

## RESEARCH PAPER

# High costs and burden of illness in acute rhinosinusitis: real-life treatment patterns and outcomes in Swedish primary care

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Received 2nd September 2011; revised 9th December 2011; accepted 12th December 2011; online 20th February 2012

### Abstract

**Background:** Few studies have investigated the impact of acute rhinosinusitis on disease-specific quality of life, and disease costs have not been studied previously in Scandinavia.

**Aims:** To study symptoms, treatment patterns, quality of life and costs in adults with acute rhinosinusitis.

**Methods:** This was an observational study in primary care. Patients aged 18–80 years seeking care for acute rhinosinusitis were evaluated using the Major Symptom Score (MSS) on days 0 and 15. Recommended and used treatments, quality of life and costs were assessed by questionnaires including EQ-5D™ and a visual analogue scale (VAS) on the same days.

**Results:** 150 patients were enrolled; 143 provided follow-up data. The proportion of MSS responders was 91%. Mean MSS decreased from 8.4 on day 0 (N=150) to 1.9 on day 15 (N=143). Patients reporting pain/discomfort and problems with usual activities decreased from 88.4% to 31.5% and from 43.2% to 1.4%, respectively, and mean VAS increased from 58.7 to 79.5. Intranasal corticosteroids were the most recommended and/or prescribed drugs. Total cost for an episode was 10,260 SEK (€1,102), of which 75% were indirect costs.

**Conclusions:** With treatment dominated by intranasal corticosteroids, a high proportion of responders and good symptom relief were seen. Acute rhinosinusitis seems to cause a high burden on quality of life and also a high cost for society.

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P Stjärne *et al.* *Prim Care Respir J* 2012; 21(2): 174-179

<http://dx.doi.org/10.4104/pcrj.2012.00011>

**Keywords** EQ-5D™, Major Symptom Score, primary care, rhinosinusitis, observational study, visual analogue scale, adults

See linked editorial by Friedlander on pg 130

The full version of this paper, with online Appendices, is available at [www.thepcrj.org](http://www.thepcrj.org)

## Introduction

Symptoms consistent with acute rhinosinusitis are common clinical problems in primary care practice. In Europe, 1–2% of all patient visits to physicians in primary care are for suspected acute sinusitis.<sup>1</sup> The high number of visits to physicians translates to a high burden on the healthcare system. In 2000 the direct costs in the United States were estimated to be nearly \$6 billion.<sup>2,3</sup>

According to current European guidelines, the recommended treatment for mild acute rhinosinusitis is symptomatic, with nasal

steroids advised in moderate cases.<sup>4,5</sup> Antibiotics should only be added if severe symptoms are present (e.g. fever >38°C, severe pain). The evidence for the efficacy of decongestants in the treatment of acute rhinosinusitis in adults is poor, and decongestants only have a grade D recommendation.<sup>4,5</sup>

According to recent estimates, bacterial infection is present in ≤50% of patients with symptoms of acute rhinosinusitis<sup>6,7</sup> and may be as low as 0.5–2%.<sup>8</sup> Several studies have shown that antibiotics commonly used in Sweden are of limited value in the management of patients with mild to moderate acute rhinosinusitis.<sup>9–11</sup> An increase in antibiotic resistance has also been observed among *Streptococcus pneumoniae* isolates globally, particularly to amoxicillin and other beta-lactam antibiotics.<sup>12</sup> Despite this, antibiotics have previously been estimated to be

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prescribed to more than 90% of adult patients seen for acute rhinosinusitis in the UK.<sup>13</sup> The proportion receiving antibiotics in Scandinavia has also been high.<sup>14</sup> However, a recent questionnaire study performed in the Netherlands showed that current treatment patterns might have changed, as the antibiotic prescription frequency in that study was 20% and 34% for mild and moderate acute rhinosinusitis, respectively.<sup>15</sup>

Few studies investigating the impact of acute rhinosinusitis on disease-specific quality of life have been performed and generic quality of life data are limited.<sup>16</sup> Furthermore, to our knowledge, the costs for acute rhinosinusitis have not previously been investigated in Scandinavia.

The purpose of this study was to investigate the treatment patterns in acute rhinosinusitis in primary care and to increase knowledge of the disease population, symptoms, quality of life and costs.

## Methods

### Study design and patients

This was a prospective non-interventional observational study in adults with acute rhinosinusitis performed at 11 Swedish primary care practices from November 2008 to December 2009. The study protocol was approved by the ethics committee in Stockholm, Sweden, and written informed consent was obtained from all patients before inclusion in the study.

