W222R, or D225N and S193F accompanied the loss of hemagglutination.^{20,21,48} A comprehensive analysis using an array of glycans clearly identified that H3 evolution during six decades has progressively changed the type of glycan recognized. Early human H3 viruses had preference for short-branched glycans containing NeuAca2,6 structures. A transition to recognize glycans containing longer branches and linear structures containing *N*-acetylactosamine (LacNAc) repeats occurred in viruses from the second and third decades. By the years 1998–2008, viruses had lost affinity for binding short glycans and gained affinity for long linear NeuAca2,6(LacNAc)₃ structures.⁴⁹ A recent expansion of these observations analyzed the evolution of human H3 viruses through different SA α 2,6 conformations. A change in binding from single short glycans to branched *N*- and

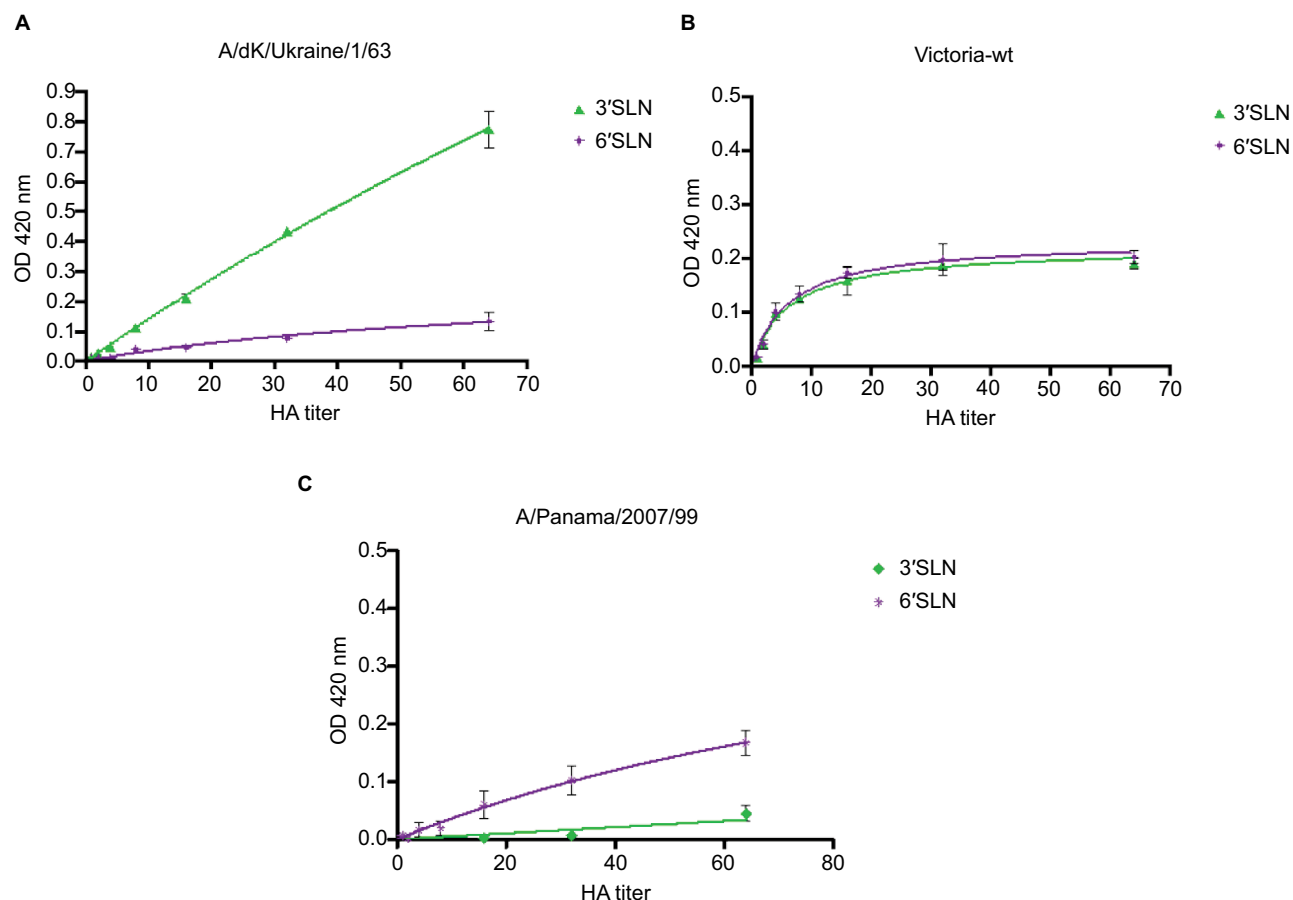


Figure 3 Binding of influenza H3 viruses representing different stages of the subtype evolution prior to and since its emergence in humans.

Notes: Viruses were analyzed by cross-link binding assay using receptor mimics 3'SLN and 6'SLN. Virus binding to polymers was detected with virus subtype-specific antiserum.⁶⁴ **(A)** Binding of an AIV (H3N8) to synthetic polymers. The virus shows preference for 3'SLN. **(B)** Binding of early human IAV (H3N2) to synthetic polymers. The virus shows dual preference for 3'SLN and 6'SLN. **(C)** More recent human H3N2 viruses switch the receptor preference for 6'SLN and reduced affinity for the avian-like polymers 3'SLN.

Abbreviations: 3'SLN, 3'-sialyl-N-acetylglucosamine trisaccharide; 6'SLN, 6'-sialyl-N-acetylglucosamine; HA, hemagglutinin.

O-glycans with three or more LacNAc repeats has been attributed to the loss of affinity for chicken RBCs. This study has proposed a bidentate binding model, where these human-type receptors with elongated branches bind to two subunits of the same HA, increasing the avidity of recent H3 viruses for the complex forms of human SA α 2,6 receptors.⁵⁰

