**Lower respiratory tract infection in the community: associations between viral aetiology and illness course**

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**ABSTRACT**

**OBJECTIVES.** This study determined associations between respiratory viruses and subsequent illness course in primary care adult patients presenting with acute cough and/or suspected lower respiratory tract infection (LRTI).

**METHODS.** A prospective European primary care study recruited adults with symptoms of lower respiratory tract infection between Nov-Apr 2007-2010. Real-time in-house polymerase chain reaction (PCR) was performed to test for six common respiratory viruses. In this secondary analysis, symptom severity (scored 1=no problem, 2=mild, 3=moderate, 4=severe) and symptom duration were compared between groups with different viral aetiologies using regression and Cox proportional hazard models, respectively. Additionally, associations between baseline viral load (cycle threshold (Ct) value) and illness course were assessed.

**RESULTS.** The PCR tested positive for a commonrespiratory virus in 1,354 of the 2,957 (45.8%) included patients. The overall mean symptom score at presentation was 2.09 (95%CI 2.07-2.11) and the median duration until resolution of moderately bad or severe symptoms was 8.70 days (interquartile range 4.50-11.00). Patients with influenza virus, human metapneumovirus (hMPV), respiratory syncytial virus (RSV), coronavirus (CoV) or rhinovirus had a significantly higher symptom score than patients with no virus isolated (0.07-0.25 points or 2.3-8.3% higher symptom score). Time to symptom resolution was longer in RSV infections (adjusted hazard ratio (AHR) 0.80, 95%CI 0.65-0.96) and hMPV infections (AHR 0.77, 95%CI 0.62-0.94) than in infections with no virus isolated. Overall, baseline viral load was associated with symptom severity (difference 0.11, 95%CI 0.06-0.16 per 10 cycles decrease in Ct value), but not with symptom duration.

**CONCLUSIONS.** In healthy, working adults from the general community presenting at the general practitioner with acute cough and/or suspected LRTI respiratory viruses other than influenza impose an illness burden comparable to influenza. Hence, the public health focus for viral respiratory tract infections should be broadened.

**INTRODUCTION**  
From the few studies describing the aetiology of acute lower respiratory tract infections (LRTIs) in primary care patients, we know that most LRTIs in the general community are caused by viral pathogens, in particular rhinovirus, influenza virus, coronavirus (CoV), respiratory syncytial virus (RSV), human metapneumovirus (hMPV), and parainfluenza virus (PiV)(1,2). The illness course of LRTIs in adults presenting in this setting - a relatively healthy, working population - is mostly self-limiting and complications are rare(3). However, with an average of 3.5 days sick leave per year, LRTIs cause a substantial socio-economic burden(3,4). In adults, influenza virus, bacteria, and viral-bacterial coinfections are assumed to cause the most severe illnesses, with most systemic symptoms, longest illness durations, and most complications(5–7). However, evidence on associations between aetiology and severity are mainly derived from hospital care settings with vulnerable patient populations(8–10). In this setting, a focus on pathogens with the highest complication rates is obvious. Quite often, however, this focus is also applied in the general community, with public health interventions as the annual influenza vaccinations targeted at the most vulnerable people with the aim of reducing the risk of complications and death(11). Although data on the impact of respiratory viruses in the primary care setting are limited due to restricted microbial testing and absence of a standardized, validated outcome measure to evaluate illness severity(12), there are studies suggesting that the burden of disease from infections due to respiratory viruses other than influenza – i.p. rhinovirus, coronavirus and RSV - may be greater overall(13). In this study, we aimed to explore the associations between respiratory viral pathogens, including viral load, and illness course in the adult primary care community, thereby opening up possibilities to base the public health focus on the impact of respiratory viruses in primary care, rather than on extrapolated data from hospital settings. This study was conducted in a large European cohort consisting of prospectively enrolled adult patients with acute cough and/or a clinical suspicion for LRTI.

