Increased detection in Australia and Singapore of a novel influenza A(H1N1)2009 variant with reduced oseltamivir and zanamivir sensitivity due to a S247N neuraminidase mutation

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A novel influenza A(H1N1)2009 variant with mildly reduced oseltamivir and zanamivir sensitivity has been detected in more than 10% of community specimens in Singapore and more than 30% of samples from northern Australia during the early months of 2011. The variant, which has also been detected in other regions of the Asia-Pacific, contains a S247N neuraminidase mutation. When combined with the H275Y mutation, as detected in an oseltamivir-treated patient, the dual S247N+H275Y mutant had extremely high oseltamivir resistance.

Introduction

The emergence and global spread in 2007/08 of an oseltamivir-resistant seasonal influenza A(H1N1) variant containing a histidine to tyrosine substitution (H275Y) in the neuraminidase (NA), demonstrated the potential for drug-resistant influenza viruses to arise and spread within the community in the absence of drug-selective pressure [1]. Since the start of the 2009 pandemic, the oseltamivir-resistant H275Y variant has only been detected on rare occasions in pandemic influenza A(H1N1)2009 community specimens (<1%), although recent reports have suggested that this frequency may be increasing [2]. Other NA mutations in influenza A(H1N1)2009 viruses have been reported to confer mildly reduced oseltamivir and/or zanamivir sensitivity, such as substitutions at the isoleucine residue at position 223 (N1 numbering), but the detection of these mutants has been very rare and has occurred mostly in isolated cases of immunocompromised individuals under long-term NA inhibitor (NAI) treatment [3,4]. Here we report the identification and increased rate of detection in community samples of a novel influenza A(H1N1)2009 variant with reduced oseltamivir and zanamivir sensitivity. The variant contained a

serine to asparagine mutation at residue 247 (S247N) of the NA, and has been detected in recent community specimens from Australia, Brunei and Singapore. Although the mutation has been described before in a small number of seasonal influenza A(H1N1) and highly pathogenic influenza A(H5N1) viruses with reduced NAI sensitivity [5,6], it has not previously been reported in influenza A(H1N1)2009 viruses, and has not occurred in any strains at the frequencies that are described here.

Materials and methods

Specimens and isolates from the Asia-Pacific region were submitted to the World Health Organization (WHO) Collaborating Centre for Reference and Research on Influenza, Melbourne, as part of the WHO Global Influenza Surveillance Network. Isolates cultured in Madin-Darby canine kidney (MDCK) cells were analysed for oseltamivir, zanamivir and peramivir sensitivity using a fluorescence-based neuraminidase inhibition assay [7]. Oseltamivir carboxylate, the active form of the ethyl ester prodrug oseltamivir phosphate, was kindly provided by Hoffmann-La Roche Ltd, Switzerland; zanamivir was kindly provided by GlaxoSmithKline, Australia; peramivir was kindly provided by BioCryst, United States. Isolates and specimens were sequenced using standard techniques at the WHO Collaborating Centre, Melbourne, except for Singaporean strains which were sequenced at the National Public Health Laboratory, Singapore. Haemagglutinin (HA) and NA sequences reported here have been deposited on GISAID (www.gisaid.org; accession numbers EPI319165 to EPI319183) or GenBank (accession numbers CY091664 to CY091724 and CY063853 to CY063854). Other HA and NA sequences were analysed from GISAID or Genbank sequence databases (see supplementary* for details: http://mendel.bii.a-star.edu.sg/SEQUENCES/H1N1/S247N/Hurt_et_al_NA-S247N_suppl).

pdf). A concatenated HA and NA nucleotide alignment was created with MAFFT (FFT-NS-2) (<http://mafft.cbrc.jp/alignment/software/>) and a maximum likelihood phylogenetic tree was generated with PhyML (<http://www.atgc-montpellier.fr/phyml/>) using the HKY85 substitution model, 4 gamma-distributed categories (shape parameter 0.46) and the approximate likelihood ratio test. The trees were visualised and annotated in MEGA5 (<http://www.megasoftware.net/>).

The S247N and dual S247N+H275Y NA mutations were investigated further by site-directed mutagenesis and reverse genetics using the eight-plasmid system. Mutations were engineered into the NA of influenza A/Auckland/1/2009(H1N1) and transfected together with plasmids containing the seven remaining segments

from the influenza A/Puerto Rico/8/34 strain, as described previously [8].

The mutations S247N and H275Y were modelled into the influenza A(H1N1)2009 neuraminidase crystal structure and the effects of the mutations on oseltamivir binding were visualised and examined using FoldX (<http://foldx.crg.es>) and Yasara (<http://www.yasara.org>).

Results

From April 2009 to December 2010, over 2,900 influenza A(H1N1)2009 influenza viruses from the Asia-Pacific region were analysed for NA sensitivity with only 23 H275Y oseltamivir-resistant viruses detected (0.8%), and virtually all of the remaining strains being fully sensitive to both oseltamivir and zanamivir. However

FIGURE 1

Detection of influenza A(H1N1)2009 S247N variants since 2009**

A. Year and frequency of detection of S247N mutants globally since 2009 (n=49). Dots are overlaid on top of each other for Singapore.

