



MINISTRY OF HEALTH
SINGAPORE

Management of Rhinosinusitis and Allergic Rhinitis

MOH Clinical Practice Guidelines 2/2010



Academy of Medicine,
Singapore



Chapter of
Otorhinolaryngologists
College of Surgeons,
Singapore



College of Family Physicians,
Singapore



Feb 2010

Levels of evidence and grades of recommendation

Levels of evidence

Level	Type of Evidence
1 ⁺⁺	High quality meta-analyses, systematic reviews of randomised controlled trials (RCTs), or RCTs with a very low risk of bias.
1 ⁺	Well conducted meta-analyses, systematic reviews of RCTs, or RCTs with a low risk of bias.
1 ⁻	Meta-analyses, systematic reviews of RCTs, or RCTs with a high risk of bias
2 ⁺⁺	High quality systematic reviews of case control or cohort studies. High quality case control or cohort studies with a very low risk of confounding or bias and a high probability that the relationship is causal
2 ⁺	Well conducted case control or cohort studies with a low risk of confounding or bias and a moderate probability that the relationship is causal
2 ⁻	Case control or cohort studies with a high risk of confounding or bias and a significant risk that the relationship is not causal
3	Non-analytic studies, e.g. case reports, case series
4	Expert opinion

Grades of recommendation

Grade	Recommendation
A	At least one meta-analysis, systematic review of RCTs, or RCT rated as 1 ⁺⁺ and directly applicable to the target population; or A body of evidence consisting principally of studies rated as 1 ⁺ , directly applicable to the target population, and demonstrating overall consistency of results
B	A body of evidence including studies rated as 2 ⁺⁺ , directly applicable to the target population, and demonstrating overall consistency of results; or Extrapolated evidence from studies rated as 1 ⁺⁺ or 1 ⁺
C	A body of evidence including studies rated as 2 ⁺ , directly applicable to the target population and demonstrating overall consistency of results; or Extrapolated evidence from studies rated as 2 ⁺⁺
D	Evidence level 3 or 4; or Extrapolated evidence from studies rated as 2 ⁺
GPP (good practice points)	Recommended best practice based on the clinical experience of the guideline development group.

CLINICAL PRACTICE GUIDELINES

**Management of Rhinosinusitis
and Allergic Rhinitis**

MOH Clinical Practice Guidelines 2/2010

Published by Ministry of Health, Singapore
16 College Road,
College of Medicine Building
Singapore 169854

Printed by Golden City Colour Printing Co. (Pte) Ltd

Copyright © 2010 by Ministry of Health, Singapore

ISBN 978-981-08-5211-5

Available on the MOH website: <http://www.moh.gov.sg/cpg>

Statement of Intent

These guidelines are not intended to serve as a standard of medical care. Standards of medical care are determined on the basis of all clinical data available for an individual case and are subject to change as scientific knowledge advances and patterns of care evolve.

The contents of this publication are guidelines to clinical practice, based on the best available evidence at the time of development. Adherence to these guidelines may not ensure a successful outcome in every case. These guidelines should neither be construed as including all proper methods of care, nor exclude other acceptable methods of care. Each physician is ultimately responsible for the management of his/her unique patient, in the light of the clinical data presented by the patient and the diagnostic and treatment options available.

Contents

	Page
Executive summary of recommendations	1
1. Introduction	15
2. Management of common cold (acute viral rhinosinusitis) and use of antibiotics in acute bacterial rhinosinusitis	17
3. Management of infective rhinosinusitis in adults	21
4. Management of infective rhinosinusitis in children	38
5. Management of allergic rhinitis	48
6. Management of rhinitis in pregnancy	61
7. Cost-effectiveness issues	66
8. Clinical quality improvement	67
References	68
Multiple choice questions	88
Workgroup members	91

Foreword

Rhinosinusitis is one of the most common health problems encountered by primary care physicians worldwide. The condition is made up of infectious rhinitis (including upper respiratory tract infections) and allergic rhinitis.

In Singapore, the 2005 Primary Care Survey¹ found that the leading condition seen at primary care clinics was upper respiratory tract infections, accounting for 29% of consultations; an earlier study² put the prevalence of rhinitis in schoolchildren at 44%. While usually self-limited, the commonness of the common cold (acute viral rhinosinusitis) means that it is a condition with significant impact on ability to function and quality of life. The common cold should also be distinguished from acute bacterial rhinosinusitis. Variation in investigation and treatment is a concern, as is inappropriate use of antibiotics.

The prevalence of allergic rhinitis in the general adult population in Singapore has been estimated at 5.5%.³ Allergic rhinitis is more common in children and prevalence may be as high as 40% in school-going children.⁴ Allergic rhinitis in children may be associated with co-morbidities including asthma, atopic dermatitis/eczema, allergic conjunctivitis, chronic sinusitis and chronic otitis media with effusion.

It is therefore timely to develop this first national guideline which incorporates the best available evidence from the scientific literature and expert consensus, to assist primary care physicians in the management of rhinosinusitis. I hope you will find the guidance useful in managing your patients.

PROFESSOR K SATKU
DIRECTOR OF MEDICAL SERVICES

¹ Primary Care Survey 2005, Integrated Health Services Division, Ministry of Health, Singapore.

² Goh DYT, Chew FT, Quek SC, Lee BW. Prevalence and severity of asthma, rhinitis, and eczema in Singapore schoolchildren. *Arch Dis Child* 1996;74:131–135.

³ Ng TP, Tan WC. Epidemiology of chronic (perennial) rhinitis in Singapore: prevalence estimates, demographic variation and clinical allergic presentation. *Ann Acad Med Singapore*. 1994 Jan;23(1):83-8.

⁴ Wright AL, Holberg CJ, Martinez FD, Halonen M, Morgan W, Taussig LM. Epidemiology of physician-diagnosed allergic rhinitis in childhood. *Pediatrics* 1994; 94:895- 901.

Executive summary of key recommendations

Details of recommendations can be found in the main text at the pages indicated.

Management of common cold (acute viral rhinosinusitis) and use of antibiotics in acute bacterial rhinosinusitis

Acute viral rhinosinusitis (common cold)

A Antibiotics are not recommended for treatment of the common cold in children or adults (pg 17).

Grade A, Level 1++

A Dextromethorphan should be considered as a treatment option for adults with cough caused by the common cold (pg 17).

Grade A, Level 1++

A Topical (intranasal) or oral nasal decongestants, used for up to three days, is recommended for adolescents and adults with the common cold (pg 17).

Grade A, Level 1+

A Topical ipratropium may be considered as a treatment option for nasal congestion in children older than six years and in adults with moderate to severe common cold (pg 18).

Grade A, Level 1+

A Codeine and other narcotics, dextromethorphan, anti-histamines, and combination anti-histamine/decongestants are not recommended to treat cough or other cold symptoms in children (pg 18).

Grade A, Level 1++

A First-generation anti-histamines and combination anti-histamine/decongestants may be considered for cough and cold symptoms in adults if the benefits outweigh the adverse effects (pg 18).

Grade A, Level 1++

A Vitamin C, zinc, and Echinacea are not recommended for active treatment of common cold due to lack of effectiveness in preventing the common cold (pg 18).

Grade A, Level 1++

Use of antibiotics in acute bacterial rhinosinusitis

Adults

A Antibiotics are not recommended for adults with non-severe acute bacterial rhinosinusitis (mild pain and temperature < 38.3 degrees centigrade) till after 10 days of symptoms from onset (pg 18).

Grade A, Level 1+

D Besides severity of illness, the patient's age, general health, cardiopulmonary status, and co-morbid conditions should be considered in deciding start of antibiotic treatment in patients with acute bacterial rhinosinusitis (pg 19).

Grade D, Level 4

A The first-line empiric antibiotic for adults with acute bacterial rhinosinusitis is amoxicillin. If the patient is allergic to amoxicillin, trimethoprim-sulfamethoxazole or macrolides may be used (pg 19).

Grade A, Level 1+

A For adults with acute bacterial rhinosinusitis, the recommended duration of appropriate oral antibiotic regime is 7 days. Clinician assessment after 7 days is recommended. Antibiotic regime can be extended to 14 days if patient's symptoms fail to resolve (pg 19).

Grade A, Level 1++

B A second-line antibiotic such as high dose amoxicillin-clavulanate, ampicillin-sulbactam or flouroquinolone should be considered in adults with acute bacterial rhinosinusitis if there is no clinical response after at least 7 days of first line antibiotics (pg 19).

Grade B, Level 2+

Children

D Appropriate antibiotic regimes are recommended for children with the following conditions:

1. Non-severe acute bacterial rhinosinusitis: In a child with protracted symptoms with asthma, chronic bronchitis or acute otitis media.
2. Severe acute bacterial rhinosinusitis: In ambulatory patients, an oral antibiotic resistant to beta-lactamase enzymes (amoxicillin-clavunate or a second generation cephalosporin such as cefuroxime axetil).

3. Severe illness or toxic condition: In a child with suspected or proven suppurative complication.
(pg 20) **Grade D, Level 4**

D Intravenous antibiotic effective against penicillin-resistant *Streptococcus pneumoniae*, beta-lactamase producing *Haemophilus influenzae* and *Moraxella catarrhalis* should be used in children with severe acute bacterial rhinosinusitis (pg 20).

Grade D, Level 4

D Amoxicillin (45 mg/kg/day, doubled if age under 2 years or with risk factors for resistance) is recommended for a child with non-severe acute bacterial rhinosinusitis with protracted symptoms. If the symptoms do not improve within 72 hours, an antibiotic against the resistant organism prevalent in the community should be considered. Azithromycin or clarithromycin as first-line therapy is recommended in penicillin allergy (pg 20).

Grade D, Level 4

Management of infective rhinosinusitis in adults

Acute rhinosinusitis

GPP Other diagnosis should be considered in adults with acute rhinosinusitis who present with unilateral symptoms of bleeding, crusting, or anosmia (pg 22).

GPP

D Immediate referral to an ENT specialist is indicated for acute rhinosinusitis in adults who present with sinister signs indicative of complications of acute intermittent rhinosinusitis. These include:

- Peri-orbital oedema
- Displaced globe
- Double vision
- Ophthalmoplegia
- Reduced visual acuity
- Severe unilateral or bilateral frontal headache
- Frontal swelling
- Signs of meningitis or focal neurological deficits

(pg 23)

Grade D, Level 4

D Plain sinus x-rays are not recommended for the diagnosis of acute rhinosinusitis in adults (pg 23).

Grade D, Level 4

Treatment of acute rhinosinusitis

D Alleviate symptoms of mild acute rhinosinusitis in adults with the following options

- Decongestants
- Nasal saline spray and/ or irrigation
- Antihistamines, only in patients with concomitant allergic rhinitis
- Analgesics

(pg 23)

Grade D, Level 4

D Treat underlying inflammatory process of moderate to severe acute rhinosinusitis in adults with:

- Intranasal steroid
- Antibiotic, empiric: 7-14 days

Alleviate symptoms with the following options:

- Decongestants
- Nasal saline spray and/ or irrigation
- Antihistamines, in patients with concomitant allergic rhinitis
- Analgesics

(pg 24)

Grade D, Level 4

GPP The workgroup recommends that patients with acute rhinosinusitis should be reviewed for **symptom resolution**. Patients whose symptoms worsen or persist despite therapy should be referred to a specialist for further evaluation and management (pg 24).

GPP

A Nasal steroid spray twice daily is recommended for adults with acute rhinosinusitis which has not resolved after 5 days of initial presentation (pg 26).

Grade A, Level 1+

A Oral corticosteroids are not recommended for adults with acute rhinosinusitis (pg 27).

Grade A, Level 1+

D Antihistamines are not recommended in the treatment of acute bacterial rhinosinusitis in adults (pg 27).

Grade D, Level 4

A Antihistamines may be used as an adjunct to antibiotic treatment in acute bacterial rhinosinusitis patients with concomitant allergic rhinitis (pg 27).

Grade A, Level 1+

D New generation oral antihistamines are preferred in adults with acute rhinosinusitis for their favourable efficacy/safety ratio and pharmacokinetics. Refrain from first generation antihistamines to avoid sedation and anti-cholinergic side effects (pg 27).

Grade D, Level 4

GPP Topical decongestants may be used for adults with acute rhinosinusitis whose symptoms fail to resolve after 10 days of initial presentation (pg 27).

GPP

GPP The duration of treatment with topical decongestants should be limited to less than 10 days to avoid rhinitis medicamentosa (pg 28).

GPP

A Nasal hypertonic saline irrigation, alone or in conjunction with other adjunctive measures, may be used to reduce symptoms and medication use in adults with frequent acute rhinosinusitis (pg 28).

Grade A, Level 1+

D Mucolytics are not recommended to be prescribed routinely for adult patients with acute rhinosinusitis (pg 29).

Grade D, Level 4

Chronic rhinosinusitis

GPP All adults with persistent and recurrent rhinosinusitis should be referred to a specialist for nasal endoscopy to assess for differential causes (pg 30).

GPP

GPP Other diagnosis should be considered in adults with chronic rhinosinusitis who present with unilateral symptoms of bleeding, crusting, or cacosmia (pg 30).

GPP

D Immediate referral to an ENT specialist is indicated for chronic rhinosinusitis in adults who present with sinister signs such as:

- Peri-orbital oedema
- Displaced globe
- Double vision
- Ophthalmoplegia
- Reduced visual acuity
- Severe unilateral or bilateral frontal headache
- Frontal swelling
- Signs of meningitis or focal neurological deficits.

(pg 31)

Grade D, Level 4

D Sinus x-rays are **not** recommended to support the diagnosis of chronic rhinitis in adults (pg 31).

Grade D, Level 4

Treatment of chronic rhinosinusitis without nasal polyps

D For chronic rhinosinusitis without nasal polyps, alleviate symptoms with the following options:

- Nasal saline irrigation

Treat underlying inflammatory process with:

- Intranasal steroid
- Antibiotic, in patients with acute exacerbation of chronic rhinosinusitis, culture directed: 10-14 days

(pg 33)

Grade D, Level 4

C Short-term oral antibiotics are recommended for acute exacerbation of chronic rhinosinusitis without nasal polyps (pg 33).

Grade C, Level 2+

A Nasal corticosteroids may be prescribed for chronic rhinosinusitis without nasal polyps (pg 33).

Grade A, Level 1+

A Nasal saline irrigation may be prescribed for chronic rhinosinusitis without nasal polyps (pg 34).

Grade A, Level 1+

GPP Oral steroids, oral/topical decongestants, mucolytics or antihistamines are not recommended in treatment of chronic rhinosinusitis without nasal polyps (pg 34).

GPP

Treatment of chronic rhinosinusitis with nasal polyps

D For chronic rhinosinusitis with nasal polyps, alleviate symptoms with the following options:

- Nasal saline irrigation
- Anti-histamines, in patients with concomitant allergic rhinitis

Treat underlying inflammatory process with:

- Intranasal steroid.

(pg 35)

Grade D, Level 4

GPP Adults with chronic rhinosinusitis with nasal polyps should be reviewed for symptom control. Patients whose symptoms worsen during or persist despite therapy should be referred to a specialist for further evaluation and management (pg 35).

GPP

C Short-term oral antibiotics are recommended to improve symptoms in acute exacerbation of chronic rhinosinusitis with nasal polyps (pg 36).

Grade C, Level 2+

C Long-term, low-dose macrolide therapy may be considered for chronic rhinosinusitis patients with nasal polyps (pg 36).

Grade C, Level 2+

GPP Management by a specialist is recommended for patients with chronic rhinosinusitis with nasal polyps being prescribed long term, low dose macrolide therapy, in view of its side effects (pg 36).

GPP

A Nasal corticosteroid therapy may be used in adults with chronic rhinosinusitis with nasal polyps (pg 36).

Grade A, Level 1+

C Antihistamines are not recommended in chronic rhinosinusitis with nasal polyps (pg 37).

Grade C, Level 2+

Management of infective rhinosinusitis in children

GPP Allergic rhinitis often coexists with paediatric acute and chronic rhinosinusitis. The history should evaluate for symptoms of allergic rhinitis and identify possible allergens (pg 39).

GPP

GPP Otoscopy should be performed routinely to exclude otitis media in paediatric acute and chronic rhinosinusitis (pg 40).

GPP

D Plain X-ray is not recommended routinely to confirm the diagnosis of rhinosinusitis in children (pg 40).

Grade D, Level 4

A Topical corticosteroids may be used in children as an adjunct to antibiotics. It can reduce the cough and nasal discharge earlier in acute bacterial rhinosinusitis (pg 43).

Grade A, Level 1+

GPP Topical decongestants should be used in children no longer than 4-5 days to avoid toxicity and rhinitis medicamentosa (pg 43).

GPP

D Saline nose drops or sprays may be considered to decrease the mucus trapping and crusting associated with acute rhinosinusitis in children (pg 43).

Grade D, Level 3

D The workgroup recommends antibiotics use only in acute exacerbation of paediatric chronic rhinosinusitis, by following the recommendations from the Consensus Meeting in Brussels, 1996:

- For chronic rhinosinusitis, especially with frequent exacerbations, 2 weeks of oral antibiotics is advised. The antibiotic is changed if there is no response within 5-7 days.
- Failing this, sinus secretions for culture or investigations to exclude recalcitrant causes are considered.
- If there is slow response, a second 2-week course can be prescribed.

- In rare cases with clear-cut improvement but persisting symptoms, a 3rd course can be given before surgery is considered.
- Parenteral antibiotic may be appropriate if oral antibiotics fail.

(pg 44)

Grade D, Level 4

C Nasal douching may be considered for paediatric chronic rhinosinusitis (pg 44).

Grade C, Level 2+

D Antral lavage, inferior meatal antrostomy (except possibly in primary ciliary dyskinesia), Caldwell-Luc operation (risks damage to un-erupted teeth) are not recommended in paediatric chronic rhinosinusitis (pg 45).

Grade D, Level 3

Management of allergic rhinitis

GPP The diagnosis of allergic rhinitis should be made based upon concordance between a typical history of allergic symptoms and diagnostic tests (pg 48).

GPP

D The workgroup recommends using the algorithm for the diagnosis and assessment of severity of allergic rhinitis proposed by ARIA 2008 (refer to Figure 6 on page 48) (pg 48).

Grade D, Level 4

GPP Besides a nasal examination for allergic rhinitis, look out for:

- Ocular signs of irritation e.g. allergic conjunctivitis; redness and rubbing of eyes indicative of itchiness.
- Chest examination to rule out concurrent asthma.

(pg 49)

GPP

D The workgroup recommends using the algorithm for the classification of allergic rhinitis proposed by ARIA 2008 (refer to Figure 7 on page 49) (pg 49).

Grade D, Level 4

D The workgroup recommends using the algorithm for the management of allergic rhinitis proposed by ARIA 2008 (refer to Figure 8 on page 51) (pg 50).

Grade D, Level 4

GPP Mattress encasings or High Efficiency Particulate Air Filters for house dust mite and pet allergy in adults with rhinitis should be part of the overall management of allergic rhinitis (pg 52).

GPP

A Second-generation oral or intranasal H1-antihistamines are recommended for the treatment of allergic rhinitis and conjunctivitis in adults and children (pg 52).

