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HOW CRITICAL IS TIMING FOR THE DIAGNOSIS OF INFLUENZA IN GENERAL PRACTICE?

THESE

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Résumé

La performance diagnostic des signes et symptômes de la grippe a principalement été étudiée dans le cadre d'études contrôlées avec des critères d'inclusion stricts.

Il apparaît nécessaire d'évaluer ces prédicteurs dans le cadre d'une consultation ambulatoire habituelle en tenant compte du délai écoulé entre le début des symptômes et la première consultation ainsi que la situation épidémiologique.

Cette étude prospective a été menée à la Policlinique Médicale Universitaire durant l'hiver 1999-2000. Les patients étaient inclus s'ils présentaient un syndrome grippal et si le praticien suspectait une infection à *Influenza*. Le médecin administrait un questionnaire puis une culture d'un frottis de gorge était réalisée afin de documenter l'infection.

201 patients ont été inclus dans l'étude. 52% avaient une culture positive pour *Influenza*. En analyse univariée, une température > 37.8° (OR 4.2 ;95% CI 2.3-7.7), une durée des symptômes < 48h (OR 3.2; 1.8-5.7), une toux (OR 3.2; 1-10.4) et des myalgies (OR 2.8; 1.0-7.5) étaient associés au diagnostic de grippe. En analyse de régression logistique, le modèle le plus performant qui prédisait la grippe était l'association d'une durée des symptômes <48h, une consultation en début d'épidémie, une température > 37.8° et une toux (sensibilité 79%, spécificité 69%, valeur prédictive positive 67%, une valeur prédictive négative de 73% et aire sous la courbe (ROC) de 0.74).

En plus des signes et symptômes prédicteurs de la grippe, le médecin de premier recours devrait prendre en compte dans son jugement la durée des symptômes avant la première consultation et le contexte épidémiologique (début, pic, fin de l'épidémie), car ces deux paramètres modifient considérablement la valeurs des prédicteurs lors de l'évaluation de la probabilité clinique d'un patient d'avoir une infection à *Influenza*.

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Abstract

Questions under study: The diagnostic performance of clinical symptoms/signs of influenza has mainly been assessed in the context of controlled studies with stringent inclusion criteria. There was a need to extend the evaluation of these predictors in the context of general practice, and according to the duration of symptoms and to the dynamics of the epidemic.

Principles: Prospective study conducted in the Medical Outpatient Clinic, winter season 1999-2000. Patients with influenza-like syndrome were included, as long as the primary care physician envisaged the diagnosis of influenza. A questionnaire was administered by the physician, a throat swab was performed and a culture to document the diagnostic of influenza.

Results: 201 patients were included in the study. 52% were culture positive for influenza. By univariate analysis, a temperature >37.8^{0}C (OR 4.2; 95% CI 2.3-7.7), a duration of symptoms <48 hours (OR 3.2; 1.8-5.7), cough (OR 3.2; 1-10.4) and myalgia (OR 2.8; 1.0-7.5) were associated with a diagnosis of influenza. In a multivariable logistic analysis, the best model predicting influenza was the association of a duration of symptom < 48 hours, medical attendance at the beginning of the epidemic (weeks 49-50), fever > 37.8 and cough, with a sensitivity of 79%, specificity of 69%, positive predictive value of 67%, negative predictive value of 73% and an area under the ROC curve of 0.74

Conclusions: Besides relevant symptoms and signs, the physician should also consider the duration of symptoms and the epidemiological context (start, peak or end of the epidemic) in his judgement, since both parameters modify considerably the value of the clinical predictors when assessing the probability of a patient to have influenza.

Key-words : influenza, clinical predictors, clinical diagnosis, epidemic, time

Introduction

The recent marketing of antiviral drugs against influenza has changed the diagnostic approach of this disease. Indeed, there is a need to quickly make the diagnosis so that the new drugs can be used appropriately [1-3]. The general practitioner needs epidemiological, clinical and/or laboratory tools to improve the reliability of the diagnosis of influenza at first attendance.

The diagnostic performance of clinical symptoms/signs of influenza has mainly been assessed in the context of controlled studies (clinical trials of new antiviral drugs) often using stringent inclusion criteria. A temperature >37.8°C, cough and sudden onset of symptoms have been identified as indicators of influenza [4-11]. In the present study, we wanted to assess these predictors in the context of general practice and more importantly, the magnitude of their variation according to the duration of symptoms and the dynamic of the epidemic (start, peak and end).

