Impact of educational training and C-reactive protein point-of-care testing on antibiotic prescribing in rural and urban family physician practices in Latvia: a randomised controlled intervention study

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Abstract

Background: Although self-limiting viral infections are predominant, children with acute infections are often prescribed antibiotics by family physicians. The aim of the study is to evaluate the impact of two interventions, namely C-reactive protein point-of-care testing and educational training, on antibiotic prescribing by family physicians.

Methods: This randomised controlled intervention study included acutely ill children consulted by 80 family physicians from urban and rural practices in Latvia. The family physicians were divided into two groups of 40. The family physicians in the intervention group received both interventions, i.e. C-reactive protein point-of-care testing and educational training, whereas the family physicians in the control group continued to dispense their standard care. The primary outcome measure was the antibiotic prescribing at the index consultation (delayed or immediate prescription) in both study groups. The secondary outcome was CRP testing per study group. Patient- and family physician-related predictors of antibiotic prescribing were analysed as associated independent variables. Practice location effect on the outcomes was specially addressed, similar to other scientific literature.

Results: In total, 2039 children with acute infections were enrolled in the study. The most common infections observed were upper and lower respiratory tract infections. Overall, 29.8% (n = 607) of the study population received antibiotic prescription. Our binary logistic regression analysis did not find a statistically significant association between antibiotic prescriptions and the implemented interventions. In the control group of family physicians, a rural location was associated with more frequent antibiotic prescribing and minimal use of CRP testing of venous blood samples. However, in the intervention group of family physicians, a rural location was associated with a higher level of C-reactive protein point-of-care testing. Furthermore, in rural areas, a significant reduction in antibiotic prescribing was observed in the intervention group compared with the control group (29.0% (n = 118) and 37.8% (n = 128), respectively, p = 0.01).

Conclusion: Our results show that the availability of C-reactive protein point-of-care testing and educational training for family physicians did not reduce antibiotic prescribing. Nevertheless, our data indicate that regional variations in antibiotic-prescribing habits exist and the implemented interventions had an effect on family physicians practices in rural areas.
**Introduction**

Acute infection in children is one of the most common reasons for attending family physicians (FP) and these visits often result in antibiotic prescription. The most frequent indication for antibiotic use is respiratory infection, with the highest incidence rate for very young children (up to 2 years old) [1] despite viral aetiology predominance in this age group that does not require specific treatment [2]. Indeed, at least 30% of antibiotics prescribed in outpatient settings are considered to be unnecessary [3].

Recent studies have shown that inappropriate antibiotic usage may be due to several reasons. These include difficulties differentiating between viral and bacterial diseases based on clinical signs alone [4, 5], fear of complications or missing serious bacterial infections, heavy workloads and even parental insistence on antibiotics being prescribed [3]. In order to optimise outpatient antibiotic prescribing, the effectiveness of several different types of interventions has been assessed. These include patient and FP education, communication training, point-of-care testing (POCT), active prescription monitoring and delayed prescribing [3]. However, as a multidirectional combination of interventions is more likely to reduce unnecessary antibiotic prescribing than a single intervention [3, 6, 7], we conducted the present study to evaluate the effect of two interventions, namely access to C-reactive protein (CRP) POCT in FP practices and educational training for FP, focused on targeted antibiotic prescribing.

CRP is an acute phase protein that can reduce diagnostic uncertainty. In recent years, it has been more widely used for POCT in routine primary care practice in several European countries [8]. The main advantages of CRP POCT are its ease of use, rapid feedback of the test result allowing immediate decision-making on whether or not to prescribe antibiotics and higher patient satisfaction with the child-friendly finger prick test instead of an invasive venous puncture [9]. In Latvia, CRP POCT is currently available in only a few FP practices and not state covered. In the main, venous blood samples are sent to laboratories for CRP detection. However, especially in rural areas of Latvia, the result might not be reported until the next working day, which is often too long a delay for acutely ill patients and consequently the decision to initiate antibacterial treatment is frequently based upon clinical examination alone. A previous meta-analysis has shown that CRP POCT is associated with a lower antibiotic prescription rate for adults with respiratory tract infections in primary care [10]. However, the findings regarding CRP POCT for children with acute illnesses are presently incomplete and controversial and thus this subject requires further analysis [11]. Additionally, interventions such as patient, parent and physician education and communication training have also been shown to have an impact on reducing antibiotic prescribing [12]. However, data on impact of such interventions on antibiotic prescription specifically by FP s is scarce.