Grade A, Level 1++

A Intranasal glucocorticosteroids are strongly recommended for the treatment of allergic rhinitis in adults and children (pg 52).

Grade A, Level 1++

D Intramuscular glucocorticosteroids and the long term use of oral preparations are not recommended for the treatment of allergic rhinitis due to safety concerns (pg 52).

Grade D, Level 3

A Topical H1-antihistamines are recommended for the treatment of allergic rhinitis and conjunctivitis. Its therapeutic effects are superior and faster than oral anti-histamines (pg 52).

Grade A, Level 1+

A Intranasal ipratropium may be considered as a treatment option for rhinorrhoea associated with allergic rhinitis (pg 52).

Grade A, Level 1+

A Topical chromones should be considered as a treatment option for allergic rhinitis and conjunctivitis. However, they are only moderately effective (pg 53).

Grade A, Level 1+

A Montelukast may be considered as a treatment option for seasonal allergic rhinitis and asthma in patients over 6 years of age. It should not be used more than 4 weeks since there is limited data of its efficacy in patients with persistent allergic rhinitis for more than 4 weeks (pg 53).

Grade A, Level 1+

C Intranasal decongestants may be used for a short period of time in patients with severe nasal obstruction caused by allergic rhinitis (pg 53).

Grade C, Level 2+

C Oral decongestants (and their combination with oral H1-antihistamines) may be considered in the treatment of allergic rhinitis in adults, but side effects are common (pg 53).

Grade C, Level 2++

GPP Education of patient and/or patient's carer on the management of allergic rhinitis should be considered as an option to maximize compliance and optimize treatment outcomes (pg 54).

GPP

Paediatric aspects of allergic rhinitis

GPP Symptoms of sneezing, nasal itching, discharge and congestion that persist longer than 2 weeks should prompt a search for a cause other than infection in children (pg 55).

GPP

GPP It is recommended to ask about family history of atopy and progression of atopy of the child (pg 55).

GPP

B Skin prick tests should be performed and interpreted reliably early in life (pg 55).

Grade B, Level 2+

GPP The principles of treatment are the same in children as in adults with allergic rhinitis, but dosages should be adapted and care should be taken to avoid the side effects involving impairment of growth and cognitive development (pg 56).

GPP

GPP Pharmacologic management for allergic rhinitis in children should be individualized and polypharmacy avoided (pg 56).

GPP

A Intranasal glucocorticosteroid with bioavailability of <1% such as fluticasone propionate or mometasone furoate should be considered as a treatment option for allergic rhinitis and allergic conjunctivitis (pg 56).

Grade A, Level 1++

B Intranasal glucocorticosteroids with high bioavailability such as betamethasone should not be used in children with allergic rhinitis due to its effect upon growth and growth velocity (pg 56).

Grade B, Level 1++

A Oral and depot glucocorticosteroid preparations should be avoided in children with allergic rhinitis due to negative effect on short term growth and growth velocity (pg 56).

Grade A, Level 1+

A Second generation H1-antihistamines such as cetirizine, levocetirizine and loratadine should be considered as a treatment option in the treatment of allergic rhinitis in children (pg 57).

Grade A, Level 1+

GPP Nasal saline drops or spray may be considered in children with allergic rhinitis to clear the nose before eating or sleeping (pg 57).

GPP

A Sublingual immunotherapy (SLIT) should be considered in children above age 5 years who have poor symptomatic control of allergic rhinitis despite maximal therapy or who cannot or will not take medication (pg 57).

Grade A, Level 1++

GPP The family and the child should be educated about the recurrent or persistent nature of the disease, allergen avoidance and avoidance of allergen triggers and respiratory tract irritants, the most important of which is tobacco smoke (pg 58).

GPP

Management of rhinitis in pregnancy

D Nasal endoscopy on a decongested nose may be considered as an option to differentiate pregnancy rhinitis from sinusitis (pg 61).

Grade D, Level 4

D Imaging studies are not recommended to make a diagnosis in rhinitis in pregnancy (pg 61).

Grade D, Level 4

D Skin prick tests are not recommended for rhinitis in pregnancy because use of potent antigens in skin testing may be associated with systemic reactions (pg 61).

Grade D, Level 4

GPP In treating rhinitis of pregnancy, all drug therapy should ideally be avoided especially in the first trimester. If drug therapy cannot be avoided then treatment will depend upon the predominant symptoms, with the topical agents as first line since they have minimal systemic exposure (pg 62).

GPP

C Cromones are safe with no known teratogenic effect but they are moderately effective. It may be given for the treatment of rhinitis in the first 3 months of pregnancy, 3-4 times daily (pg 62).

Grade C, Level 2+

C If cromones are ineffective and poorly tolerated, they should be replaced with anti-histamines. Chlorpheniramine and tripeleminamine are the anti-histamines of choice for pregnant women with rhinitis. Cetirizine and loratadine may be considered after the first trimester (pg 62).

Grade C, Level 2+

C Intranasal steroids should be prescribed as an alternative to or in combination with anti-histamines for severe cases of rhinitis in pregnancy (pg 62).

Grade C, Level 2+

C Budesonide is the only recommended intranasal steroid for rhinitis in pregnancy (pg 62)

Grade C, Level 2+

C Topical decongestants like oxymetazoline may be considered as second-line therapy for short-term relief and when no other safer alternatives are available for the treatment of rhinitis in pregnancy (pg 63).

Grade C, Level 2+

C Oral decongestants are not recommended for rhinitis in pregnancy (pg 63).

Grade C, Level 2+

C Leukotriene Modifiers are not recommended for allergic rhinitis in pregnancy (pg 63).

Grade C, Level 2+

A Amoxicillin is the drug of choice for pregnant patients with rhinitis who are not allergic to penicillin (pg 63).

Grade A, Level 1+

D Amoxicillin-clavulanate or cephalosporin may be given to pregnant women with rhinitis not responding to amoxicillin (pg 63).

Grade D, Level 3

C Metronidazole should be used in rhinitis in pregnancy caused by anaerobic pathogens (pg 63).

Grade C, Level 2+

D Immunotherapy is not recommended for rhinitis in pregnancy. However, it may be continued if the maintenance phase has been reached (pg 63).

Grade D, Level 4

1 Introduction

1.1 Background information

The two most common diagnoses of rhinitis encountered in clinical practice in Singapore are infectious rhinitis, including upper respiratory tract infections, and allergic rhinitis. These two conditions form the main focus of this clinical practice guideline.

Rhinitis is defined as an inflammation of the lining of the nose and is characterized by nasal symptoms including anterior or posterior rhinorrhea, sneezing, nasal blockage and/or itching of the nose. These symptoms occur during two or more consecutive days for more than one hour on most days.

Sinusitis and rhinitis usually coexist and are concurrent in most individuals; thus, the correct terminology for sinusitis is **rhinosinusitis**.

These clinical practice guidelines aim to help clinicians manage rhinosinusitis based on the best available evidence as well as expert opinion in areas where studies are lacking.

1.2 Development of guidelines

Clinical practice guidelines from the World Health Organization, USA and the European Union were evaluated and local data when available were included into this CPG. Recommendations are based on locally available prescriptions and procedures. This workgroup was made up of otorhinolaryngologists with a special interest in rhinology and paediatric otorhinolaryngology, pediatricians and a general practitioner.

1.3 Objectives

The main objective of these guidelines is to provide evidence based management strategies for the diagnosis and treatment of the two most common causes of rhinosinusitis i.e. infective rhinosinusitis and allergic rhinitis at the primary care level and guidelines for specialist referral.

1.4 Review of guidelines

Evidence based clinical guidelines are only as current as the evidence that supports them. Users must keep in mind that new evidence could supersede recommendations in these guidelines. The workgroup advises that these guidelines be scheduled for review five years after publication or if new evidence appears that requires substantive changes to the recommendations.

2 Management of common cold (acute viral rhinosinusitis) and use of antibiotics in acute bacterial rhinosinusitis

There is a very high incidence of acute viral rhinosinusitis (common cold). Adults, in general, suffer 2 to 5 colds per year and school children 6 to 8 colds per year. The most common agents are Rhinovirus (24%) and Influenza virus (11%).¹ Bacterial infections, which may be mild and often resolve spontaneously, complicate only 0.5% to 2% of acute viral rhinosinusitis.²

Differentiating between a bacterial (acute bacterial rhinosinusitis) and viral (viral rhinosinusitis) etiology is largely symptom and duration based.

2.1 Acute viral rhinosinusitis (common cold)

In acute viral rhinosinusitis, the duration of symptoms is usually less than 10 days.

A common cold is characterized by sore throat, malaise and low-grade fever at onset. These symptoms resolve within a few days and are followed by nasal congestion, rhinorrhea, and cough within 24 to 48 hours after onset of the first symptoms. The second set of symptoms are what prompt most patients to see a physician for relief.³ Symptoms usually peak around day 3 or 4 and begin to resolve by day 7.⁴ Nasal discharge, appearing at the peak of illness, can become thick and purulent and may be misdiagnosed as a bacterial sinus infection.¹

A Antibiotics are not recommended for treatment of the common cold in children or adults.^{5,6}

Grade A, Level 1++

A Dextromethorphan should be considered as a treatment option for adults with cough caused by the common cold.^{7,8}

Grade A, Level 1++

A Topical (intranasal) or oral nasal decongestants, used for up to three days, is recommended for adolescents and adults with the common cold.^{9,10}

Grade A, Level 1+

A Topical ipratropium may be considered as a treatment option for nasal congestion in children older than six years and in adults with moderate to severe common cold.¹¹

Grade A, Level 1+

A Codeine and other narcotics, dextromethorphan, anti-histamines, and combination anti-histamine/decongestants are not recommended to treat cough or other cold symptoms in children.^{7,12-14}

Grade A, Level 1++

A First-generation anti-histamines and combination anti-histamine/decongestants may be considered for cough and cold symptoms in adults if the benefits outweigh the adverse effects.¹⁵

Grade A, Level 1++

A Vitamin C, zinc, and Echinacea are not recommended for active treatment of common cold due to lack of effectiveness in preventing the common cold.^{15,16,16a}

Grade A, Level 1++

However, among available complementary treatments, vitamin C prophylaxis may decrease the severity and duration of cold symptoms.^{15,16}

2.2 Use of antibiotics in acute bacterial rhinosinusitis

When symptoms persist for more than 10 days, or double worsening occurs in which symptoms initially improve, but then worsen within a 10-day period, bacterial sinusitis is presumed.¹⁷

ADULTS

A Antibiotics are not recommended for adults with non-severe acute bacterial rhinosinusitis (mild pain and temperature < 38.3 degrees centigrade) till after 10 days of symptoms from onset.^{18,19}

Grade A, Level 1+

The rationale for observing acute bacterial rhinosinusitis is based upon a high percentage of spontaneous improvement when patients receive placebo in randomized controlled trials and the modest incremental benefit from antibiotic therapy.^{18,19}

Using this time course of symptoms as a guide reduces the inappropriate use of antibiotics for viral illness and reduces the development of antibiotic resistant organisms and side effects associated with antibiotic use.²⁰

D Besides severity of illness, the patient's age, general health, cardiopulmonary status, and co-morbid conditions should be considered in deciding start of antibiotic treatment in patients with acute bacterial rhinosinusitis.¹⁷

Grade D, Level 4

A The first-line empiric antibiotic for adults with acute bacterial rhinosinusitis is amoxicillin.^{19,21-25} If the patient is allergic to amoxicillin, trimethoprim-sulfamethoxazole^{18,23,24,26,27} or macrolides may be used.

Grade A, Level 1+

A For adults with acute bacterial rhinosinusitis, the recommended duration of appropriate oral antibiotic regime is 7 days. Clinician assessment after 7 days is recommended. Antibiotic regime can be extended to 14 days if patient's symptoms fail to resolve.²⁸

Grade A, Level 1++

B A second-line antibiotic such as high dose amoxicillin-clavulanate, ampicillin-sulbactam or flouroquinolone should be considered in adults with acute bacterial rhinosinusitis if there is no clinical response after at least 7 days of first line antibiotics.^{29,30}

Grade B, Level 2+

Short-course oral antibiotic regime has comparable effectiveness to a longer course of therapy in acute bacterial rhinosinusitis. Shorter duration of treatment, particularly for patients without severe disease and complicating factors, might lead to fewer adverse events, better patient compliance, lower rates of resistance development and fewer costs.²⁸

CHILDREN

D Appropriate antibiotic regimes are recommended for children with the following conditions:³¹

1. Non-severe acute bacterial rhinosinusitis: In a child with protracted symptoms with asthma, chronic bronchitis or acute otitis media.
2. Severe acute bacterial rhinosinusitis: In ambulatory patients, an oral antibiotic resistant to beta-lactamase enzymes (amoxicillin-clavunate or a second generation cephalosporin such as cefuroxime axetil).
3. Severe illness or toxic condition: In a child with suspected or proven suppurative complication.

Grade D, Level 4

D Intravenous antibiotics effective against penicillin-resistant *Streptococcus pneumoniae*, beta-lactamase producing *Haemophilus influenzae* and *Moraxella catarrhalis* should be used in children with severe acute bacterial rhinosinusitis.

Grade D, Level 4

D Amoxicillin (45 mg/kg/day, doubled if age under 2 years or with risk factors for resistance) is recommended for a child with non-severe acute bacterial rhinosinusitis with protracted symptoms. If the symptoms do not improve within 72 hours, an antibiotic against the resistant organism prevalent in the community should be considered. Azithromycin or clarithromycin as first-line therapy is recommended in penicillin allergy.³²

Grade D, Level 4

3 Management of infective rhinosinusitis in adults

3.1 Acute rhinosinusitis

3.1.1 Diagnosis

The diagnosis of acute rhinosinusitis is based on symptoms. Nasal examination may yield supporting signs. Radiographic imaging is unnecessary in patients who meet the diagnostic criteria, unless there is suspicion of complications or an alternative diagnosis.

Symptoms and duration

A sudden onset of two or more of the symptoms:³³

- blockage, congestion, or stuffiness;
- nasal discharge or post nasal drip, often mucopurulent;
- facial pain or pressure, headache, and
- reduction/loss of smell

Acute/ intermittent rhinosinusitis

The duration of two or more of above symptoms which last less than **12 weeks with** symptom-free intervals.³³

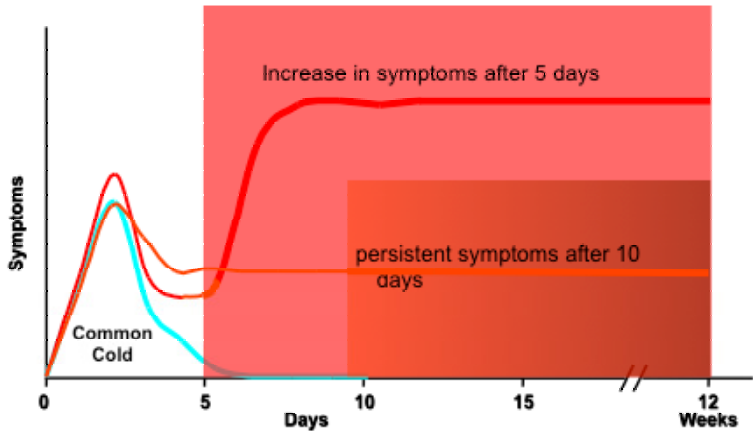
Common cold/acute viral rhinosinusitis

The duration of two or more of above symptoms which last **less than 10 days** (without worsening or persistence of symptoms).³³ (Figure 1, pg 22)

Acute non-viral rhinosinusitis

Increase in symptoms after 5 days or persistent symptoms after 10 days with less than 12 weeks duration.³³ (Figure 1, pg 22)

Figure 1 Increase in symptoms after 5 days or persistent symptoms after 10 days with less than 12 weeks duration



Adapted from Fokkens et al. EP⁺OS Guidelines. *Rhinol Suppl.* 2005;18:1.

Signs³³

- Nasal: swelling, redness, deformity
- Oropharyngeal: postnasal drip
- Oral: dental infection
- Otologic: otitis media

Anterior rhinoscopy remains the basic tool in primary care to determine the existence of pathology, but alone is limited to examining the anterior portion of the sinonasal passages.

Nasal endoscopy helps identify oedema, inflammation, mucopurulent discharge, scarring, crusting, and nasal polyps.

GPP Other diagnosis should be considered in adults with acute rhinosinusitis who present with unilateral symptoms of bleeding, crusting, or cacosmia.

GPP

D Immediate referral to an ENT specialist is indicated for acute rhinosinusitis in adults who present with sinister signs indicative of complications of acute intermittent rhinosinusitis. These include³³:

- Peri-orbital oedema
- Displaced globe
- Double vision
- Ophthalmoplegia
- Reduced visual acuity
- Severe unilateral or bilateral frontal headache
- Frontal swelling
- Signs of meningitis or focal neurological deficits

Grade D, Level 4

Imaging

D Plain sinus x-rays are not recommended for the diagnosis of acute rhinosinusitis in adults.³³⁻³⁵

Grade D, Level 4

Plain sinus x-rays have 76% sensitivity and 79% specificity in diagnosing acute non-viral rhinosinusitis, giving rise to significant false positive and negative results.²³

3.1.2 Principle of treatment

The treatment guidelines are evidence-based and dictated by the severity of rhinosinusitis. The goals of medical treatment are:

- Alleviate symptoms
- Treat underlying inflammatory process and return sinuses back to health; and
- Prevent the development of acute complications

Mild acute rhinosinusitis³³

D Alleviate symptoms of mild acute rhinosinusitis in adults with the following options (Figure 2, pg 25)

- Decongestants
- Nasal saline spray and/ or irrigation
- Antihistamines, only in patients with concomitant allergic rhinitis
- Analgesics

Grade D, Level 4

Moderate and severe acute rhinosinusitis³³

D Treat underlying inflammatory process of moderate to severe acute rhinosinusitis in adults with: (Figure 2, pg 25)

- Intranasal steroid
- Antibiotic, empiric: 7-14 days

Alleviate symptoms with the following options:

- Decongestants
- Nasal saline spray and/ or irrigation
- Antihistamines, in patients with concomitant allergic rhinitis
- Analgesics

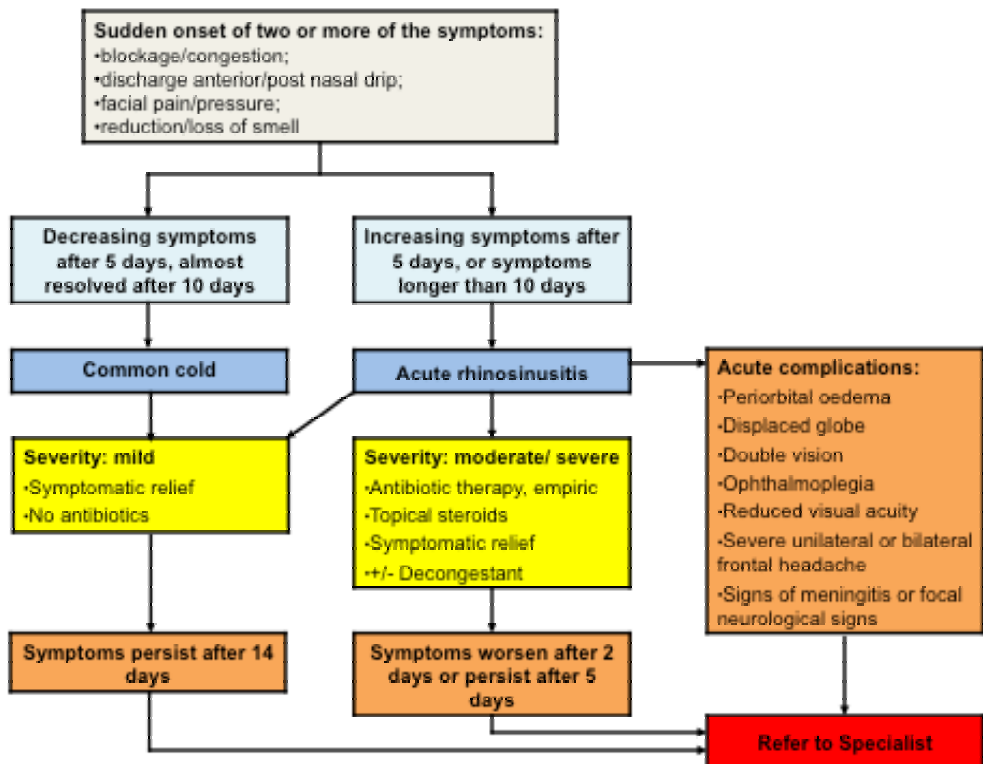
Grade D, Level 4

Specific recommendations on these treatments follow.