Methods

Design:

Prospective study conducted during the winter season 1999-2000 at the Medical Outpatient Clinic, University of Lausanne, Switzerland, a primary care centre that serves an urban population of approximately 150,000 inhabitants. The study was conducted within a national surveillance programme of influenza epidemics in Switzerland called Sentinella [12]. All along the year, naso-pharyngal swabs are collected and tested for influenza and other infectious diseases from several institutions (outpatient clinics) and private practices in Switzerland, in ordrer to detect and monitor epidemics outbreaks. The proportion of medical consultations for influenza-like illness (% MC-ILI) is also reported, and is used to describe the dynamic of epidemic of influenza (the threshold to define an epidemic is 1.5%).

Patients and procedure:

Patients were recruited in the study by the physician on duty, if he/she felt that the symptoms or signs were compatible with a diagnosis of influenza. There were no specific criteria for inclusion or exclusion (such as those of Sentinella surveillance in Switzerland, for example) in order to avoid selection bias (patient with a high pre-test probability) and to reflect the real practice. After oral consent, the physician administered a questionnaire to the patient to collect demographic (i.e. age, sex) and clinical data (i.e. symptoms of cough, sore throat,

rhinitis, myalgia, headache, fatigue, chills/sweating, as well as the duration of symptoms from onset to medical attendance and signs, mainly axillary temperature).A threshold value of 37.8° was used to define fever as in most of the other studies [3].A throat-swab was performed and sent in medium (Leibowitz, BSA, bicarbonate, hybrimax and gentamycin) to the reference laboratory for the Sentinella Surveillance Program (IKMI, St-Gallen, Switzerland) for a MDCK culture in order to identify influenza A and B viruses.

Data analysis:

To measure the association between the explanatory variables (duration of the symptoms, the period of the consultation, axillary temperature of >37.8°C, cough, sore throat, rhinitis, myalgia, headache, fatigue, chills/sweating), and the outcome variable (presence of influenza A or B in throat swab culture), we estimated the odds ratio with the program CIA SOFTWARE version 2.0.0 from BMJ using univariate analysis. For each of these variables , we estimated the sensitivity (Se), specificity (Sp), positive and negative predictive values (PPV/NPV).

We then built an multivariable logistic model using STATA 8.2 software, starting with a simple model including only duration of symptoms and period of consultation. The categories for the duration of symptoms were < 24 h, 24-48h and > 48h. The categories for the time-period of consultation were: week 49-51 (pre-epidemic), 52-1 (peak of the epidemic) and > 1 (post-epidemic) [based on the proportion of medical consultations due to flu-like syndromes estimated by the Swiss infectious disease surveillance system (Sentinella)]. We added step by step those clinical variables with odds ratio higher than one. We retained in the model variables for which the estimated odds ratio was >1 (p-value <0.05). Going backwards, we proceeded to a simplification in the definition of the categorical variables, using the deviance statistic to judge the loss of diagnostic power. The ROC curves of both the initial and the final models were computed, as well as the sensitivity, specificity, PPV and NPV.

Results

This study was conducted from December 1999 to February 2000. 222 patients were included in the study and 21 patients with incomplete data or where no throat-swab had been done were excluded from the analysis, so it remained 201 patients. 104 of 201 (52%) had a positive throat-swab for influenza, of which 103 for influenza A and 1 for influenza B. The mean age was similar in the group with positive culture (mean = 34.3 years, SD = 13) than the one with negative culture (mean = 34.3, SD = 12). Height patients were aged more than 60 years and

half of them were positive for influenza. The demographical characteristics as well as the prevalence of symptoms and signs among cases of influenza vs controls are summarised in table 1.

By univariate analysis, temperature >37.8°, cough, duration of symptoms <48 hours before consultation and myalgia were associated with a diagnosis of influenza.

Table 2 shows the diagnostic performance (PPV, NPV, sensitivity and specificity) of clinical variables for the diagnosis of influenza.