The study protocol did not provide any restrictions for the treatment of acute rhinosinusitis. At the discretion of the investigator, medical treatment was prescribed in accordance with local standard practice, i.e. the chosen treatment by the investigator of each centre in this study for each patient with acute rhinosinusitis, as defined by the inclusion/exclusion criteria.

The primary objective and endpoint was to estimate the proportion of responders according to the Major Symptom Score (MSS) for patients with acute rhinosinusitis. A responder was predefined in the study protocol as a patient who improved in the MSS by at least 30% from day 0 to day 15 (preferably  $\pm 2$  days). Secondary objectives and endpoints were: (1) to estimate the changes in MSS and individual item scores from day 0 to day 15 (preferably  $\pm 2$  days); (2) to describe current drug treatment patterns; (3) to describe the health-related quality of life status

associated with acute rhinosinusitis and related to the severity of the symptoms; and (4) to estimate the total costs related to episodes of acute rhinosinusitis and to the severity of the symptoms.

The study design is shown in Figure 1. Patients were recruited from the pool of patients attending the practice seeking medical treatment on an outpatient basis. Inclusion was based on duration of symptoms ( $\geq 7$  but  $\leq 28$  days) and severity of acute rhinosinusitis (MSS  $\geq 5$  but  $\leq 12$ ; see Questionnaires and data collection forms, available as online appendices at [www.thepcrj.org](http://www.thepcrj.org)).<sup>10</sup> Exclusion criteria included signs and symptoms suggestive of fulminant bacterial acute rhinosinusitis (fever  $\geq 38.3^\circ\text{C}$  or  $\geq 38.5^\circ\text{C}$  if a digital thermometer was not available, persistent severe unilateral facial or tooth pain, facial swelling, dental involvement or a worsening of symptoms after initial improvement).

On the inclusion day (day 0), questionnaires were completed by the patients, and demographic data, relevant medical history and recommended and/or prescribed medications were recorded by the investigator in a case report form. On day 15 (preferably  $\pm 2$  days), a telephone follow-up was conducted by an independent medically-qualified person.

### Questionnaires and data collection

#### Major Symptom Score (MSS)

The MSS<sup>17</sup> was the sum of scores for the five major symptoms rhinorrhoea, postnasal drip, nasal congestion/stuffiness, sinus headache and facial pain/pressure/tenderness (see Appendix 1, available online at [www.thepcrj.org](http://www.thepcrj.org)). Each symptom was graded as 0 (none), 1 (mild), 2 (moderate) or 3 (severe). The patients were asked to grade symptoms related to acute rhinosinusitis that they had experienced during 12 h prior to the visit on day 0 and to the follow-up telephone call on day 15. The questionnaire was reviewed by the investigator on day 0

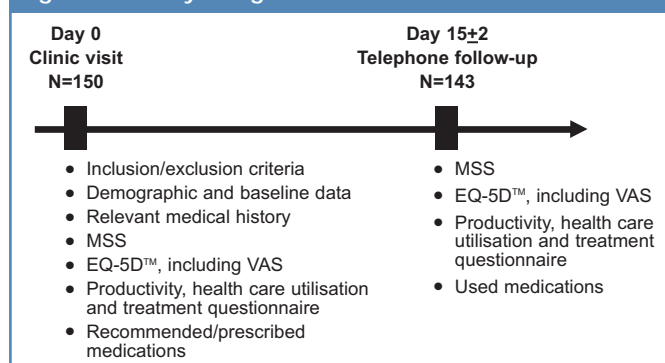
#### Quality of life

Quality of life was assessed by the EQ-5D<sup>TM</sup> (see Appendix 2 online).<sup>18</sup> The patients were asked to grade their health status on days 0 and 15. Mobility, self-care, usual activities, pain/discomfort and anxiety/depression were graded as 'none/no problems', 'moderate/some problems' or 'severe/extreme problems'. The patients also indicated their health status on a visual analogue scale (VAS) where 0 indicated worst conceivable status and 100 indicated best conceivable status.