**METHODS**

*Design and study population*

This prospective study in primary care is part of the GRACE study (Genomics to combat Resistance against Antibiotics in Community-acquired LRTI in Europe). Participants were recruited between November 2007 and April 2010 by general practitioners (GPs) from 16 primary care networks in 11 European countries (*Supplementary Figure 1*). Patients aged ≥18 years presenting with acute cough (duration of ≤28 days) and/or suspected LRTI, were asked to participate in this study, i.e. to fill out study materials and provide written informed consent(14). Exclusion criteria were pregnancy, breast-feeding, any serious immunocompromised condition and antibiotic use in the previous month(14). About one third of these patients agreed to being randomised to either the intervention (amoxicillin) or placebo arm of the original randomized controlled trial(14). Remaining patients were not randomly assigned, but were included in the observational part of the study(1). In the current study, both trial and observational patients were analysed together, but patients without PCR and/or serology results on viral aetiology (all due to practical reasons) were excluded. Ethical approval was obtained for all participating networks.

*Clinical measurements*

For the collection of clinical data on the day of presentation (baseline), standardized case report forms (CRFs) were used. GPs completed the CRF on the following 12 symptoms rated by the patients using a 4-point Likert-scale (1=no problem, 2=mild, 3=moderate, 4=severe): cough, sputum production, shortness of breath, wheeze, blocked or runny nose, fever, chest pain, muscle aching, headache, disturbed sleep, feeling generally unwell, and interference with normal daily activities. Additionally, the symptoms confusion/disorientation and diarrhoea were rated. Following initial presentation, patients were asked to fill out a symptom diary at home on a daily basis until they had no more symptoms or until the end of follow-up (day 28). Patients were asked to rate the same 12 symptoms by using a 7-point Likert-scale (0=normal, 1=very little problem, 2=slight problem, 3=moderately bad, 4=bad, 5=very bad, 6=as bad as it could be). This diary was internally reliable, valid, and sensitive to change for acute LRTI(15).

*Microbiological measurements*

At baseline, two nasopharyngeal flocked swabs were taken by trained staff within 24 hours after recruitment and before any antimicrobial treatment had started. Swabs were placed in universal transport medium immediately, frozen locally, and transported on dry ice to the central laboratory (University of Antwerp). Real-time in-house polymerase chain reaction (RT-PCR) testing was performed either as four multiplex RT-PCRs (combining INF-A, INF-B, and RSV; PIV1-4; HRV, hMPV, and the EAV internal control; and finally the human CoV: 229E, OC43, NL63, and HKU1), or as monoplex (all other viruses) (16). RNA/DNA extractions and amplification methods were described previously(1,16). Based on the results from our study comparing the prevalence of viral pathogens between symptomatic and asymptomatic matched controls(1), we evaluated rhinoviruses, influenza viruses, coronaviruses, RSV, hMPV and PiV. Since (pan-)adenovirus (1.3% vs. 1.1%, p=0.33), bocavirus (0.6% vs. 0.8%, p=0.43) and WU/KI polyomaviruses (2.2% vs. 2.5%, p=0.02) were not detected more frequently in symptomatic patients than in controls, they were not considered pathogenic respiratory viruses and therefore excluded from our analyses(1). A cycle threshold (Ct) value - an inverse, logarithmic, quantitative measurement of viral load – below 45 was chosen as cut-off for a positive result. We adjusted our analyses for bacterial infections, which were defined as having at least one of the following pathogens detected in a sputum or nasopharyngeal sample: *Streptococcus* species*,* Gram-negative species,or *Aspergillus* (fungus). Commensals and *Candida* species were considered contaminants for which analyses were not adjusted.Microbiologists who determined the results were blinded to clinical information.

*Outcome parameters*

We focused on two main outcome parameters: symptom severity at presentation and illness duration. Symptom severity was measured as the mean CRF score for all 12 symptoms (scored 1-4) at baseline(14,17–19). Illness duration was defined as the duration until absence of any symptoms rated moderately bad or severe (score 3 or above) in the symptom diary following initial presentation(14,17–19). Additionally, the severity of all individual symptoms was analysed, dichotomizing symptom severity at no/mild/moderate versus severe.