B. Detection of S247N mutants in Darwin and Singapore between October 2010 and April 2011 (n=19). S247N mutants are indicated by light blue circles and wildtype S247 viruses are indicated by dark blue circles.

since December 2010, 22 influenza A(H1N1)2009 viruses containing a novel S247N NA mutation (S246N based on N2 numbering), have been detected in both clinical specimens and isolates from the Asia-Pacific region. The majority of the S247N variants detected since December 2010 (n=19) have occurred in two clusters, one in the Darwin region in northern Australia and the other in Singapore (Figure 1A). The variant has also been detected in Western Australia and Brunei (in this study), and based on data from the public sequence databases has occurred, albeit rarely, in other locations such as

the United States, Europe and Asia since 2009 (Figure 1A). During the first three months of 2011, 28 influenza A(H1N1)2009 strains were sampled from the Darwin region, of which nine contained the S247N mutation, while in Singapore 10 out of 80 viruses sampled since December 2010 had the S247N NA mutation (Figure 1B).

Of the 22 S247N variants detected, nine were cultured and in an NA inhibition assay showed a mean six-fold reduction in oseltamivir sensitivity, a three-fold reduction in zanamivir sensitivity, and no significant

TABLE

Neuraminidase inhibitor sensitivity of naturally occurring and recombinant S247N and S247N+H275Y influenza A(H1N1)2009 variants, December 2010–March 2011

Virus	Mutation	Specimen date	Zanamivir		Oseltamivir carboxylate		Peramivir	
			Mean IC ₅₀ ± SD (nM)	x-fold difference vs wildtype	Mean IC ₅₀ ± SD (nM)	x-fold difference vs wildtype	Mean IC ₅₀ ± SD (nM)	x-fold difference vs wildtype
Mean of sensitive influenza A(H1N1) 2009 viruses (n=3,169)	-	-	0.28±0.15	-	0.45±0.35	-	0.20±0.10 ^a	-
Mean of S247N influenza A(H1N1) 2009 viruses ^b (n=9)	S247N	-	0.85±0.10	3	2.68±0.61	6	0.21±0.02	1
Naturally occurring strains								
A/Singapore/GP4565/2010	S247N	10 Dec 2010	0.96±0.06	3	3.21±0.50	7	0.21±0.02	1
A/Singapore/GP4588/2010	S247N	15 Dec 2010	0.81±0.08	3	3.02±0.54	7	0.21±0.02	1
A/Darwin/2/2001	S247N	13 Jan 2011	0.88±0.10	3	3.20±0.47	7	0.23±0.01	1
A/Brunei/1/2011	S247N	13 Jan 2011	0.86±0.08	3	1.62±0.11	4	0.26±0.03	1
A/Darwin/10/2011	S247N	9 Feb 2011	0.79±0.08	3	2.41±0.47	5	0.19±0.01	1
A/Perth/30/2011 ^c	S247N	1 Mar 2011	0.80±0.01	3	2.34±0.21	5	0.22±0.01	1
A/Perth/29/2011 ^c	S247N+H275Y	8 Mar 2011	1.30±0.08	5	2,646.81±293.55	5,880	66.88±4.19	334
A/Darwin/70/2011	S247N	14 Mar 2011	1.04±0.10	4	3.49±0.45	8	0.21±0.01	1
A/Darwin/74/2011	S247N	16 Mar 2011	0.68±0.03	2	2.15±0.23	5	0.19±0.01	1
A/Darwin/75/2011	S247N	17 Mar 2011	0.82±0.04	3	2.69±0.11	6	0.21±0.02	1
Recombinant strains ^d								
RG-wildtype	-	-	0.24±0.05	-	0.30±0.20	-	0.09±0.01	-
RG-S247N	S247N	-	0.57±0.10	2	0.96±0.48	3	0.18±0.02	2
RG-S247N+H275Y	S247N+H275Y	-	1.25±0.17	5	2,149.96±309.02	7,073	60.91±5.88	704
RG-H275Y	H275Y	-	0.26±0.03	1	195.02±21.05	642	19.72±1.42	228

RG: reverse genetics; SD: standard deviation.

^a Mean and standard deviation of peramivir IC₅₀ values based on analysis of n=273 isolates.

^b Mean does not include the dual S247N+H275Y mutant.

^c Viruses were from the same patient before and after oseltamivir treatment (refer to the text for more details).

^d Recombinant strains were derived using site-directed mutagenesis and reverse genetics.

FIGURE 2

Phylogenetic trees of concatenated haemagglutinin and neuraminidase sequences from influenza A(H1N1)2009 S247N variants detected December 2010-March 2011

Dark blue dots: S247N variants identified in this study (n=22); light blue dots: S247N variants previously reported on Genbank or GISAID sequence databases.

Branches and designations are coloured according to their sample collection date. The nucleotide sequences for these strains were concatenated such that a single sequence contains nucleotide sequences encoding both the haemagglutinin and neuraminidase proteins in order to increase the evolutionary signal for strain differences of these almost identical sequences.