In this study as the second intervention, we included educational training of FP s based on new recommendations for the management of respiratory infections as well as fever for children that have been recently introduced in Latvia. Furthermore, although the Happy Audit study has previously reported that interventions focused on patient or FP education and CRP POCT for adults reduce inappropriate antibiotic prescriptions [13–15], the effectiveness of these two types of interventions on antibiotic prescribing has not been evaluated in a paediatric population.

**Materials and methods**

This randomised controlled intervention study was conducted in Latvia between November 2019 and February 2020. The aim of the study was to evaluate the effect of the combination of two interventions – access to CRP POCT in FP practices and educational training for FP – on the antibiotic prescribing rate of FP for acutely ill children. Patient- and FP-related predictors of antibiotic prescribing were also analysed.

**Participating family physicians**

There are approximately 360,000 children (age 0–17 years) in Latvia and their health needs are served by an estimated 1268 FP. FP are self-employed, usually independently located and serve an extensive age range of children. For the purposes of this study, we had access to 40 CRP POCT devices, enabling 80 FP to be recruited. The FP selection process has previously been reported [16]. Briefly, the participating FP were recruited using two approaches. First, from the country’s 1268 FP, by means of an Excel random-number generator, we selected 160 FP (the expected response rate was 50%) across different geographically located practices (urban and rural areas) and sent invitations to participate in the study via both email and paper-based letter form. Unfortunately, the response rate was lower than expected and only 38 participants were recruited using this approach. Secondly, we directly addressed FP at a meeting of the
Latvian Family Physicians Association and consequently achieved the requisite number of 80 participants. It was expected that each FP might see about 30 suitable patients during the duration of the study.

Interventions
FP were stratified according to practice location and each stratum was divided into two groups of 40 FP using random numbers generated by MS Excel Random Number function. Practices in the intervention group received both interventions (CRP POCT and educational training). Specifically, each FP in this group received a CRP POCT device for use during the duration of the study and was individually tutored by diagnostic test company on how to perform the CRP test during a face-to-face meeting, and ongoing support by the company was available to the FP. In addition, FP were contacted proactively by the study team to address any issues. We used the Orion Diagnostica QuikRead go CRP POCT system for the quantitative determination of CRP in blood with a sample volume of 20μl obtained via a finger prick. This system has a measuring range of 5–200mg/L and the result is available within 2 minutes. As CRP cut-off levels for children in primary care are currently undetermined [17], the FP did not receive any guidelines on the interpretation of results. FP in the intervention group were allowed to order a CRP test based on individual indications if they believed the result would help them make a more informed decision on antibiotic necessity after a clinical assessment. Other tests such as rapid strep test, urine dipstick test or laboratory analyses were also available as usual; however, the availability of testing was different between urban and rural practices.

FP in the intervention group also received educational training based on new recommendations for the management of respiratory infections and fever in children introduced in 2019 in Latvia. The key topics were:

- child with fever – evaluation, precautionary level system and management,
- child with upper and lower respiratory infection – evaluation and management,
- principles of antibiotic resistance and safer prescribing of antibiotics.

This intervention involved one four-hour training seminar, followed by educational materials in video and printed format. FP also received parent information booklets about managing children with fever at home and signs to look out for that indicate a FP should be contacted.

FP in the control group received no interventions and continued to dispense their standard care.

Participating children
FP were asked to record the data of consecutive children (1 month up to 17 years old) who attended their practice for a face-to-face visit with current clinical signs of an acute infection that had been present for less than 5 days. Patients were excluded if they were aged under 1 month, had received antibiotics prior to the visit or were already in the reconvalescent stage of disease.