*Statistical analysis*

At baseline, the variables of interest and confounders had less than 1% missing values. We accounted for missing values using a multiple imputation model including baseline characteristics, predictors and outcome variables by which one imputed dataset was created. Patients without follow-up data were excluded from the analysis on illness duration. Baseline characteristics were reported descriptively as N and percentage, mean and standard deviation (SD) or median and interquartile range (IQR) as appropriate. The association between symptom severity at baseline and the presence of specific viruses and the number of detected viruses was analysed with linear regression models. The assumptions for the model (distributional assumptions, homoscedasticity) were assessed with residual plots(20). Results were expressed as differences in mean symptom severity with a 95% confidence interval (CI) and p-values. In an additional step, we evaluated the association between individual symptoms and the presence of detected viruses. Results were expressed as odds ratios (OR) with a 95% CI and p-values. The association between presence of detected viruses and the duration until absence of symptoms rated moderately bad or severe were analysed with cox proportional hazard models. For the latter analysis, patients were censored at the end of follow-up or if less than ten symptoms were filled out in the symptom diary. If patients already met the event criteria at baseline (n=104), we defined their time to event as one day. Results were expressed as hazard ratios (HR) with a 95% CI and p-values. For all analyses, we adjusted for the following potential confounders defined beforehand: age, gender, pulmonary comorbidities (asthma, COPD and other chronic lung diseases), hearth failure, current or former smoking, influenza vaccination during the preceding fall or winter, duration of symptoms before presentation (in days), detection of one or more bacteria and detection of a second respiratory virus. Statistical analyses were performed using SPSS v.25.0 for Windows and the “survival” and “survminer” packages in R v.4.

**RESULTS**

*Study population*

We included 2,957 adult patients (*Figure 1*). Demographics and clinical symptoms at presentation are presented in *Table 1*. Patients had a median age of 50 years (IQR 36-63), 1,195 (40.4%) were male and 1,603 (54.2%) were a former or current smoker. The overall mean symptom score at presentation was 2.09 (95%CI 2.07-2.11). Respiratory viruses (1,411) were detected in 1,354 patient samples (*Figure 2*). The proportion of influenza virus positive patients was lower among patients who received the annual influenza vaccination during the preceding fall/winter (38/707, 5.4%) than among non-vaccinated patients (259/2250, 11.5%) (p<0.001). Follow-up data were available for 2,393 patients (80.9%). Baseline disease characteristics did not differ between patients who did (n=2,393), or did not (n=564) fill out a symptom diary. Of all 2,393 patients included in the symptom duration analysis, 2,186 patients (91.3%) documented resolution of symptoms rated moderately bad or severe before the end of follow-up, with a median duration of 6.00 days (IQR 4.00-11.00 days). At presentation, only two patients were prescribed antiviral medication (oseltamivir).