The effect of receptor-binding variations in human H3 viruses through all these years has represented a scientific challenge for virus isolation, antigenic characterization, and vaccine strain selection. A transition in in vitro systems has required moving from only chicken to a broad source as turkey, guinea pig, or human RBCs, without any success. Thus, virus neutralization assays are now more commonly used in addition to hemagglutination inhibition assays to characterize antigenicity and antibody responses.⁵¹

Avian viruses: H5N1 and H7Nx

Perhaps the most extensive study on the glycobiology of influenza virus has been done in AIV as a strategy to gain

knowledge on the receptor-binding properties of these viruses that might acquire human-to-human transmission properties. In general, AIV, including those that are highly pathogenic in birds, do not replicate efficiently or cause disease in humans. Prior to the influenza virus H5N1 isolated in Hong Kong in 1997, the only reported natural infections in humans by avian viruses are two cases of conjunctivitis associated with avian H7 viruses, one of which was an infection with a seal virus of avian origin.^{52,53}

While influenza's primary site of infection in humans is the upper respiratory tract, the primary infection site in birds is their digestive tract. In contrast to human upper airways, avian guts predominantly express SA α 2,3 receptors. Studies on viruses isolated early during human pandemics, when the HA gene has only just crossed over from an avian source, clearly demonstrate these differences. Sequence analysis of viruses isolated early during the pandemic of 1957 and 1968 revealed that changes at conserved residues of the RBS were associated with human adaptation. Early H2N2

viruses isolated from humans in 1957 possessed avian-like amino acids Q226 G228 (H3 numbering) at the HA RBS; in vitro assays showed high affinity to the SA α 2,3-linked ligand. However, soon after its introduction in humans, the virus acquired a single amino acid change at 226Q→L and the receptor specificity of these viruses changed toward SA α 2,6 binding. A complete human adaptation step occurred when the virus also mutated at the second position Q226L G228S.^{1,8,54} A similar adaptation process occurred for H3 viruses from the 1968 pandemic, once the virus was fully adapted to transmit efficiently between humans.⁸