Sample size
We presumed that the frequency of antibiotic prescribing in the intervention group compared to the control group was 34 and 42%, respectively, as per Martínez-González et al. [18]. Thus, according to Fleiss et al. [19], with 80% power and α level 5%, our study required 571 patients in each study group.

Data collection
FP collected data in anonymised form. The variables recorded included whether or not an antibiotic was prescribed, patient demographics, and diagnosis based on a pre-defined list (upper respiratory infections (common cold, rhinosinusitis, otitis, pharyngitis, tonsillitis, stomatitis, laryngitis), lower respiratory infections (bronchitis, bronchiolitis, pneumonia), gastrointestinal infections, urinary tract infections, skin and soft tissue infections and joint and bone infections) and the diagnostic tests undertaken prior to the initiation of antibiotic treatment (e.g. CRP POCT, CRP measurement from a venous blood sample, full blood count, urine dipstick test and microscopy, Group A streptococcal rapid antigen test, rapid influenza diagnostic test, bacteriological cultures, X-ray).

The primary outcome measure was the antibiotic prescribing at the index consultation (delayed or immediate prescription) in both study groups. The secondary outcome was CRP testing per study group. Patient- and family physician- related predictors of antibiotic prescribing were analysed as associated independent variables. Practice location effect on the outcomes was specially addressed, similar to other scientific literature.

Statistical analyses
Descriptive statistics, such as means (with standard deviations) and medians (with interquartile range (IQR)) for continuous variables and proportions for categorical variables, were calculated. For determination of the statistical significance of differences in the proportions of dependent variables between subgroups of independent variables, the Chi-square test was used. Normal distribution was tested using the Kolmogorov-Smirnov test. To identify factors associated with antibiotic prescription or CRP testing, binary logistic regression was used.
Results were considered as statistically significant if $p < 0.05$. Data processing was performed using IBM SPSS Statistics (Statistical Package for the Social Sciences, Version 23.0).

**Results**

Initially, 80 FP started the study; however, 5 FP from the control group declined to participate further after randomisation. The mean age of the FP was 51.9 years and the majority were female. Considerable heterogeneity existed regarding the length of time working in FP practice (ranging from 1 year to 52 years) and the number of paediatric patients registered at practices (ranging from 48 to 1843 children). Table 1 details the characteristics of the FP in the intervention and control groups. There were no significant differences between the two groups regarding the age, sex and work experience of the FP, number of registered paediatric patients and practice location.

| Variables                        | Intervention group ($n = 40$) | Control group ($n = 35$) |
|----------------------------------|------------------------------|--------------------------|
| Age (years)                      |                              |                          |
| Median                           | 52.5 (IQR 46.3–59.8)         | 53.0 (IQR 46.0–61.0)     |
| Sex                              |                              |                          |
| Male                             | 1 (2.5%)                     | 1 (2.9%)                 |
| Female                           | 39 (97.5%)                   | 34 (97.1%)               |
| Work experience (years)          |                              |                          |
| Mean                             | 25.4 (SD 13.1)               | 24.6 (SD 11.9)           |
| Proportion of children on patient list (%) |          |                          |
| Median                           | 24.3 (IQR 16.7–43.4)         | 24.2 (IQR 16.9–38.1)     |
| Location                         |                              |                          |
| Rural areas                      | 14 (35.0%)                   | 10 (28.6%)               |
| Regional cities                  | 10 (25.0%)                   | 8 (22.9%)                |
| Capital of Latvia                | 16 (40.0%)                   | 17 (48.6%)               |

**Fig. 1** Flowchart of the study's recruitment process. FP: family physician.
During the three-month study period, 2347 patients were recruited; however, 308 patients were excluded due to symptom duration of more than 5 days or missing information concerning diagnoses (Fig. 1). Therefore, a total of 2039 patients met the inclusion criteria (1153 patients in the intervention group and 886 in the control group). The mean number of included patients per FP was 27.2.

