Surveillance of respiratory syncytial virus infections in adults, Austria, 2017 to 2019

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Respiratory syncytial virus (RSV) testing is generally available in most care centres, but it is rarely performed because clinicians’ seldom suspect RSV to be the underlying pathogen in adults with respiratory disease. Here, we evaluate the impact of broad combined influenza/RSV testing on the clinical practice. Overall, 103 patients were tested positively for RSV. Our study indicates that positively tested patients were mostly of advanced age and suffered from chronic diseases. Mortality was significant in our cohort and higher in patients with advanced age. Further, we report a significant increase in detected RSV cases but also in detection rate. Together, these findings suggest that implementation of a combined influenza/RSV testing led to a significant increase in detection rate, supported clinicians establishing the correct diagnosis and allowed a safe and controlled handling of RSV patients.

The respiratory syncytial virus (RSV), a single-stranded RNA virus, is primarily known as a major cause of lower respiratory tract infection in infants and young children1. In the recent past, numerous studies have demonstrated significant disease burden in adults, especially if immunocompromised, suffering from an oncological disease, chronic obstructive pulmonary disease, severe asthma or congestive heart failure2,3. Shi et al. estimated a global number of 336,000 hospitalized patients per year due to RSV in the group of older adults (≥ 65 years)4.

Previously, an overwhelming amount of studies reported similar rates of hospitalization and mortality of influenza virus A and B compared to RSV5–8. Similar clinical presentation of both viruses and comparable seasonal characteristics makes a differential diagnosis based on clinical or epidemiological characteristics nearly impossible without adequate diagnostics9. Moreover, both viruses can be transmitted by aerosol droplets or direct contact to the virus, such as over fomites10,11. Hence, prevention protocols have to be implemented to protect health care workers (HCWs) and prevent disease transmission within health care facilities.

Despite overwhelming evidence that RSV is associated with a significant disease burden in adults, clinicians’ awareness is still low12,13. At the Vienna General Hospital, 2776 samples were tested for Influenza A and B, however only 387 samples were tested for RSV during season 2017/2018. In autumn 2018, a combined flu/RSV test was introduced at the Vienna General Hospital. Hence, patients with suspected respiratory tract infection tested for influenza were automatically tested for RSV and treated according to strict infection control and prevention strategies (supplementary Table 1).

Here, we set out to describe RSV disease burden in hospitalized adults at a university hospital, highlight the impact on the clinical practice and elucidate pitfalls of misinterpretation with other respiratory viruses.

Methods

This retrospective cohort analysis assesses the influence of a newly implemented combined influenza/RSV RT-PCR test on the clinical practice at a university hospital. The study protocol was approved by the Ethics Committee of the Medical University of Vienna, Austria (ECS 1523/2019) and all study-related procedures were conducted according to the declaration of Helsinki. This manuscript is reported following the RECORD statement14.

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Data collection and analysis. Firstly, all samples of patients with suspected influence-like illness admitted to the General Hospital during October 2017 to April 2019 were included. Detection rates of viruses (RSV, influenza A and influenza B) were described in dependence of the surveillance period. Secondly, all RSV positively tested patients during October 2017 to April 2019 were included into the data analysis. Figure 1 demonstrates the investigated collective and performed analysis. For the data analysis demographic parameters (age, sex, body-mass index, co-morbidities and history of smoking), microbiological data (urinary antigen testing, blood-culture, bronchoalveolar lavage fluid culture), radiological data (evidence of pneumonia), laboratory parameters (haemoglobin, thrombocytes, leucocytes, creatinine, bilirubin and C-reactive protein), treatment procedures (admission to hospital, administration of antibiotics, admission to ICU) and mortality data were retrospectively extracted from the electronic patient information system. Firstly, diagnostic approach of RSV was compared between the surveillance period 2018 (period before combined influenza/RSV testing), defined as week 40 of 2017 to week 30 of 2018, and the surveillance period 2019 (period with influenza/RSV testing in place), defined as week 40 of 2018 to week 30 of 2019. Secondly, demographic data, handling of the patients and outcome were described. Thirdly, differences of the handling of RSV patients were assessed in dependence of clinicians’ knowledge of the RSV status (RSV status known versus RSV status not known). Therefore, an additional cohort was formed by retrospectively assessing 191 sputum samples of patients admitted to the hospital due to respiratory symptoms during the encompassed peak of 2018 RSV season in Vienna (week 8 to 12) but not tested for RSV infection, for RSV infection. RSV peak was defined in accordance to RSV peak published by Austria’s Sentinel Physician Surveillance Network15,16.