#### Costs

The productivity, healthcare utilisation and treatment questionnaire completed on days 0 and 15 captured information on (1) outpatient visit and institutional care; (2) travel to the doctor; (3) employment status and absence due to illness; and (4) use of medications for acute rhinosinusitis (see Appendix 3 online). For the day 0 questionnaire, patients were asked to consider a recall period of 7 days. Thus the cost of 'one episode' of acute rhinosinusitis was all costs due to acute rhinosinusitis 7 days before the inclusion visit and 15 days after, a total of 22 days. To give productivity, transport and healthcare utilisation

Figure 1. Study design



monetary values, unit costs were assigned. For cost of medication, as a principle, the cost of the smallest pack(s) corresponding to the reported use was used. An exception was made for painkillers (non-steroidal anti-inflammatory drugs, paracetamol, etc) as it was assumed that patients would have a supply of painkillers at home and thus only the daily cost of painkillers was used to estimate cost. Unit costs for healthcare resource use, transport and medication were based on published price lists. Unit cost for productivity was derived from the mean annual income. Direct cost was the sum of costs for outpatient visits and institutional care, travel to the doctor and use of medications. Transportation costs from patients' visits up until and on day 0 were included. Indirect cost was the sum of costs for loss of productivity as derived from answers to questions on employment status, reduced productivity and absence due to illness. Total cost was the sum of direct and indirect costs.

The questionnaire was completed by the patient after information about the disease by the investigator.

### Statistical methods

All data were summarised by means of descriptive statistics and a 95% confidence interval (CI) for the proportion of responders was calculated. Pearson correlation was used to analyse EQ-5D™, VAS and total cost relationship to MSS.

For the presentation of descriptive statistics of EQ-5D™ and cost variables, the MSS symptom severity classes were defined as low (MSS 5–6), mid (MSS 7–9) and high (MSS 10–12).

## Results

### Studied population

A total of 150 patients were enrolled, of whom 143 provided data at follow-up (including six patients with MSS outside the prespecified MSS range).

Demographic data and main background characteristics are summarised in Table 1. The mean age was 45.1 years and 76.7% of the patients were female; 20.4% of the patients were current smokers and 2.7% were former smokers. The most common current or past medical history conditions were asthma (29.9%) and seasonal allergic rhinitis (26.5%). Sleep was impaired in approximately half of the patients due to the current episode of acute rhinosinusitis.

### Primary endpoint

The proportion of responders (i.e. the proportion of patients in whom MSS improved by at least 30% from day 0 to day 15) was 90.9% (95% CI: 85.0% – 95.1%).

### Secondary endpoints

#### *Mean change in MSS from day 0 to day 15*

Mean (SD) MSS decreased from 8.4 (2.3) on day 0 to 1.9 (2.2) on day 15. The mean (SD) change from baseline to day 15 in MSS was –6.4 (2.8). Ten patients showed no or limited change in MSS (0, 1 or –1). The mean (SD) change in individual MSS items were –1.5 (1.0) for sinus headache, –1.5 (1.0) for nasal congestion/ stuffiness, –1.3 (1.0) for facial

**Table 1. Demographic data and main background characteristics**

Variable	Statistic	Result
<i>Demographic data</i>		
Age (years)	No. of observations	150
	Mean (SD)	45.1 (14.5)
	Min, Max	18, 80
Gender	No. of observations	150
	Male	35 (23.3%)
	Female	115 (76.7%)
Smoker	No. of observations	147
	Non	113 (76.9%)
	Yes, current	30 (20.4%)
	Yes, former	4 (2.7%)
<i>Medical history</i>		
Asthma	No. of observations	147
	No	103 (70.1%)
	Yes	44 (29.9%)
Seasonal allergic rhinitis	No. of observations	147
	No	108 (73.5%)
	Yes	39 (26.5%)
Perennial allergic rhinitis	No. of observations	147
	No	132 (89.8%)
	Yes	15 (10.2%)
Sleep impairment*	No. of observations	146
	No	62 (42.5%)
	Yes	84 (57.5%)
*The question on day 0 was: "Is the patient's sleep impaired due to the current episode of acute rhinosinusitis?"		
SD=standard deviation.		

pain/pressure/tenderness, –1.2 (0.9) for postnasal drip and –0.9 (1.1) for rhinorrhoea.

### *Recommended and/or prescribed medications and use of medications*

Intranasal corticosteroids were the most recommended/prescribed medications (91%), followed by antibiotics (60%) and decongestant tablets (27%) (Table 2 and Figure 2).

