SULTANATE OF OMAN

OMAN RESPIRATORY SOCIETY & OMAN FAMCO SOCIETY

In association with

MINISTRY OF HEALTH

GUIDELINES FOR THE MANAGEMENT OF ASTHMA

SECOND EDITION 2009[©]







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PREFACE

Despite the advances in understanding asthma pathophysiology and the availability of effective treatment, asthma remains a major burden worldwide to individuals, communities and health systems.

This is the second edition of the national asthma guidelines, covering adults and children. In addition it includes "Educational" and "Special Considerations" sections to assist health care providers in the management of asthma.

The first guideline was produced by the Ministry of Health in 1998. The current guideline is a combined effort by Oman Respiratory Society and Oman FAMCO Society.

We hope that all concerned will implement these guidelines to achieve better asthma control.

James ~

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PREFACE

Asthma is a major cause of chronic morbidity throughout the world and there is evidence that its prevalence has increased considerably over the past few decades, especially in children. It is a disorder that affects physical, physiological, psychological and socioeconomic functions. Because of the social disabilities that relate both directly and indirectly to the persons affected, improving health status and thereby improving quality of life are the principle goals in asthma care.

Asthma can be controlled through appropriate prevention and management strategies. This manual outlines the principles and policies of the Ministry of Health in providing asthma care services at all levels of health care.

The aim of this guideline is to provide the approaches and techniques in early detection and diagnosis and to standardize the treatment patterns among all the health care providers.

Ministry of Health strongly endorses and hereby adopts these management guidelines. In recognition to the great efforts done, The Ministry would like to express its appreciation and gratitude to Oman Respiratory Society and Oman FAMCO Society for their initiative in updating the guideline.

Dr. Ahmed bin Mohammed Al-Saidi

The Undersecretary of Health Affairs

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ABBREVIATION LIST

Abbreviation	Terminology			
AIA	Aspirin-Induced Asthma			
A&E	Accident and Emergency			
BPD	Bronchopulmonary Dysplasia			
CO ₂	Carbon Dioxide			
COPD	Chronic Obstructive Pulmonary Disease			
COX-1	Cyclooxygenase-1			
ED	Emergency Department			
EIA	Exercise-Induced Asthma			
FEV1	Forced Expiratory Volume In One Second			
FVC	Forced Vital Capacity			
FAMCO	Family and Community Medicine			
FM & PH	Family Medicine And Public Health			
GERD	Gastro Esophageal Reflux			
GINA	The Global Initiative For Asthma			
Н	Hour			
HD	High Dependency			
ICS	Inhaled Corticosteroids			
ICU	Intensive Care Unit			
IgE	Immunoglobulin E			
INH	Inhalation			
IV	Intra-Venous			
LABA	Long Acting β_2 Agonist			
Max	Maximum			
Min	Minute			
NSAIDs	Non Steroidal Anti-Inflammatory Drugs			
PaCO2	Partial Pressure of Carbon Dioxide			
PEF	Peak Expiratory Flow			
PEFM	Peak Expiratory Flow Meter			
PHC	Primary Health Care			
PICU	Pediatric Intensive Care Unit			
Pred	Predicted			
PRN	When Needed			
SABA	Short Acting β_2 . Agonists			
SaO2	Oxygen Saturation			
SQU	Sultan Qaboos University			
SQUH	Sultan Qaboos University Hospital			
VCD	Vocal Cord Dysfunction			
, 02				

INTRODUCTION

Asthma and other allergies are common chronic conditions with very high socioeconomic burden in all societies. It is estimated that asthma affects more than 300 million people worldwide and it causes approximately 240,000 deaths per year. In Oman the prevalence of asthma in children and adolescents is not only high (10% in young children and 20% in adolescents), but also is associated with severe symptoms. Although, there are no data on the prevalence of asthma in adults, its burden on patients and the society is likely to be high. Based on the findings in children, asthma is under-diagnosed, and under-treated in Oman.