Diagnostic assays performed. Tests were performed differently depending on the season. (1) in season 2018 testing for influenza virus A and B was performed by RT-PCR (in-house protocol established after Ward et al. or Xpert Xpress Flu, Cepheid) and testing for RSV was performed by (MagNA Pure 96 system, TaqMan-based RT-PCR using the LightCycler Multiplex RNA Virus Master [Roche Diagnostics, Rotkreuz, Switzerland]17. RT-PCR was performed using the routine test for respiratory viruses in-house established after Fry et al., but not always in parallel18. (2) in season 2019 RSV infection was diagnosed by a combined Flu/RSV RT-PCR (Xpert Xpress Flu/
RSV XC, Cepheid). (3) 191 samples, not tested for RSV infection during the encompassed peak of 2018 RSV sea-
son in Vienna (week 8 to 12), were reassessed for RSV by RT-PCR with the above mentioned routine protocol.

**Statistical analysis.** Statistical analysis was performed using commercially available computer software
SPSS Statistics 20 (IBM, USA) and figures were produced in R (R Core Team, 2014) using the package ggplot2
(Wickham 2016)\(^1\)\(^9\),\(^2\)\(^0\).

Baseline characteristics of participants are summarized as frequencies and proportions for categorical data
and as means and standard deviations or medians and interquartile for metric data. The hypothesis testing was
performed by Chi-Square test for categorical variables and Student’s t tests or Mann–Whitney tests for metric
data. To illustrate correlations between categorical variables, a Pearson-correlation is performed for normally
distributed variables, as well as Spearman’s Rho for non-parametric variables. Statistical significance was defined
as p < 0.05.

**Ethics declaration.** The study protocol of this retrospective analysis was approved by the Ethics Committee
of the Medical University of Vienna, Austria (ECS 1523/2019) and all study-related procedures were conducted
according to the declaration of Helsinki. Due to the retrospective study design, informed consent was waived by
the Ethics Committee of the Medical University Vienna.

**Results**

**Analysis of respiratory samples collected during season 2018 and 2019.** A total 5196 respiratory
samples were collected, 2660 samples during season 2018 and 2536 samples during season 2019. Performed
tests and results are demonstrated in Fig. 2. Interestingly, there was a significant difference in detection rate
between the seasons for all of the reported viruses. We detected a significant increase (p < 0.001) in influenza A
in season 2019 (326/2390 [13.6%]) compared to season 2018 (192/2583 [7.4%]). Prevalence of influenza B was
significantly lower (p < 0.001) in season 2019 (1/2390 [0.04%]) compared to 2018 (461/2583 [17.8%]). Further
the newly implemented diagnostic strategy led to an increase (p = 0.08) in RSV infection rates in 2019 (111/2403
[4.6%]) compared to season 2018 (10/391 [2.6%]).