*Association between respiratory viruses and symptom severity*

We evaluated the severity of symptoms at presentation for patients with CoV, hMPV, influenza virus, PiV, rhinovirus and RSV, as compared to patients without these viruses, with adjustment for confounders, bacteria and co-viruses. Influenza virus, hMPV, RSV, CoV and rhinovirus were significantly associated with, respectively, 0.25 (95%CI 0.19-0.31), 0.16 (95%CI 0.07-0.26), 0.12 (95%CI 0.04-0.21), 0.09 (95%CI 0.02-0.16) and 0.07 (95%CI 0.02-0.12) points higher symptom scores at presentation as compared to patients without detected virus (*Table 2*). Among patients in whom a virus was detected, a ten cycles lower Ct value – i.e. a higher viral load – measured at presentation, was associated with a 0.11 (95%CI 0.06-0.16) point higher mean symptom severity as compared to patients without detected virus. After stratification for viral aetiology, we only observed an association between viral load and symptom severity for rhinovirus (increase of 0.12 per 10 cycles reduction in Ct value, 95%CI 0.04-0.20) and for RSV (increase of 0.16 per 10 cycles reduction in Ct value, 95%CI 0.01-0.30). When looking at differences in the severity of individual symptoms of these viruses (*Figure 3*), influenza virus was independently associated with severe fever (OR 6.3, 95%CI 4.0-9.8), headache (OR 3.1, 95%CI 2.2-4.5), chest pain (OR 2.0, 95%CI 1.3-3.2), muscle pain (OR 2.5, 95%CI 1.6-3.9), disturbed sleep (OR 1.4, 95%CI 1.1-1.9), being generally unwell (OR 2.5, 95%CI 1.8-3.5), and interference with daily activities (OR 2.5, 95%CI 1.8-3.5). RSV was associated with severe headache (OR 2.0, 95%CI 1.2-3.5), disturbed sleep (OR 1.7, 95%CI 1.1-2.5) and a runny nose (OR 2.9, 95%CI 1.9-4.4). hMPV was associated with severe dyspnoea (OR 2.0, 95%CI 1.0-3.7) and headache (OR 2.0, 95%CI 1.1-3.7). Rhinovirus was associated with severe wheeze (OR 1.6, 95%CI 1.0-2.6), a runny nose (OR 1.6, 95%CI 1.2-2.1) and negatively associated with severe cough (OR 0.8, 95%CI 0.6-0.9). CoV was associated with a severe runny nose (OR 2.0, 95%CI 1.4-3.0) and negatively associated with severe chest pain (OR 0.3, 95%CI 0.1-0.9).

*Association between respiratory viruses and illness duration*

After adjustment for bacterial coinfections, baseline symptom severity and other potential confounders, patients with detected viral pathogen(s) had no significantly different HR (0.93, 95%CI 0.86-1.02) for resolution of moderately bad or severe symptoms compared to patients in which no virus was detected (*Table 3*). We also assessed the duration until resolution of moderately bad or severe symptoms for the six individual viruses as compared to patients without a detected virus (*Figure 4*). Patients with RSV had an adjusted hazard ratio (AHR) of 0.80 (95%CI 0.65-0.96) and patients with hMPV an AHR of 0.77 (95%CI 0.62-0.94) for symptom resolution, indicating a significant longer symptom duration as compared to patients without RSV and hMPV, respectively. All other viral pathogens showed no significant differences in AHRs. Among patients in whom a virus was detected, there was no association between baseline viral load and duration of moderately bad or severe symptoms (AHR per unit lower Ct value 1.01, 95%CI 0.99-1.02). After stratification for viral aetiology, no significant associations were found between viral load and symptom duration.

**DISCUSSION**

Adult patients visiting the GP with acute cough or suspected LRTI due to influenza virus, hMPV, RSV, CoV or rhinovirus had a 0.07-0.25 points (or 2-8%) higher mean symptom severity score (range 1-4) at presentation as compared to patients presenting with acute cough or suspected LRTI without detection of one of these respiratory viruses. In translation, patients with RSV - who have a 0.12 point (4%) higher symptom score at presentation than patients in whom no virus is detected - rate one or two symptoms severe instead of moderate, moderate instead of mild, or mild instead of absent. Additionally, RSV and hMPV were associated with a longer duration of moderately bad or severe symptoms, which might be linked to the pattern of immune response to these viruses(21). For all respiratory virus together, a higher viral load measured at presentation, was significantly associated with a higher symptom severity. This was caused by significant associations between viral load and symptom severity for rhinovirus and RSV. There was no association between viral load and the duration of moderately bad or severe symptoms.

*Clinical implications*

This study does not provide direct clinically actionable insight. However, although we do not provide recommendations on clinical management or treatment, we do think that the large number of patients included in this study provides important information which can be used to prioritize different respiratory viruses in the primary care setting. Currently, public health resources in the general community are guided by the aim to prevent complications in the most vulnerable people, and are focused almost exclusively on influenza(11,22,23). From a socio-economic perspective, however, targeting public health resources only at influenza virus neglects the substantial illness course in the community caused by other respiratory viruses. From our results we conclude that RSV and hMPV impose a disease burden that compares well to that of influenza virus and should therefore receive more attention in the primary care setting, e.g. by supporting the development and implementation of prevention approaches like vaccines(24-26).