### *EQ-5D™ dimensions and VAS*

On day 0, pain/discomfort was reported by 88.4% of patients and problems with usual activities by 43.2%. Extreme pain/discomfort

**Table 2. Treatment pattern of acute rhinosinusitis**

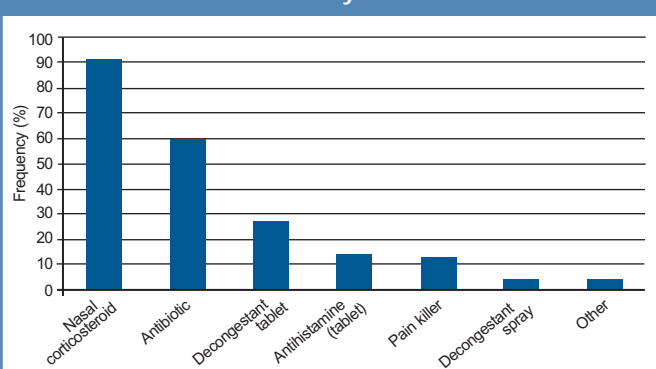
Category	Recommended and/or prescribed (N)*	Used (N)†
Nasal corticosteroid	137	127
Antibiotic	90	103
Decongestant tablet	41	49
Antihistamines (tablet)	22	6
Pain killer	21	21
Decongestant spray	8	12
Other‡	7	24

\*Data from day 0 in the case report form (N=150).

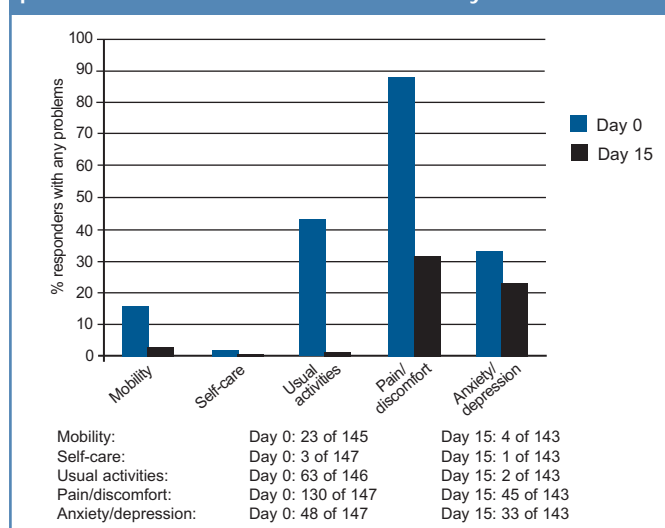
†Data from day 15 in the productivity, healthcare utilisation and treatment questionnaire (N=143).

‡For the category 'other', it was possible to fill in two different treatments; if two treatments were filled in, 'other' was counted only once per patient.

**Figure 2. Recommended and/or prescribed medication for acute rhinosinusitis on day 0**



**Figure 3. Self-reported health status (EQ-5D™) of patients with acute rhinosinusitis on days 0 and 15**



was reported by 10.9%, and 10.9% were also unable to perform their usual activities. Both dimensions showed improvement on day 15 when 31.5% of patients reported moderate/extreme pain and 1.4% reported having problems with usual activities. Self-care was the dimension least affected by acute rhinosinusitis (Figure 3).

Analysis of the EQ-5D™ data based on MSS symptom severity classes indicated that patients with higher MSS experienced extreme problems in the EQ-5D™ dimensions, mainly pain/discomfort and usual activities, to a greater extent than patients with lower MSS.

The mean (SD) VAS score improved from 58.7 (18.9) on day 0 to 79.5 (19.1) on day 15 (Table 3). The Pearson correlation coefficient between VAS and MSS on day 15 was  $-0.37$ . The Pearson correlation coefficient between VAS and MSS for the individual item scores on day 15 was  $-0.28$  for facial pain or tenderness on palpation,  $-0.32$  for sinus headache and  $-0.37$  for nasal congestion/stuffiness.

#### Costs (direct and indirect costs)

The mean total cost for one episode of acute rhinosinusitis was 10,260 SEK (€1,102 at May 2011, 1 SEK = €0.11), of which

**Table 3. Change in VAS in patients with rhinosinusitis from day 0 to day 15**

Day	No. of patients	Mean (SD) VAS
Day 0	146	58.7 (18.9)
Day 15	143	79.5 (19.1)
Day 15 – Day 0	142	21.0 (22.6)

SD=standard deviation, VAS=visual analogue scale.

**Table 4. Costs of an episode of acute rhinosinusitis**

Statistic	Direct cost	Indirect cost	Total cost
No. of observations	143	143	143
Mean (SD)	2,478 (1,256)	7,781 (9,639)	10,260 (10,104)
Median	1,981	3,456	5,993
Q1, Q3	1,867, 2,438	0, 13,943	2,325, 16,166
Min, Max	1,614, 10,099	0, 44,258	1,728, 54,357

Unit=SEK; 1 SEK= €0.11 (May 2011).

