Seasonal variations in use and outcome of rapid antigen detection tests and cultures in pharyngotonsillitis: a register study in primary care

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Abstract

Background: Diagnosis and treatment of pharyngotonsillitis are commonly focused on group A streptococci (GAS), although the disease is often associated with other pathogens. While the incidence of pharyngotonsillitis is known to vary with season, seasonal variations in the prevalence of potential pathogens are sparsely explored. The aim of this study was to explore any seasonal variations in the use and outcome of rapid antigen detection tests (RADTs) for GAS and throat cultures among patients diagnosed with pharyngotonsillitis in primary care.

Methods: We retrieved and combined retrospective data from the electronic medical record system and the laboratory information system in Kronoberg County, Sweden. Primary care visits resulting in a diagnosis of tonsillitis or pharyngitis were included, covering the period 2013–2016. The monthly rate of visits was measured, along with the use and outcome of RADTs for GAS and throat cultures obtained on the date of diagnosis. The variations between calendar months were then analysed.

Results: We found variations between calendar months, not only in the mean rate of visits resulting in a diagnosis of pharyngotonsillitis (p < 0.001), but in the mean proportion of RADTs being positive for GAS among the diagnosed (p < 0.001), and in the mean proportion of visits associated with a throat culture (p < 0.001). A lower mean rate of visits in August and September coincided with a lower proportion of RADTs being positive for GAS among them, which correlated with a higher proportion of visits associated with a throat culture.

Conclusions: This study suggests that the role of GAS in pharyngotonsillitis in Sweden is less prominent in August and September than during the rest of the year.

Keywords: Sore throat, Tonsillitis, Pharyngitis, Seasonal variation, Streptococcus pyogenes, Streptococcus dysgalactiae, Fusobacterium necrophorum

Background

A sore throat is a common condition, resulting in considerable numbers of primary care visits and prescriptions of antibiotics. According to the Primary care Record of Infections in Sweden (PRIS), 11% of the prescriptions of antibiotics are issued for sore throats (the vast majority being diagnosed with either pharyngitis or tonsillitis) [1], but consultation rates and prescription percentages...
visits in this article refers to this definition. From the EMRs, we extracted the date of diagnosis, the personal identification number, if a rapid antigen detection test (RADT) for GAS had been used for that patient and date, and if used, the result of that RADT. Any results from throat cultures obtained from that patient and date, were extracted from the laboratory information system. Two types of standardized throat cultures were available at the time, one targeting beta-haemolytic streptococci (A, C and G) only, and the other one also including selective anaerobic incubation to detect *Fusobacterium necrophorum*. As to duplicates of visits (more than one visit on the same date) in our dataset, we were not able to tell if they represented real duplicates or only duplicated data transfers, as most of them were identical regarding reporting unit/PHCC, and when so, also identical regarding RADT use and results. For technical reasons, we chose to count what appeared to be the first visit, but it should be noted that it does not necessarily reflect true chronology due to data limitations. However, the number of duplicates in the dataset was low (179 of 21,363), and the true number of duplicates possibly even lower. As to duplicates of cultures, positive results were included.

In late 2013, the microbiological laboratory switched from Lancefield classification of streptococci to binomial nomenclature, with the introduction of MALDITOF (matrix-assisted laser desorption/ionization with time-of-flight mass spectrometer). For the analyses in this article, we chose to treat group A beta-haemolytic streptococci (GAS) and *S. pyogenes* as interchangeable, and similarly, to collectively treat group C (GCS) and G (GGS) streptococci as equivalent to *S. dysgalactiae* subsp. equisimilis.