*Strengths and limitations*

Despite the fact that we had a large cohort in which data were collected in a standardized manner, and outcome measures were in line with previous studies(14,17–19), there are several potential sources of bias that might limit the validity of our results. Firstly, it is possible that non-agreement of patients to participate in this observational study was not random. The extent to which this selection might be present is uncertain since we have no information on the number and characteristics of patients who declined participation. Secondly, the use of medication, such as antibiotic treatment, antiviral treatment, (over-the-counter) symptomatic treatment, and prophylactic antibiotics with antiviral effects (as azithromycin) might have influenced outcomes . We consider it unlikely that receiving antibiotics caused biased results, since the in-study amoxicillin trial showed no differences in outcomes between the intervention and placebo group(14). Since only two patients in our cohort were prescribed antivirals (oseltamivir), we consider the effect of antiviral treatment also negligible. Unavailability of data on the use of prophylactic antibiotics and symptomatic medication made adjustment for these factors impossible. Thirdly, there might be bias in the self-report of symptoms by patients. However, previous studies showed a high internal reliability, validity and sensitivity of the symptom diary we used(15). Also, since the 95/207 (46%) patients who did not meet the event criteria and who did not fill out their symptom diary completely were censored for the analysis, we do not expect selection bias due to loss of follow-up. Fourthly, the required sample size for the prospective observational cohort was not determined on the specific requirements of the current study. Hence, inconclusive or non-significant results can therefore not be considered definite to prove the absence of associations. We specifically choose not to correct for multiple testing, as this correction may further hamper statistical power, especially for viruses only detected in a limited number of patients. Fifthly, the relatively low overall percentage of detected viruses might have been caused by the inclusion of patients with quite long duration of symptoms. Since respiratory fluids are renewed quickly in the patient, viral pathogens in patients with longer duration of symptoms might therefore not have been detectable anymore. Finally, a higher viral load was associated with a higher symptom severity at presentation. Looking at specific viruses we only found this association for RSV and rhinovirus, which confirms previous studies(27–29). However, the interpretation of single viral load measurements is difficult. Not only are viral loads of respiratory viruses highly dependent on variation in sampling location and technique, they also rise and drop rapidly and it is known that symptoms mostly follow the viral load(30,31).

In conclusion, in this study among relatively healthy adult patients presenting in a primary care setting with acute cough and/or a suspected LRTI, influenza virus, hMPV, RSV, CoV and rhinovirus were associated with an increased symptom severity at presentation as compared to patients without a detected virus. In this general community population, RSV and hMPV were associated with a longer duration of moderately bad or severe symptoms. This study emphasizes that public health policies as vaccinations and awareness among GPs should not remain focused on influenza virus exclusively, but should also include other common respiratory viruses like RSV and hMPV that pose a high socio-economic burden to the general adult community.

**TRANSPARANCY DECLARATION**

Conflict of interest: the authors declare that they have no conflict of interest. Funding: the study was part of the European Union FP6 funded Network of Excellence GRACE. Orion Diagnostics provided the QuikRead instruments and kits for this study. The study sponsors played no role in the study design; in the collection, analysis, and interpretation of data; in the writing of the report; or in the decision to submit the paper for publication. Ethical approval: all procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. Informed consent: informed consent was obtained from all individual participants included in the study.

**AUTHOR CONTRIBUTIONS**

The larger GRACE observational study was designed by CCB, TJMV, PL, DC and HG, and sampling protocols by MI, CL, KL and HG. MI, CL, PL, TV and HG supervised the day-to-day management at study sites. PCR and serological analyses were performed by KL, AMVL, CL, KZ, ECJC, MV and FC. Data were analysed by LMV, RB, NPAZ, BDLB, JJO and FC. The manuscript was designed and drafted by LMV, NPAZ and FC, and was reviewed by all authors.