**Descriptive analysis of RSV patients.** In total 103 patients with RSV infection were included into the
analysis. Baseline characteristics are highlighted in Table 1. The median age of our cohort was 57 years and
female and male patients were equally affected. Most of the patients suffered from comorbidities, such as cardiac
illness (n = 54), pulmonary illness (n = 33), oncological disease (n = 24), type 2 diabetes mellitus (n = 20) and
terminal dialysis-dependent kidney insufficiency (n = 13). 19 patients were solid-organ transplant recipients.
Nearly half of the cohort (43.7%) required in-hospital care. Risk factors for hospitalization in RSV patients
were, advanced age (> 65a) (p < 0.001), smokers (p < 0.001), comorbidities (p < 0.001), clinical or radiological
signs for pneumonia (p = 0.009) and signs for superinfection (0.014) (supplementary Table 2). Inpatients had
significant higher CRP values then patients outpatients (inpatients 1.3 mg/dl [0.52–2.91], outpatients 5.25 mg/dl [1.22–11.81]; p = 0.001) (supplementary table 3). Complications during hospitalization were most frequently worsening of pulmonary disease, such as COPD exacerbation in 16 patients, followed by cardiac decompensation in two patients. Superinfections were reported in 5 patients. *Streptococcus pneumoniae* detected by rapid urine antigen testing was the most common pathogen, followed by *Klebsiella pneumoniae* detected in blood culture (n = 1), *Enterococcus faecalis* detected in blood culture (n = 1) and *Aspergillus fumigatus* detected in a bronchoalveolar lavage fluid culture (n = 1).

Antimicrobial treatment was initiated in 29 of 103 (28.2%) patients. Decision for antimicrobial treatment was based on several factors as demonstrated in Fig. 3. Further, antimicrobial treatment was more likely to administered in patients with higher CRP values (no antibiotics prescribed CRP = 1.28 mg/dl [0.52–2.9], antibiotics prescribed CRP = 9 mg/dl [4.56–17.3]; p < 0.001).

Of the patients admitted for in-patient care 15.56% (7/45) needed further treatment and an ICU ward. Patients with preexisting pulmonary (p = 0.03), complications of RSV infection (p = 0.001), pneumonia (p = 0.002) and bacterial superinfection (p = 0.036) were more likely to be admitted to ICU. Overall 2.9% of the patients died. Age over 65 was associated with higher odds for death. Table 2 demonstrated odds for hospitalization, ICU admission and mortality in dependence of several factors, such as age, smoking status, cardiac disease, pulmonary disease, type 2 diabetes mellitus, dialysis, oncological disease, solid-organ transplant recipient, complications aside RSV, pneumonia and superinfections.

**Assessment of the impact of clinicians’ knowledge of RSV status on clinical practice.** Finally, we reassessed 191 samples respiratory samples, negatively tested for influenza during RSV high peak during season 2018 but not tested for RSV. Indeed, we revealed 10 additional RSV cases. Handling of those ten patients (patients with unknown RSV infection) was compared to the handling of all other RSV patients (patients with known RSV infection). We did not detect differences in rates of hospitalization (patients with known RSV infection = 45/103 [43.7%], patients with unknown RSV infection = 4/10 [40%], p = 1.000) or frequency of antimicrobial administration (patients with known RSV infection = 29/103, 28.2%, patients with unknown RSV infection = 2/8, 20%, p = 0.725).

**Discussion**

This study was conducted to assess the influence of a newly implemented combined flu/RSV test at the biggest tertiary care center in Austria. As expected, introduction of the test revealed that RSV is responsible for a substantial fraction of severe respiratory illnesses in adults during influenza season. Mortality in our cohort was

| Characteristics | RSV patients (n = 103) |
|-----------------|-----------------------|
| Age—year median (Q1–Q3) | 57 (40–73) |
| Female sex—no. (%) | 54 (52.4) |
| BMI median (Q1–Q3) | 25.3 (21–31.1) |
| Smoking—no. (%) | 32 (31.1) |
| **Comorbidities—no. (%)** |       |
| Respiratory illness | 33 (32) |
| Cardiac illness | 54 (52.4) |
| Type 2 diabetes mellitus | 20 (19.4) |
| Dialysis | 13 (12.6) |
| Oncological disease | 24 (23.3) |
| SOT | 19 (18.4) |
| Clinical or radiological findings for pneumonia—no. (%) | 18 (17.5) |
| Respiratory complications—no. (%) | 20 (19.4) |
| Cardiac complications—no. (%) | 2 (1.9) |
| **Superinfections—no. (%)** |       |
| Positive blood culture | 2 (1.9) |
| Rapid urine antigen testing for *S. pneumoniae* | 3 (2.9) |
| Positive broncho-alveolar lavage fluid culture | 1 (1) |
| Antibiotics administered | 29 (28.2) |
| Hospital admission | 45 (43.7) |
| ICU admission—no. (%) | 7 (6.8) |
| Length of stay—median (Q1–Q3) | 13 (4–23) |
| Death—no. (%) | 3 (2.9) |