GPP The group recommends that patients with acute rhinosinusitis should be reviewed for **symptom resolution**. Patients whose symptoms worsen or persist despite therapy should be referred to a specialist for further evaluation and management.

GPP

Figure 2 Management scheme for primary care for adults with acute rhinosinusitis



Treatment for adults with acute rhinosinusitis

Antibiotics

The use of antibiotics in acute bacterial rhinosinusitis has been validated by meta-analysis. Penicillin group antibiotics have been shown to be superior to placebo. Newer non-penicillin antibiotics (macrolide or cephalosporin) have similar rates of cure or improvement when compared to penicillin, amoxicillin or amoxicillin-clavulanate.¹⁹

The choice of empiric antibiotic is determined by the prevalence and resistance patterns of bacteria based on local and published data. A local series of intracranial and orbital complications of acute sinusitis, found 67% and 75% of patients to have bacteria sensitive to amoxicillin and amoxicillin-clavulanate, respectively, whereas 96% were sensitive to a combination of amoxicillin-clavulanate and cloxacillin.³⁷

Refer to chapter 2 for recommendations on antibiotics for adults with acute bacterial rhinosinusitis.

Nasal Corticosteroids

A Nasal steroid spray twice daily is recommended for adults with acute rhinosinusitis which has not resolved after 5 days of initial presentation.³⁸⁻⁴⁴

Grade A, Level 1+

The use of nasal steroid spray has been shown to help with symptoms of acute rhinosinusitis. As monotherapy, twice daily dosing was superior to amoxicillin and placebo in acute bacterial rhinosinusitis. Once daily dosing showed superiority to placebo but not to amoxicillin.³⁸

The use of nasal steroid spray as an adjunct to antibiotics in the treatment of acute bacterial rhinosinusitis results in significant symptom improvement.³⁹⁻⁴⁴

Oral Corticosteroids

A Oral corticosteroids are not recommended for adults with acute rhinosinusitis.^{45,46}

Grade A, Level 1+

There is limited data on the use of oral steroids in acute rhinosinusitis treatment. Two studies have shown that although there was initial symptom relief during the treatment, there was no significant difference between the treatment groups at the end of a 10 and 14 day course of antibiotics with or without short course of oral steroids.^{45,46}

Anti-histamines

D Antihistamines are not recommended in the treatment of acute bacterial rhinosinusitis in adults.^{26,47,48}

Grade D, Level 4

In adults, antihistamines might worsen the congestion by drying the mucosa.

A Antihistamines may be used as an adjunct to antibiotic treatment in acute bacterial rhinosinusitis patients with concomitant allergic rhinitis.⁴⁹

Grade A, Level 1+

Nasal obstruction and sneezing were reduced as compared to controls in a clinical trial.⁴⁹

D New generation oral antihistamines are preferred in adults with acute rhinosinusitis for their favourable efficacy/safety ratio and pharmacokinetics. Refrain from first generation antihistamines to avoid sedation and anti-cholinergic side effects.⁴⁸

Grade D, Level 4

Topical Decongestants

GPP Topical decongestants may be used for adults with acute rhinosinusitis whose symptoms fail to resolve after 10 days of initial presentation.

GPP

After 10 days, unresolved acute bacterial rhinosinusitis is presumed to be significant.

Topical decongestants have significant effect on congestion of the inferior turbinate and ostiomeatal complex patency (middle turbinates and infundibular mucosa).

Clinical experience shows that drainage of persistent mucopus in established acute bacterial rhinosinusitis from the middle meatus is facilitated by topical decongestion. Intranasal decongestants act more rapidly and more effectively than oral decongestants.

GPP The duration of treatment with topical decongestants should be limited to less than 10 days to avoid rhinitis medicamentosa.

GPP

Nasal Saline Spray and/or Irrigation

A Nasal hypertonic saline irrigation, alone or in conjunction with other adjunctive measures, may be used to reduce symptoms and medication use in adults with frequent acute rhinosinusitis.⁵⁰

Grade A, Level 1+

Hypertonic saline irrigation showed improvement in quality of life, decreased symptoms and decreased medication use in patients with frequent acute rhinosinusitis.⁵⁰

Compared to isotonic saline, buffered hypertonic saline irrigation may have a superior anti-inflammatory effect and ability to thin mucous, and transiently improve nasal mucociliary clearance in healthy volunteers.¹³ Moreover, hypertonic saline irrigation has been shown to improve mucociliary transport in chronic rhinosinusitis, while isotonic saline seemed to have a better effect on acute rhinosinusitis.⁵²

However, hypertonic saline spray did not improve subjective symptom scores (congestion, secretion, and headache) and the duration to symptom resolution in patients with the common cold and acute rhinosinusitis, compared with isotonic saline, and no treatment. In this study, rhinosinusitis patients also received antibiotics.³⁷

Mucolytics

D Mucolytics are not recommended to be prescribed routinely for adult patients with acute rhinosinusitis.⁴⁸

Grade D, Level 4

Table 1 Treatment for adults with acute rhinosinusitis

Therapy	Recommendation
Oral antibiotics	Yes, after 5 days or in mod/severe acute rhinosinusitis
Nasal corticosteroid	Yes
Topical steroid and oral antibiotic combined	Yes
Oral corticosteroid	No
Oral anti-histamines	Yes, only in allergic patients
Saline irrigation	Yes, as symptomatic relief
Decongestant	Yes, as symptomatic relief
Mucolytic	No

3.2 Chronic rhinosinusitis

3.2.1 Definition

Chronic rhinosinusitis is a group of disorders characterized by inflammation of the mucosa of the nose and paranasal sinuses of at least 12 consecutive weeks.³³

3.2.2 Diagnosis

Symptoms present longer than 12 weeks.

Two or more symptoms, one of which should be either nasal blockage/obstruction/congestion or nasal discharge (anterior/posterior nasal drip):

± Facial pain/pressure

± Reduction or loss of smell³³

Common factors associated with chronic rhinosinusitis:

- Micro-organisms – viral, bacterial, fungal
- Allergy
- Nasal anatomical variants causing obstruction
- Mucociliary dysfunction – congenital and acquired
- Immunodeficiency
- Noxious chemicals and pollutants, including cigarette smoke
- Dental disease
- Asthma
- Aspirin sensitivity

This list is by no means exhaustive.

GPP All adults with persistent and recurrent rhinosinusitis should be referred to a specialist for nasal endoscopy to assess for differential causes.

GPP

GPP Other diagnosis should be considered in adults with chronic rhinosinusitis who present with unilateral symptoms of bleeding, crusting, or cacosmia.

GPP

D Immediate referral to an ENT specialist is indicated for chronic rhinosinusitis in adults who present with sinister signs³³ such as:

- Peri-orbital oedema
- Displaced globe
- Double vision
- Ophthalmoplegia
- Reduced visual acuity
- Severe unilateral or bilateral frontal headache
- Frontal swelling
- Signs of meningitis or focal neurological deficits

Grade D, Level 4

Imaging

D Sinus x-rays are **not** recommended to support the diagnosis of chronic rhinitis in adults.³³

Grade D, Level 4

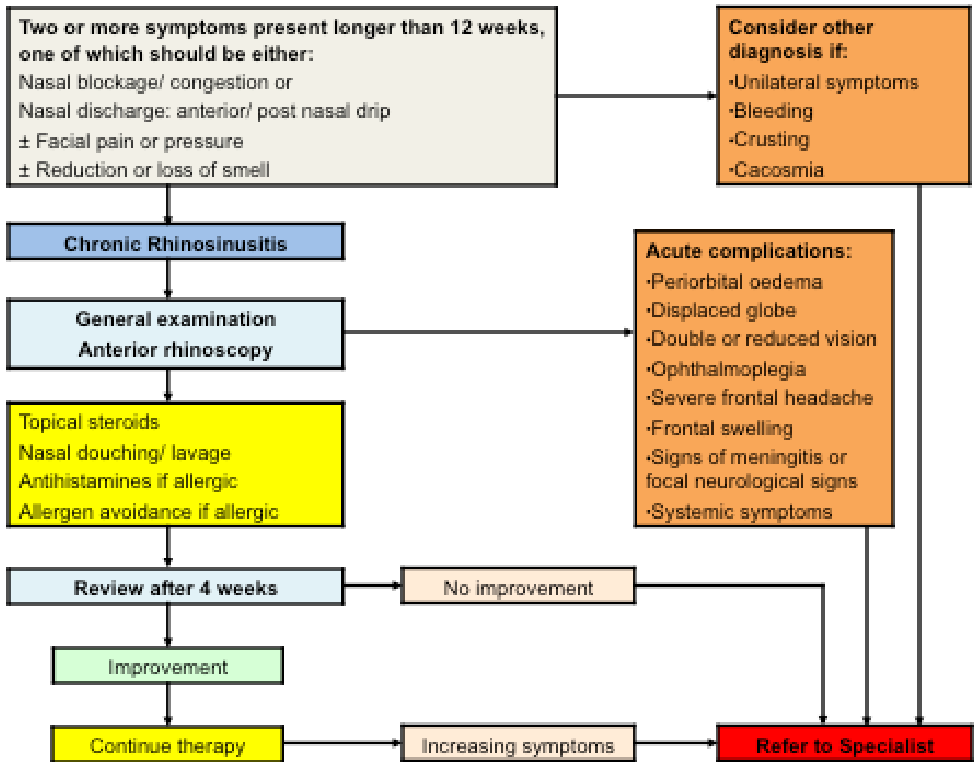
Specialist management of chronic rhinosinusitis includes serial endoscopic surveillance, microbial isolation, allergy testing, immunodeficiency screening, long-term pharmacotherapy, CT scan evaluation of the paranasal sinuses, and endoscopic sinus surgery.

3.2.3 Principle of treatment

The goals of medical treatment are:

- Alleviate symptoms
- Treat underlying inflammatory process and return sinuses back to health; and
- Prevent the development of acute complications

Figure 3 Management scheme for primary care for adults with chronic rhinosinusitis



Treatment of chronic rhinosinusitis without nasal polyps

D For chronic rhinosinusitis without nasal polyps, alleviate symptoms with the following options:

- Nasal saline irrigation

Treat underlying inflammatory process with:

- Intranasal steroid
- Antibiotic, in patients with acute exacerbation of chronic rhinosinusitis, culture directed: 10-14 days³³

Grade D, Level 4

Short-term oral antibiotic therapy

C Short-term oral antibiotics are recommended for acute exacerbation of chronic rhinosinusitis without nasal polyps.⁵³⁻⁵⁵

Grade C, Level 2+

In chronic rhinosinusitis, the role of bacteria is still not well established. Bacteria present may be infective, as super antigens or just as colonizers of the sinuses. The use of short-term antibiotics in chronic rhinosinusitis without nasal polyps has not been validated by meta-analysis, or placebo-controlled studies available. However, several open trials have shown that oral antibiotics improve the symptomatology of acute exacerbation of chronic rhinosinusitis.⁵³⁻⁵⁵ Studies have shown clinical responses to short-term antibiotics of up to 14 days duration, in 56% to 95% of patients.^{53,54}

Long-term oral macrolide therapy

Use of long-term, low-dose macrolide therapy has been shown to be beneficial with improvement of symptoms of 60% to 80% only in the Japanese population.^{56,57} The exact mechanism of action is still unknown.

In selected cases, when topical steroids and a short course of antibiotics have failed, long term low dose macrolides have been reported to have been effective in a subpopulation of patients with chronic rhinosinusitis*.

* Wallwork et al. A double-blind, randomized, placebo-controlled trial of macrolide in the treatment of chronic rhinosinusitis. *Laryngoscope*. 2006 Feb;116(2):189-93.

* Ragab SM. Evaluation of the medical and surgical treatment of chronic rhinosinusitis: a prospective, randomised, controlled trial. *Laryngoscope*. 2004 May;114(5):923-30.

Nasal corticosteroids

A Nasal corticosteroids may be prescribed for chronic rhinosinusitis without nasal polyps.^{56,57}

Grade A, Level 1+

The use of nasal steroids has shown significant improvement in nasal symptom score and inspiratory peak flow.^{56,57}

Nasal saline irrigation

A Nasal saline irrigation may be prescribed for chronic rhinosinusitis without nasal polyps.

Grade A, Level 1+

The use of isotonic saline irrigation for 7 days in the treatment of chronic rhinosinusitis showed improvement in subjective complaints, endoscopic findings, and radiological results.⁵⁸ The use of hypertonic saline irrigation, over standard treatment, showed improvement in endoscopic appearances, quality of life⁵⁹, sinus-related symptoms, and decreased medication use⁶⁰ in patients with chronic rhinosinusitis.

GPP Oral steroids, oral/topical decongestants, mucolytics or antihistamines are not recommended in treatment of chronic rhinosinusitis without nasal polyps.

GPP

Oral corticosteroids

There is no evidence-based data to support the use of oral steroids in the treatment of chronic rhinosinusitis without nasal polyps.

Anti-histamines

There are no controlled trials to show evidence of beneficial effects of anti-histamines therapy for chronic rhinosinusitis without nasal polyps.

Mucolytics

Though mucolytics are commonly prescribed as an adjunct to antibiotic treatment, there is insufficient data to draw a conclusion on the benefits of mucolytics in acute rhinosinusitis. There are no controlled trials to suggest that mucolytics are beneficial in the treatment of chronic rhinosinusitis without nasal polyps.

Decongestants

The use of decongestants in treatment of chronic rhinosinusitis without nasal polyps has not been evaluated in a randomized controlled trial.

Table 2 Treatment for adults with chronic rhinosinusitis without nasal polyps

Therapy	Recommendation
Short-term oral antibiotic therapy	Yes, in acute exacerbation of chronic rhinosinusitis
Long-term macrolide therapy	Yes
Nasal corticosteroid therapy	Yes
Oral steroid therapy	No
Nasal saline irrigation	Yes
Oral/topical decongestants	No
Mucolytics	No
Anti-histamines	No

Treatment of chronic rhinosinusitis with nasal polyps

D For chronic rhinosinusitis with nasal polyps, alleviate symptoms with the following options:

- Nasal saline irrigation
- Anti-histamines, in patients with concomitant allergic rhinitis

Treat underlying inflammatory process with:

- Intranasal steroid.³³

Grade D, Level 4

GPP Adults with chronic rhinosinusitis with nasal polyps should be reviewed for symptom control. Patients whose symptoms worsen during or persist despite therapy should be referred to a specialist for further evaluation and management.

GPP

Short-term oral antibiotic therapy

C Short-term oral antibiotics are recommended to improve symptoms in acute exacerbation of chronic rhinosinusitis with nasal polyps.

Grade C, Level 2+

There are no placebo-controlled trials with evidence of beneficial effects of the use of antibiotics in treatment of chronic rhinosinusitis with nasal polyps. However, several open trials have reported that short-term oral antibiotics improve the symptomatology of acute exacerbation of chronic rhinosinusitis.⁵³⁻⁵⁵

Long-term oral macrolide therapy

C Long-term, low-dose macrolide therapy may be considered for chronic rhinosinusitis patients with nasal polyps.^{56,57,61}

Grade C, Level 2+

Benefits have been reported with long-term, low-dose macrolide therapy for chronic rhinosinusitis with nasal polyps, in patients when corticosteroids fail.^{61,62}

GPP Management by a specialist is recommended for patients with chronic rhinosinusitis with nasal polyps being prescribed long term, low dose macrolide therapy, in view of its side effects.

GPP

Nasal Corticosteroids

A Nasal corticosteroid therapy may be used in adults with chronic rhinosinusitis with nasal polyps.⁶³⁻⁶⁸

Grade A, Level 1+

Placebo controlled studies on nasal corticosteroids in the treatment of chronic rhinosinusitis with nasal polyps have demonstrated over periods of 4-26 weeks, reduced nasal symptom scores, improved peak nasal inspiratory flow, and decreased polyp size.⁶³⁻⁶⁸ However, the effect on improvement in the sense of smell is inconsistent.

Oral corticosteroids

There are no studies performed on single treatment with oral corticosteroids for chronic rhinosinusitis with nasal polyps. Furthermore, placebo-controlled, and dose-effect studies are also

lacking. However, there is clinical acceptance from several open studies that oral corticosteroids are effective in shrinkage of polyps, reducing nasal symptoms, and improving the sense of smell.^{63,69-71}

Although widely prescribed pre-operatively, there is an absence of experimental data to support the use of oral corticosteroids. Oral corticosteroid therapy, especially if long-term, has potential side effects and should preferably be administered and monitored under specialist care.

Antihistamines

C Antihistamines are not recommended in chronic rhinosinusitis with nasal polyps.

Grade C, Level 2+

Cetirizine does not have an effect on polyp size.⁷²

Table 3 Treatment for adults with chronic rhinosinusitis with nasal polyps

Therapy	Recommendation
Short-term oral antibiotic therapy	Yes, in acute exacerbation of chronic rhinosinusitis
Long-term macrolide therapy	Yes
Topical steroid therapy	Yes
Oral steroid therapy	Yes
Nasal saline irrigation	Yes, as symptomatic relief
Oral/topical decongestants	No
Mucolytics	No
Systemic anti-histamines	No, except for allergy

4 Management of infective rhinosinusitis in children

4.1 Introduction

Rhinosinusitis is common but challenging to manage in children. Issues of special concern are how to obtain an accurate diagnosis, the extent of investigations required, and the relative lack of high level evidence on effectiveness and safety of long-term medical and surgical therapy.

This section on Paediatric Rhinosinusitis follows closely the recommendations of the European Position Paper on Rhinosinusitis and Nasal Polyposis 2007.³³

4.2 Epidemiology

In an MRI study of a non-ENT paediatric population, the prevalence of children with signs of sinusitis was 45%. This increases to 80% with bilateral mucosal swelling on rhinoscopy, and 100% with purulent secretions.⁷³

Children at higher risk include:

1. Those 2-8 years of age, when an immature immunity system and adenoid hypertrophy are more common.
2. Those with allergic rhinitis.
3. Those attending day care centres.