**Statistics**

Age quartiles were calculated, after adjusting age to the number of full years lived. Three age groups were defined, to control for age as a potentially interfering factor. These age groups were chosen due to previous knowledge of age being of relevance to aetiology [8, 11]. For descriptive statistics, the rate of visits was measured per calendar month, with the numbers weighted to the length of the month by dividing by the number of days in the month and multiplying by 30. Data from all 4 years were combined, for further analyses. Seasonal variation was first analysed with chi-squared goodness of fit tests for detection of overall differences between all calendar months (with aggregated data from all 4 years, if not stated otherwise in the text). The magnitude of any detected seasonality was then estimated with peak-to-low ratios between the months with the highest and the lowest rates or proportions, as exemplified by Christiansen et al. [12]. The confidence level was set to 95%. Confidence intervals
for proportions were calculated using Wilson’s method. Confidence intervals for ratios (ratios of proportions) were calculated according to Daly, 1998, with p-values from z-tests according to Sheskin, 2004. Correlations were analysed using Pearson’s bivariate correlation. The analyses were performed using Excel, version 16.16.2 (Microsoft, Redmond, WA) and SPSS, version 23.0.0.0 (IBM, Armonk, NY).

Results
Visits with a diagnosis of pharyngotonsillitis
Throughout the 4 years of the study, a total of 21,184 visits with a diagnosis of tonsillitis or pharyngitis were identified, after exclusion of 179 duplicates (more than one visit on the same date). The median age was 20 years. Of all visits, 56% (95% CI 56–57) were made by men. However, this gender difference was not seen in the youngest age group (0–14 years). The rate of visits per month showed seasonal variation when comparing means for all calendar months (p < 0.001). The highest mean was seen in December, and the lowest in September, with a peak-to-low rate ratio of 1.40 (95% CI 1.22–1.60, p < 0.001). The second lowest mean was seen in August, with a corresponding rate ratio (December/August) of 1.36 (95% CI 1.19–1.55, p < 0.001). However, peak and low months were not the same for all the years studied (Fig. 1). For instance, the beginning of 2013 and the end of 2016 had higher numbers, thus skewing the mean. In the age group of 15–29 years, the mean pattern was somewhat different, with a peak in August. Data for all years are shown in Table 1, including rate ratios between peak and low months. Summarized characteristics per calendar month are presented in Tables 2, 3, 4, 5. Non-aggregated data are also shown in Additional file 1: Table 1.

Table 1 Visits with a diagnosis of pharyngotonsillitis 2013–2016, shown as rate per 1000 persons and month

|       | Jan | Feb | Mar | Apr | May | Jun | Jul | Aug | Sep | Oct | Nov | Dec | p   | RR peak/low (95% CI) | p   |
|-------|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|---------------------|-----|
| 2013  | 3.81| 3.71| 3.36| 3.22| 2.90| 2.62| 2.61| 2.32| 1.93| 2.55| 2.24| 2.67| <0.001| 1.98 (1.74–2.26), p<0.001|
| 2014  | 2.33| 2.74| 2.37| 1.97| 2.27| 2.44| 2.17| 2.01| 2.07| 2.47| 2.16| 2.50| <0.001| 1.39 (1.21–1.59), p<0.001|
| 2015  | 2.15| 2.29| 2.26| 2.12| 1.99| 2.35| 1.87| 1.89| 1.91| 2.15| 2.28| 2.48| <0.001| 1.32 (1.15–1.53), p<0.001|
| 2016  | 2.54| 2.60| 2.50| 2.20| 2.26| 2.57| 2.13| 2.18| 2.26| 2.50| 2.98| 3.76| <0.001| 1.76 (1.56–2.00), p<0.001|
| Mean  | 2.71| 2.83| 2.62| 2.38| 2.36| 2.49| 2.20| 2.10| 2.04| 2.42| 2.42| 2.86| <0.001| 1.40 (1.22–1.60), p<0.001|

All ages. The numbers are weighted to the length of the month (divided by the number of days and multiplied by 30)