**REFERENCES**

1. Ieven M, Coenen S, Loens K, Lammens C, Coenjaerts F, Vanderstraeten A, et al. Aetiology of lower respiratory tract infection in adults in primary care: a prospective study in 11 European countries. Clin Microbiol Infect. 2018;24(11):1158–63.

2. Woodhead M, Blasi F, Ewig S, Garau J, Huchon G, Ieven M, et al. Guidelines for the management of adult lower respiratory tract infections - Summary. Clin Microbiol Infect. 2011;17(SUPPL. 6):1–24.

3. Verheij, T; Hermans, J; Kaptein, A; Mulder J. Acute bronchitis: course of symptoms and restrictions in patients’ daily activities. Scand J Prim Heal Care. 1995;13(1):8–12.

4. Fragaszy EB, Warren-Gash C, White PJ, Zambon M, Edmunds WJ, Nguyen-Van-Tam JS, et al. Effects of seasonal and pandemic influenza on health-related quality of life, work and school absence in England: Results from the Flu Watch cohort study. Influenza Other Respi Viruses. 2018;12(1):171–82.

5. Pavia AT. What is the Role of Respiratory Viruses in Community Acquired Pneumonia; What is the Best Therapy for Influenza and Other Viral Causes of CAP? Infect Dis Clin North Am. 2014;27(1):157–75.

6. Voiriot G, Visseaux B, Cohen J, Nguyen LBL, Neuville M, Morbieu C, et al. Viral-bacterial coinfection affects the presentation and alters the prognosis of severe community-acquired pneumonia. Crit Care. 2016;20(1):1–9.

7. Qu JX, Gu L, Pu ZH, Yu XM, Liu YM, Li R, et al. Viral etiology of community-acquired pneumonia among adolescents and adults with mild or moderate severity and its relation to age and severity. BMC Infect Dis. 2015;15(1):1–8.

8. Sokolow LZ, Naleway AL, Li DK, Shifflett P, Reynolds S, Henninger ML, et al. Severity of influenza and noninfluenza acute respiratory illness among pregnant women, 2010-2012. Am J Obstet Gynecol. 2015;212(2):202.e1-202.e11.

9. Treanor J, Falsey A. Respiratory viral infections in the elderly. Antiviral Res. 1999;44(2):79–102.

10. Gaunt ER, Harvala H, McIntyre C, Templeton KE, Simmonds P. Disease burden of the most commonly detected respiratory viruses in hospitalized patients calculated using the disability adjusted life year (DALY) model. J Clin Virol. 2011;52(3):215–21.

11. Demicheli V, Jefferson T, Di Pietrantonj C, Ferroni E, Thorning S, Thomas RE, et al. Vaccines for preventing influenza in the elderly (Review). Cochrane Database Syst Rev. 2018;2:CD004876.

12. Rath B, Conrad T, Myles P, Alchikh M, Ma X, Hoppe C, et al. Influenza and other respiratory viruses: standardizing disease severity in surveillance and clinical trials. Expert Rev Anti Infect Ther. 2017;15(6):545–68.

13. Nicholson KG, Kent J, Hammersley V, Cancio E. Acute viral infections of upper respiratory tract in elderly people living in the community: comparative, prospective, population based study of disease burden. BMJ. 2011;315(7115):1060–4.

14. Little P, Stuart B, Moore M, Coenen S, Butler CC, Godycki-Cwirko M, et al. Amoxicillin for acute lower-respiratory-tract infection in primary care when pneumonia is not suspected: A 12-country, randomised, placebo-controlled trial. Lancet Infect Dis. 2013;13(2):123–9.

15. Watson L, Little P, Moore M, Warner G, Williamson I. Validation study of a diary for use in acute lower respiratory tract infection. Fam Pract. 2001;18(5):553–4.