The mean age of the patients was 6.1 years. Boys comprised 50.9% of the study participants. The mean duration of illness was 2.6 days. Only 8.7% of patients had a chronic disease and bronchial asthma was the most common (84.8%). Table 2 summarises the characteristics of the patients in both study groups. There were small imbalances between the groups regarding age, duration of symptoms, frequency of chronic disease and vaccination status.

No significant difference was found in the distribution of diagnoses between the two groups. The most common infections were upper respiratory tract infections (78.3% (n = 1597) of patients) and lower respiratory tract infections (18.8% (n = 384)). Gastrointestinal (1.8% (n = 36)), urinary tract (0.9% (n = 18)), skin and soft tissue (0.1% (n = 3)), and bone and joint (0.05% (n = 1)) infections featured to a much lesser extent.

Overall, 29.8% (n = 607) of the study population received antibiotic prescription. The proportions of patients treated with antibiotics for each type of infection are shown in Fig. 2.

Comparing the two study groups, 27.8% (n = 320) episodes prompted antibiotic prescription in the intervention group, whereas it was 32.4% (n = 287) in the control group. This difference was statistically significant (p < 0.02).

Table 3 shows the patient- and FP-related predictors of antibiotic prescribing, also including the interventions as a single factor. Our data showed that antibiotic prescribing was significantly associated with younger children (adjusted odds ratio (aOR) for children aged 10–14 years vs. 0–4 years was 0.62, p = 0.002), middle-aged FP (aOR for FP aged 41–50 years vs. 30–40 years was 1.76, p = 0.002), a rural location of the FP practice (aOR was 1.42, p = 0.01 compared to the capital city) and a larger number of registered paediatric patients (aOR was 1.68, p < 0.001 and 1.59, p = 0.02 for 501–1000 and 1001+ patients, respectively, compared to < 500 patients).

Our binary logistic regression analysis did not find a statistically significant association between antibiotic prescriptions and the implemented interventions. However, subgroup analysis by location of practice showed a significant reduction in antibiotic prescribing in the intervention group compared to the control group in rural areas (29.0% (n = 118) and 37.8% (n = 128), respectively, p = 0.01). The proportions of antibiotic prescriptions in relation to location of practice in both study groups are presented in Table 4.

The CRP level was frequently measured in the intervention group; 72.4% (n = 835) of episodes, with CRP POCT being preferred over standard laboratory testing of a venous blood sample (99.0% (n = 827) and 1.0% (n = 8), respectively). For 8 patients with CRP POCT testing of a venous blood sample was also provided. In contrast, the CRP level was measured in just 8.8% (n = 78) of episodes in the control group, where only venous blood sample testing was available. Furthermore, 79.4% of antibiotic prescriptions were preceded by CRP testing in the intervention group compared with only 12.5% in the control group.

| Table 2 Characteristics of patients according to the study groups |
|---------------------------------------------------------------|
| Variables          | Intervention group (n = 1153) | Control group (n = 886) |
| Age (years)        |                               |                        |
| Median             | 5.0 (IQR 3.0–9.0)             | 5.0 (IQR 2.0–8.0)      |
| 0–4 years          | 484 (42.6%)                   | 431 (49.4%)            |
| 5–9 years          | 383 (33.7%)                   | 279 (32.0%)            |
| 10–14 years        | 186 (16.4%)                   | 120 (13.8%)            |
| 15–17 years        | 82 (7.2%)                     | 42 (4.8%)              |
| Sex                |                               |                        |
| Boys               | 591 (51.6%)                   | 440 (50.0%)            |
| Girls              | 555 (48.4%)                   | 440 (50.0%)            |
| Duration of illness (days) | 3.0 (IQR 2.0–4.0) | 3.0 (IQR 2.0–4.0)      |
| Chronic disease    |                               |                        |
| Yes                | 75 (6.5%)                     | 102 (11.5%)            |
| No                 | 1078 (93.5%)                  | 784 (88.5%)            |
| Full vaccination   | 1046 (92.7%)                  | 820 (95.0%)            |
| Partial vaccination| 69 (6.1%)                     | 40 (4.6%)              |
| No vaccination     | 13 (1.2%)                     | 3 (0.3%)               |
| Diagnoses          |                               |                        |
| Upper respiratory infection | 922 (80.0%) | 675 (76.2%) |
| Lower respiratory infection | 204 (17.7%) | 180 (20.3%) |
| Gastrointestinal infection | 17 (1.5%)     | 19 (2.1%)            |
| Urinary tract infection | 8 (0.7%)          | 10 (1.1%)             |
| Skin and soft tissue infection | 2 (0.2%)     | 1 (0.1%)             |
| Bone and joint infection | 0 (0.0%)            | 1 (0.1%)               |
| Ambulatory patients | 1136 (98.5%)                | 879 (99.2%)            |
| Referred to hospital | 17 (1.5%)               | 7 (0.8%)               |

