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Research Article





Evaluation of Antibiotic Prescription for Upper Respiratory Tract Infections in the Community Pharmacy Setting

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Abstract

Objectives: To empirically classify the etiology of URTI; to study the prescription pattern for antibiotics in URTI; and to analyze the necessity and adequacy of prescribed therapy.

Study design: Observational non analytical cross-sectional study.

Study sites: Two community pharmacies located in Almada and in Elvas.

Study population: Patients with an antibiotic prescription for a URTI (self use) and age greater than or equal to 18 years.

Methods: Clinical and therapeutic data were collected during patient observation performed by a training pharmacist, and complemented with information collected through a questionnaire administered face-to-face to patients meeting the inclusion criteria and agreeing to participate. Empirical classifications were developed, one to ascertain the probability of URTI, based on Centor criteria, and another, to judge the need for antibiotic therapy, based on additional criteria considering subpopulations known to be at higher risk. Data was analyzed using the software SPSS, version 20.0. **Results:** The sample included 22 patients (27.3% recruited in Almada and 72.7% in Elvas). The most frequently prescribed antibiotic class was macrolides (54.5%), followed by penicillins (36.4%). In the first group, clarithromycin prevailed (66.7%) and in the second group, the more common was the association amoxicillin/clavulanic acid (75%). Considering the empirical classification developed, it was estimated that only one patient (4.5%) presented signs and symptoms suggestive of URTI with probable bacterial aetiology, and only three patients (13.6%) had an indication for antimicrobial therapy.

Conclusion: The majority of patients observed were classified as having infections with apparently non-bacterial aetiology, for which the prescription of antibiotic would have been probably needless. This alerts to the overuse of antibiotics in the community setting, particularly for URTI, and its contribution to resistance.

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Introduction

Upper respiratory tract infections (URTI) are highly incident infectious diseases, which affect adults 2-5 times per year^[1]. These infections include common cold, sore throat, sinusitis, laryngitis and otitis media and may involve several regions of the upper respiratory tract simultaneously.

URTI comprises seasonal infections, many prevailing during winter season^[2]. Several factors may predispose an individual to acquire these diseases, including his immune state, exposure to pollution or smoking and asthma^[3,4]. Although some studies have confirmed the impact of allergy on rhinosinusitis resulting from a higher number of receptors for rhinovirus on allergic patients, the role of allergy as a predisposing factor for URTI is controversial^[3]. Furthermore, gastroesophageal reflux disease (GERD) may also induce respiratory symptoms (eg. chronic cough), resulting from bronchoconstriction through vagal reflex or by aspiration. Even though GERD can lead to misdiagnosis, it is also considered as a predisposing factor for URTI triggered by chemical injury^[5].



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Although it is relatively difficult to establish a differential diagnosis in pharyngitis, the Centor criteria are highly recognized and commonly used during patient observation. These criteria consider the signs and symptoms most commonly involved in pharyngitis, which can be summarised by: presence of pharyngeal exudates, anterior cervical adenopathy, fever and the absence of cough. Consequently, in the presence of at least three of these features, it is possible to predict that the infection is probably of bacterial origin^[6].

Rhinosinusitis can be classified as acute sinusitis, when the duration of the symptoms is less than one month; subacute, when the symptoms persist for twelve weeks, and chronic when exceeding twelve weeks. Sinusitis may also be classified as recurrent when the individual presents four or more episodes per year^[7].

Antibiotic treatment instituted in these URTI is essentially empirical since the microbiological diagnosis is rarely made, due to the time limitations, difficulty in obtaining clinical samples (sinusitis) and costs of rapid tests (e.g. rapid antigen detection test for group A streptococcus)^[8,9]. Taking into account that URTI are frequently caused by respiratory viruses, antibiotic therapy proves to be unnecessary in most of the cases. Nevertheless, antimicrobials are used mainly to prevent potential complications that may arise from non-antibiotic indication. In Portugal, primary care prescription focuses mainly on beta-lactamic antibiotics, especially the association amoxicillin/clavulanic acid^[10]. Antibacterial therapy may be beneficial to avoid the development of suppurative complications such as tonsillitis, sinusitis, otitis media and cellulite, however delayed antibiotic therapy can be similarly effective in preventing complications when compared to immediate and empirical prescription^[6,11,12].

Furthermore, antimicrobials are indicated in certain circumstances, such as patients who have complications (e.g. pneumonia and mastoiditis), who have a higher risk of developing complications (e.g. renal impairment) and elderly over 65 with cough and presenting two or more Centor criteria, or elderly over 80 with diabetes mellitus, hospital admission in the preceding year, history of heart failure, or under oral glucocorticoid treatment^[13].