16. Loens K, Van Loon AM, Coenjaerts F, Van Aarle Y, Goossens H, Wallace P, et al. Performance of different mono- and multiplex nucleic acid amplification tests on a multipathogen external quality assessment panel. J Clin Microbiol. 2012;50(3):977–87.

17. Bruyndonckx R, Stuart B, Little P, Hens N, Ieven M, Butler CC, et al. Amoxicillin for acute lower respiratory tract infection in primary care: subgroup analysis by bacterial and viral aetiology. Clin Microbiol Infect. 2018;24(8):871–6.

18. Teepe J, Little P, Elshof N, Broekhuizen BDL, Moore M, Stuart B, et al. Amoxicillin for clinically unsuspected pneumonia in primary care: subgroup analysis. Eur Respir J. 2016;47(1):327-30.

19. Moore M, Stuart B, Coenen S, Butler CC, Goossens H, Verheij TJM, et al. Amoxicillin for acute lower respiratory tract infection in primary care: Subgroup analysis of potential high-risk groups. Br J Gen Pract. 2014;64(619):75–80.

20. Harrell F. Regression Modeling Strategies. With Applications to Linear Models, Logistic and Ordinal Regression, and Survival Analysis. 2015, p. 33-35.

21. Ascough S, Paterson S, Chiu C. Induction and subversion of human protective immunity: Contrasting influenza and respiratory syncytial virus. Front Immunol. 2018;9:323

22. Moncion K, Young K, Tunis M, Rempel S, Stirling R, Zhao L. Effectiveness of hand hygiene practices in preventing influenza virus infection in the community setting: A systematic review. Canada Commun Dis Rep. 2019;45(1):12–23.

23. Doyon-Plourde P, Fakih I, Tadount F, Fortin É, Quach C. Impact of influenza vaccination on healthcare utilization – A systematic review. Vaccine. 2019;37(24):3179-89.

24. Zhu T, Zhang C, Yu L, Chen J, Qiu H, Lyu W, et al. The preventive effect of vaccine prophylaxis on severe respiratory syncytial virus infection: A meta-analysis. Virol Sin. 2015;30(5):371–8.

25. Ren J, Phan T, Bao X. Recent vaccine development for human metapneumovirus. J Gen Virol. 2015;96(7):1515–20.

26. Papadopoulos NG, Megremis S, Kitsioulis NA, Vangelatou O, West P, Xepapadaki P. Promising approaches for the treatment and prevention of viral respiratory illnesses. J Allergy Clin Immunol. 2017;140(4):921–32.

27. Fuller JA, Njenga MK, Bigogo G, Aura B, Ope MO, Nderitu L, et al. Association of the CT values of real-time PCR of viral upper respiratory tract infection with clinical severity, Kenya. J Med Virol. 2013;85(5):924–32.

28. Feikin DR, Fu W, Park DE, Shi Q, Higdon MM, Baggett HC, et al. Is Higher Viral Load in the Upper Respiratory Tract Associated With Severe Pneumonia? Findings From the PERCH Study. Clin Infect Dis. 2017;64(suppl\_3):S337–46.

29. Waghmare A, Kuypers JM, Xie H, Leisenring W, Campbell AP, Jerome KR, et al. Viral load in hematopoietic cell transplant recipients infected with human rhinovirus correlates with burden of symptoms. Biol Blood Marrow Transplant. 2015;21(2):S317–8.

30. Bagga B, Woods CW, Veldman TH, Gilbert A, Mann A, Balaratnam G, Lambkin-Williams R, Oxford JS, McClain MT, Wilkinson T, Nich DJ. Comparing influenza and RSV viral and disease dynamics in experimentally infected adults predicts clinical effectiveness of RSV antivirals. Antivir Ther. 2013;18(6):785–91.

31. Garcia-Mauriño C, Moore-Clingenpeel M, Thomas J, Mertz S, Cohen DM, Ramilo O, et al. Viral Load Dynamics and Clinical Disease Severity in Infants With Respiratory Syncytial Virus Infection. J Infect Dis. 2018;219:1207–15.