* Denominators may vary due to the missing values.
binary logistic regression analysis found a significant association between higher usage of CRP testing and rural location of FP practice in the intervention group (aOR was 2.51, \( p < 0.001 \)). However, the relationship in the control group was the opposite, with patients more likely to have their CRP level tested in laboratories in urban areas than in rural areas (aOR was 0.05, \( p < 0.001 \)) (Table 5).

The majority of patients who underwent CRP testing – 78.9% (\( n = 727 \)) – were found to have a CRP level < 20 mg/L, whereas 15.3% (\( n = 141 \)) had a level between 20.01 and 50 mg/L, 4.5% (\( n = 41 \)) between 50.1 and 99 mg/L, and 1.3% (\( n = 12 \)) > 100 mg/L. Of note, 28.4% (\( n = 317 \)) of all patients who did not undergo CRP testing were prescribed antibiotics by FP. Furthermore, of the patients with a CRP level < 20 mg/L, 19.9% also received antibiotics. Overall, antibiotic prescribing increased with increasing measured CRP level (\( p < 0.0001 \)).

**Discussion**

The aim of this study was to evaluate the effect of CRP POCT in FP practices and educational training for FP on antibiotic prescribing for children in primary care – a two-intervention combination that has not previously been studied in a paediatric population. CRP POCT is already routinely used in primary care by several other countries [8]. A meta-analysis has previously shown that CRP POCT significantly reduces immediate antibiotic prescribing for adults compared with standard care [18]; however, evidence for the use of CRP POCT to guide antibiotic prescribing for children is currently lacking. Nonetheless, a reduction in antibiotic prescribing for children has been reported in studies in which guidance for CRP result interpretation was provided [20–22]. Educational training of primary health care providers or parents on appropriate antibiotic usage has been shown to be effective in changing the antibiotic-prescribing behaviour [23, 24]. Multifaceted interventions, including educational training for physicians and patients, communication skills training and the introduction of POCT into clinical practice, have the potential to reduce inappropriate antibiotic usage even further [7, 15].

Although we did not find a statistically significant association between antibiotic prescriptions and the implemented interventions in regression analysis, analysing the data by location of practice revealed that the interventions significantly reduced antibiotic usage in rural areas (29.0% in the intervention group versus 37.8% in the control group). To the best of our knowledge, this is the first time the effectiveness of CRP POCT and educational
training on antibiotic prescribing has been analysed by practice location in a paediatric population.

The overall effect was smaller than anticipated when comparing the intervention and control groups; however, this should be seen in the light of an overall low antibiotic prescribing rate. Other studies have pointed out that interventions may be more beneficial in cases of generally higher prescribing [25]. The observed antibiotic prescribing rate was lower than expected in both study groups (27.8% in the intervention group and 32.4%)}
in the control group). Dumpis et al. previously reported an antibiotic prescribing rate of 42% in Latvia and 53% in Lithuania [26]. It is possible that during the study period our recruited FP were more inclined to avoid prescribing antibiotics. However, despite a low general antibiotic prescribing rate, significant variations were detected between rural and urban areas. Higher overall antibiotic prescribing rates were associated with practices located in rural areas (aOR was 1.42, \( p = 0.01 \)) compared with the capital city. This finding is consistent with those of previous studies where rurality was found to be a risk factor for inappropriate and more frequent prescribing [3, 27, 28]. More extensive use of antibiotics in rural areas may be explained by fewer FP per head of population, increased workload, limited access to laboratory testing with consequent diagnostic uncertainty [28] and fears of missing secondary bacterial infections which may occur when patients are unable to access medical care [29].