This study has focused on pharyngitis and sinusitis because these are the leading causes of acute morbidity in adults in primary care and, the majority of cases for which antimicrobials are prescribed^[2]. Also, URTI are differently expressed in children due to their susceptibility to bacterial otitis media, associated with Eustachian tube dysfunction and immaturity of the immune system^[2-4].

Thereby, the main objective of this study was to evaluate the need for antibiotic treatment, considering the presence of symptoms and previously known risk factors. Two research questions were formulated: "Is the aetiology of the infection, in case of antibiotic prescription, probably of bacterial origin?" and, "Is the prescribed antibiotic therapy for the treatment of URTI, necessary?"

Methods

Study design

This study was implemented through a cross-sectional study (observational non-analytical) that included a survey with clinical observation of the patients. Data gathered by both means allowed empirical classification of the infection aetiology and the need for antibiotic prescription. Data collection was undertaken between November and December of 2014.

Population and sample

The target population included patients aged 18 or older, which represent 79.76% of the total resident population in Portugal^[14]. Paediatric patients were excluded due to specific clinical and treatment issues. Considering that recruitment was pursued in community pharmacies with a minimum capitation of 3500 inhabitants^[15], the study population considered was 2792 inhabitants. Estimation of sample size was calculated considering a population of 2792 inhabitants, a prevalence of the phenomena (p) to be 2.36%, a confidence interval (CI) of 95% and a beta error (d) of 3%. Thus, it was estimated that it would be necessary to recruit a sample (m) of 95 patients. To define the prevalence value (p), it was considered that the prevalence of patients presenting URTI is $15.7\%^{[16]}$; from which 22% (ranging from 11 to 33%) seek health care^[17], resulting in a prescription rate of antibiotic therapy of $68.2\%^{[13]}$.

Eligibility criteria for patient recruitment were as following: Inclusion criteria:

• Patients presenting themselves at the pharmacy where the study was being carried out during the study period;

- Patients presenting a prescription for antibiotic therapy;
- Patients stating to have an URTI;
- Exclusion criteria were:
- Patients refusing to be submitted to a simple physical observation;
- Individuals with a prescription for others;
- Individuals aged ≤ 18 years.

Data collection

Data collection was performed during a face-to-face interview following a questionnaire specifically designed for this purpose (appendix 1). This interview included a clinical observation of the patient performed in a private consultation area. This tool comprised three sections: I) sociodemographic characterization, II) characterization of the patient's present condition (self-reported signs and symptoms) and medical history, and III) clinical observation of signs (observed by the pharmacist). Section II of the questionnaire was developed following the Code of Good Pharmaceutical Practice^[18], and included information about the symptoms, duration thereof, presence of relevant comorbidities and/or prescription of other medicines that could affect disease progression or therapeutic selection. Patient observation included the measurement of body temperature, observation of the pharynx and tonsils, palpation of the neck (detection of cervical adenopathy), palpation of the maxillary and frontal sinuses (pain evaluation) and the visualization of some signs such as rhinorrhoea and sneezing, as suggested by Centor criteria. Both questionnaire and patient observation enabled a complete information collection and were used in the establishment of the probable aetiology of infection and consequent need for antibiotic treatment^[19]. For quality assurance purposes, the principal investigator was submitted to a preliminary training period, which consisted on the clinical observation of a group of volunteer students, with and without symptoms of URTI, under the supervision of two experienced clinical pharmacists.



Table 1: Empirical classification of the aetiology of infection

	Parameters	Assigned weighting	Empirical classification of the etiology of infection
Sore throat	Cervical lymphadenopathy	Yes = 2 No = 0	
	Pharyngeal exudate	Yes = 2 No = 0	
	Cough	Yes = 0 No = 1	
	Severe malaise	Yes = 2 No = 0	 ≤3: Probable viral actiology 4-5: Uncertain outcome
Rhinosinusitis	Duration of symptoms	$\geq 10 \text{ days} = 2 < 10 \text{ days} = 0$	≥ 6 : Probable bacterial aetiology
	Face pain/sensibility	Yes = 1 No = 0	
	Purulent nasal secretions	Yes = 1 No = 0	
General	Fever	$\geq 39^{\circ}C = 2 < 39^{\circ}C = 0$	