4.3 Bacteriology

The common pathogens in acute rhinosinusitis are *Streptococcus pneumoniae*, *Haemophilus influenzae*, and *Moraxella catarrhalis*. These pathogens isolated from the maxillary sinuses correlated well with middle meatal (83%), but not nasopharyngeal cultures.⁷⁴

4.4 Diagnosis

A detailed medical history and clinical examination is important.

4.5 History

4.5.1 Acute rhinosinusitis

Definition

Sudden onset of two or more symptoms, one of which should be either nasal blockage or nasal discharge (anterior/posterior nasal drip), with or without facial pain/pressure, and with or without reduction of smell. Fever and pain are more often associated with acute rhinosinusitis than chronic rhinosinusitis. There is complete resolution of symptoms within 12 weeks.

Differences between acute rhinosinusitis and common cold

Acute rhinosinusitis is suspected when³²:

1. Signs and symptoms of a cold persist beyond 10 days (nasal discharge and day-time cough worsening at night).
2. A cold seems more severe than usual (high fever, copious purulent discharge, peri-orbital oedema and pain).
3. A cold worsens after several days of improvement (with or without fever).

4.5.2 Chronic rhinosinusitis

Definition

Symptoms are the same as acute rhinosinusitis, but there is no complete resolution of symptoms by 12 weeks. Acute-on-chronic exacerbations are possible.

GPP Allergic rhinitis often coexists with paediatric acute and chronic rhinosinusitis. The history should evaluate for symptoms of allergic rhinitis and identify possible allergens.

GPP

4.6 Clinical examination

Paediatric acute rhinosinusitis or chronic rhinosinusitis is frequently associated with otitis media with effusion. Recurrent otitis episodes can impact hearing, speech, language development, and may require placement of grommet tubes.

GPP Otoscopy should be performed routinely to exclude otitis media in paediatric acute and chronic rhinosinusitis.

GPP

Complications of acute rhinosinusitis can develop rapidly in a child. Eye swelling from orbital infection can occur without pain in the eye or any history of rhinosinusitis.⁷⁵ Complications like osteomyelitis, intracranial infection and cavernous sinus thrombosis are also possible. Any change in sensorium, extreme lethargy or spiking fevers should prompt a suspicion of intracranial infection.

4.7 Investigations

1. Culture and sensitivity analysis

Swabs of nasal discharge from middle meatus or maxillary antrum are obtained in⁷⁶:

- severe illness or toxicity
- acute illness failing 48-72 hours of medical therapy
- immunocompromised patients
- intra-orbital or intracranial abscess complications

Careful decongestion of the middle meatal area usually allows mucopus from the sinus to be cultured without the need for antral puncture. In children younger than 7 years old, the floor of the maxillary sinus remains above the level of the floor of the nose, increasing the risk of damaging the tooth roots during an antral puncture.

2. Imaging

D Plain X-ray is not recommended routinely to confirm the diagnosis of rhinosinusitis in children.^{33,77}

Grade D, Level 4

Van der Veken et al reported that plain x-ray is not sensitive, with unnecessary radiation exposure.⁷⁸ Transillumination is also not useful under 10 years of age.⁷⁷

3. The following conditions are associated with recalcitrant cases.

a. Allergic rhinitis

The association between allergy and chronic rhinosinusitis remains controversial. However, children with allergy have more upper respiratory tract infections.⁷⁹

Children with a positive history of allergy are further evaluated for inhalant and food allergies.⁸⁰

b. Immunocompromised state

Recurrent or chronic rhinosinusitis is the most common clinical presentation of common variable immunodeficiencies.⁸¹ Primary deficiencies include secretory IgA and IgG subclass antibody deficiencies. Secondary immunocompromise related to organ transplants, AIDS, malignancies and drugs is possible. In immunocompromise states, unusual or resistant micro-organisms and fungi are more common.

c. Laryngopharyngeal reflux

Laryngopharyngeal reflux can cause chronic cough, hoarseness and even stridor. In 30 children with chronic sinus disease, 63% had esophageal reflux and 32% had nasopharyngeal reflux.⁸²

The 24-hour pH probe is the gold standard diagnostic tool for acid reflux, but will miss a diagnosis of alkaline reflux.

d. Cystic Fibrosis

Cystic fibrosis is uncommon in Singapore. It is more common in non-Hispanic Caucasians and Ashkenazi Jews. Sinusitis and nasal polyps were found in more than 50% of patients with cystic fibrosis.⁸³ Nasal polyposis in chronic rhinosinusitis should prompt a suspicion of cystic fibrosis.

e. **Primary Ciliary Dyskinesia**

It is also rare in Asians. However, rhinitis at birth in an otherwise well child, atypical recalcitrant asthma, chronic wet cough, very severe gastro-oesophageal reflux, bronchiectasis, rhinosinusitis, chronic severe secretory otitis media (especially post grommet tubes), and Kartegener's syndrome (situs inversus, bronchiectasis and chronic rhinosinusitis) should prompt a suspicion of primary ciliary dyskinesia.⁸⁴

4.8 Pharmacotherapy

Principles of medical therapy are:

1. To control symptoms
2. To reduce underlying nose and sinus inflammation
3. To eradicate pathogens

The workgroup adopted recommendations from the European Position Paper on Rhinosinusitis and Nasal Polyposis (EPOS). However, decisions for individual patients should be specifically tailored.

4.9 Management of paediatric acute rhinosinusitis

Usually, only symptomatic treatment is needed.

1. Antibiotics

Refer to chapter 2 for recommendations on antibiotics in children with acute bacterial rhinosinusitis.

Cochrane meta-analysis⁸⁵ of antibiotics for persistent nasal discharge concluded that antibiotics for 10 days reduced the persistence of acute rhinosinusitis in the short to medium term. For 8 children treated, only 1 additional child would be cured (NNT 8, 95% CI 5 to 29). No long term benefits were documented.⁸⁶

2. Topical corticosteroids

A Topical corticosteroids may be used in children as an adjunct to antibiotics. It can reduce the cough and nasal discharge earlier in acute bacterial rhinosinusitis.⁸⁷

Grade A, Level 1+

3. Topical or oral decongestants

GPP Topical decongestants should be used in children no longer than 4-5 days to avoid toxicity and rhinitis medicamentosa.

GPP

No benefit has been shown, though Alpha-2 agonists xylometazoline and oxymetazoline are commonly used in acute rhinitis.

A double-blind, randomized controlled trial showed no additive effect of adding oral decongestant-antihistamine to amoxicillin.⁸⁸

4. Nasal douching

D Saline nose drops or sprays may be considered to decrease the mucus trapping and crusting associated with acute rhinosinusitis in children.⁷⁹

Grade D, Level 3

Figure 4 (pg 46) showed the summary of treatment evidence and recommendations for children with acute rhinosinusitis³³

4.10 Management of paediatric chronic rhinosinusitis

Most cases of paediatric chronic rhinosinusitis resolve spontaneously.⁸⁹

There is a significant paucity of data on specific treatment of chronic rhinosinusitis in children.

1. Antibiotics

The benefit of antibiotics has been shown only in acute bacterial rhinosinusitis as described earlier. The only study addressing antibiotics in paediatric chronic rhinosinusitis⁹⁰ did not show any benefit.

D The workgroup recommends antibiotics use only in acute exacerbation of paediatric chronic rhinosinusitis, by following the recommendations from the Consensus Meeting in Brussels, 1996³¹:

- For chronic rhinosinusitis, especially with frequent exacerbations, 2 weeks of oral antibiotics is advised. The antibiotic is changed if there is no response within 5-7 days.
- Failing this, sinus secretions for culture or investigations to exclude recalcitrant causes are considered.
- If there is slow response, a second 2-week course can be prescribed.
- In rare cases with clear-cut improvement but persisting symptoms, a 3rd course can be given before surgery is considered.
- Parenteral antibiotic may be appropriate if oral antibiotics fail.

Grade D, Level 4

2. Topical corticosteroids

Many studies show that topical corticosteroids are effective and safe in children with rhinitis, and it is assumed that this similarly applies for chronic rhinosinusitis. There is no data on the efficacy of topical corticosteroids in paediatric chronic rhinosinusitis specifically.

3. Nasal douching

C Nasal douching may be considered for paediatric chronic rhinosinusitis.^{91,92}

Grade C, Level 2+

Nasal douching was effective in 2 small randomized studies. Hypertonic or normal saline were equally effective.^{91,92}

4. Reflux therapy

Gastro-oesophageal reflux (GER) is treated in chronic rhinosinusitis before any surgical intervention. In children with chronic rhinosinusitis and gastro-oesophageal reflux proven by 24-hour pH monitoring, reflux therapy improved sinus disease.⁸² Reflux therapy avoided surgery in 89 % of children with chronic rhinosinusitis.⁹³

4.11 Surgical therapy

A. Functional Endoscopic Sinus Surgery (FESS)

A meta-analysis showed positive outcomes of 88-92 % with average follow-up of 3.7 years, concluding that FESS is a safe and effective treatment for chronic rhinosinusitis.⁹⁴ However, outcomes were based on symptomatic relief, not endoscopic examination or CT scan.

B. Adenoidectomy

This is still controversial in paediatric chronic rhinosinusitis. Recently, it has been suggested that adenoidectomy is safe and effective, and should be performed before functional endoscopic sinus surgery (FESS), especially in younger children with obstructive symptoms.⁹⁵

Outcomes are based on symptomatic relief, not endoscopic examination or CT scan. A meta-analysis showed positive outcomes of 88-92 % with average follow-up of 3.7 years, concluding that FESS is a safe and effective treatment for chronic rhinosinusitis.⁹⁴

C. Other interventions

D Antral lavage, inferior meatal antrostomy (except possibly in primary ciliary dyskinesia), Caldwell-Luc operation (risks damage to un-erupted teeth)^{74,79,96} are not recommended in paediatric chronic rhinosinusitis.

Grade D, Level 3

Figure 5 (pg 47) showed the summary of clinical management scheme for children with chronic rhinosinusitis.³³

Figure 4 Summary of treatment evidence and recommendations for children with acute rhinosinusitis³³

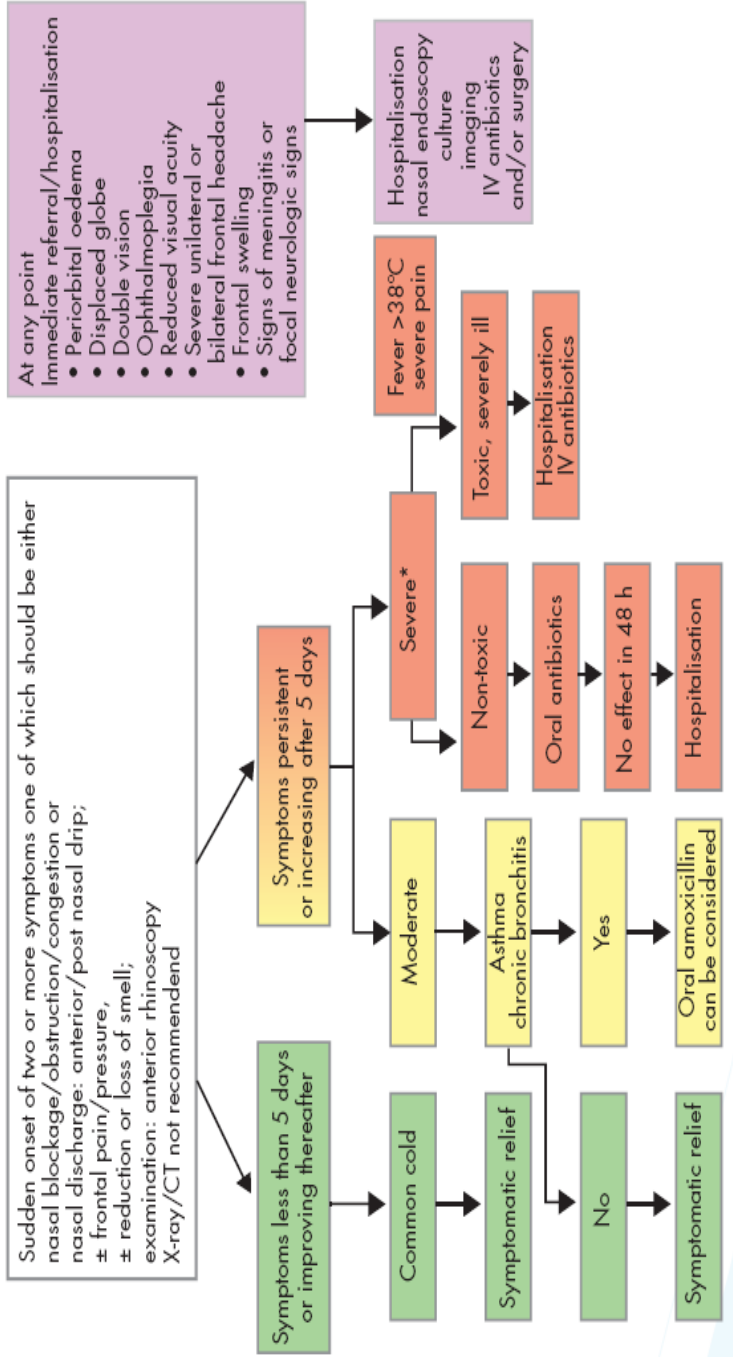
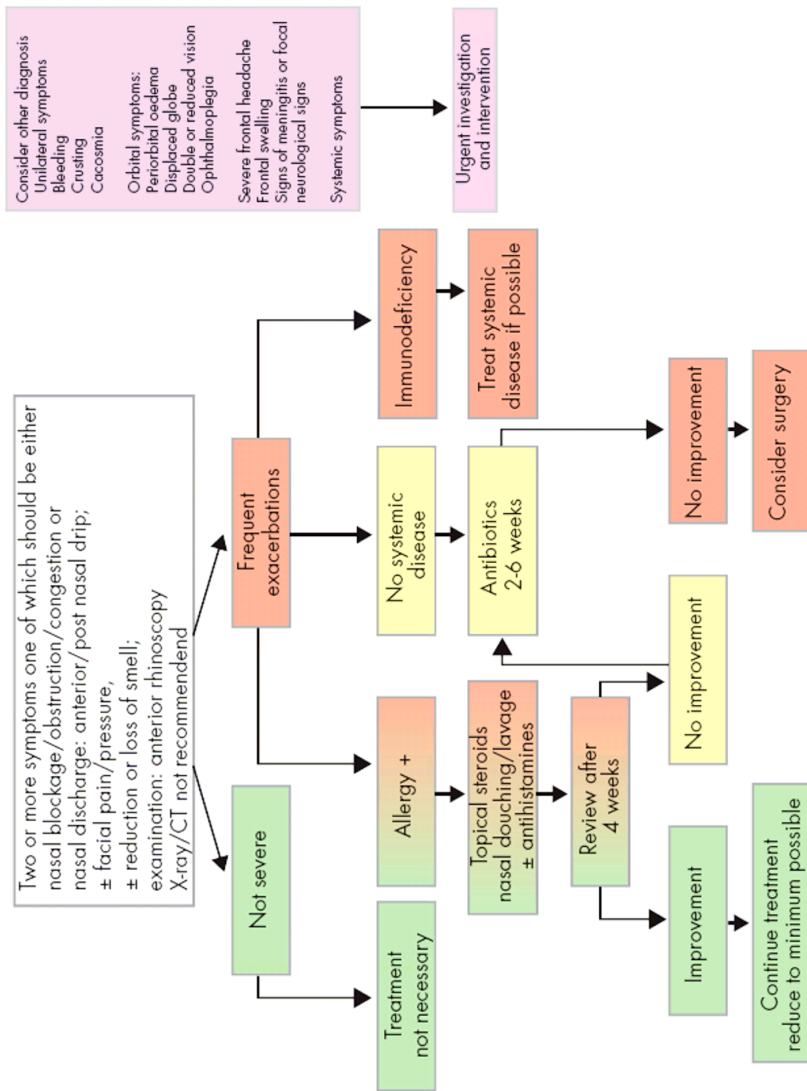


Figure 5 Summary of clinical management scheme for chronic rhinosinusitis in children³³



5 Management of allergic rhinitis

5.1 Diagnosis

Allergic rhinitis is a symptomatic disorder of the nose, induced after allergen exposure by an IgE mediated inflammation of the membranes lining the nose, resulting in the cardinal symptoms of sneezing, nasal obstruction, anterior or posterior rhinorrhoea and/or itching.

These symptoms occur during two or more consecutive days for more than one hour in most days.

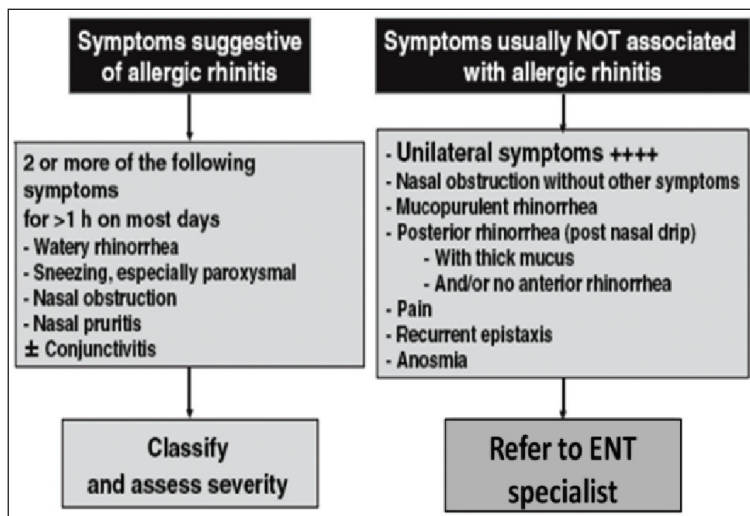
GPP The diagnosis of allergic rhinitis should be made based upon concordance between a typical history of allergic symptoms and diagnostic tests.

GPP

D The workgroup recommends using the algorithm for the diagnosis and assessment of severity of allergic rhinitis proposed by ARIA 2008 (refer to Figure 6).

Grade D, Level 4

Figure 6 Symptoms of allergic rhinitis



(Adapted with permission from ARIA 2008 Update).