H5N1 viruses

The human infections caused by H5N1 chicken viruses in Hong Kong in 1997 demonstrated for the first time that AIV could infect humans directly without intermediate evolution. H5 viruses isolated between 1997 and 2004 showed 100% identity in the RBS; however, viruses from early 2005 have several amino acid changes in the HA, some of which are located in the vicinity of the RBS.⁵⁵ Phylogenetic analysis of H5 viruses isolated from humans and chickens since 1997 shows that 10 clades (0–9) were distributed in different geographic regions of Asia.⁵⁶ Clade 2 became predominant with an extensive intercontinental expansion of the Asian clade 2.3.4.4.^{57,58} The expansion of the A/Goose/Guangdong/1/1996 lineage included outbreaks of H5N8 viruses in China and South Korea in 2014 and the first detection in the American continent, Canada, and USA of a highly pathogenic reassortant H5N2 virus.⁵⁹

A novel H5N6 virus emerged in China in 2014 causing human infections; this novel genotype has replaced the H5N1 as the dominant subtype in southern China. These AIV are known to be the only subtype of clade 2.3.4.4 that is able to cross the host species barrier and infect humans with documented exposure to infected poultry.^{60,61} Nonetheless, all H5 AIV isolated to date from birds, mammals, or humans show preferential binding to SA α 2,3-linked receptors.^{62–67} As described with the recent human H3N2 viruses, a progression on the type of SA α 2,3-linked receptors recognized by the avian H5 has been observed, resulting from the high rate of antigenic and genetic diversity in H5N1 viruses. Mutations K222Q and S227R are predominant in the RBS of virus HAs from the actual clade 2.3.4.4, conferring a change in receptor-binding specificity and preference for fucosylated receptors and limited capacity for replication and transmission in mammals.^{68,69}

Interestingly and of relevance to the potential risk to humans, the chances that avian H5N1 viruses will give rise

to a pandemic have been deeply analyzed. Viruses and the effect of amino acid changes at the RBS have been tested under several in vitro and in vivo systems, and it is generally accepted that a double mutation at residues 226Q→L and 228G→S changed the receptor-binding activity of the avian virus to a human phenotype. The single mutation Q226L seems not to play an important role for the switch on receptor recognition by the H5 HA, even though it was paramount for the H2N2 and H3N2 viruses.⁸ Also, although it makes some change to the SA binding, the single mutant G228S by itself does not affect the complete switch. Then, it may be possible that Q226L is the driving force for the intermediate G228R mutate onward to a G228S.⁷⁰

Despite all the knowledge generated in the last 20 years about the H5N1 viruses, it is still unknown why these viruses have not yet become pandemic to humans. The most recent evidence using the ferret model suggested that for the avian H5N1 virus to become transmissible between humans, several mutations at the HA RBS are required, including those previously described in residues 226 and 228.^{71–73} Considering what is already known about the virus and how long it has been circulating in humans, it might be that H5N1 viruses are still far away from an evolutionary adaptation in the human host. However, it should be considered that some other internal genes also contribute to host adaptation; for instance, the PB2 gene appears to play a central role in the ability of IAV to replicate in different hosts. The phenotype of efficient replication in mammalian cells seems to result from an amino acid difference at position 627 (amino acid change E627K, lysine in human strains and glutamate in avian viruses).^{74,75}

H7Nx viruses

Avian influenza infections in humans are not just of H5N1 subtype. Avian viruses of the H7N9 subtype have also become a threat to humans. These AIV, which are restricted to China, infected humans for the first time in 2013; since then, a total of 1557 laboratory-confirmed cases of human infections including at least 605 deaths have been reported to the World Health Organization.⁷⁶ Soon, as the H7N9 viruses were first isolated from the humans, the determinants of pathogenicity were analyzed; it is of great relevance that most of the H7N9 viruses isolated possessed a genetic marker of human adaptation in the amino acid residue 226Q→L,⁷⁷ which, in pandemic H2 and H3 viruses from 1957 to 1968, allowed the virus to transmit between humans.⁸ Infection studies in different animal models, including mice, macaques, and guinea pigs, shed light on the capacity of H7N9 viruses to replicate