Table 2: Empirical classification of the need for antibiotics

	Parameters	Assigned weighting	Empirical classification of the need for antibiotics			
Sore throat	Cervical lymphadenopathy	Yes = 2 No = 0				
	Pharyngeal exudate	Yes = 2 No = 0				
	Cough	Yes $=0$ No $=1$				
	Severe malaise	Yes = 2 No = 0				
Rhinosinusitis	Duration of symptoms	≥ 10 days=2 <10 days = 0	≤3: No need for antibiotic 4-5: Uncertain outcome			
	Face pain/sensibility	Yes = 1 No = 0	≥ 6 : Need for antibiotic			
	Purulent nasal secretions	Yes = 1 No = 0				
General	Fever	$\geq 39^{\circ}C = 2 < 39^{\circ}C = 0$				
	Age	< 65 years = $0 \ge 65$ years = $1 \ge 80$ years = 2				
	Corticosteroid therapy	Yes = 1 No = 0				

Tool development

Based on diagnostic and treatment guidelines, a scale was developed to empirically classify the aetiology of infections (Table 1) and the need for antibiotic therapy (Table 2)^[11,13,20]. A weighting scale was defined according to the severity of signs and symptoms.

Ethics and confidentiality

This study complied with the established ethical and legal requirements, including collection of patient informed consents and the authorisation of the technical directors of the community pharmacies where the study was implemented. Patients' were anonymised and the identity of the pharmacies involved was preserved.

Data analysis

Data were analyzed using the Statistical Package for Social Sciences software (SPSS), version 20, including descriptive univariate analysis of the categorical (absolute and relative frequencies) and scalar variables [(dispersion measures (range and standard deviation) and measures of central tendency (mean and median)]. Bivariate analysis was conducted using non-parametric tests (simple chi-square test and Fisher's exact test), considering a 95% confidence interval.

Results

The sample included 22 patients, implying that only 23.16% of the intended samples were recruited (95 patients). These were recruited in two different locations, 6 patients (27.3%) in Almada, Lisbon Metropolitan Area, and the remaining 16 (72.7%) in Elvas, Alentejo.

Socio demographic characterization of the sample

Most recruited patients were female, with a mean and median age corresponding to 38 years ($\sigma = 16.92$) and 35, respectively, ranging from 19 to 78 years. Nearly half the sample had higher education (45.5%), although educational level in Elvas was lower.

Clinical and therapeutic characterization

On average, patients presented six symptoms ranging from two to eleven, as described in (Figure 1).

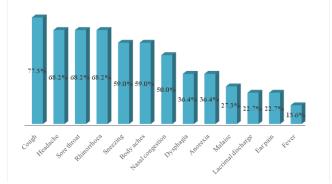


Figure 1: Symptoms described by patients

During clinical observation, only one patient presented with fever (4.5%). The sign more often present was erythema of the pharynx (n=9; 40.9%). Pharyngeal exudates was present in 9.1% of cases (n=2). Only one patient (4.5%) revealed pain or tenderness in maxillary and frontal sinuses. Palpation of the neck resulted in the identification of 5 patients presenting anterior adenopathy (22.7%) (n=5) and one presenting posterior adenopathy (4.5%). The remaining observations revealed: rhinorrhoea in 7 patients (31.8%), sneezing in 2 (9.1%) and one



patient had signs of ocular pathology (4.5%).

On the other hand, history of allergy and GERD were described in 7 (31.8%) and 2 (9.1%) patients, respectively.

The most often prescribed antibiotic was the macrolide, clarithromycin (n=8; 36.4%) (Figure 2).

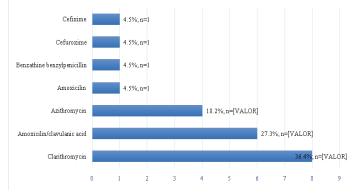


Figure 2: Prescribed antibiotics for the treatment of URTI

Empirical classification of the aetiology of URTI

Using the developed classification for aetiology of URTI (table 1) six patients (27.2%) were considered to present an infection with questionable aetiology (rating between 4 and 5), and only one patient (4.5%) was considered to have a possible bacterial infection. The remaining sample (68.2%, n=15) had an empirical classification lower or equal to 3 points, which suggests the aetiology of the infection was probably viral. The prescribing pattern of antibiotics focused on amoxicillin/clavulanic acid, clarithromycin and benzylpenicillin (Table 3).