Cl confidence interval

* P-value from chi-squared goodness of fit test for overall differences in rates between all months

b Rate ratio between the month with the highest rate and the month with the lowest rate, and corresponding p-value from z-test (Sheskin, 2004)
Table 2  Summary of visits with a diagnosis of pharyngotonsillitis, per calendar month (2013–2016 combined). All ages

|                     | Jan  | Feb  | Mar  | Apr  | May | Jun  | Jul  | Aug  | Sep  | Oct  | Nov  | Dec  |
|---------------------|------|------|------|------|-----|------|------|------|------|------|------|------|
| Mean number of weighted visits* | 479  | 502  | 465  | 422  | 419 | 443  | 390  | 374  | 363  | 430  | 431  | 510  | 436  |
| Median age, years (IQR) | 20 (37–8) | 19 (35–7) | 18 (35–7) | 20 (36–8) | 19 (36–8) | 21 (37–9) | 21 (36–11) | 19 (34–10) | 20 (34–8) | 18 (32–7) | 19 (35–8) | 20 (35–8) |
| Women, % of visits (95% CI) | 46 (44–48) | 46 (43–48) | 42 (40–44) | 43 (40–45) | 41 (39–44) | 39 (37–42) | 45 (43–48) | 44 (42–47) | 45 (42–47) | 45 (41–45) | 43 (41–45) | 44 (43–44) |
| RADT used, % of visits (95% CI) | 64 (62–66) | 65 (63–67) | 65 (63–67) | 66 (64–68) | 65 (63–67) | 69 (66–71) | 66 (63–68) | 64 (61–66) | 62 (60–65) | 63 (61–66) | 66 (63–68) | 67 (65–69) | 67 (65–66) |
| Positive RADTs/all RADTs, % (95% CI) | 68 (65–70) | 68 (65–70) | 68 (65–71) | 69 (67–72) | 68 (66–71) | 66 (63–69) | 60 (57–63) | 54 (51–57) | 56 (53–59) | 61 (58–64) | 66 (63–68) | 68 (65–70) | 65 (64–66) |
| Culture obtained, % of visits (95% CI) | 48 (40–59) | 5.5 (4.6–6.6) | 5.7 (4.8–6.8) | 5.7 (4.7–6.9) | 6.1 (5.1–7.4) | 6.3 (5.3–7.6) | 6.2 (5.1–7.5) | 8.8 (7.5–10.3) | 8.5 (7.2–10.0) | 6.4 (5.4–7.6) | 6.6 (5.5–7.9) | 4.5 (3.7–5.4) | 6.2 (5.8–6.5) |

IQR interquartile range; CI confidence interval; RADT rapid antigen detection test for group A streptococci

*The numbers are weighted to the length of the month (divided by the number of days and multiplied by 30)
Table 3  Summary of visits with a diagnosis of pharyngotonsillitis, per calendar month (2013–2016 combined). Age 0–14 years

|                  | Jan | Feb | Mar | Apr | May | Jun | Jul | Aug | Sep | Oct | Nov | Dec |
|------------------|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|
| Mean number of weighted visits* | 180 | 198 | 187 | 193 | 163 | 169 | 126 | 107 | 111 | 160 | 166 | 202 | 163 |
| Women, % of visits (95% CI) | 56  | 52  | 50  | 51  | 49  | 49  | 51  | 48  | 51  | 51  | 54  | 52  | 53  | 51  |
| RADT used, % of visits (95% CI) | 69  | 69  | 69  | 71  | 67  | 71  | 67  | 67  | 64  | 66  | 70  | 69  | 68  |
| Positive RADTs/all RADTs, % (95% CI) | 80  | 79  | 80  | 85  | 82  | 78  | 73  | 73  | 69  | 72  | 76  | 81  | 78  |
| Culture obtained, % of visits (95% CI) | 2.7 | 2.8 | 3.4 | 4.0 | 3.6 | 3.4 | 3.8 | 5.2 | 4.5 | 4.1 | 3.1 | 1.9 | 3.4 |

CI confidence interval; RADT rapid antigen detection test for group A streptococci