We observed a markedly higher level of CRP testing in the intervention group (72.4% compared to the control group (8.8%). Furthermore, antibiotic prescriptions were preceded by CRP testing far more frequently in the intervention group. Moreover, in line with previous studies [9], FP in the intervention group almost exclusively used POCT (99.0%) for CRP level measurement rather than laboratory testing of a venous blood sample. This highlights user friendliness, patient satisfaction and clinical utility as the main advantages of CRP POCT. The majority of FP did not have any previous experience of using CRP POCT and so perhaps were more interested in trying it during the study period, thus resulting in the observed high usage level in the intervention group. Having said that, other studies have reported that CRP testing is also widespread for self-limiting viral infections in countries where POCT is available [30]. For comparison, in Sweden the level of CRP is measured in up to 50% of all consultations for respiratory infections [25]. The very low CRP testing level in the control group, especially in rural areas, may be due to differences in the availability of testing and timing of reporting of test results from central laboratories.

Although we observed a low rate of antibiotic prescribing, unnecessary antibiotic prescribing still occurs. Antibiotics were often prescribed at the early stage of disease as we only included patients with a symptom duration of less than 5 days (median duration 3 days). Overall, the duration of symptoms was not associated with higher antibiotic prescribing. We found that antibiotics were still widely used for self-limiting upper respiratory tract infections (26.0% in the control group and 23.1% in the intervention group), despite the majority of tested patients having a low CRP concentration and consequently not requiring antibacterial treatment. Moreover, for 19.9% of patients with a CRP level < 20 mg/L, antibiotics were prescribed. These data indicate that FP require more experience and guidance interpreting CRP level results.

### Strengths and limitations
The FP response rate was lower than expected from random selection and it is possible that the participating FP may have been more active and inclined to perform well. Nevertheless, our study included FP with different paediatric patient counts, work experience and practice localities, thus providing widespread information about the country. Moreover, as CRP POCT is not currently integrated into primary care in Latvia, it was valuable to evaluate its effect.

Consistent with other studies [31], we observed a lower antibiotic prescribing rate than expected. This may have been due to knowledge regarding the aim of the study.

### Table 4 Antibiotic prescribing according to FP practice location in the two groups

| Location of practice | Intervention group | Control group | P  |
|----------------------|--------------------|---------------|----|
| Rural areas          | 118 (29.0)         | 128 (37.8)    | 0.01 |
| Regional cities      | 69 (26.1)          | 50 (32.1)     | 0.19 |
| Capital of Latvia    | 133 (27.6)         | 109 (27.9)    | 0.93 |

| Location of practice | Adjusted OR* (95% CI) | Adjusted OR* (95% CI) | P  |
|----------------------|-----------------------|-----------------------|----|
| Rural areas          | 2.51 (1.57–4.00)      | 0.05 (0.02–0.17)      | < 0.001 |
| Regional cities      | 0.39 (0.27–0.57)      | 0.99 (0.53–1.86)      | 0.99 |
| Capital of Latvia    | 1                     | 1                     | 1   |

*Adjusted OR: adjusted odds ratio – adjusted for patient-related factors (age, sex, duration of symptoms) and FP-related factors (age, sex, work experience, number of registered paediatric patients), except the age of FP due to the collinearity with the duration of the career of FP