Evaluation of the need for antibiotic therapy

Applying the second classification system, developed (table 2) for evaluating the need for antibiotic therapy, we found four patients to have a questionable result (18.2%), with classifications between 4 and 5, and only three patients (13.6%) evidenced the need for antibiotic treatment, presenting a rating

above 6. Therefore, the remaining fifteen patients (68.2%) had ratings below 3, suggesting antibiotics were not required in these cases. Antibiotics prescribed for patients with probable need for antimicrobials were amoxicillin/clavulanic acid and clarithromycin (Table 4).

Discussion

Using the empirical classification of the aetiology of infection and considering risk factors (such as age over 65), we assumed to be possible to ascertain the need for antibiotic therapy. Results suggest that most patients had viral infections and only one evidenced signs of bacterial infection. As expected, this study demonstrated a generalised overuse of antimicrobials, evidenced by the finding that only 13.6% of patients had a rating supporting antibiotic treatment^[17]. Some studies suggest that this over prescription may be a result of patients' expectations about treatment, pressuring physicians to institute the treatment patients believe to be more effective for URTI^[9]. This suggests that is it important for healthcare professionals to explain patients that URTI are self-limiting infections and the overuse of antibiotics increases the emergence of resistances.

Clarithromycin and azithromycin (both belonging to macrolide class) were prescribed in over half of the patients. However, according to the "Infectious Diseases Society of America" (IDSA) guidelines, this class of antimicrobials is considered second-line therapy in bacterial pharyngitis (except in penicillin-allergic patients)^[20]. Therefore, the treatment with macrolides in rhinosinusitis must be discouraged due to the higher resistance rates of *Streptococcus Pneumoniae*. In addition, although benzylpenicillin is considered first-line therapy in bacterial pharyngitis, it was only prescribed to one patient, who presented an infection classified to be of doubtful aetiology. On the other hand, *amoxicillin*/clavulanic acid, which is considered first-line treatment in rhinosinusitis, was often prescribed in this study^[21].

Empirical classification of the etiology of the infection	Amoxicillin/ clavulanate n (%)	Clarithro- mycin n (%)	Benzathine benzylpeni- cillin n (%)	Amoxicil- lin n (%)	Azithromy- cin n (%)	Cefu- roxime n (%)	Cefixime n (%)	Total n (%)	
Probable viral infection (classification: ≤3)	3 (50,0)	5 (62,5)	0 (0,0)	1 (4,5)	4 (100,0)	1(100,0)	1 (100,0)	15 (68,2)	
Doubtful etiology (classifica- tion: 4-5)	3 (50,0)	2 (25,0)	1 (100,0)	0 (0,0)	0 (0,0)	0 (0,0)	0 (0,0)	6 (27,2)	
Probable bacterial infection (classification: 8)	0 (0,0)	1 (12,5)	0 (0,0)	0 (0,0)	0 (0,0)	0 (0,0)	0 (0,0)	1 (4,5)	
Total	6 (27,3)	8 (36,4)	1 (4,5)	1 (4,5)	4 (18,2)	1 (4,5)	1 (4,5)	22 (100,0)	

Classification of the need for antibiotics	Amoxicillin/ clavulanate n (%)	Claritromy- cin n (%)	Azitromy- cin n (%)	Benzathine ben- zylpenicillin n (%)	Amoxicil- lin n (%)	Cefurox- ime n (%)	Cefixime n (%)	Total n (%)
Probably unnecessary (classification: ≤3)	3 (50.0)	5 (62.5)	4 (100.0)	0 (0.0)	1 (100.0)	1 (100.0)	1 (100.0)	15 (68.2)
Doubtful need (classi- fication: 4-5)	1 (16.7)	2 (25.0)	0 (0.0)	1 (100.0)	0 (0.0)	0 (0.0)	0 (0.0)	4 (18.2)
Probable need (classifi- cation: ≥6)	2 (33.3)	1 (12.5)	0 (0.0)	0 (0.0)	0 (0.0)	0 (0.0)	0 (0.0)	3 (13.6)
Total	6 (27.3)	8 (36.4)	4 (18.2)	1 (4.5)	1 (4.5)	4 (18.2)	1 (4.5)	22 (100.0)

Previous studies have found that in Portugal the prescription pattern for antibiotics in respiratory infections focuses on the beta-lactamics/penicillins class, especially the combination of amoxicillin/clavulanic acid and cephalosporins, followed by macrolides^[10,16]. Our findings support this predominance in one of the regions studied–Elvas. In Almada, most patients were prescribed a macrolide, followed by beta-lactamics.