*The numbers are weighted to the length of the month (divided by the number of days and multiplied by 30)
Table 4  Summary of visits with a diagnosis of pharyngotonsillitis, per calendar month (2013–2016 combined). Age 15–29 years

|                    | Jan   | Feb   | Mar   | Apr   | May   | Jun   | Jul   | Aug   | Sep   | Oct   | Nov   | Dec   | ?    |
|--------------------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-----|
| Mean number of weighted visits* | 130   | 120   | 124   | 113   | 112   | 122   | 122   | 143   | 128   | 137   | 126   | 130   | 125 |
| Women, % of visits (95% CI) | 42 (38–46) | 42 (37–46) | 34 (30–38) | 39 (35–44) | 40 (36–45) | 35 (31–39) | 44 (40–49) | 43 (39–47) | 43 (38–46) | 42 (33–42) | 39 (35–43) | 40 (39–41) |
| RADT used, % of visits (95% CI) | 59 (55–63) | 62 (58–66) | 61 (57–64) | 60 (55–64) | 64 (59–66) | 67 (63–71) | 66 (62–70) | 61 (57–65) | 61 (57–65) | 64 (59–68) | 62 (57–66) | 62 (57–66) | 62 |
| Positive RADTs/all RADTs, % (95% CI) | 48 (43–54) | 50 (44–55) | 49 (44–55) | 48 (42–54) | 48 (43–54) | 50 (44–55) | 46 (40–51) | 36 (31–41) | 40 (35–46) | 44 (39–49) | 45 (44–55) | 49 (40–51) | 46 (44–48) |
| Culture obtained, % of visits (95% CI) | 8.0 (6.0–10.6) | 8.9 (6.7–11.7) | 9.2 (7.0–12.0) | 8.6 (6.4–11.5) | 9.5 (7.1–12.2) | 9.7 (7.1–12.2) | 10.2 (8.0–12.9) | 12.6 (10.1–15.7) | 12.6 (10.0–15.5) | 11.9 (9.4–14.9) | 7.8 (5.8–10.4) | 9.5 (8.8–10.2) |

CI = confidence interval; RADT = rapid antigen detection test for group A streptococci

*The numbers are weighted to the length of the month (divided by the number of days and multiplied by 30)
|                  | Jan  | Feb  | Mar  | Apr  | May  | Jun  | Jul  | Aug  | Sep  | Oct  | Nov  | Dec  | %  |
|------------------|------|------|------|------|------|------|------|------|------|------|------|------|----|
| **Mean number of weighted visits*** | 170  | 155  | 155  | 144  | 144  | 156  | 143  | 124  | 104  | 133  | 121  | 178  | 144|
| **Women, % of visits** (95% CI) | 40   | 42   | 39   | 37   | 34   | 34   | 41   | 42   | 39   | 37   | 36   | 36   | 38 |
| **RADT used, % of visits** (95% CI) | 63   | 64   | 63   | 65   | 63   | 68   | 63   | 64   | 62   | 60   | 64   | 68   | 64 |
| **Positive RADTs/all RADTs, %** (95% CI) | 67   | 67   | 67   | 67   | 67   | 67   | 61   | 56   | 61   | 66   | 66   | 68   | 65 |
| **Culture obtained, % of visits** (95% CI) | 4.7  | 5.9  | 5.8  | 5.2  | 6.4  | 6.8  | 5.3  | 10.4 | 7.9  | 7.6  | 6.0  | 4.9  | 6.3 |

CI confidence interval; RADT rapid antigen detection test for group A streptococci

*The numbers are weighted to the length of the month (divided by the number of days and multiplied by 30).
Use of rapid antigen detection tests for group A streptococci

A RADT for GAS was used in 65% (95% CI 65–66) of the visits, totalling 13,817 tests. The proportion of visits where a RADT was used did show a seasonal variation comparing means for all calendar months (p = 0.015). The highest proportion was seen in June (69%, 95% CI 66–71) and the lowest in September (62%, 95% CI 60–65), with a peak-to-low ratio of 1.10 (95% CI 1.05–1.16, p < 0.001). Non-aggregated data are also shown in Additional file 1: Table 2.