Direct cost is the sum of costs for doctor visit and institutional care, travel to doctor's visit, use of medications. Indirect cost is the sum of costs for fall in productivity (derived from employment status and absence due to illness). Total cost is the sum of direct and indirect costs.

SD=standard deviation

mean direct costs were 2,478 SEK (€266) (Table 4). The inter-individual variability in indirect costs was large (minimum 0 and maximum 44,258 SEK (€4,752)).

There was no trend in direct, indirect or total costs based on MSS symptom severity classes. Mean total costs ranged from 9,663 SEK (€1,037) for low severity (MSS 5–6) to 10,154 SEK (€1,090) for high severity (MSS 10–12).

The Pearson correlation coefficients between direct, indirect and total costs and MSS were 0.00, 0.11 and 0.11, respectively, on day 0, and 0.18, 0.11 and 0.13, respectively, on day 15.

## Discussion

### Main findings

In this observational study in primary care, nine out of 10 adult patients with acute rhinosinusitis showed significant improvement in symptoms 15 days after a visit to a primary care physician where patients were treated according to local practice, and 91% of patients were recommended and/or prescribed intranasal corticosteroids.

As the diagnosis 'acute rhinosinusitis' does not exist in the official Swedish registry of diagnoses, based on WHO International Classification of Diseases (ICD-10), it was not possible to perform a retrospective or prospective registry study in this case. We thus needed to perform a prospective observational study to collect data.

The study design was therefore based on the EP<sup>3</sup>OS 2007 recommendations for study definitions of acute rhinosinusitis and outcomes in primary care.<sup>4,5</sup> The inclusion criteria were based on data from an earlier clinical study,<sup>10</sup> international guidelines<sup>4,5</sup> and Swedish clinical practice. This approach tried to exclude

'common colds' by excluding patients with a symptom duration shorter than 7 days and an MSS <5, as well as excluding severe acute rhinosinusitis by not including patients with MSS >12 and by applying the exclusion criteria.

The results from the analyses of the primary endpoint showed a very high proportion of responders, with about 90% of patients demonstrating a clinically relevant improvement during the time period investigated. The results further showed that there was good symptom relief within the studied time period, as mean MSS decreased from 8.4 at baseline to 1.9 on day 15. This was a greater improvement than seen in the randomised placebo-controlled clinical study by Meltzer *et al.* in which MSS was used as a primary endpoint to compare mometasone furoate nasal spray with amoxicillin and placebo.<sup>10</sup> However, it should be noted that our study is not easily comparable with the study by Meltzer *et al.* as it was observational and was not designed to compare efficacy and safety of treatments for acute rhinosinusitis. Furthermore, MSS was used in a different way in our study from that in the study by Meltzer *et al.*

The treatments prescribed and used during the study period consisted mainly of intranasal corticosteroids and/or antibiotics. The finding of the surprisingly high use of intranasal steroids is an indication of a change in the clinical treatment of acute rhinosinusitis in primary care, consistent with the EP<sup>3</sup>OS evidence-based guidelines.<sup>4,5</sup>

The slight decrease in the number of patients using nasal corticosteroids and the corresponding increase in patients using antibiotics during the 15 days of study remains to be explained. One theory could be that some patients did not respond to initial nasal steroid treatment and were switched to antibiotics.

### Interpretation of findings in relation to previously published work

A comparison of the results from the current study with a recently published French survey of 397 general practitioners (GPs), summarising data from 1,585 patients, showed that the use of intranasal steroids was higher (91% vs. 38.7%) in our study and the use of oral antibiotics was lower (60% vs. 86.5%).<sup>19</sup> However, the study populations differed in characteristics as the French questionnaire survey was based on physician diagnosis of acute maxillary sinusitis with signs of a bacterial infection. A recent study performed in the Netherlands using questionnaires sent to GPs showed that oral antibiotics were prescribed by approximately one-third of the practitioners for mild and moderate acute rhinosinusitis.<sup>15</sup> In the current study, we found that antibiotic treatment was common, despite national programmes to reduce its use.<sup>20</sup> The most recent Cochrane update emphasised that antibiotics have a limited treatment effect in acute rhinosinusitis and stated that most cases will resolve without antibiotics within 2 weeks.<sup>21</sup> Our interpretation of the data is that, in contrast to medical evidence and recent guidelines, there still seems to be a general view among GPs that mild to moderate acute rhinosinusitis is the result of a bacterial infection.