GPP Besides a nasal examination for allergic rhinitis, look out for:

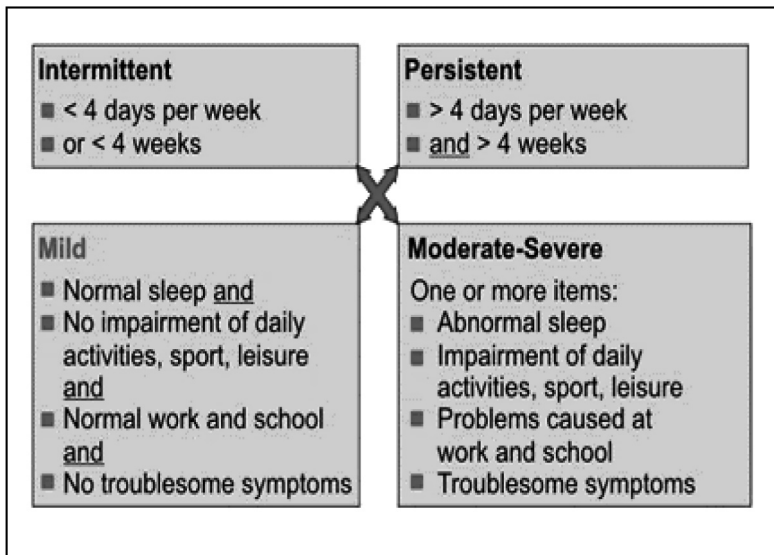
- Ocular signs of irritation e.g. allergic conjunctivitis; redness and rubbing of eyes indicative of itchiness.
- Chest examination to rule out concurrent asthma.

GPP

D The workgroup recommends using the algorithm for the classification of allergic rhinitis proposed by ARIA 2008 (refer to Figure 7 below).

Grade D, Level 4

Figure 7 Classification and severity of allergic rhinitis



(Reproduced with permission from ARIA 2008 Update)

5.2 Management

This encompasses the following measures:

- Pharmacotherapy
- Avoidance of allergen
- Allergen specific immunotherapy
- Patient education

In the treatment of allergic rhinitis, one should consider the severity and duration of the disease, the patient's preference, the safety measures, as well as the efficacy, availability and cost of medications.

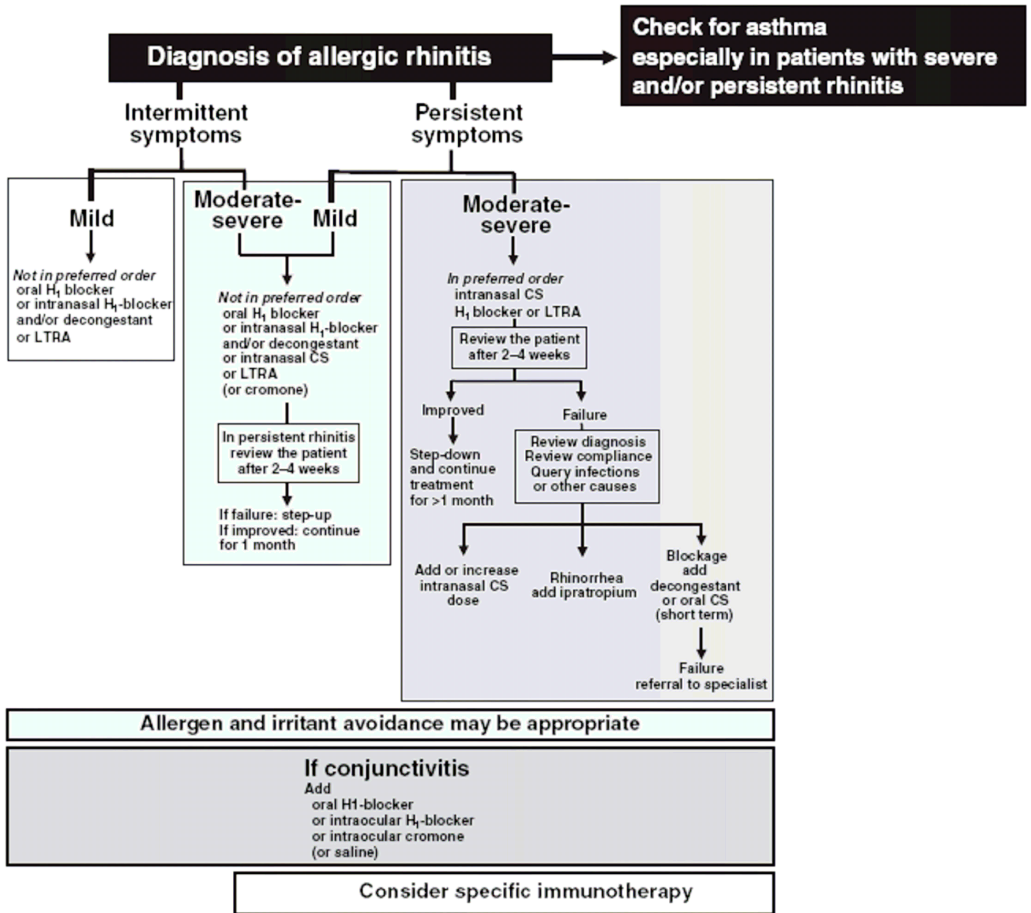
A stepwise approach depending on the severity and duration of rhinitis is proposed.

Not all patients with moderate/severe allergic rhinitis are controlled despite optimal pharmacotherapy.

D The workgroup recommends using the algorithm for the management of allergic rhinitis proposed by ARIA 2008 (refer to Figure 8, pg 51).

Grade D, Level 4

Figure 8 Algorithm for management of allergic rhinitis



(Reproduced with permission from ARIA 2008 Update)

5.2.1 Environmental control (inhalant-allergen avoidance)

GPP Mattress encasings or High Efficiency Particulate Air Filters for house dust mite and pet allergy in adults with rhinitis should be part of the overall management of allergic rhinitis.

GPP

A range of inhalant allergens has been associated with allergic rhinitis, of which house dust mite (HDM) is the most important locally and most investigated. Although the general consensus is that allergen avoidance should lead to an improvement of symptoms, studies have shown that the majority of single preventive measures of indoor allergen control fail to achieve a clinically relevant improvement of asthma and rhinitis.

5.2.2 Pharmacotherapy of allergic rhinitis

A Second-generation oral or intranasal H1-antihistamines are recommended for the treatment of allergic rhinitis and conjunctivitis in adults and children.⁹⁷⁻⁹⁹

Grade A, Level 1++

First-generation oral H1-antihistamines tend to have sedative effects and should be used with care.

A Intranasal glucocorticosteroids are strongly recommended for the treatment of allergic rhinitis in adults and children.¹⁰⁰⁻¹⁰²

Grade A, Level 1++

D Intramuscular glucocorticosteroids and the long term use of oral preparations are not recommended for the treatment of allergic rhinitis due to safety concerns.¹⁰³

Grade D, Level 3

A Topical H1-antihistamines are recommended for the treatment of allergic rhinitis and conjunctivitis. Its therapeutic effects are superior and faster than oral anti-histamines.^{104,105}

Grade A, Level 1+

A Intranasal ipratropium may be considered as a treatment option for rhinorrhoea associated with allergic rhinitis.^{106,107}

Grade A, Level 1+

A Topical cromones should be considered as a treatment option for allergic rhinitis and conjunctivitis. However, they are only moderately effective.¹⁰⁸⁻¹¹⁴

Grade A, Level 1+

A Montelukast may be considered as a treatment option for seasonal allergic rhinitis and asthma in patients over 6 years of age. It should not be used more than 4 weeks since there is limited data of its efficacy in patients with persistent allergic rhinitis for more than 4 weeks.¹¹⁵⁻¹²³

Grade A, Level 1+

C Intranasal decongestants may be used for a short period of time in patients with severe nasal obstruction caused by allergic rhinitis.¹²⁴⁻¹³⁰

Grade C, Level 2+

C Oral decongestants (and their combination with oral H1-antihistamines) may be considered in the treatment of allergic rhinitis in adults, but side effects are common.¹²

Grade C, Level 2++

Complementary/ alternative medicine appear to be satisfactory for allergic rhinitis patients, but it lacks evidence based level of recommendations.¹³¹⁻¹³⁶

5.2.3 Allergen-specific immunotherapy: therapeutic vaccines for allergic diseases

Allergen-specific immunotherapy is the practice of administering gradually increasing quantities of an allergen extract to an allergic subject to ameliorate the symptoms associated with the subsequent exposure to the causative allergen.

It induces clinical and immunologic tolerance, has long-term efficacy, improves the quality of life, and may prevent the progression of allergic disease.

There is sound evidence that subcutaneous immunotherapy using inhalant allergens is clinically effective in the treatment of allergic rhinitis and asthma for pollen and mite allergy in both adults and

children, but it is burdened by the risks of side effects which may be life-threatening.

Sublingual immunotherapy is effective for the treatment of pollen and mite allergy in adults and children.

5.2.4 Education

GPP Education of patient and/or patient's carer on the management of allergic rhinitis should be considered as an option to maximize compliance and optimize treatment outcomes.

GPP

5.2.5 Surgical intervention

Surgery is not a modality for treatment of allergic rhinitis. However, it may be used as an adjunctive intervention in few highly-selected patients such as the relief of nasal obstruction due to persistent turbinate hypertrophy, cartilaginous or bony obstruction of the nasal airways or secondary sinus disease.

5.3 Paediatric aspects of allergic rhinitis

Allergic rhinitis is the most prevalent chronic allergic disease in children. It can significantly affect the child's quality of life, and may exacerbate a number of common co-morbidities, including asthma and sinusitis.

Allergic and non-allergic rhinitis are often difficult to differentiate based on symptoms. 50% of childhood rhinitis is induced by allergy.¹³⁷

Inhalant allergens may play an important role in the early development of asthma. However, in preschool children, in contrast to older children, allergic rhinitis occurs at the same time or later than asthma. Sensitization to indoor allergens occurs early in life. Not all children with an allergic sensitization will have atopic disease or develop symptoms after exposure to an allergen.

It is important to note that epidemiology, diagnosis, and treatment of paediatric allergic rhinitis significantly differ between preschool and older children.

The prevalence of allergic rhinitis in preschool children is 4%.¹³⁸ By the age of 6, doctor-diagnosed allergic rhinitis may occur in more than 40% of children.¹³⁹

5.3.1 Diagnosis

GPP Symptoms of sneezing, nasal itching, discharge and congestion that persist longer than 2 weeks should prompt a search for a cause other than infection in children.

GPP

Children with moderate/severe allergic rhinitis may develop noisy breathing, repeated throat clearing, snoring and sleep apnea, loss of olfaction and taste, have allergic salute or an allergic transverse nasal crease, malaise and disturbed nocturnal sleep with subsequent daytime fatigue. Co-morbidities associated with allergic rhinitis in children include asthma, atopic dermatitis/eczema, allergic conjunctivitis, chronic sinusitis and chronic otitis media with effusion.

GPP It is recommended to ask about family history of atopy and progression of atopy of the child.

GPP

B Skin prick tests should be performed and interpreted reliably early in life.¹⁴⁰

Grade B, Level 2+

Positive tests to food allergens in infancy may predict a later development of sensitization to inhaled allergens.

5.3.2 Treatment

The goal of treatment is to control the symptoms, improve the child's ability to function and prevent the complications of allergic rhinitis.

Allergic rhinitis and asthma are commonly present together in preschool and school children. It is therefore important to carefully

assess the side effects of treatments, especially in children with both rhinitis and asthma.

GPP The principles of treatment are the same in children as in adults with allergic rhinitis, but dosages should be adapted and care should be taken to avoid the side effects involving impairment of growth and cognitive development.

GPP

GPP Pharmacologic management for allergic rhinitis in children should be individualized and polypharmacy avoided.

GPP

5.3.2.1 Pharmacologic treatment

Glucocorticosteroids

A Intranasal glucocorticosteroid with bioavailability of <1% such as fluticasone propionate or mometasone furoate should be considered as a treatment option for allergic rhinitis and allergic conjunctivitis in children.¹⁴¹⁻¹⁴⁴

Grade A, Level 1++

Mometasone furoate is available for children of 2 years and above. Fluticasone propionate is approved for children aged 4 years and older and other intranasal glucocorticosteroids may be used in those over the age of 5 years.^{145,146}

B Intranasal glucocorticosteroids with high bioavailability such as betamethasone should not be used in children with allergic rhinitis due to its effect upon growth and growth velocity.¹⁴⁵⁻¹⁴⁷

Grade B, Level 1++

Present day intranasal glucocorticosteroids do not appear to have an effect on the hypothalamic-pituitary-adrenal-axis in children.¹⁴⁸

A Oral and depot glucocorticosteroid preparations should be avoided in children with allergic rhinitis due to negative effect on short term growth and growth velocity.^{141,149}

Grade A, Level 1+

H₁-antihistamines

First-generation oral H₁-antihistamines, which are often included in the formulations of oral decongestants, have central nervous system side effects, including sedation, fatigue, paradoxical hyperactivity, insomnia, irritability and may further reduce the cognitive function of children with allergic rhinitis.

A Second generation H₁-antihistamines such as cetirizine, levocetirizine and loratadine should be considered as a treatment option in the treatment of allergic rhinitis in children.¹⁵⁰⁻¹⁵²

Grade A, Level 1+

Second-generation H₁-antihistamines are effective and safe in the treatment of allergic rhinitis in children, in particular, cetirizine, levocetirizine and loratadine.¹⁵⁰⁻¹⁵²

Intranasal H₁-antihistamines like levocabastine and azelastine have rapid onset of action and few adverse effects in children with allergic rhino-conjunctivitis.¹⁵²⁻¹⁵⁵

Disodium cromoglycate is safe and effective for allergic rhino-conjunctivitis in children. However, a dosage of four to six times a day is required for cromoglycate, and compliance with treatment is often difficult.^{154,156,157}

GPP Nasal saline drops or spray may be considered in children with allergic rhinitis to clear the nose before eating or sleeping.

GPP

Immunotherapy

Allergen-specific immunotherapy can reduce allergic rhinitis symptoms, alter natural course of the disease and induce long term clinical remission.

A Sublingual immunotherapy (SLIT) should be considered in children above age 5 years who have poor symptomatic control of allergic rhinitis despite maximal therapy or who cannot or will not take medication.¹⁵⁸⁻¹⁶⁰

Grade A, Level 1++

Studies¹⁵⁸⁻¹⁶⁰ have shown that it is effective in young children with allergic rhinitis with only mild and transient local side effects. It may also possibly prevent a later development of asthma.

It can be administered safely at home, has greater acceptance by parents and children, and is well adapted for poly-sensitized patients (mites & pollen).

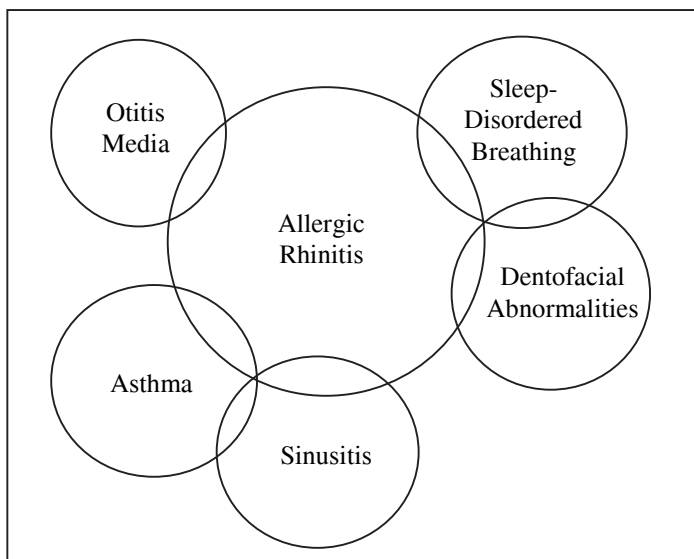
The medication dosing for paediatrics can be found in Appendix 1 (pgs 59-60).

5.3.2.2 Non-pharmacological treatment

GPP The family and the child should be educated about the recurrent or persistent nature of the disease, allergen avoidance and avoidance of allergen triggers and respiratory tract irritants, the most important of which is tobacco smoke.

GPP

Figure 9 Link between allergic rhinitis and other chronic disorders and complications



Appendix 1 Medication dosing for pediatrics

Medication	Age range				
	<12 months	12-23 months	2-5 years	6-11 years	≥12 years
Oral H ₁ anti-histamines	Cetirizine 6-12 mo: 2.5 mg qd	Cetirizine 2.5 mg qd or bid	Cetirizine 2.5 or 5 mg qd or 2.5 mg bid	Cetirizine 5 or 10 mg qd	Cetirizine 5 or 10 mg qd
H ₁ anti-histamines nasal spray		Loratadine 5 mg qd	Loratadine 5 mg qd	Loratadine 10 mg qd	Desloratadine 5 mg qd Fexofenadine 60 mg qd or 180 mg qd
	Corticosteroid nasal spray		Azelastine 5-11 yrs: 1 spray each nostril bid	Azelastine 5-11 yrs: 1 spray each nostril bid	Azelastine 5-11 yrs: 1 spray each nostril bid
			Fluticasone propionate 4-11 yrs: 1 or 2 sprays each nostril qd	Fluticasone propionate 4-11 yrs: 1 or 2 sprays each nostril qd	Budesonide 6-12 yrs: 1 or 2 sprays each nostril qd
			Mometasone furoate 1 spray each nostril qd	Budesonide 6-12 yrs: 1 or 2 sprays each nostril qd	Budesonide >12 yrs: 1-4 sprays each nostril qd
				Flunisolide 6-14 yrs: 1 spray each nostril tid or 2 sprays each nostril bid	Flunisolide >14 yrs: 2 sprays each nostril bid-tid
				Fluticasone propionate 4-11 yrs: 1 or 2 sprays each nostril qd	Fluticasone propionate 1 or 2 sprays each nostril qd or 1 spray each nostril bid

6

Management of rhinitis in pregnancy

Rhinitis may occur in 30% of pregnant women.¹⁶¹

1. Allergic rhinitis: the most common cause of rhinitis in pregnancy. It occurs in 18-30% of women in their childbearing age.¹⁶²

2. Gestational or hormonal rhinitis: defined as nasal congestion for at least 6 weeks of pregnancy without signs of infection, tumor growth or known allergic causes. It occurs in approximately one-fifth of pregnancies during any gestational weeks and resolves within two weeks after the delivery.¹⁶³

3. Infective rhinitis: only occurs in 1.5% of pregnant women and commonly presents as nasal congestion with purulent nasal discharge. Common pathogens identified are *Hemophilus influenzae* and *Streptococcus pneumoniae*.

Diet may affect the prevalence of rhinitis in pregnancy. Fish, N-3 polyunsaturated fat, soya and isoflavones may reduce its prevalence.

Smoking, sensitization to house dust mites, and chronic sinusitis increase prevalence of rhinitis in pregnancy.

6.1 Diagnosis

D Nasal endoscopy on a decongested nose may be considered as an option to differentiate pregnancy rhinitis from sinusitis.¹⁶³

Grade D, Level 4

D Imaging studies are not recommended to make a diagnosis in rhinitis in pregnancy.³³

Grade D, Level 4

D Skin prick tests are not recommended for rhinitis in pregnancy because use of potent antigens in skin testing may be associated with systemic reactions.¹⁶⁴

Grade D, Level 4

Treatment

GPP In treating rhinitis of pregnancy, all drug therapy should ideally be avoided especially in the first trimester. If drug therapy cannot be avoided then treatment will depend upon the predominant symptoms, with the topical agents as first line since they have minimal systemic exposure.