Results of rapid antigen detection tests for group A streptococci

The mean proportion of RADTs being positive for GAS (positive RADTs/all RADTs) was 65% (95% CI 64–66). It showed seasonal variation comparing means for all calendar months (p < 0.001). The highest mean proportion was seen in April (69%, 95% CI 67–72) and the lowest in August (54%, 95% CI 51–57), with a peak-to-low ratio of 1.28 (95% CI 1.20–1.38, p < 0.001). The second lowest mean proportion was seen in September (56%, 95% CI 53–59), with a corresponding ratio (April/September) of 1.24 (95% CI 1.15–1.33, p < 0.001). This seasonal pattern was seen in all age groups (Tables 2, 3, 4, 5). Data from all years are shown in Fig. 2.

Use of throat cultures

In total, 1,304 throat cultures were obtained from the patient cohort, after exclusion of eight duplicates (more than one culture from the same patient and date). The mean proportion of visits where a culture was obtained was 62% (95% CI 5.8–6.5), and when comparing means for all calendar months, it showed significant seasonal variation (p < 0.001). The highest mean proportion was seen in August (8.8%, 95% CI 7.5–10.3) and the lowest in December (4.5%, 95% CI 3.7–5.4), with a peak-to-low ratio of 1.97 (95% CI 1.53–2.55, p < 0.001). The second highest mean proportion was seen in September (56%, 95% CI 53–59), with a corresponding ratio (April/September) of 1.24 (95% CI 1.15–1.33, p < 0.001). This seasonal pattern was seen in all age groups (Tables 2, 3, 4, 5). Data from all years are shown in Fig. 2.

Results of throat cultures

Most cultures were negative (61%, 95% CI 58–63), and this proportion did not vary significantly comparing means for all calendar months (p = 0.86). Of all cultures, 18% (95% CI 16–20) were positive for GAS, with the highest mean proportion in April (29%, 95% CI 21–39) and the lowest in September (11%, 95% CI 6–17), but the overall variation between the calendar months, comparing means for all months, was not statistically significant (p = 0.055). GCS or GGS were detected in 13% (95% CI 11–15) of the cultures, with the highest (non-significant) mean proportion in September (19%, 95% CI 13–27). F. necrophorum was detected in 16% (95% CI 13–19) of the cultures targeting this species, with the highest (non-significant) mean proportion in October (23%, 95% CI 14–36). Of all positive cultures (n = 513), 21 were positive for more than one of the analysed bacteria. Almost all of these (n = 20) were positive for F. necrophorum, and more often in combination with GCS or GGS (n = 14) than with GAS (n = 6). The proportions of positive cultures are shown in Fig. 3.

Discussion

In this study, more light has been shed on the seasonal variations in the aetiology of pharyngotonsilitis in primary care. As expected, a seasonal variation was seen in the mean rate of visits with a diagnosis of pharyngotonsilitis comparing calendar months, with the highest mean observed in December and the lowest in September. More interestingly, a lower mean rate of visits in August and September coincided with a lower proportion of RADTs being positive for GAS among them, which correlated with a higher proportion of visits associated with a throat culture. The high number of visits and RADTs analysed, during 4 consecutive years, provides strong evidence for a lower likelihood of GAS involvement in August and September than during the rest of the year.

Subgroup analysis indicated a somewhat different visiting behaviour for adolescents and young adults (15–29 years), with a mean peak of visits in August instead of December. However, this age group showed the same pattern of a lower proportion of RADTs being positive for GAS in August and September, thus making age
a less likely explanatory factor for this variation in GAS involvement. The peak of visits in August among adolescents and young adults was unexpected, and it remains to be explained what factors are involved in this difference between age groups. Furthermore, men were overrepresented from the age of 15 years and above, which contrasts with what is generally seen in primary care visits [13].