We started to construct the multivariable logistic model with the two variables: time-period of consultation and duration of symptoms before first medical attendance, each divided in 3 categories as described in the data analysis section. In this situation the area under the ROC curve was 0.69 with a prediction rule of 0.5, Se was 64% (IC 95% 54-74), Sp 64% (54-73), PPV 66%(56-75) and NPV 63% (52-72) (see figure 1). We then added step by step the variables with a estimated OR > 1, temperature > 37.8, cough, and myalgia. The latter symptom added nothing to the power of the model and was thus withdrawn. After that, we replaced temperature > 37.8° by continuous temperature measurement. We then simplified the 3 time-period categories into two(week 49-50 and week 51-5) and also the 3 categories of duration of symptoms into two (< 48h and >48h) without changing the power of our model. Finally, the model with 2 categories for time-period of consultation and duration of symptoms, continuous temperature and cough had an area under the ROC curve of 0.76 with a Se of 74% (65-82), Sp of 69% (59-78), PPV of 72% (62-80) and NPV of 71% (61-80) (see figure 2). When we replaced continuous temperature by categorical temperature > 37.8, the area under the ROC curve was 0.74 with a Se of 80% (70-87), Sp of 59% (48-69), PPV of 67% (58-76) and NPV of 73% (62-82) (see figure 3).

Discussion

The present study shows that the best model for the prediction of influenza in clinical practice is the association of a duration between symptom onset and first medical consultation <48 hours, medical attendance at the beginning of the epidemic, a temperature >37.8°C, and cough. The model was even better when we used temperature as a continuous measurement, meaning that the highest the temperature was, the better the prediction. However in practice it is much easier to use a fixed threshold (> 37.8°).

Temperature > 37.8° and cough were good clinical predictors of influenza, which is in line with the results described in previous studies conducted in selected populations aimed at assessing the safety and efficacy of antiviral drugs [4-11,13]. This means that primary care physicians can also safely use these predictors to guide their practice.

According to our results, the time must also be considered in presence of clinical signs and symptoms suggesting influenza. Indeed, a duration of symptoms less than 48 hours, and first medical attendance at the beginning of the epidemic were also good predictors for the positive diagnostic of influenza in the presence of fever and cough .The primary care physician should therefore consider the duration of symptoms and the time of the epidemic when assessing the probability of his/her patient to have a diagnosis of influenza in the presence of a known clinical predictor.

In summary, the probability of having influenza is highest when the patient attends rapidly after symptom onset, at the beginning of the epidemic and in the presence of a temperature >37.8°C + cough. At the peak of the epidemic, almost all patients have influenza, irrespective of their symptoms and signs. The clinical predictors, as well as the rapid diagnostic tests (due to the important variability of specificity of these assays [14-16]) loose thus their usefulness at that time. The time period of consultation during and around the epidemic influencing the prediction of influenza highlights the necessity for the clinician to consider the epidemiological context at the time of consultation, when estimating the probability of his/her patient to have influenza.

The identification of clinical predictors of influenza, as well as a fair estimation of their variability in time, should help to establish clinical scores that could be used by the general practitioner to optimise the care of patients in terms of rapid diagnostic tests use and antiviral therapy initiation.

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Table 1: demographical characteristics and prevalence of symptoms and signs among cases of influenza (culture +) vs controls (culture-)

Characteristics	% (number) of patients with positive culture (n=104)	% (number) patients with negative culture (n=97)	Odds Ratio*	CI 95 %
Female sex	53 (55)	46 (44)	1.3	0,8 – 2.3
Duration of symptoms before medical attendance < 48h	66 (69)	38 (37)	3.2	1.8 – 5.7
Time-period of consultation weeks 49-50 (pre-epidemic) weeks 51-5 (epidemic and post-epidemic)	26 (27) 74 (77)	28 (27) 72 (70)	2.3 ref	1.2-4.5 1.1-9.9
Temperature >37.8°C	74 (77)	40 (39)	4.2	2.3 – 7.7
Cough	96 (100)	89 (86)	3.2	1.0 – 10.4
Temperature > 37,8 and cough	72 (75)	37 (36)	4.4	2.4 – 7.9
Sore throat	75 (78)	75 (73)	1	0.5 – 1.9
Myalgia	94 (98)	86 (83)	2.8	1.0 – 7.5
Rhinitis	81 (84)	81 (79)	1	0.5 – 1.9
Headache	85 (88)	84 (81)	1.1	0.5 – 2.3
Fatigue	91 (95)	92 (89)	1	0.4 – 2.6
Chills/sweating	88 (91)	77 (75)	2.1	1.0 – 4.4