Cough was a prevailing symptom described by patients, which may indicate that these infections were not restricted to the upper respiratory tract, also affecting the lower tract. However, taking into account the high prevalence of smoking habits in this sample and considering tobacco to be the leading cause of chronic cough, this symptom can result from lifestyle related factors. Furthermore, the presence of smoking habits is a predisposing factor for URTI, so these patients are more susceptible to acquire these illnesses^[22]. It is noteworthy that the prevalence of smoking in the sample was more than twice higher (45.5%) than the prevalence of smoking habits in Portugal (20.8%)^[23].

Despite the absence of allergic and GERD patients in our sample, the relationship between these conditions and URTI has been noticed by other authors^[4,5]. However, an information bias must be considered because there could have been a misinterpretation or misunderstanding regarding the questions made during the interview. Also, most of the patients were from Elvas, Alentejo, where there is an overall lower educational level which can justify the lack of awareness about the history of allergy and GERD.

This study revealed some limitations, including the limited sample size, in comparison with the estimated 95 patients, which invalidate the extrapolation of results. This may be explained by the fact that when the study began, URTI were not at the peak season. Additionally, many people presenting a prescription were not the sick patient, precluding their recruitment. It may be assumed that the patients able to acquire their own medication had the less severe clinical situations, hence the high prevalence of probable viral infections aetiology. The exclusion of patients without antimicrobial prescription prevented to estimate the need for antibiotic therapy in patients who resort to the pharmacy when respiratory symptoms appear. Nonetheless, this was considered to be the best methodological option, since the presence of the principal investigator could have influenced the standard pharmaceutical intervention. The limited study period and human resources also contributed to the limited sample size. It is also appropriate to reflect on the difference of the sample size between the two locations, which can be explained by several reasons. First, the greater availability to participate in the study in Elvas; equally important was the fact that the recruitment time was different in the two locations: in Elvas, it was primarily held during shift schedules (at night and weekend). Third, it should be noted that, according to data from the General Health Directorate, the ambulatory antibiotic consumption is higher in Elvas than in other regional areas, so it can be expected an easier recruitment at this location^[24]. Finally, the study began in Almada, when URTI were not yet in seasonal peak. Consequently, due to the asymmetric sample size in these places, the comparative analysis of the data was limited, and any characterization of the sample recruited in Almada can also be compromised. However, this study was based on developing clinical competencies of a pharmacy student and as such, had to be feasible and compatible with classes, explaining the methodological option taken.

The methodology used to classify the aetiology of infections and assess the need and appropriateness of the therapy is questionable for various reasons. For example, the presence of fever may have been underestimated because many patients were under antipyretic medication effect. It should also be borne in mind that the questionnaire did not include information concerning beta-lactamic antibiotic allergy, so it is not possible to establish if some of the prevailing prescription of macrolides is due to the presence of allergic patients. The scale was tested but not validated, which means that the sensitivity and specificity are unknown. Nonetheless, these scales are pioneers, developed through international guidelines, so it would be an asset to validate them on a representative sample, in the future. Despite having received a differentiated training the principal investigator was, at the time, a practicing pharmacist and his experience in observation was certainly lower than of a doctor and consequently some ratings may be subject to error or discussion. Nonetheless, we believe it was a valuable experience for the student to develop his clinical competencies further, and also for the pharmacy to have the opportunity to provide patients with an advanced service, currently not offered in most pharmacies (a complete clinical evaluation including observation when presenting minor symptoms). This pitfall of the system is not restricted to pharmacies but generalised in primary care, where doctors for many reasons do not have the opportunity to conduct a thorough and complete evaluation of the signs and symptoms, as mentioned by some patients during this study^[9].

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Despite the aforementioned limitations, this study focuses on a relevant topic, introducing an innovative approach to pharmaceutical interventions, and presents results consistent with previous studies, highlighting the overuse of antimicrobials, particularly in sore throat^[10]. This suggests that the campaigns developed to promote the rational use of antibiotics, although important, are insufficient to change behaviours and could benefit from pharmacists' intervention, a professional privileged by its trained skills and unique position in the healthcare system. These interventions could be fostered by studies like this.

Conclusion

Antibiotic treatment prescribed in most clinical situations was probably unnecessary, since the analysed URTI did not suggest a bacterial aetiology. Clarithromycin and amoxicillin/ clavulanic acid revealed a high indication. Hence, it is essential to adopt corrective measures in primary care to promote the rational use of antimicrobials, in order to fight the emergence of resistance and further decrease treatment costs.

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