As to other studies of seasonal variations in the aetiology of pharyngotonsillitis, a lower likelihood of GAS involvement among sore throat cases in the northern temperate summer than in the colder winter has previously been hypothesized [9]. The incidence of GAS-associated pharyngotonsillitis in children has also been shown to be higher when temperatures are lower (in a Mediterranean climate), and when the number of school-free days is reduced [14]. Recently, seasonal variations in the percentage of RADT positivity have been reported among children with symptoms of acute pharyngitis in a paediatric hospital setting in Portugal, with a decline in August, similar to our findings [15]. A seasonal pattern has also been suggested for GAS among peritonsillar abscesses, with one study of Danish hospital patients showing a higher prevalence of GAS during the winter and the spring [16].

The findings in our study of a lower likelihood of GAS involvement in pharyngotonsillitis in August and September than during the rest of the year, seem to confirm that a seasonality of aetiology is present. An overall decline in GAS transmission due to summer holidays (mid-June to mid-August in Sweden) and higher temperatures (with less time spent indoors) appear to be plausible factors behind this pattern, but it still needs to be shown why the same factors would not cause a similar decline in other pathogens. Seasonal variations in the prevalence of other beta-haemolytic streptococci in pharyngotonsillitis have previously been found among symptomatic children, with higher non-GAS recovery rates during warm weather [17]. A contrasting pattern has been reported from Hungary, with a peak incidence of GCS/GGS in pharyngeal isolates during the colder months (January–March) [18]. As to virus-associated cases of sore throat in children, these have been shown to be more common during the Scottish winter [19].

For comparison, seasonal variations in Swedish cases of invasive GAS infection have been described in a report from the Public Health Agency of Sweden, with the incidence being at its highest in the 1st months of the year [20]. Similar reports of invasive infections, with peak incidence of invasive GAS in the winter (December–February) have been reported both from Norway and Hungary, with contrasting incidence peaks of invasive GCS/GGS in the summer (June–August) [18, 21].

Given the limited number of cultures in our study, no conclusions could be made from their results with regard to seasonal variations. The low number of cultures was expected, since the current guidelines do not encourage throat cultures as part of routine diagnosis. Hence, they can be thought of as actively selected cases (for instance treatment failure, relapsing episodes, unexpected clinical presentations or RADT results). It is still unclear whether the decline in GAS in August and September is offset by non-GAS bacteria or if other pathogens are involved. In our study, we did not detect any significant variation.
between calendar months regarding the proportion of negative cultures.

Another factor, potentially obscuring patterns of seasonal variations in the aetiology, is the seasonal variations in the asymptomatic pharyngeal colonization by potential pathogens. Asymptomatic colonization by GAS has been shown to vary with age, time and concurrent infections in the environment [3]. A small, previous study has also indicated higher prevalence of asymptomatic GAS during the Swedish summer (mid-July to mid-September) than during the winter (mid-January to mid-February) [9]. In the same study, no significant difference was seen in the prevalence of GAS between symptomatic and asymptomatic children < 16 years of age during the summer, which could question the overall applicability of RADTs during this time of the year. However, other studies have indicated the opposite, that the asymptomatic prevalence of GAS and other beta-haemolytic streptococci is independent of season [22, 23].

Obviously, GAS are not the only pathogens causing pharyngotonsillitis, but the significance of non-GAS bacteria and other pathogens needs to be better understood. GCS and GGS are more commonly found among pharyngotonsillitis patients than in asymptomatic controls [8], and GCS have been thought to be of more relevance than GGS [3], but the pathogenic importance of GCS and GGS in pharyngotonsillitis is much less clear than that of GAS [7, 8, 24]. \( \text{F. necrophorum} \) too is more commonly found among patients with pharyngotonsillitis than in asymptomatic controls [8], and more so among teenagers and young adults [11]. However, there is yet no strong evidence for causality, and neither is it known if antibiotic treatment can alleviate symptoms or reduce the risk of complications [25], such as development of the rare Lemierre’s syndrome [26]. Thus, it remains unclear to what extent symptomatic non-GAS cases could benefit from antibiotic treatment. In general, the absolute benefits of antibiotics for sore throats are considered modest, and the number needed to treat is high, to prevent suppurative and non-suppurative complications in high-income countries [27].