### Table 5 Binary logistic regression analysis of the effect of location of the FP practice on CRP testing in the two groups

| Intervention group | CRP testing n (%) | Adjusted OR* (95% CI) | P  |
|--------------------|-------------------|-----------------------|----|
| Location of practice | CRP testing n (%) | Adjusted OR* (95% CI) | P  |
| Rural areas        | 370 (90.9)        | 2.51 (1.57–4.00)      | < 0.001 |
| Regional cities    | 145 (54.9)        | 0.39 (0.27–0.57)      | < 0.001 |
| Capital of Latvia  | 328 (68.0)        | 1                     | 1   |

| Control group | CRP testing n (%) | Adjusted OR* (95% CI) | P  |
|---------------|-------------------|-----------------------|----|
| Location of practice | CRP testing n (%) | Adjusted OR* (95% CI) | P  |
| Rural areas        | 3 (0.9)           | 0.05 (0.02–0.17)      | < 0.001 |
| Regional cities    | 20 (12.8)         | 0.99 (0.53–1.86)      | 0.99 |
| Capital of Latvia  | 55 (14.1)         | 1                     | 1   |

*Adjusted OR: adjusted odds ratio – adjusted for patient-related factors (age, sex, duration of symptoms) and FP-related factors (age, sex, work experience, number of registered paediatric patients), except the age of FP due to the collinearity with the duration of the career of FP
influencing participants’ prescribing habits. However, this factor would be expected to affect both study groups equally.

To reflect real daily practice, we asked the FP not to use POCT for all patients but only for those they were unsure about prescribing antibiotics for and felt that additional testing would help their decision-making. This is in line with other studies where testing was not recommended for all patients but restricted to those believed to be at higher risk following clinical assessment [21]. Previous studies have shown that for patient’s risk following clinical assessment, additional testing would help their decision-making. Nevertheless, the FP ordered CRP testing frequently, possibly because the test was not accessible in their practice prior to the study and they were interested in trying it.

A limitation of this study is that FP did not include all patients with acute infection episodes who visited FP during the study period, but still the target number of patients was achieved. Also, we don’t have follow up data of patient’s recovery, hospitalization or subsequent antibiotic prescribing.

Conclusion
Our results show that the availability of CRP POCT and educational training for FP did not reduce antibiotic prescribing. Nevertheless, our data indicate that regional variations in antibiotic-prescribing habits exist and the implemented interventions had an effect on FP practices in rural areas.

In the absence of CRP POCT, especially in rural areas, patients undergo minimal CRP testing prior to antibiotic prescribing, consequently leading to initiation of unwarranted antibacterial treatment.

Abbreviations
FP: Family physicians; POCT: Point-of-care testing; CRP: C-reactive protein; AB: Antibiotic.

Acknowledgements
We would like to thank all the family physicians who participated in the study, as this study could not have been performed without their valuable input.

Authors’ contributions
Conceptualization, Z.L. and J.P.; methodology, Z.L. and J.P.; formal analysis, Z.L.; investigation, Z.L., VS, resources, Z.L., VS, and J.P.; data curation, Z.L. and A.K-U.; writing—original draft preparation, Z.L.; writing—review and editing, J.P. and A.K-U.; visualization, Z.L. and A.K-U.; supervision, J.P.; project administration, J.P. All authors reviewed the manuscript. The author(s) read and approved the final manuscript.

Funding
This study was supported through an educational grant from Riga Stradins University.

Availability of data and materials
The data presented in this study can be viewed on a separate dataset spreadsheet: https://docs.google.com/spreadsheets/d/1kpJRWe91b1S_mwHYbERLCLvbr4t6VVG.

Declarations
Ethics approval and consent to participate
The study was conducted according to the guidelines of the Declaration of Helsinki and was approved by the Research Ethics Committee of Riga Stradins University, approval no. 6–3/5/21 (received: 30 May 2019). Patient consent forms was not relevant to our study according to decision of the the Research Ethics Committee of Riga Stradins University because only the participating FP were exposed to the study interventions, but patients did not undergo any intervention and CRP testing was used only for clinical indications as in routine practice. Patients were informed about the objective of the project, and they were told that specific anonymous clinical information related to the consultation was collected for the study purposes.

Consent for publication
Not applicable.

Competing interests
The authors declare no conflicts of interest.

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Received: 21 April 2022 Accepted: 13 September 2022
Published online: 21 September 2022

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