GPP

A. Supportive therapy

Supportive therapy includes simple treatment measures like avoidance of allergens, head elevation, nasal douching and reassurance that the condition is self-limiting. Nasal douching uses saline wash to help remove mucus from the nasal passageways improving discomfort and breathing. Saline washes also help lubricate the nasal mucosa.¹⁶⁵

B. Pharmacotherapy

C Cromones are safe with no known teratogenic effect but they are moderately effective. It may be given for the treatment of rhinitis in the first 3 months of pregnancy, 3-4 times daily.^{166,167}

Grade C, Level 2+

C If cromones are ineffective and poorly tolerated, they should be replaced with anti-histamines. Chlorpheniramine and tripeleennamine are the anti-histamines of choice for pregnant women with rhinitis. Cetirizine and loratadine may be considered after the first trimester.¹⁶⁸⁻¹⁷¹

Grade C, Level 2+

C Intranasal steroids should be prescribed as an alternative to or in combination with anti-histamines for severe cases of rhinitis in pregnancy.¹⁶⁶

Grade C, Level 2+

C Budesonide is the only recommended intranasal steroid for rhinitis in pregnancy.^{172,173}

Grade C, Level 2+

C Topical decongestants like oxymetazoline may be considered as second-line therapy for short-term relief and when no other safer alternatives are available for the treatment of rhinitis in pregnancy.¹⁷⁴

Grade C, Level 2+

Studies on pregnant women using this drug showed no report of association with congenital abnormalities.^{175,176}

C Oral decongestants are not recommended for rhinitis in pregnancy.^{177,178}

Grade C, Level 2+

C Leukotriene Modifiers are not recommended for allergic rhinitis in pregnancy.¹⁷⁹

Grade C, Level 2+

The safety of leukotriene modifiers during pregnancy is not well established.

Antibiotics should only be used in infective rhinitis or sinusitis which is severe (painful, temperature $>38.3^{\circ}$ C) or persists after 10 days of symptoms from onset.¹⁸⁰

A Amoxicillin is the drug of choice for pregnant patients with rhinitis who are not allergic to penicillin.¹⁸¹

Grade A, Level 1+

D Amoxicillin-clavulanate or cephalosporin may be given to pregnant women with rhinitis not responding to amoxicillin.¹⁸²

Grade D, Level 3

C Metronidazole should be used in rhinitis in pregnancy caused by anaerobic pathogens.¹⁸³

Grade C, Level 2+

D Immunotherapy is not recommended for rhinitis in pregnancy. However, it may be continued if the maintenance phase has been reached.^{161,165,184}

Grade D, Level 4

Table 4 shows the list of drugs commonly used for the treatment of sinusitis and their US FDA Pregnancy Risk Category.

Table 4 Drugs commonly used in the treatment of rhinosinusitis in pregnancy¹⁸⁵

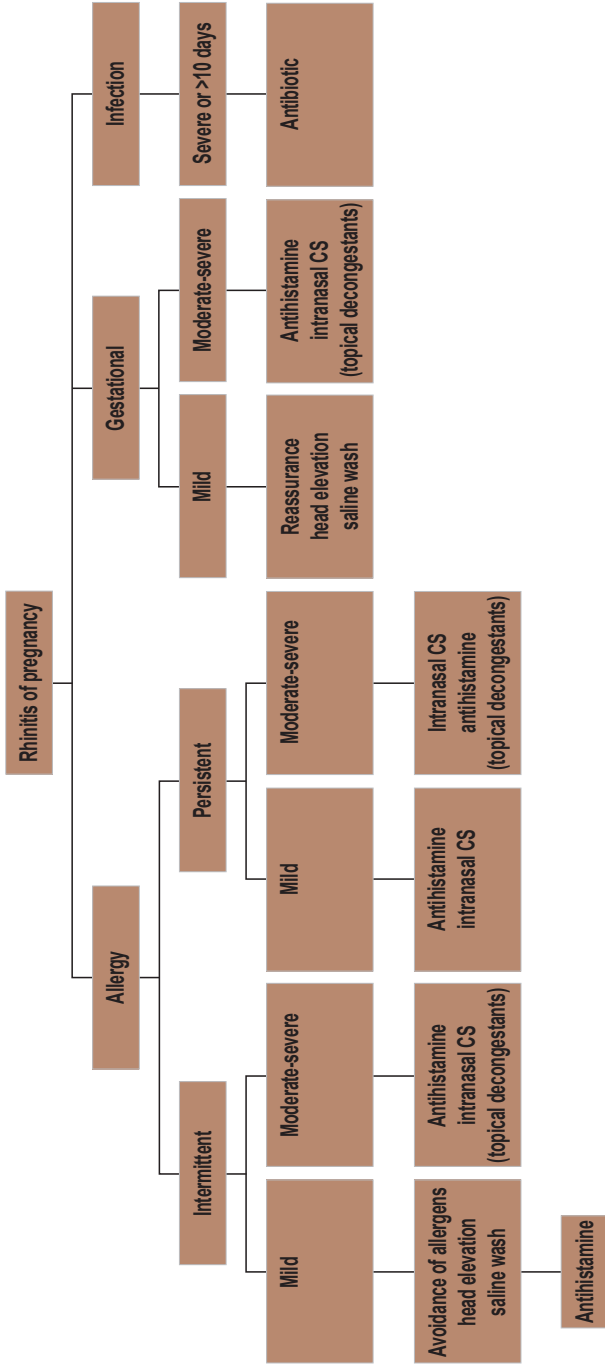
ANTIBIOTICS	FDA Pregnancy Risk Category
Ampicillin-sulbactam	B
Amoxicillin	B
Amoxicillin-clavulanate	B
Cefuroxime	B
Clarithromycin	C
Clindamycin	B
Erythromycin	B
Metronidazole	B

Table 5 shows the FDA Pregnancy Risk Category, an assessment of the risk of fetal injury when using certain drugs. These categories will help the physician in making a correct treatment decision. It is a must-know prior to giving any drugs to pregnant patients.

Table 5 US Food and Drug Administration Pregnancy Risk Category¹⁸⁶

Category A	Adequate and well-controlled human studies have failed to demonstrate a risk to the fetus. Possibility of fetal harm seems remote
Category B	Controlled studies done on animals in reproduction do not indicate risk to the fetus. No adequate and well-controlled studies done on pregnant women.
Category C	Studies in animals show adverse effect and toxicity on fetus. No adequate and well-controlled studies done on pregnant women. But the benefits of these drugs may out weigh the potential risks in humans.
Category D	Positive evidence of human fetal risk exists, but benefits may outweigh the risks in certain situations.
Category X	Studies in animals or human beings have demonstrated fetal abnormalities. The risk of the use of drug in pregnant women clearly outweighs possible benefit. Contraindicated in pregnant women.

Figure 10 Treatment of rhinitis of pregnancy



Abbreviation:
CS - Corticosteroids

Allergic rhinitis imposes a substantial economic burden on society with indirect costs of productivity loss being larger than the direct health costs. It has been estimated that the burden of illness cost for allergic rhinitis ranges from US 2 to 5 billion dollars in USA.¹⁸⁶

The many variables in the study of cost effectiveness of allergic rhinitis management such as identification of allergic rhinitis patients, differences in cost assignment, and difficulties in assigning indirect costs such as reduced productivity preclude formal cost effectiveness evaluations that compare incremental costs and benefits of alternative treatment strategies.¹⁸⁷

Although there are presently no strong cost-effectiveness arguments available comparing each specific treatment option in allergic rhinitis, management in general is important in reducing a substantial economic burden on society.¹⁸⁸

The following clinical quality improvement parameters, based on recommendations in these guidelines, are proposed:

Management of rhinosinusitis in adults

1. Percentage of patients who had plain sinus X-ray to diagnose rhinosinusitis. (Page 26)
2. Percentage of acute rhinosinusitis patients who received nasal steroid spray prior to commencement of antibiotics. (Page 29)

Management of rhinosinusitis in children

3. Percentage of children who had plain sinus X-ray to diagnose rhinosinusitis. (Page 43)

References

1. American Academy of Pediatrics. Subcommittee on Management of Sinusitis and Committee on Quality Improvement. Clinical practice guideline: Management of sinusitis. *Pediatrics* 2001;108: 798-808.
2. Gwaltney JM Jr. Acute community-acquired sinusitis. *Clin Infect Dis* 1996; 23:1209-1223.
3. Woodwell DA, Cherry DK. National ambulatory medical care survey:002 summary. *Adv Data* 2004;346:1-44.
4. Heikkinen T, Jarvinen A. The common cold. *Lancet* 2003;361:51-9.
5. Arroll B, Kenealy T. Antibiotics for the common cold and acute purulent rhinitis. *Cochrane Database Syst Rev* 2005;(3):CD000247.
6. Fahey T, Stocks N, Thomas T. Systematic review of the treatment of upper respiratory tract infection. *Arch Dis Child* 1998;79:225-30.
7. Schroeder K, Fahey T. Over-the-counter medications for acute cough in children and adults in ambulatory settings. *Cochrane Database Syst Rev* 2004;(4):CD001831.
8. Pavesi L, Subburaj S, Porter-Shaw K. Application and validation of a computerized cough acquisition system for objective monitoring of acute cough: a meta-analysis. *Chest* 2001;120:1121-8.
9. Eccles R, Jawad MS, Jawad SS, Angello JT, Druce HM. Efficacy and safety of single and multiple doses of pseudoephedrine in the treatment of nasal congestion associated with common cold. *Am J Rhinol* 2005;19:25-31.
10. Del Mar C, Glasziou P. Upper respiratory tract infection. *Clin Evid* 2003;10:1747-56.
11. Hayden FG, Diamond L, Wood PB, Korts DC, Wecker MT. Effectiveness and safety of intranasal ipratropium bromide in common colds. A randomized, double-blind, placebo-controlled trial. *Ann Intern Med* 1996;125:89-97.

12. Sutter AI, Lemiengre M, Campbell H, Mackinnon HF. Antihistamines for the common cold. *Cochrane Database Syst Rev* 2003;(3): CD001267
13. Paul IM, Yoder KE, Crowell KR, Shaffer ML, McMillan HS, Carlson LC, et al. Effect of dextromethorphan, diphenhydramine, and placebo on nocturnal cough and sleep quality for coughing children and their parents. *Pediatrics* 2004;114:E85-90. Accessed July 25, 2006, at: <http://pediatrics.aappublications.org/cgi/content/full/114/1/e85>.
14. Schroeder K, Fahey T. Should we advise parents to administer over the counter cough medicines for acute cough? Systematic review of randomised controlled trials. *Arch Dis Child* 2002;86:170-5.
15. Harri Hemilä, Elizabeth Chalker, Barbara Treacy, Bob Douglas. Vitamin C for preventing and treating the common cold. *Cochrane Database of Systematic Reviews*, Issue 4, 2009
16. Linde K, Barrett B, Wölkart K, Bauer R, Melchart D. Echinacea for preventing and treating the common cold. *Cochrane Database Syst Rev* 2006;(1):CD000530.
- 16a. Simasek M, Blandino D. Treatment of common cold. *Journal of American Family Physician* 2007: 75:4.
17. Rosenfeld RM, Andes D, Bhattacharyya N. Clinical practice guideline: Adult sinusitis. *Otolaryngology-Head and Neck Surgery* 2007; 137 (Suppl):S1–S31.
18. de Ferranti SD, Ioannidis JPA, Lau J, et al. Are amoxicillin and folate inhibitors as effective as other antibiotics for acute sinusitis? A meta-analysis. *BMJ* 1998;317:632–7.
19. Williams JW Jr, Aguilar C, Cornell J, et al. Antibiotics for acute maxillary sinusitis. *The Cochrane Database of Systematic Reviews* 2003, Issue 2. Art No: CD000243. DOI: 10.1002/14651858. CD000243
20. Pearlman AN, Conley DB. Review of current guidelines related to the diagnosis and treatment of rhinosinusitis. *Curr Opin Otolaryngol Head Neck Surg.* 2008 Jun;16(3):226-30.

21. Anon JB, Jacobs MR, Poole MD, et al; Sinus And Allergy Health Partnership. Antimicrobial treatment guidelines for acute bacterial rhinosinusitis. *Otolaryngol Head Neck Surg* 2004;130(Suppl):1– 45.
22. Snow V, Mottur-Pilson C, Hickner JM. Principles of appropriate antibiotic use for acute sinusitis in adults. *Ann Intern Med* 2001;134: 495–7.
23. Lau J, Zucker D, Engels EA, et al. Diagnosis and treatment of acute bacterial rhinosinusitis. Evidence Report/Technology Assessment No. 9 (Contract 290-08-0019 to the New England Medical Center). Rockville (MD): Agency for Health Care Policy and Research; March 1999.
24. de Bock GH, Dekker FW, Stolk J. Antimicrobial treatment in acute maxillary sinusitis: a meta-analysis. *J Clin Epidemiol* 1997;50:881–90.
25. Low DE, Desrosiers M, McSherry J, et al. A practical guide for the diagnosis and treatment of acute sinusitis. *CMAJ* 1997;156(Suppl 6):1–14.
26. Slavin RG, Spector SL, Bernstein IL, et al. The diagnosis and anagement of sinusitis: a practice parameter update. *J Allergy Clin Immunol* 2005;116(6 Suppl):S13– 47.
27. Institute for Clinical Systems Improvement. Acute sinusitis in adults. Bloomingdale (MN): Institute for Clinical Systems Improvement; May 2004.
28. Falagas ME, Karageorgopoulos DE, Grammatikos AP, Matthaiou DK. Effectiveness and safety of short vs. long duration of antibiotic therapy for acute bacterial sinusitis: a meta-analysis of randomized trials. *Br J Clin Pharmacol*. 2009 Feb;67(2):161-171. Epub 2008 Sep 19.
29. Craig W. Pharmacokinetic/pharmacodynamic parameters: rationale for antibacterial dosing of mice and men. *Clin Infect Dis* 1998;26:1–12.

30. Ambrose PG, Grasela DM, Grasela TH, et al. Pharmacodynamics of fluoroquinolones against *Streptococcus pneumoniae* in patients with community-acquired respiratory tract infections. *Antimicrob Agents Chemother* 2001;45:2793–7.
31. Clement PA, Bluestone CD, Gordts F, Lusk RP, Otten FW, Goossens H, et al. Management of rhinosinusitis in children: consensus meeting, Brussels, Belgium, September 13, 1996. *Arch Otolaryngol Head Neck Surg.* 1998 Jan; 124(1):31-4.
32. Wald ER. Beginning antibiotics for acute rhinosinusitis and choosing the right treatment. *Clin Rev Allergy Immunol.* 2006 Jun; 30(3):143-52.
33. European Position Paper on Rhinosinusitis and Nasal Polyposis. *Rhinology*, Supplement 20, 2007; www.rhinologyjournal.com; www.eaaci.net.
34. Jonas I, Mann W. Misleading x-ray diagnosis due to maxillary sinus asymmetries (author’s transl). *Laryngol Rhinol Otol (Stuttg).* 1976;55(11):905-13.
35. Comparison of plain radiographs and coronal CT scans in infants and children with recurrent sinusitis. *AJR Am J Roentgenol.* 1989;153(6):1259-64.
36. deleted.
37. Hwang SY, Tan KK. *Ann Otol Rhinol Laryngol.* 2007 May;116(5):381-5. *Streptococcus viridans* has a leading role in rhinosinusitis complications.
38. Meltzer EO, Bachert C, Staudinger H. Treating acute rhinosinusitis: comparing efficacy and safety of mometasone furoate nasal spray, amoxicillin, and placebo. *J Allergy Clin Immunol.* 2005 Dec;116(6):1289-95.
39. Qvarnberg Y, Kantola O, Salo J, Toivanen M, Valtonen H, Vuori E. Influence of topical steroid treatment on maxillary sinusitis. *Rhinology.* 1992;30(2):103-12.

40. Meltzer EO, Orgel HA, Backhaus JW, Busse WW, Druce HM, Metzger WJ, Mitchell DQ, Selner JC, Shapiro GG, Van Bavel JH, et al. Intranasal flunisolide spray as an adjunct to oral antibiotic therapy for sinusitis. *J Allergy Clin Immunol.* 1993 Dec;92(6):812-23.
41. Barlan IB, Erkan E, Bakir M, Berrak S, Başaran MM. Intranasal budesonide spray as an adjunct to oral antibiotic therapy for acute sinusitis in children. *Ann Allergy Asthma Immunol.* 1997 Jun;78(6):598-601.
42. Meltzer EO, Charous BL, Busse WW, Zinreich SJ, Lorber RR, Danzig MR. Added relief in the treatment of acute recurrent sinusitis with adjunctive mometasone furoate nasal spray. The Nasonex Sinusitis Group. *J Allergy Clin Immunol.* 2000;106(4):630-7.
43. Dolor RJ, Witsell DL, Hellkamp AS, Williams JW Jr., Califf RM, Simel DL. Comparison of cefuroxime with or without intranasal fluticasone for the treatment of rhinosinusitis. The CAFFS Trial: a randomized controlled trial. *Jama.* 2001;286(24):3097-105.
44. Nayak AS, Settipane GA, Pedinoff A, Charous BL, Meltzer EO, Busse WW, et al. Effective dose range of mometasone furoate nasal spray in the treatment of acute rhinosinusitis. *Ann Allergy Asthma Immunol.* 2002;89(3):271-8.
45. Gehanno P, Beauvillain C, Bobin S, Chobaut JC, Desautly A, Dubreuil C, et al. Short therapy with amoxicillin-clavulanate and corticosteroids in acute sinusitis: results of a multicentre study in adults. *Scand J Infect Dis.* 2000;32(6):679-84.
46. Klossek JM, Desmont-Gohler C, Deslandes B, Coriat F, Bordure P, Dubreuil C, et al. Treatment of functional signs of acute maxillary rhinosinusitis in adults. Efficacy and tolerance of administration of oral prednisone for 3 days. *Presse Med.* 2004 Mar 13;33(5):303-9.
47. W. Fokkens, V. Lund and C. Bachert *et al.*, EAACI position paper on rhinosinusitis and nasal polyps: executive summary, *Allergy* 60 (2005), pp. 583-601.
48. R.S. Zeiger, Prospects for ancillary treatment of sinusitis in the 1990's, *J Allergy Clin Immunol* 90 (1992), pp. 478-495.

49. Braun JJ, Alabert JP, Michel FB, Quiniou M, Rat C, Cougnard J, et al. Adjunct effect of loratadine in the treatment of acute sinusitis in patients with allergic rhinitis. *Allergy*. 1997;52(6):650-5.
50. Rabago D, Zgierska A, Mundt M, et al. Efficacy of daily hypertonic saline nasal irrigation among patients with sinusitis: a randomized controlled trial. *J Fam Pract* 2002;51:1049-55.
51. Talbot AR, Herr TM, Parsons DS. Mucociliary clearance and buffered hypertonic saline solution. *Laryngoscope*. 1997 Apr;107(4):500-3.
52. Ural A, Oktemer TK, Kizil Y, Ileri F, Uslu S.J. Impact of isotonic and hypertonic saline solutions on mucociliary activity in various nasal pathologies: clinical study. *Laryngol Otol*. 2008 Oct 28:1-5. [Epub ahead of print]
53. Namyslowski G, Misiolek M, Czecior E, et al. Comparison of the efficacy and tolerability of amoxicillin/clavulanic acid 875 mg b.i.d. with cefuroxime 500 mg b.i.d. in the treatment of chronic and acute exacerbation of chronic sinusitis in adults. *J Chemother* 2002;14(5):508-17.
54. Legent F, Bordure P, Beauvillain C, et al. A double-blind comparison of ciprofloxacin and amoxicillin/clavulanic acid in the treatment of chronic sinusitis. *Chemotherapy* 1994;40(Suppl. 1):8-15.
55. Subramanian HN, Schechtman KB, Hamilos DL. A retrospective analysis of treatment outcomes and time to relapse after intensive medical treatment of chronic sinusitis. *Am J Rhinol* 2002;16:303-12.
56. Hashiba M, Baba S. Efficacy of long-term administration of clarithromycin in the treatment of intractable chronic sinusitis. *Acta Otolaryngol Suppl* 1996;525:73-8.
57. Nishi K, Mizuguchi M, Tachibana H, et al. Effect of clarithromycin on symptoms and mucociliary transport in patients with sino-bronchial syndrome. *Nihon Kyobu Shikkan Gakkai Zasshi* 1995;33(12):1392-1400.