FIGURE 3

The structural effect of the S247N and S247N+H275Y mutations in the neuraminidase of influenza A(H1N1)

A

B

C

A. Wildtype.

B. Dual S247N+H275Y mutant.

C. The merged image shows the superimposition of the wildtype and the double mutant relative to bound oseltamivir (pink, from PDB:3c10). The crystal structure of influenza A(H1N1)2009 neuraminidase (PDB:3nss, chain A) was used to model the double mutations S247N and H275Y using Yasara.

Hydrogen bonds among the important residues are shown in yellow. Side chains of residues undergoing conformational changes upon the double mutation are shown in cyan.

reduction in peramivir sensitivity compared to the mean IC_{50} (concentration required to inhibit 50% of NA activity) of sensitive influenza A(H1N1)2009 viruses (Table).

In Perth, Western Australia, an immunocompromised patient was found on 1 March 2011 to be infected with an influenza virus that contained the S247N NA mutation (A/Perth/30/2011) (Table). Oseltamivir treatment was commenced two days after detecting the initial S247N variant (3 March). A sputum specimen collected five days later (8 March) contained an influenza virus with both the S247N and H275Y NA mutations (A/Perth/29/2011) (Table). Despite commencement of intravenous zanamivir the patient died on 16 March from multiple organ failure. The isolate with the dual S247N and H275Y mutations had an oseltamivir IC_{50} nearly 6,000-fold higher than sensitive viruses and 10-fold higher than seen for influenza A(H1N1)2009 viruses with the H275Y mutation alone (Table). Clonal analysis of the virus population from the sputum specimen collected after oseltamivir treatment confirmed that both the S247N and H275Y mutations occurred together in the NA. The effect of the S247N and dual S247N and H275Y mutations on NAI sensitivity was confirmed by reverse genetics experiments (Table).

Sequence analysis of the S247N variants from the recent Darwin and Singapore clusters revealed that the majority belonged to the same genetic lineage (Figure 2). The two strains from the immunocompromised patient in Perth were also genetically similar to the Darwin and Singapore strains, but the virus from Brunei and one strain from Singapore were placed in a separate clade together with one other S247N strain from the United States (Figure 2).

Structural analyses indicate that the S247N mutation could change the hydrogen bonding network and side chain conformation of residue E277 (E276 by N2 numbering) reflecting similar but weaker alterations known from the H275Y mutation. When adding H275Y, E277 gets pushed further into the drug binding pocket which is believed to weaken oseltamivir binding (Figure 3).

Discussion

Previous studies have reported the effect of the S247N mutation on NAI sensitivity in N1 neuraminidases. The levels of oseltamivir and zanamivir sensitivity reported for the S247N mutation in a pre-pandemic seasonal influenza A(H1N1) virus were similar to those reported here for the mutation in influenza A(H1N1)2009 strains [5]. However a greater reduction in oseltamivir sensitivity was reported for a highly pathogenic influenza A(H5N1) strain with a S247N mutation compared to the influenza A(H1N1)2009 strains reported here (24-fold vs 6-fold reduction compared to their respective wildtypes) [6]. The same study also reported on an

influenza A(H5N1) virus with the S247N and two additional mutations, I223L and K150N (N1 numbering). This triple mutant virus had a greater reduction in oseltamivir sensitivity compared to the S247N mutation alone (77-fold vs 24-fold reduction), further demonstrating that mutations at other key residues such as I223, in combination with S247N, can cumulatively decrease NAI sensitivity. Given that a number of studies have recently reported I223 mutations conferring NAI resistance in influenza A(H1N1)2009 strains [3,4,9,10], and that data from the United Kingdom show an increased frequency of H275Y mutants in community samples [2], the likelihood of mutation combinations S247N+I223X and S247N+H275Y is considerably increased if the S247N influenza A(H1N1)2009 variant continues to spread.

Pharmacokinetic data would suggest that the maximum drug levels achieved via the recommended dose easily exceed the observed IC_{50} values of the S247N mutant [11], and therefore the variant is unlikely to be clinically resistant. However, it is noteworthy that recent clinical studies have demonstrated a reduced oseltamivir efficacy for normal influenza B viruses [12] which have IC_{50} values only 6-fold higher than that of the S247N influenza A(H1N1)2009 variants (mean oseltamivir $IC_{50} \pm SD$ for influenza B viruses from 2010 and 2011: $15.5 \text{ nM} \pm 11.3$ ($n=557$)). Data from Singapore and Darwin demonstrate that the S247N variant is able to circulate widely and therefore does not appear to have compromised viral fitness or transmissibility. If the S247N variant spreads globally, the greatest concern is that other NA mutations which may have previously caused only mild reductions in NAI susceptibility (e.g. mutations at the I223 residue) could instead cumulatively decrease NAI sensitivity to levels that may be clinically significant and affect treatment efficacy. Laboratories should consider screening currently circulating specimens and isolates for the S247N NA mutation to determine whether the variant is spreading into other regions.

***Note:** Supplementary information made available by the authors on an independent website is not edited by *Eurosurveillance*, and *Eurosurveillance* is not responsible for the content.

****Erratum:** The map in Figure 1 was misaligned with the dots. This was corrected on 10 June 2011.

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