Table 1. Demonstrates the baseline demographics of RSV patients. BMI, body-mass-index; SOT, solid-organ transplantation; ICU, intensive care unit.
high, and was even higher in patients with advanced age. Despite early and broad testing of our patients we were not able to safely discern any influence on admission rates to hospital or administration of antimicrobial agents. Previously published studies demonstrated a particularly high disease burden of RSV disease burden in patients aged > 75 years. Consistent with previous reports, our patients were mostly of advanced age and suffered from chronic diseases, such as respiratory illness, cardiac illness, type 2 diabetes mellitus, oncological disease, terminal dialysis-dependent kidney insufficiency and solid-organ transplant recipients, bacterial superinfections (bacterial growth in blood cultures) and age > 65a.

![Figure 3. Demonstrates factors influencing administration of antibiotics in patients with RSV infections.](image)

**Table 2.** Highlights odds-ratios for different variables depending on outcome parameters hospitalization, ICU admission and mortality. SOT, solid-organ transplantation; ICU, intensive care unit.

|                | Hospitalization | ICU admission | Mortality |
|----------------|-----------------|---------------|-----------|
|                | OR (95% CI)     | p-value       | OR (95% CI) | p-value   | OR (95% CI) | p-value   |
| Age > 65a      | 5.25 (2.2–12.5) | <0.001        | 2.43 (0.51–11.51) | 0.42 | 1.13 (1–12.74) | 0.048 |
| Smoking—yes    | 9.11 (3.4–24.38) | <0.001        | 1.73 (0.36–8.24) | 0.67 | 1.11 (0.1–12.74) | 1 |
| Cardiac illness| 4.03 (1.75–9.28) | 0.001        | 6 (0.2–51.74) | 0.115 | 1.85 (0.16–21.02) | 1 |
| Respiratory illness | 3.35 (1.41–7.96) | 0.006        | 6.07 (1.11–33.16) | 0.033 | 4.45 (0.39–50.95) | 0.24 |
| DM type II     | 2.96 (1.07–8.21) | 0.044        | 1.73 (0.31–9.66) | 0.62 | 9.11 (0.78–106.01) | 0.096 |
| Dialysis       | 3.38 (0.97–11.79) | 0.07         | 0.87 (0.8–0.94) | 0.59 | 0.87 (0.81–0.94) | 1 |
| Oncological illness | 2.17 (0.86–5.49) | 0.108        | 2.68 (0.56–12.92) | 0.349 | 7.09 (0.61–81.9) | 0.135 |
| SOT            | 1.82 (0.66–5.02) | 0.31         | 0.68 (0.08–5.97) | 1 | 0.8 (0.72–0.89) | 1 |
| Complications  | –               | –            | 21.43 (2.44–187.98) | 0.001 | 6 (0.52–69.03) | 0.167 |
| Pneumonia      | 4.3 (1.4–13.21) | 0.009        | 15.96 (2.8–91.02) | 0.002 | 10.5 (0.9–122.81) | 0.078 |
| Superinfection | –               | –            | 12.4 (1.67–91.87) | 0.036 | 12 (0.89–161.65) | 0.14 |

Despite an increasing number of reports highlighting the disease burden of RSV in adults, awareness of many health care providers to adults seems to be low. The clinical presentation of patients with RSV and influenza is non-specific and similar, which makes it almost impossible to differentiate between the viruses based on symptoms alone. Furthermore, RSV and influenza virus A and B epidemics show similar seasonality with most synchronized peaks in temperate regions. Chronological sequence of the viruses’ peak incidence differs depending on the country and is not necessarily consistent from year-to-year. In Austria, the RSV epidemic starts later than influenza virus A. As a result, clinicians in emergency rooms and ambulances had to separate patients with respiratory tract infections due to RSV and influenza quickly and reliably to prevent nosocomial transmission. We demonstrate that a combined test led to an expected increase in performed tests and detected cases, but also to an increase in detection rate from 2.6 to 4.6%. We believe that the demonstrated increase in
detection rate emphasizes these diagnostic difficulties and supports the application of combined PCR tests in the clinical practice.