In the Dutch study, decongestants were the most commonly prescribed treatments for both mild (91%) and moderate (83%) acute rhinosinusitis and were the first choice treatment in both cases.<sup>15</sup> The use of decongestants was approximately three times more common than in our study, which is interesting as the evidence level is low for the efficacy of decongestants in mild to moderate disease.<sup>4,5</sup> However, only one-third of the GPs in the Dutch study considered prescribing intranasal corticosteroids in mild or moderate acute rhinosinusitis.

The quality of life measure chosen by us was a generic and short validated method, the EQ-5D<sup>TM</sup>. As this is a non-interventional study, any 'intervention' (including time for filling out questionnaires) outside common clinical practice, taking an unreasonably long time for the patient (or the primary care centre) could have jeopardised the approval by the Ethics Committee.

The data suggest a high burden of acute rhinosinusitis on quality of life parameters, an area that has not so far been well studied. At baseline, pain/discomfort was reported by 88.4% of the patients and problems with usual activities by 43.2%. Both dimensions showed improvement on day 15. Investigation of the correlation between MSS and quality of life showed that increased MSS to a large extent gave worse problems in the EQ-5D<sup>TM</sup>, mostly pain/discomfort and usual activities. The mean health status measured using the VAS score improved from day 0 to day 15 and there was some correlation between EQ-5D<sup>TM</sup>, VAS and MSS on day 15. The data are in line with the results from a randomised placebo-controlled study by Bachert and Meltzer which demonstrated significant improvement in disease-specific quality of life among effectively treated patients with acute rhinosinusitis.<sup>16</sup> However, the SNOT-20 questionnaire used in that study was originally not developed for acute rhinosinusitis. Quality of life deserves further investigation and is a recommended assessment to be included in clinical studies of acute rhinosinusitis.<sup>22</sup>

The mean total cost for an episode of acute mild to moderate rhinosinusitis in patients seeking primary care was 10,260 SEK (approximately €1,102 at May 2011). The direct costs (medications, visits to the physician) constituted only about 25% of the total costs, indicating that the main costs for acute rhinosinusitis are due to loss in productivity (indirect costs). There was a large difference between mean and median numbers in the cost analysis, and a few patients reported very high costs. In cost of illness studies it is not uncommon to find large variability in patient level cost estimates.<sup>23,24</sup> When the cost was correlated to the MSS, some correlation between direct, indirect and total costs and MSS on day 15 could be shown. There was also some correlation for indirect and total costs with MSS on day 0.

This is the first observational study in Scandinavia to address costs for acute rhinosinusitis in primary care. A recent Swedish questionnaire study of allergic rhinitis and the common cold used a similar method to estimate productivity loss and found a mean productivity loss of €653 per worker per year, adding up to an

indirect cost of €2.7 billion for society as a whole.<sup>25</sup> Although acute rhinosinusitis is arguably less prevalent than allergic rhinitis and the common cold, the high cost per episode suggests that acute rhinosinusitis has a considerable financial impact on society.

### Limitations and strengths of this study

Weaknesses of the study are the recognised problems of observational studies as well as, in our case, a small population size and the use of a non-validated productivity, healthcare utilisation and treatment questionnaire. Having a pharmaceutical company as a sponsor of the study could potentially influence the treatment choices, even though local Swedish guidelines and European legislation on conduct of non-interventional studies were followed.

On the other hand, the strengths of our study are the primary care 'real-life' setting and the prospective design with independent follow-up. Furthermore, we used validated questions on quality of life (EQ-5D™) and symptom score (MSS).

### Conclusions

In conclusion, with a pharmacological treatment pattern dominated by intranasal corticosteroids, a high proportion of responders and good symptom relief were seen within the study time period. In addition, our data suggest a high burden of acute rhinosinusitis on quality of life and also a high cost for society, two areas that have not been well studied in Europe.

**Handling editor** Osman Mohammed Yusuf

**Statistical review** Gopal Netuveli

**Acknowledgements** The authors thank Patrick Svarvar (Merck) for input to the study design, Catarina Jansson-Blixt (TFS Sweden) for statistical support, Sari von Reedtz (MSD) for supporting the study report, Johanna Andlin (MSD) for supporting the study protocol and Hanna Liedman (TFS Sweden) for editorial support. The authors also thank all the investigators in the study; Rickard Ekesbo, Dalby; Ines Vinge, Lidingö; Bo Polhem, Uddevalla; Ingrid Johansson, Fjugesta; Tomas Lindskog, Växjö; Christer Dahlqvist, Limhamn; Per Helleke, Göteborg; Kjell Larsson, Arvidsjaur; Anne-Sofie Lindberg, Bro; Maria Papachristou Örebro; and Luisa Escuder, Bromma, all in Sweden.