58. Lund VJ, Black JH, Szabo LZ, Schrewelius C, Akerlund A. Efficacy and tolerability of budesonide aqueous nasal spray in chronic rhinosinusitis patients. *Rhinology* 2004;42:57-62.
59. Lavigne F, Cameron L, Renzi PM, et al. Intranasal administration of topical budesonide to allergic patients with chronic sinusitis following surgery. *Laryngoscope* 2002;112(5):858-64.
60. Bachmann G, Hommel G, Michel O. Effect of irrigation of the nose with isotonic salt solution on adult patients with chronic paranasal sinus disease. *Eur Arch Otorhinolaryngol* 2000;257(10):537-41.
61. Ichimura K, Shimazaki Y, Ishibashi T, et al. Effect of new macrolide roxithromycin upon nasal polyps associated with chronic sinusitis. *Auris Nasus Larynx* 1996;23:48-56.
62. Ragab SM, Lund VJ, Scadding G. Evaluation of the medical and surgical treatment of chronic rhinosinusitis; a prospective, randomised, controlled trial. *Laryngoscope* 2004;114(5):923-30.
63. Lildholdt T, Rundcrantz H, Bende M, et al. Glucocorticoid treatment for nasal polyps. The use of topical budesonide powder, intramuscular betamethasone, and surgical treatment. *Ach Otolaryngol Head Neck Surg* 1997;123(6):595-600.
64. Holmberg K, Juliusson S, Balder B, et al. Fluticasone propionate aqueous nasal spray in the treatment of nasal polyposis. *Ann Allergy Asthma Immunol* 1997;78(3):270-6.
65. Lund VJ, Flood J, Sykes AP, et al. Effect of fluticasone in severe polyposis. *Arch Otolaryngol Head Neck Surg* 1998;124(5):513-8.
66. Tos M, Svendstrup F, Arndal H, et al. Efficacy of an aqueous and a powder formulation of nasal budesonide compared in patients with nasal polyps. *Am J Rhinol* 1998;12:183-189.
67. Keith P, Nieminen J, Hollingworth K, et al. Efficacy and tolerability of fluticasone propionate nasal drops 400 microgram once daily compared with placebo for the treatment of bilateral polyposis in adults. *Clin Exp Allergy* 2000;30(10):1460-8.

68. Penttila M, Poulsen P, Hollingworth K, et al. Dose-related efficacy and tolerability of fluticasone propionate nasal drops 400 microg once daily and twice daily in the treatment of bilateral nasal polyposis: a placebo-controlled randomized study in adult patients. *Clin Exp Allergy* 2000;30(1):94-102.
69. van Camp C, Clement PA. Results of oral steroid treatment in nasal polyposis. *Rhinology* 1994;32(1):5-9.
70. Damm M, Jungehulsing M, Eckel HE, et al. Effects of systemic steroid treatment in chronic polypoid rhinosinusitis evaluated by magnetic resonance imaging. *Otolaryngol Head Neck Surg* 1999;120(4):517-23.
71. Benitez P, Alobid I, de Haro Jet al. A short course of oral prednisone followed by intranasal budesonide is an effective treatment of severe nasal polyps. *Laryngoscope* 2006;116:770-775.
72. Haye R, Aanesen JP, Burtin B, et al. The effect of cetirizine on symptoms and signs of nasal polyposis. *J Laryngol Otol* 1998;112:1042-1046.
73. Gordts F, Clement PA, Destryker A, Desprechins B, Kaufman L. Prevalence of sinusitis signs on MRI in a non-ENT paediatric population. *Rhinology*.1997 Dec; 35 (4):154-7.
74. Orobello PW, Jr., Park RI, Belcher LJ, Eggleston P, Lederman HM, Banks JR, et al. Microbiology of chronic sinusitis in children. *Arch Otolaryngol Head Neck Surg*. 1991; 117(9): 980-3.
75. Oxford LE, McClay J. Complications of acute sinusitis in children. *Otolaryngology - Head & Neck Surgery*. 2005; 133(1):32-7
76. Clement PA, Bluestone CD, Gordts F, Lusk RP, Otten FW, Goossens H, et al. Management of rhinosinusitis in children. *Int J Pediatr Otorhinolaryngol*. 1999 Oct 5; 49 Suppl 1:S95-100.
77. Otten FW, Grote JJ. The diagnostic value of transillumination for maxillary sinusitis in children. *Int J Pediatr Otorhinolaryngol*. 1989;18(1):9-11.

78. van der Veken PJ, Clement PA, Buisseret T, Desprechins B, Kaufman L, Derde MP. CT-scan study of the incidence of sinus involvement and nasal anatomic variations in 196 children. *Rhinology*. 1990;28(3):177-84.
79. Manning SC. Pediatric sinusitis. *Otolaryngol Clin North Am*. 1993; 26(4):623-38.
80. Ciprandi G, Tosca MA, Fasce L. Allergic children have more numerous and severe respiratory infections than non-allergic children. *Pediatr Allergy Immunol*. 2006 Aug; 17(5):389-91.
81. Polmar SH. The role of the immunologist in sinus disease. *J Allergy Clin Immunol*. 1992; 90(3 Pt 2):511-4.
82. Phipps CD, Wood WE, Gibson WS, Cochran WJ. Gastroesophageal reflux contributing to chronic sinus disease in children: a prospective analysis. *Arch Otolaryngol Head Neck Surg*. 2000; 126(7):831-6.
83. Yung MW, Gould J, Upton GJ. Nasal polyposis in children with cystic fibrosis: a long-term follow-up study. *Ann Otol Rhinol Laryngol*. 2002; 111 (12 Pt 1):1081-6.
84. Sleigh MA. Primary ciliary dyskinesia. *Lancet*. 1981 Aug 29; 2 (8244):476.
85. Morris P, Leach A. Antibiotics for persistent nasal discharge (rhinosinusitis) in children (Cochrane Review). *Cochrane Database Syst Rev*. 2002(4):CD001094.
86. Balatsouras DG, Korres S, Rallis E, Eliopoulos P, Ferekidis E. Twice-daily dosing of loracarbef 15 mg/kg versus 30 mg/kg in the treatment of children with acute sinusitis. *Drugs Exp Clin Res*. 2005; 31 Suppl:1-5.
87. Yilmaz G, Varan B, Yilmaz T, Gurakan B. Intranasal budesonide spray as an adjunct to oral antibiotic therapy for acute sinusitis in children. *Eur Arch Otorhinolaryngol*. 2000; 257(5):256-9.
88. McCormick DP, John SD, Swischuk LE, Uchida T. A double blind, placebo-controlled trial of decongestant-antihistamine for the

- treatment of sinusitis in children. *Clin Pediatr (Phila)*. 1996; 35(9):457-60.
89. Poole MD. Pediatric endoscopic sinus surgery: the conservative view. *Ear Nose Throat J*. 1994 Apr; 73(4):221-7.
 90. Otten FW, Grote JJ. Treatment of chronic maxillary sinusitis in children. *Int J Pediatr Otorhinolaryngol*. 1988; 15(3):269-78.
 91. Van Bever HP, Bosmans J, Stevens WJ. Nebulization treatment with saline compared to bromhexine in treating chronic sinusitis in asthmatic children. *Allergy*. 1987; 42(1):33-6.
 92. Shoseyov D, Bibi H, Shai P, Shoseyov N, Shazberg G, Hurvitz H. Treatment with hypertonic saline versus normal saline nasal wash of pediatric chronic sinusitis. *J Allergy Clin Immunol*. 1998;101(5):602-5.
 93. Bothwell MR, Parsons DS, Talbot A, Barbero GJ, Wilder BI. Outcome of reflux therapy on pediatric chronic sinusitis. *Otolaryngol Head Neck Surg*. 1999; 121(3):255-62.
 94. Hebert RL, 2nd, Bent JP, 3rd. Meta-analysis of outcomes of pediatric functional endoscopic sinus surgery. *Laryngoscope*. 1998; 108(6):796-9.
 95. Ungkanont K, Damrongsak S. Effect of adenoidectomy in children with complex problems of rhinosinusitis and associated diseases. *Int J Pediatr Otorhinolaryngol*. 2004 Apr; 68(4):447-51.
 96. Lusk RP, Lazar RH, Muntz HR. The diagnosis and treatment of recurrent and chronic sinusitis in children. *Pediatr Clin North Am*. 1989; 36(6):1411-21.
 97. Vuurman EF, van Veggel LM, Uiterwijk MM, Leutner D, O'Hanlon JF. Seasonal allergic rhinitis and antihistamine effects on children's learning. *Ann Allergy* 1993; 71:121-6.
 98. Sheikh A, Khan-Wasti S, Price D, Smeeth L, Fletcher M, Walker S, Schwarzer G, Bassler D, Mitra A, Ducharme FM, Forster J. Ketotifen alone or as additional medication for long-term control of asthma and

- wheeze in children. *Cochrane Database Syst Rev* 2004; CD001384.
treatment: the effect on cells and cytokines in nasal allergic
inflammation. *Am J Rhinol* 1998; 12:21–6.
99. Nelson HS. Prospects for antihistamines in the treatment of asthma. *J Allergy Clin Immunol* 2003; 112:S96–100. nose drops. In rhinological disease betamethasone.
 100. Yanez A, Rodrigo GJ. Intranasal corticosteroids versus topical H1 receptor antagonists for the treatment of allergic rhinitis: a systematic review with meta-analysis. *Ann Allergy Asthma Immunol* 2002; 89:479–84.
 101. Weiner JM, Abramson MJ, Puy RM. Intranasal corticosteroids versus oral H1 receptor antagonists in allergic rhinitis: systematic review of randomised controlled trials. *BMJ* 1998; 317:1624–9.
 102. Wilson AM, O’Byrne PM, Parameswaran K. Leukotriene receptor antagonists for allergic rhinitis: a systematic review and meta-analysis. *Am J Med* 2004; 116:338–44.
 103. Nasser SM, Ewan PW. Lesson of the week: depot corticosteroid treatment for hay fever causing avascular necrosis of both hips. *BMJ* 2001; 322:1589–91.
 104. McNeely W, Wiseman LR. Intranasal azelastine. A review of its efficacy in the management of allergic rhinitis. *Drugs* 1998; 56:91–114.
 105. Portnoy JM, Van OT, Williams PB. Evidence-based strategies for treatment of allergic rhinitis. *Curr Allergy Asthma Rep* 2004; 4:439–46.
 106. Grossman J, Banov C, Boggs P et al. Use of ipratropium bromide nasal spray in chronic treatment of nonallergic perennial rhinitis, alone and in combination with other perennial rhinitis medications. *J Allergy Clin Immunol* 1995; 95:1123–7.
 107. Tan R, Corren J. Optimum treatment of rhinitis in the elderly. *Drugs Aging* 1995; 7:168–75.

108. Welsh PW, Stricker WE, Chu CP, Naessens JM, Reese ME, Reed CE, et al. Efficacy of beclomethasone nasal solution, flunisolide, and cromolyn in relieving symptoms of ragweed allergy. *Mayo Clin Proc* 1987;62:125-134.
109. Meltza EO. Efficacy and patient satisfaction with cromolyn sodium nasal solution in the treatment of seasonal allergic rhinitis: a placebo controlled study. *Clin Ther* 2002;24:942-952.
110. Druce HM, Goldstein S, Melamed I. Multicenter placebo-controlled study of nedocromil sodium 1% nasal solution in ragweed seasonal allergic rhinitis. *Ann Allergy* 1990;65:212-216.
111. Siplla P, Sorri M, Pukande J. Double-blind comparison of nedocromil sodium (1% nasal spray) and placebo in rhinitis caused by birch pollen. *Clin Otolaryngol* 1987;12:365-370.
112. Sculler DE, Selcow JE, Joos TH, Hannawy PJ, Hirsch SR, Schwartz HJ, et al. A multicenter trial of nedocromil sodium, 1% nasal solution, compared with cromolyn sodium and placebo in ragweed seasonal allergic rhinitis. *J Allergy Clin Immunol* 1990;6:554-561.
113. Magyar P, Gyori Z, Mark Z, Hutás I. The protective effect of N-acetyl-aspartyl-glutamate (NAAGA) against nasal obstruction provoked by antigen in allergic rhinitis. *Allergy* 1993;48:631-633.
114. James IG, Campbell LM, Harrison JM, Fell PJ, Ellers-Lenz B, Petold U. Comparison of the efficacy and tolerability of topically administered azelastine, sodium cromoglycate and placebo in the treatment of seasonal allergic conjunctivitis and rhino-conjunctivitis. *Curr Med Res Opin* 2003;19:313-320.
115. Meltzer E, Malmstrom K, Lu S, Brenner B, Wei L, Weinstein S, et al. Concomitant montelukast and loratadine as treatment for seasonal allergic rhinitis: placebo-controlled clinical trial. *J Allergy Clin Immunol* 2000;105:917-922.
116. Philip G, Malmstrom K, Hampel FC, Weinstein SF, LaForce CF, Ratner PH, et al. Montelukast for treating seasonal allergic rhinitis: a randomized, double-blind, placebocontrolled trial performed in the spring. *Clin Exp Allergy* 2002;32:1020-1028.

117. Patel P, Philip G, Yang W, Call R, Horak F, LaForce C, et al. Randomized, double-blind, placebo-controlled study of montelukast for treating perennial allergic rhinitis. *Ann Allergy Asthma Immunol* 2005;95:551–557.
118. Chervinsky P, Philip G, Malice MP, Bardelas J, Nayak A, Marchal JL, et al. Montelukast for treating fall allergic rhinitis: effect of pollen exposure in 3 studies. *Ann Allergy Asthma Immunol* 2004;92:367–373.
119. Nayak AS, Banov C, Corren J, Feinstein BK, Floreani A, Friedman BF, et al. Once-daily mometasone furoate dry powder inhaler in the treatment of patients with persistent asthma. *Cochrane Database Syst Rev* 2000;2:417–424.
120. van Adelsberg J, Philip G, Pedinoff AJ, Meltzer EO, Ratner PH, Menten J, et al. Montelukast improves symptoms of seasonal allergic rhinitis over a 4-week treatment period. *Allergy* 2003;58:1268–1276.
121. van Adelsberg J, Philip G, LaForce CF, Weinstein SF, Menten J, Malice MP, et al. Randomized controlled trial evaluating the clinical benefit of montelukast for treating spring seasonal allergic rhinitis. *Ann Allergy Asthma Immunol* 2003;90:214–222.
122. Philip G, Nayak AS, Berger WE, Leynadier F, Vrijens F, Dass SB, et al. The effect of montelukast on rhinitis symptoms in patients with asthma and seasonal allergic rhinitis. *Curr Med Res Opin* 2004;20:1549–1558.
123. Busse WW, Casale TB, Dykewicz MS, Meltzer EO, Bird SR, Hustad CM, et al. Efficacy of montelukast during the allergy season in patients with chronic asthma and seasonal aeroallergen sensitivity. *Ann Allergy Asthma Immunol* 2006;96:60–68.
124. Johnson DA, Hricik JG. The pharmacology of alpha-adrenergic decongestants. *Pharmacotherapy* 1993;13:110S–115S; discussion 43S–46S.
125. Johannssen V, Maune S, Werner JA, Rudert H, Ziegler A. Alpha 1-receptors at pre-capillary resistance vessels of the human nasal mucosa. *Rhinology* 1997;35:161–165.

126. Graf P, Hallen H, Juto JE. Fourweek use of oxymetazoline nasal spray (Nezeril) once daily at night induces rebound swelling and nasal hyperreactivity. *Acta Otolaryngol* 1995;115:71–75.
127. Graf P, Hallen H. Effect on the nasal mucosa of long-term treatment with oxymetazoline, benzalkonium chloride, and placebo nasal sprays. *Laryngoscope* 1996;106:605–609.
128. Graf P. Rhinitis medicamentosa: aspects of pathophysiology and treatment. *Allergy* 1997;52(Suppl. 40):28–34.
129. Scadding GK. Rhinitis medicamentosa [editorial]. *Clin Exp Allergy* 1995;25:391–394.
130. Graf P. Rhinitis medicamentosa: a review of causes and treatment. *Treat Respir Med* 2005;4:21–29.
131. Kung YY, Chen YC, Hwang SJ, Chen TJ, Chen FP. The prescriptions frequencies and patterns of Chinese herbal medicine for allergic rhinitis in Taiwan. *Allergy* 2006;61:1316–1318.
132. Passalacqua G, Bousquet PJ, Carlsen KH, Kemp J, Lockey RF, Niggemann B, et al. ARIA update: I. Systematic review of complementary and alternative medicine for rhinitis and asthma. *J Allergy Clin Immunol* 2006;117:1054–1062.
133. Linde K, Jonas WB, Melchart D, Willich S. The methodological quality of randomized controlled trials of homeopathy, herbal medicines and acupuncture. *Int J Epidemiol* 2001;30:526–531.
134. Linde K, Hondras M, Vickers A, Riet Gt G, Melchart D. Systematic reviews of complementary therapies – an annotated bibliography. Part 3: Homeopathy. *BMC Complement Altern Med* 2001;1:4.
135. Linde K, ter Riet G, Hondras M, Vickers A, Saller R, Melchart D. Systematic reviews of complementary therapies – an annotated bibliography. Part 2: Herbal medicine. *BMC Complement Altern Med* 2001;1:5.
136. Linde K, Vickers A, Hondras M, ter Riet G, Thormahlen J, Berman B, et al. Systematic reviews of complementary therapies – an annotated

- bibliography. Part 1: Acupuncture. *BMC Complement Altern Med* 2001;1:3.
137. Gentile D, Shapiro G, Sloner D. Allergic rhinitis. In: Leung D, Sampson H, Geha R, Szeffler S, editors. *Pediatric allergy. Principles and practice*. St. Louis, Missouri: Mosby, 2003:287–297.
 138. Blackwell DL, Tonthat L. Summary health statistics for U.S. children: National Health Interview Survey, 1999. *Vital Health Stat* 10, 2003:1-50.
 139. Wright AL, Holberg CJ, Martinez FD, Halonen M, Morgan W, Taussig LM. Epidemiology of physician-diagnosed allergic rhinitis in childhood. *Pediatrics* 1994; 94:895- 901.
 140. Menardo JL, Bousquet J, Rodiere M, Astruc J, Michel FB. Skin test reactivity in infancy. *J Allergy Clin Immunol* 1985;75:646–651.
 141. Daley-Yates PT, Richards DH. Relationship between systemic corticosteroid exposure and growth velocity: development and validation. *Clin Ther* 2004;26:1905–1919.
 142. Fink RS, Pierre LN, Daley-Yates PT, Richards DH, Gibson A, Honour JW. Hypothalamic-pituitary-adrenal axis function after inhaled corticosteroids: unreliability of urinary free cortisol estimation. *J Clin Endocrinol Metab* 2002;87: 4541–4546.
 143. Schenkel E, Skoner D, Bronsky E, Miller S, Pearlman D, Rooklin A, et al. Absence of growth retardation in children with perennial allergic rhinitis following 1 year treatment with mometasone furoate aqueous nasal spray. *Pediatrics* 2000;101:e22.
 144. Allen DB. Do intranasal corticosteroids affect childhood growth? *Allergy* 2000;55(Suppl. 62):15–18. Ngamphaiboon J, Thepchatri A, Chatchatee P, Chumdermpadetsuk S. Fluticasone propionate aqueous.
 145. Brannan MD, Herron JM, Affrime MB. Safety and tolerability of once-daily mometasone furoate aqueous nasal spray in children. *Clin Ther* 1997;19:1330–1339.