* by univariate analysis

Table 2 : diagnostic performance: Positive Predictive Value (PPV), sensitivity (Se) and specificity (Sp)) of clinical variables for the diagnosis of influenza

symptoms and signs	PPV	NPV	Se	Sp
Cough	54	73	96	11
Sore throat	52	48	75	25
Rhinitis	52	47	81	19
Myalgia	54	70	94	14
Headache	52	50	85	17
Fatigue	52	47	91	8
Duration of symptoms < 48h	65	63	66	62
Chills/sweating	55	63	88	23
Temperature >37,8	66	68	74	60
Cough + Temp. > 37,8	68	68	72	63

Figure 1 ROC curve, model with the two variables: time-period of consultation (weeks 49-51, 52-1 and >1) and duration of symptoms before first medical attendance (< 24 h, 24-48h, >48h).

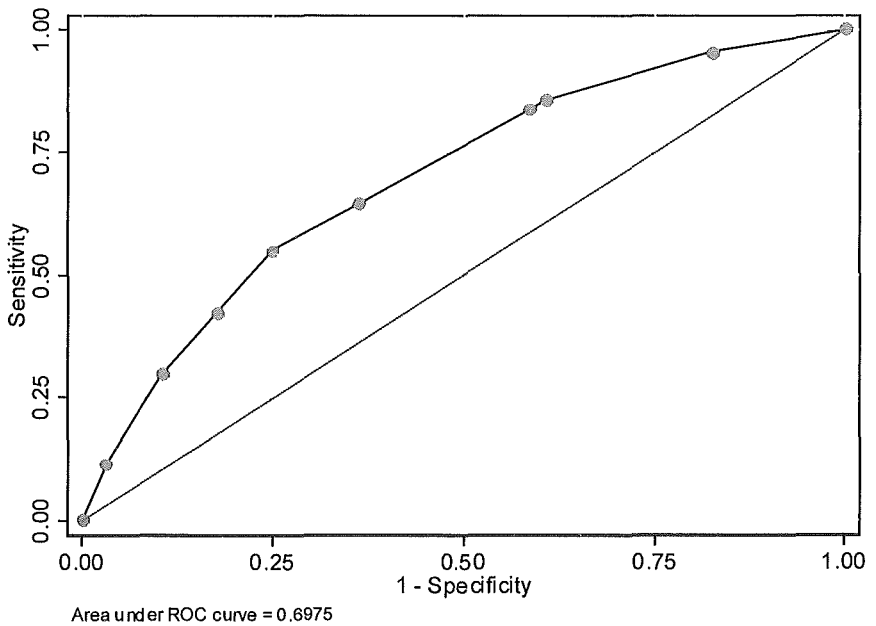

Figure 2 ROC curve, model with the four variables: time-period of consultation (weeks 49-50, 51-5), duration of symptoms before first medical attendance (< 48h, >48h), continuous temperature and cough.

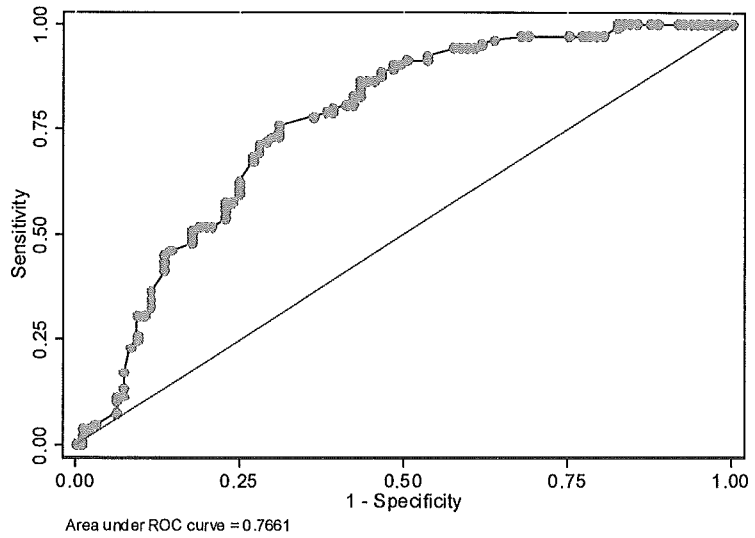

Figure 3 ROC curve, model with the four variables: time-period of consultation (weeks 49-50, 51-5), duration of symptoms before first medical attendance (< 48h, >48h), temperature > 37.8° and cough.