**Conflicts of interest** BS and P Odebäck have received honoraria for educational activities from MSD, GSK and AstraZeneca. PS has received grants for studies and honoraria for educational activities from MSD, GSK, AstraZeneca and Novartis. P Olsson was employed by MSD during the conduct of the study and is currently employed by Boehringer-Ingelheim. JL is an employee of MSD and owns stock options and shares in Merck & Co. BS is an Associate Editor of the *PCRJ*, but was not involved in the editorial review of, nor the decision to publish, this article.

**Contributorship** All authors contributed equally to this paper.

**Funding** MSD (Sweden) AB funded this study, as well as the editorial support by Hanna Liedman.

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Appendix 1. MSS questionnaire

**QUESTIONNAIRE ON SYMPTOMS OF  
ACUTE RHINOSINUSITIS**

Patient number: \_\_\_\_\_

Day 0 Day 15 (+/-2)	Date: 200____ Month Day
------------------------	----------------------------

## Appendix 1. MSS questionnaire continued

**PLEASE SEE THE INSTRUCTIONS**

How have you been affected by the following symptoms during the last 12 hours?				
Facial pain, pressure or tenderness on palpation	0	1	2	3
Sinus headache	0	1	2	3
Rhinorrhea	0	1	2	3
Post-nasal drip	0	1	2	3
Nasal congestion/stuffiness	0	1	2	3



## Appendix 1. MSS questionnaire continued

**INSTRUCTIONS FOR:****QUESTIONNAIRE ON SYMPTOMS OF ACUTE RHINOSINUSITIS**

- Use a blue/black ball point pen
- Write legibly
- Tick only one box that corresponds to symptom severity

## Rhinosinusitis symptoms

**Facial pain, pressure, tenderness on palpation over the paranasal sinuses**

0 = None	No symptom evident
1 = Mild	Symptoms easily tolerated
2 = Moderate	Moderate symptoms that are bothersome but tolerable
3 = Severe	Severe symptoms hard to tolerate; causes interference with activities of daily living and/or sleeping

**Sinus headache (forehead, eye-region and temple)**

0 = None	No headache
1 = Mild	Mild headache, allowing normal activity
2 = Moderate	Disturbing but not prohibiting normal activity, bed rest is not necessary
3 = Severe	Severe headache; normal activity has to be discontinued, bed rest may be necessary

**Rhinorrhea (nasal discharge/runny nose)**

0 = None	No runny nose
1 = Mild	Snuffles; occasional wiping and/or nose blowing
2 = Moderate	Frequent wiping and/or nose blowing; frequently interrupts talking, is annoying
3 = Severe	Very frequent wiping and/or nose blowing. Very annoying and causes interference with activities of daily living and/or sleeping.

**Post-nasal drip (mucus in the throat)**

0 = None	No post-nasal drip
1 = Mild	Snuffles, occasional throat clearing
2 = Moderate	Frequent throat clearing; frequently interrupts talking, is annoying
3 = Severe	Very frequent throat clearing; constantly interrupts talking and causes interference with activities of daily living and/or sleeping

**Nasal congestion/stuffiness**

0 = None	No congestion, or same as before
1 = Mild	Slight block in one or both nostrils; nasal air flow somewhat impeded and annoying, no or only infrequent mouth breathing
2 = Moderate	Moderate block in one or both nostrils, nasal air flow noticeably impaired, is annoying, frequent mouth breathing
3 = Severe	Both nostrils completely blocked, very annoying, need to breath through the mouth all or almost all of the time

Appendix 2.



**Health Questionnaire**  
**English version for the UK**  
**(validated for Ireland)**

SAMPLE

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Appendix 2. continued

By placing a tick in one box in each group below, please indicate which statements best describe your own health state today.

**Mobility**

- I have no problems in walking about
- I have some problems in walking about
- I am confined to bed

**Self-Care**

- I have no problems with self-care
- I have some problems washing or dressing myself
- I am unable to wash or dress myself

**Usual Activities** (e.g. work, study, housework, family or leisure activities)

- I have no problems with performing my usual activities
- I have some problems with performing my usual activities
- I am unable to perform my usual activities

**Pain/Discomfort**

- I have no pain or discomfort
- I have moderate pain or discomfort
- I have extreme pain or discomfort

**Anxiety/Depression**

- I am not anxious or depressed
- I am moderately anxious or depressed
- I am extremely anxious or depressed

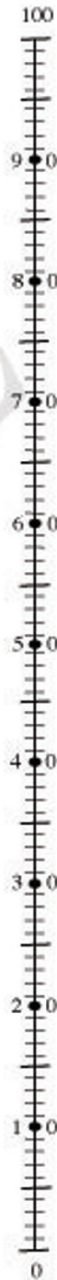
Appendix 2. continued

To help people say how good or bad a health state is, we have drawn a scale (rather like a thermometer) on which the best state you can imagine is marked 100 and the worst state you can imagine is marked 0.