in different tissues of the upper and lower respiratory airways, which is lethal in mice.^{78,79} Important to the pandemic potential that these viruses represent, besides the presence of mutation Q226L, several transmission studies using the ferret model have demonstrated limited transmission by respiratory droplets.^{78–82} Receptor-binding analysis of H7N9 viruses using solid-phase binding assays or the glycan array have shown that only the presence of 226L in these viruses is not sufficient for a complete switch from avian SA α 2,3 to the human SA α 2,6 receptors.^{83–87} However, if the virus acquires additional mutations, V186G/K-K193T-G228S or V186N-N224K-G228S, it has the capacity to recognize and bind to human trachea to the same extent as a human H3 virus and switch the receptor specificity from avian to human type.^{88,89} The evolutive constraints that maintain H7N9 viruses as a threat to human health are still unknown; implementation of extreme surveillance measures and risk assessment tools will alert in the event of a virus capable of crossing the host range barrier.

Not least important these avian viruses of the H7N7 subtype are also a threat to human health. Worldwide, isolated cases or outbreaks of H7N7 have been reported from the Netherlands and Italy.^{90,91} In humans, infections with avian viruses H7N7 are clinically associated to self-limiting conjunctivitis, and no person-to-person transmission has been reported. Experimental work using the ferret animal model showed that H7 viruses isolated from the American lineage have already acquired the phenotype of human receptor recognition by binding to several SA α 2,6-linked glycans and losing affinity for the avian SA α 2,3-linked carbohydrates. Moreover, an avian H7N7 isolated from an individual with respiratory disease was transmitted to naïve control ferrets only by direct contact.^{92,93} Infection of eye conjunctiva by the avian H7N7 virus correlates with the predominance of SA α 2,3.⁹⁴

H7N3 viruses are also associated with conjunctivitis when humans are infected. Historically, these viruses have caused human infection in Italy in 2002, in Canada in 2004, and in England in 2006.^{91,95,96} Recently, two human cases of associated conjunctivitis were reported in two Mexican workers during an outbreak of highly pathogenic avian H7N3 in poultry flocks.⁹⁷ The receptor-binding affinity of H7N3 viruses from the Eurasian and North American lineages showed increased binding of SA α 2,6 glycans, exhibiting the characteristics of a human influenza virus without losing affinity for the avian SA α 2,3 receptor. However, besides the fact that viruses replicate efficiently in the upper and lower respiratory tracts of ferrets including the recent Mexican

viruses from 2012 human cases, transmission between the ferrets still occurs only by direct contact.^{78,92}

The most recent event of zoonotic infections with influenza viruses of avian origin was reported in companion animals. An AIV of the H7N2 subtype was isolated from a person infected through direct contact with sick domestic cats. Viruses isolated from the person and the cat showed 99.9% identity, and they were phylogenetically related to H7N2 viruses isolated from poultry in the USA. The glycan array analysis showed in these viruses preference for SA α 2,3 glycans, though there was some affinity for SA α 2,6 human-like receptors. Even though this seems an isolated event of AIV trespassing the human host range, it certainly reminds us the plasticity of the IAV to cross the species barrier.⁹⁸

Concluding remarks

Bearing in mind the accumulated evidence about H1, H2, and H3 viruses that caused the pandemics in the 20th and 21st centuries, it seems very probable that in addition to reassortment, a prerequisite for a pandemic virus is that its HA protein must adapt to bind SA α 2,6 rather than SA α 2,3. The human airway epithelium is a complex organ of different cell types and functions, and is also the target by which the influenza viruses gain entry to their hosts and from where they are transmitted onward to a new host. However, binding to the target cell is a limiting step for efficient virus replication in the host, and human influenza viruses are presumably adapted to optimize their interactions with this organ.

Dissecting the receptor-binding phenotype of influenza virus has become a primary task for researchers and world health authorities around the globe. The World Health Organization published a Tool for Influenza Pandemic Risk Assessment for those countries that have experienced human infections with animal influenza viruses. This information can help scientists better understand the risk these viruses pose to human health and can help support development of tools and strategies for prevention and treatment.

Disclosure

The author reports no conflicts of interest in this work.

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