

Clinical technique sets new standard for speed in battle to prevent pandemic infection

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A new diagnosis technique developed by researchers at the RIKEN Omics Science Center (OSC) has succeeded in detecting influenza virus infection in only 40 minutes and with one hundred times the sensitivity of conventional methods. Clinical research conducted in 2009 and 2010 confirms the new technique accurately identified the 2009 pandemic (pdm) influenza virus in Japanese patients less than 24 hours after fever onset, much faster than standard diagnostic tests.

The human-to-human transmission of new, highly pathogenic strains of [influenza](#) virus poses today a major threat to human health and to the security of global society. With its rapid global spread, the 2009 pandemic (pdm) [influenza virus](#) reminded the world of this threat, resulting in an estimated 18,000 deaths worldwide. In Japan, infected patients over the winter season of 2009 accounted for a staggering 16% of the total population.

Tackling the challenge of such global pandemics requires new technology for rapid [clinical diagnosis](#). To answer this need, Toshihisa Ishikawa and colleagues at the RIKEN OSC developed the RT-SmartAmp assay, a technique to rapidly detect the 2009 pdm influenza A(H1N1) virus from patient swab samples. By combining both reverse transcriptase (RT) and isothermal DNA amplification reactions in one step, the RT-SmartAmp assay does away with the need for RNA extraction and PCR reaction. The researchers adapted the RT-SmartAmp technique using a fluorescent primer to specifically detect the 2009 pdm influenza A(H1N1) virus within 40 minutes, without cross-reacting with the seasonal A(H1N1), A(H3N2), or B-type (Victoria) viruses.

The effectiveness of the RT-SmartAmp method was confirmed in clinical research carried out at Japanese hospitals during the period of October

2009 to January 2010, where it outperformed standard diagnosis tests in both speed and sensitivity. Of a total 255 clinical samples, 140 (54.9%) were identified as 2009 pdm A(H1N1)-positive by RT-SmartAmp, compared to only 110 (43.1%) detected by standard [diagnostic tests](#). In 72.8% of all 140 infection-positive cases, the RT-SmartAmp assay detected the presence of the pdm influenza virus within 24 hours of fever onset.

Taken together, these results set a new standard for infection diagnosis speed, providing a highly-effective tool for rapidly detecting sub-types of the H5N1 virus and oseltamivir-resistant influenza viruses and promising support in the battle to prevent global pandemic infection.

The study is published today in *PLoS ONE*.

Provided by RIKEN

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