146. Cutler D, Banfield C, Affrime M. Safety of mometasone furoate nasal spray in children with allergic rhinitis as young as 2 years of age: a randomized controlled trial. *Pediatr Asthma, Allergy Immunol* 2006;19:146–153.
147. Agertoft L, Pedersen S. Short-term lower leg growth rate in children with rhinitis treated with intranasal mometasone furoate and budesonide [In Process Citation]. *J Allergy Clin Immunol* 1999;104:948–952.
148. Bousquet J, Khaltaev N. et al. Allergic rhinitis and its Impact on Asthma (ARIA) 2008 Update. *Allergy* 2008; 63 (Suppl 86): 1-160.
149. Wolthers OD, Pedersen S. Shortterm growth in children with allergic rhinitis treated with oral antihistamine, depot and intranasal glucocorticosteroids [see comments]. *Acta Paediatr* 1993;82:635–640. *Immunol* 2004;15:261–266.
150. Simons FE, Silas P, Portnoy JM, Catuogno J, Chapman D, Olufade AO, et al. Safety of cetirizine in infants 6 to 11 months of age: a randomized, double-blind, placebo-controlled study. *J Allergy Clin Immunol* 2003;111:1244–1248.
151. Grimfeld A, Holgate ST, Canonica GW, Bonini S, Borres MP, Adam D, et al. Prophylactic management of children at risk for recurrent upper respiratory infections: the Preventia I Study. *Clin Exp Allergy* 2004;34:1665–1672.
152. Simons FER, Group ObotEPoAiACES. Safety of levocetirizine treatment in young atopic children. A 18-month study. *Pediatr Allergy Immunol*. 2007;18:535–542.
153. Herman D, Garay R, Le-Gal M. A randomized double-blind placebo controlled study of azelastine nasal spray in children with perennial rhinitis. *Int J Pediatr Otorhinolaryngol* 1997;39:1–8.
154. Vermeulen J, Mercer M. Comparison of the efficacy and tolerability of topical levocabastine and sodium cromoglycate in the treatment of seasonal allergic rhinoconjunctivitis in children. *Pediatr Allergy Immunol* 1994;5:209–213.

155. Sabbah A, Marzetto M. Azelastine eye drops in the treatment of seasonal allergic conjunctivitis or rhinoconjunctivitis in young children. *Curr Med Res Opin* 1998;14:161–170.
156. Engstrom I, Oberger E, Blyckert A, Kraepelien S. Disodium cromoglycate in the treatment of seasonal allergic rhinoconjunctivitis in children. *Ann Allergy* 1971;29:505–509.
157. Sensi LG, Seri A, Siracusa A, Pertici L, Marcucci F. Allergic rhinitis in children: effects of fluticasone and disodium cromoglycate on nasal eosinophil cationic protein. *Clin Exp Allergy* 1997;27:270-276.
158. Agostinis F, Tellarini L, Canonica GW, Falagiani P, Passalacqua G. Safety of sublingual immunotherapy with a monomeric allergoid in very young children. *Allergy* 2005;60:133.
159. Baena-Cagnani CE, Passalacqua G, Baena-Cagnani RC, Croce VH, Canonica WG. Sublingual immunotherapy in pediatric patients: beyond clinical efficacy. *Curr Opin Allergy Clin Immunol* 2005;5:173-177.
160. Penagos M, Compalati E, Tarantini F, Baena-Cagnani R, Huerta J, Passalacqua G, et al. Efficacy of sublingual immunotherapy in the treatment of allergic rhinitis in pediatric patients 3 to 18 years of age: a meta-analysis of randomized, placebocontrolled, double-blind trials. *Ann Allergy Asthma Immunol* 2006;97:141-148.
161. Gani F, Braida A, Lombardi C, Del Giudice A, Senna GE. Rhinitis in pregnancy. *Eur Ann Allergy Clin Immunol.* 2003;35(8):306-313.
162. Somoskovi A, Bartfai Z, Tamasi L. Population-based case-control study of allergic rhinitis during pregnancy for birth outcomes. *European Journal of Obstetrics and Gynecology and Reproductive Biology.* 2007;131:21-27.
163. Ellegard EK. The etiology and management of pregnancy rhinitis. *Am J Respir Med* 2003;2:469-475.
164. Nathan R. The burden of allergic rhinitis. *Allergy and Asthma Proceedings.* 2007;28(1):3-9.

165. Scadding G, Mirakian R. National Knowledge Week for Rhinitis 2007-Rhinitis and Pregnancy. National Health Library. 2007.
166. Gilbert C, Mazzotta P, Loebstein R, Koren G. Fetal safety of drugs used in the treatment of allergic rhinitis: a critical review. *Drug Saf.* 2005;28(8):707-719.
167. Demoly P, Piette V, Daures JP. Treatment of allergic rhinitis during pregnancy. *Drugs.* 2003;63:1813-1820.
168. Einaron A, Bailey B, Jung G, Spizzirri D, Baillie M, Koren G. Prospective controlled study of hydroxyzine and cetirizine in pregnancy. *Ann Allergy Asthma Immunol.* 1997 Feb;78(2):183-6.
169. Schoendorfer, Corinna Weber and Schaefer, Christof. The safety of cetirizine during pregnancy. A prospective observational cohort study. Berlin Institute for Clinical Teratology and Drug Risk Assessment in Pregnancy. 13 May 2008.
170. Diav-Citrin O, Schetman S, Aharonovich A, Moerman L, Arnon J. Pregnancy outcome after gestational exposure to loratadine or antihistamines:a prospective controlled cohort study. *J Allergy Clin Immunol.* 2003;111:1239-1243.
171. Moretti ME, Caprara D, Coutinho CJ, Bar-oz B, Berkovitch M, Addis A. Fetal safety of loratadine use in the first trimester of pregnancy:a multicenter study. *J Allergy Clin Immunol.* 2003;111:479-483.
172. Gluck PA, Gluck JC. A review of pregnancy outcomes after exposure to orally inhaled or intranasal budesonide. *Curr Med Res Opin.* 2005;21(7):1075-1084.
173. Källén BA, Otterblad Olausson P. Maternal drug use in early pregnancy and infant cardiovascular defect. *Reprod Toxicol* 2003; 17: 255-61.
174. Incaudo GA. Diagnosis and treatment of allergic rhinitis and sinusitis during pregnancy and lactation. *Clin Rev Allergy Immunol* 2004; 27: 159-77.

175. Rayburn WF, Anderson JC, Smith CV, et al. Uterine and fetal Doppler flow changes from a single dose of a long-acting intranasal decongestant. *Obstet Gynecol* 1990;76(2):180.
176. Baxi LV, Gindoff PR, Pregonzer GJ, et al. Fetal heart rate changes following maternal administration of a nasal decongestant. *Am J Obstet Gynecol* 1985;153(7):799–800.
177. Kallen B, Ollausson P. Use of oral decongestants during pregnancy and delivery outcome. *Am J Obs Gyne.* 2006;194:480-485.
178. Toll K, Graf P. Phenylpropanolamine's decongestive effect on the nasal mucosa of pregnant women with nasal stuffiness. *Rhinology.* 2006;44(4):274-277.
179. Gilbert C, Mazzotta P, Loebstein R, et al. Fetal safety of drugs used in the treatment of allergic rhinitis: a critical review. *Drug Saf* 2005; 28: 707-19.
180. Williams JW Jr, Aguilar C, Cornell J, et al. Antibiotics for acute maxillary sinusitis. *The Cochrane Database of Systematic Reviews* 2003, Issue 2. Art No: CD000243. DOI: 10.1002/14651858.CD000243.
181. Heinonen OP, Slone D, Shapiro S. *Birth Defects and Drugs in Pregnancy.* Littleton, MA:Publishing Sciences Group, 1977.
182. Czeizel AE, Rockenbauer M, Sorensen H., and Olsen J. Augmentin treatment during pregnancy and the prevalence of congenital abnormalities: A population-based study. *Eur Jour Obs and Gyn and Repro Bio.* 2001; 97:188-192.
183. Sorenzen HT., Larsen H., Jensen ES et al. Safety of metronidazole during pregnancy: a cohort study of risk of congenital abnormalities, preterm delivery and low birth weight in 124 women. *Journal of Antimicrobial Chemotherapy.* 1999; 44:847-855.
184. Metzger WJ. Indications for allergen immunotherapy during pregnancy. *Compr Ther.* 1990;16(3):17-26.

185. Einarson A, Shuhaiber S, Koren G: Effects of antibacterials on the unborn child: what is known and how should this influence prescribing. *Paediatr Drugs*. 2001;3(11):803-16.
186. Food and Drug Administration. Federal Register 1980;44:37434-67.
187. Reed, S. D., T. A. Lee, et al. (2004). "The economic burden of allergic rhinitis: a critical evaluation of the literature." *Pharmacoeconomics* 22(6): 345-61.
188. Simoens, S. and G. Laekeman (2009). "Pharmacotherapy of allergic rhinitis: a pharmaco-economic approach." *Allergy* 64(1): 85-95.

Self-assessment (MCQs)

After reading the Clinical Practice Guidelines, you can claim one CME point under Category 3A (Self-Study) of the SMC Online CME System. Alternatively, you can claim one CME point under Category 3B (Distance Learning - Verifiable Self Assessment) if you answer at least 60% of the following MCQs correctly. You can submit your answers through the SMJ website at this link: <http://smj.sma.org.sg/cme/smj/index.html>. The answers will be published in the SMJ May 2010 issue and at the MOH webpage for these guidelines after the period for submitting the answers is over.

Instruction: Choose “True” or “False”.

	True	False
1. The following suggest a diagnosis of acute bacterial rhinosinusitis instead of a common cold:		
A) More severe symptoms than usual.	<input type="checkbox"/>	<input type="checkbox"/>
B) Cold symptoms lasting more than 10 days.	<input type="checkbox"/>	<input type="checkbox"/>
C) Low grade fever.	<input type="checkbox"/>	<input type="checkbox"/>
D) Symptoms worsen after several days of improvement.	<input type="checkbox"/>	<input type="checkbox"/>
2. Which of the following statements is FALSE on paediatric rhinosinusitis?		
A) Eye swelling from orbital infection can occur without pain in the eye or history of rhinosinusitis.	<input type="checkbox"/>	<input type="checkbox"/>
B) Plain X ray is sensitive in the diagnosis of acute rhinosinusitis.	<input type="checkbox"/>	<input type="checkbox"/>
C) Recalcitrant cases of rhinosinusitis require an exclusion of laryngopharyngeal reflux.	<input type="checkbox"/>	<input type="checkbox"/>
D) Frequent exacerbations of chronic rhinosinusitis may benefit from 2 weeks of oral antibiotics.	<input type="checkbox"/>	<input type="checkbox"/>
3. In patients with persistent allergic rhinitis		
A) the most common aeroallergen locally is house dust mites.	<input type="checkbox"/>	<input type="checkbox"/>
B) evaluation for asthma should be performed.	<input type="checkbox"/>	<input type="checkbox"/>
C) measurement of total IgE is useful in the diagnosis.	<input type="checkbox"/>	<input type="checkbox"/>
D) long-term use of oral glucocorticosteroids is not recommended due to safety concerns.	<input type="checkbox"/>	<input type="checkbox"/>

	True	False
4. With regards to antibiotics and acute rhinosinusitis;		
A) In general, adults suffer 6 to 8 colds per year.	<input type="checkbox"/>	<input type="checkbox"/>
B) Antibiotics need not be started in patients with acute rhinosinusitis until after 10 days from onset of symptoms unless symptoms are severe.	<input type="checkbox"/>	<input type="checkbox"/>
C) Greenish nasal discharge suggestive of bacterial sinusitis always requires antibiotic treatment.	<input type="checkbox"/>	<input type="checkbox"/>
D) The recommended duration of use of antibiotics is 14 days.	<input type="checkbox"/>	<input type="checkbox"/>
5. In the diagnosis of acute infective rhinosinusitis in adults,		
A) fever is a diagnostic criteria.	<input type="checkbox"/>	<input type="checkbox"/>
B) radiological imaging is not needed to make the diagnosis.	<input type="checkbox"/>	<input type="checkbox"/>
C) eye swelling requires immediate specialist referral.	<input type="checkbox"/>	<input type="checkbox"/>
D) symptoms should resolve within 5 days.	<input type="checkbox"/>	<input type="checkbox"/>
6. For treatment of acute bacterial rhinosinusitis,		
A) anti-histamines are indicated in all patients.	<input type="checkbox"/>	<input type="checkbox"/>
B) fluoroquinolones should not be used as first line antibiotics.	<input type="checkbox"/>	<input type="checkbox"/>
C) nasal corticosteroid spray has not been shown to reduce symptoms.	<input type="checkbox"/>	<input type="checkbox"/>
D) oral steroids should be used for all patients.	<input type="checkbox"/>	<input type="checkbox"/>
7. Drugs recommended for use in rhinitis in pregnancy include:		
A) Budesonide	<input type="checkbox"/>	<input type="checkbox"/>
B) Oral decongestants	<input type="checkbox"/>	<input type="checkbox"/>
C) Leukotriene Modifiers	<input type="checkbox"/>	<input type="checkbox"/>
D) Cefuroxime	<input type="checkbox"/>	<input type="checkbox"/>
8. Recommended treatment for chronic sinusitis with nasal polyps in adults include		
A) Antibiotics	<input type="checkbox"/>	<input type="checkbox"/>
B) Nasal corticosteroid therapy	<input type="checkbox"/>	<input type="checkbox"/>
C) Oral Steroids	<input type="checkbox"/>	<input type="checkbox"/>
D) Mucolytics	<input type="checkbox"/>	<input type="checkbox"/>

	True	False
9. Which of the following symptoms together with persistent nasal congestion or discoloured nasal discharge lasting for more than 3 months suggest the diagnosis of chronic sinusitis in adults?		
A) Visual changes	<input type="checkbox"/>	<input type="checkbox"/>
B) Epistaxis	<input type="checkbox"/>	<input type="checkbox"/>
C) Loss of smell	<input type="checkbox"/>	<input type="checkbox"/>
D) Headache	<input type="checkbox"/>	<input type="checkbox"/>
10. Regarding paediatric allergic rhinitis:		
A) In pre-school children, allergic rhinitis occurs at the same time as asthma	<input type="checkbox"/>	<input type="checkbox"/>
B) Allergic rhinitis often come to light when preschoolers are being treated for co-morbidities, e.g. chronic otitis media with effusion	<input type="checkbox"/>	<input type="checkbox"/>
C) The aim of treatment in paediatric allergic rhinitis is to cure the disease.	<input type="checkbox"/>	<input type="checkbox"/>
D) All intranasal steroid sprays are safe for use in children.	<input type="checkbox"/>	<input type="checkbox"/>

Workgroup members

The members of the workgroup, who were appointed in their personal professional capacity, are:

Chairman

Clin A/Prof Siow Jin Keat
Clinical Associate Professor
National University of Singapore;
Senior Consultant
Dept of Otorhinolaryngology
Tan Tock Seng Hospital

Members

Clin A/Prof Abhilash Balakrishnan
Clinical Associate Professor
National University of Singapore;
Senior Consultant
Dept. of Otolaryngology
Singapore General Hospital and KK
Women's and Children's Hospital

A/Prof Lynne Lim
Senior Consultant
Dept of Otolaryngology-Head &
Neck Surgery
National University Hospital and
National University of Singapore

A/Prof Wang De Yun
Research Director
Dept of Otolaryngology-Head &
Neck Surgery
National University Health System

Dr Leong Jern-Lin
Consultant
ASCENT Ear Nose Throat Specialist
Group
Mount Elizabeth Medical Centre

Clin A/Prof Henry Tan
Deputy Chairman
Division of Surgery,
Head and Senior Consultant
Dept of Otolaryngology (Paediatric
Otolaryngology)
KK Women's and Children's
Hospital

Dr Anita Menon
Consultant
Infectious Disease Service
Dept of Paediatric Medicine
KK Women's and Children's
Hospital

Dr Chao Siew Shuen
Senior Consultant
Dept of Otolaryngology-Head &
Neck Surgery (Rhinology)
National University Hospital

Dr Julian Lee Cheow Yew
Senior Consultant
Dept of Otolaryngology
Tan Tock Seng Hospital

Members

Dr Jason Hwang Siew Yoong
Consultant
Dept of Otorhinolaryngology
Changi General Hospital

Clin A/Prof Dharmbir S Sethi
Clinical Associate Professor
National University of Singapore;
Senior Consultant
Dept of Otolaryngology
Singapore General Hospital

Dr Chan Kwai Onn
Consultant
K O Chan ENT Sinus & Sleep
Centre

Dr Nada Ali Alshaikh
Clinical Fellow
Dept of Otorhinolaryngology
Tan Tock Seng Hospital

A/Prof Goh Lee Gan
Associate Professor
Dept of COFM
Yong Loo Lin School of Medicine;
President
College of Family
Physicians, Singapore

Subsidiary editors:

Dr Pwee Keng Ho
Deputy Director (Health Technology Assessment)
Health Services Research & Evaluation Division
Ministry of Health

Dr Sandi Chit Lwin
Manager (Health Technology Assessment)
Health Services Research & Evaluation Division
Ministry of Health

Acknowledgement:

Dr Edwin Chan Shih-Yen
Head, Epidemiology
Singapore Clinical Research Institute
Assoc Professor, Duke-NUS Graduate Medical School, Singapore
Director, Singapore Branch, Australasian Cochrane Centre;
Head (Evidence-based Medicine)
Health Services Research & Evaluation Division
Ministry of Health

This page has been intentionally left blank

This page has been intentionally left blank

This page has been intentionally left blank

Ministry of Health, Singapore

College of Medicine Building

16 College Road

Singapore 169854

TEL (65) 6325 9220

FAX (65) 6224 1677

WEB www.moh.gov.sg

ISBN 978-981-08-5211-5