The study has methodological limitations. For example, no validation of diagnoses was made, as no individual EMRs were read. Several previous studies have suggested that the adherence to diagnostic guidelines in clinical practice is low [28, 29]. It could be suspected that the outcome of RADTs for GAS among patients with a sore throat is in practice affecting the choice of diagnosis,
altering the “true” proportion of positive/negative RADTs among patients with pharyngotonsillitis. However, our study does not describe the general population of sore throats, but one of visits resulting in a diagnosis of acute tonsillitis or pharyngitis. Because of the Swedish health care system, with publicly financed primary care to everyone, and with compulsory diagnosing of all visits, our data can be expected to represent the vast majority of diagnoses of pharyngotonsillitis in our county during the period studied. Another limitation is our time span for duplicate exclusion. Choosing a longer time span, aiming at finding disease episodes rather than visits, would have been an alternative approach. This could have reduced the risk of overestimation of cases due to “second opinions”, for instance after getting a negative RADT. However, setting an optimal time span for what to exclude would not be easy, and a longer time span could have led to exclusion of cultures representing treatment failure that we wanted to find. Thus, we chose to accept this risk of “dilution”, which should be kept in mind also when interpreting the proportion of visits with RADT being used and the proportion of RADTs being positive.

Conclusions
Among the visits with a diagnosis of pharyngotonsillitis, variation between calendar months was seen as to likelihood of GAS involvement. A lower proportion of RADTs being positive for GAS was seen in August and September, correlating with a higher use of throat cultures. The role of group A streptococci in pharyngotonsillitis in Sweden appears to be less prominent in August and September than during the rest of the year.

Abbreviations
CI: Confidence interval; EMR: Electronic medical record; GAS/GCS/GGS: Group A/C/G streptococci; ICD-10: International classification of diseases, tenth revision; IQR: Interquartile range; PHCC: Primary health care centre; RADT: Rapid antigen detection test.

Supplementary Information
The online version contains supplementary material available at https://doi.org/10.1186/s12879-021-06774-5.

Additional file 1: Table 1. Number of visits in primary care resulting in a diagnosis of pharyngotonsillitis (all ages). Table 2. Number of rapid antigen detection tests for group A streptococci used at visits with a diagnosis of pharyngotonsillitis. Table 3. Number of throat cultures (and positive results) at visits with a diagnosis of pharyngotonsillitis.

Acknowledgements
Not applicable.

Authors’ contributions
MA wrote the original draft and performed the analyses and visualizations of data. KH and MS conceptualized the overarching research aims. OC and KH were responsible for the data collection. OC, KH and JP contributed to the curation of data. KH was also responsible for the funding acquisition, the project administration and the supervision. All authors read and approved the final manuscript.

Funding
This work was supported by Region Kronoberg, Sweden. The funder had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Availability of data and materials
The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Declarations
Ethics approval and consent to participate
This retrospective study was part of a larger project focusing on the aetiology of pharyngotonsillitis in primary care. It was approved, and informed consent was waived, by the Regional Ethical Review Board, Linköping University (Dnr. 2016/529–31). Although personal identification numbers were used in order to match the datasets, these were subsequently anonymized, and no personal EMRs were read during the study. Permissions for data collection were obtained from all the managers of the PHCCs, and the data were stored on servers with restricted access, in compliance with good clinical research practice. Any potential harm to the integrity of the patients was considered negligible, and the data collection had no impact on past or future access to health care.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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Received: 18 April 2021 Accepted: 27 September 2021
Published online: 26 October 2021

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