We would like you to indicate on this scale how good or bad your own health is today, in your opinion. Please do this by drawing a line from the box below to whichever point on the scale indicates how good or bad your health state is today.

Your own  
health state  
today

Best  
imaginable  
health state



Worst  
imaginable  
health state

Appendix 3. Productivity, healthcare utilisation and treatment questionnaire

**Questionnaire for patients with acute sinusitis**

Patient number: — — — —

Date of visit: Year 20\_\_ Month \_\_\_\_\_ Day \_\_\_\_\_

**If you feel that anything in the questionnaire is unclear, or if you do not understand what it refers to, please ask a doctor or nurse for help.**

(By sinusitis is meant acute sinusitis)

## I. Doctor visit and institutional care

1. Have you **during the last week** visited a doctor or other caregiver as a result of your sinusitis?

- a  yes      b  no (go to question 3)

2. **If yes**, which of the following doctors or other caregivers have you visited?  
More than one alternative may be selected

- a  ear, nose and throat doctor, please state number of visits .....
- b  general practitioner, community health clinic, please state number of visits .....
- c  doctor's urgent care clinic, please state number of visits .....
- d  doctor emergency room, please state number of visits .....
- e  district nurse or other nurse, please state number of visits .....
- f  other caregiver, please state number of visits .....

3. Have you **during the last week** been admitted to a **hospital** as a result of your sinusitis?

- a  yes      b  no (go to question 5)

4. **If yes**, how many **nights** in total were you in the hospital as a result of your **sinusitis or other infection or inflammation in your respiratory passages** during the past week?

Number of nights: .....

Appendix 3. Productivity, healthcare utilisation and treatment questionnaire continued

**II Travel to doctor's visit**

**5. How many kilometers** have you travelled today from your home/work to make this visit to the doctor?

- a  0 - 2 kilometers
- b  2 - 4 kilometers
- c  4 - 6 kilometers
- d  6 or more kilometers

**6. What was your primary means of transportation** today, in order to make this visit to the doctor?

- a  Bicycled/walked
- b  Car
- c  Bus/train
- d  Other .....

**III Employment status and absence due to illness**

**7. What is your primary employment status?**

- a  Work full-time
- b  Work part-time
- c  Study full-time
- d  Study part-time
- e  Other (e.g. unemployed, parental leave, disability retired, retired, on sick leave for reason other than sinusitis)

**8. Have you been absent from your work/studies as a result of sinusitis during the past week?**

- a  yes
- b  no (go to question 10)

**9. If yes, how many days during the past week have you reported absent due to illness from work/studies as a result of sinusitis?**

Number of days: .....

**10. Have you during the past week had problems with sinusitis and still worked/studied?**

- a  yes
- b  no (go to question 13)

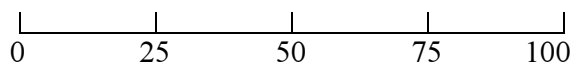
## Appendix 3. Productivity, healthcare utilisation and treatment questionnaire continued

**11. If yes, estimate how many days during the past week you have had problems with sinusitis and still worked/studied.**

Number of days: .....

**12. How would you rate your ability to perform at work/studies during those days you have had problems with sinusitis and still worked/studied? 0 = no ability  
100 = complete ability to perform**

(mark an X on the scale)



#### IV Use of medications for sinusitis

**13. Have you during the past week used any medication/natural remedy/self-treatment as a result of your sinusitis?**

a  yes

b  no

**14. If yes, which of the following preparations have you used?**

Mark an X for the preparations you used (several alternatives can be stated)

a  Antibiotic tablets/liquid                      Number of days .....

b  Cortisone nasal spray/drops/powder                      Number of days .....

c  Decongestant tablets                      Number of days .....

d  Decongestant spray/pipette                      Number of days .....

e  Allergy tablets/antihistamine tablets/                      Number of days .....

liquid/dissolving

f  Pain-relief tablets/liquid                      Number of days .....

g  Other medication/natural remedy

indicate type ..... Number of days .....

indicate type ..... Number of days .....