Broad testing did change the management of patients. Positively tested patients were admitted to distinct wards and had to remain in quarantine for 5 days. Further, health care workers in contact with those patients had to wear personal protective equipment. Herewith we achieved a safe and controlled handling of RSV patients. Although, the implemented strategies suggest a reduction of nosocomial infection, the study design does not allow causative conclusions. Apart from the implemented measures, the impact of the test on the clinical reasoning is difficult to quantify. In theory, detection of viral pathogens responsible for pneumonia should reduce administration of antimicrobial treatment and prevent nosocomial infections. Coinciding data from previously published retrospective studies showed that increased viral testing only partially altered antimicrobial treatment\textsuperscript {23,24}. Here, we demonstrate that reasoning for ab treatment depends on multiple factors such as an elevated CRP, clinical and radiological indication for pneumonia, as well as age of the patient rather than on the clinicians’ knowledge about RSV/test result.

Due to increasing availability and potency of viral diagnostic tools, viruses are increasingly detected as a cause for pneumonia world-wide. Clinical influence of broad viral testing was often discussed controversially, mostly due to three factors. Firstly, apart from neuraminidase inhibitors for influenza virus infections no specific antiviral options exists\textsuperscript {25}. Secondly, attempts to provide convincing evidence that broad antiviral testing reduces rates of antimicrobial treatment were unsuccessful\textsuperscript {23,24}. Thirdly, interpretation of naso-pharyngeal swabs poses a challenge of its own as the detected virus could be responsible for co-existing upper respiratory tract colonization or an actual pneumonia pathogen, making diagnosis difficult\textsuperscript {25}. The above points should not be interpreted in the sense that testing is unnecessary, but rather that further research is urgently necessary to improve diagnostic procedures and the availability of treatment options. Currently, however, the biggest argument for broad-based testing remains the prevention of the spread of disease and protection of HCWs.

We acknowledge several limitations. First, our control group of patients with unknown RSV status was relatively small, as we only detected 10 additional cases. Hence, we only compared 10 patients from season 2018 to 103 cases in season 2019 in terms of clinical practice and reasoning, which is a limiting factor of our study. However, it is plausible clinicians’ select antimicrobial treatment based on multiple factors such as CRP, indication for pneumonia, age and not only on a test result. Furthermore, the assessed 191 samples corresponded to all samples during the suspected peak of RSV season in 2018, week 8 to 12. Hence, analysis of other periods would have been associated with increased efforts regrading costs and time. Secondly, we did not broadly test patients after the isolation period and cannot ultimately conclude if implemented strategies proved feasible in reducing risk of transmission after the isolation period. Thirdly, the present study was conducted retrospectively. Hence, study results were depended on the quality of documentation done by clinicians and does not allow conclusion about consequences not documented as, such as source of infection (nosocomial versus community acquired) and quality of care performed.

In conclusion, we demonstrated that although RSV disease is a well described cause for pneumonia in adults. Implementation of a combined influenza/RSV test led to a significant increase in detection rate, supported clinicians establishing the correct diagnosis and allowed a safe and controlled handling of RSV patients.

Data availability
The datasets generated during and/or analyzed during the current study are available from the corresponding author on reasonable request.

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**Author contributions**

L.S. and C.S. wrote the main text, J.S. and L.S. contributed for data acquisition, A.N.H. and R.S. performed the diagnostic assays, L.S. and F.L. prepared Figs. 2, 3 and S.T., M.K. and M.R. contributed for statistical analysis and interpretation of the data. All authors reviewed the manuscript.

**Competing interests**

The authors declare no competing interests.

**Additional information**

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