Inadequate control of asthma continues to be a serious problem all over the world despite advances in understanding the inflammatory basis of asthma and well established disease management guidelines. Patients with inadequately controlled asthma are at a high risk of serious morbidity and mortality with consequent high economic cost to the population.

Primary care physicians are the main care providers for asthma patients and they can achieve complete control of asthma in the majority of their patients. However, this requires knowledge and understanding of asthma and its risk factors as well as adherence to the best practice management guidelines

PART ONE ASTHMA MANAGEMENT IN ADULTS

1. Definition of Asthma

"Asthma is a chronic inflammatory disorder of the airways in which many cells and cellular elements play a role. The chronic inflammation is associated with airway hyperresponsiveness that leads to recurrent episodes of wheezing, breathlessness, chest tightness, and coughing, particularly at night or in the early morning. These episodes are usually associated with widespread, but variable, airflow obstruction within the lung that is often reversible either spontaneously or with treatment". (*GINA 2007*)

2. When to Suspect Asthma

Breathlessness may be due to pulmonary, cardiac, hematological, metabolic or to diseases of other organs. The presence of multiple key indicators increases the probability of asthma, but Spirometry is needed to establish a diagnosis. The following factors help in differentiating asthma from other conditions:

- Wheezing: (A lack of wheezing and a normal chest examination do not exclude asthma)
- History of any of the following:
 - Cough (worse particularly at night)
 - Recurrent wheeze
 - Recurrent difficulty in breathing
 - Recurrent chest tightness
- Symptoms occur or worsen in the presence of:
 - Exercise
 - Viral infection
 - Inhalant allergens (e.g. animal fur or hair, house-dust mites, molds, pollen)
 - Irritants (tobacco or wood smoke, airborne chemicals)
 - Changes in weather
 - Strong emotional expression (laughing or crying hard)
 - Stress
 - Menstrual cycles
- Symptoms occur or worsen at night, awakening the patient from sleep

3. Medical History:

A detailed history of symptoms, aggravating factors, environmental factors, presence of the illness in the family, social aspects and patterns of exacerbation has to be taken during the first few visits. The impact of the disease on the person and the family and their perception of the disease are often needed in order to plan a proper treatment strategy.

a. Symptoms

- Coughing, Wheezing, Shortness of breath, Chest tightness and Sputum production

b. Pattern of Symptoms

- Perennial, seasonal, or both
- Continuous, episodic, or both
- Onset, duration, frequency, (number of days or nights per week or months)
- Diurnal variations, especially nocturnal and early morning

c. Precipitating and/or Aggravating Factors

- Viral respiratory infections
- Environmental allergens, indoor (e.g. mold, house-dust mite, cockroach, animal dander or secretory products) and outdoor (e.g. pollen)
- Characteristics of home including age, location, cooling and heating system, woodburning stove (Barbecue)
- Humidifiers, carpeting over concrete, presence of molds or mildew, presence of pets with fur or hair, characteristics of rooms where patient spends time (e.g. bedroom and living room with attention to bedding, floor covering, stuffed furniture)
- Smoking (patient and others at home or work)
- Exercise
- Environmental changes (e.g. moving to new home; going on vacation; and/or alterations in workplace, work processes, or materials used)
- Irritants (e.g. tobacco smoke, strong odors, air pollutants)
- Occupational chemicals, dusts and particulates, vapors, gases, and aerosols
- Emotions (e.g. stress, crying or laughing hard, fear, anger, frustration)
- Incense, perfume and *bukhoor*
- Drugs (e.g. aspirin and other nonsteroidal anti-inflammatory drugs, beta-blockers including eye drops)
- Food, food additives and preservatives (e.g. sulfites)
- Changes in weather, or exposure to cold air

- Endocrine factors (e.g. menses, pregnancy, thyroid disease)
- Comorbid conditions (e.g. sinusitis, rhinitis, gastroesophageal reflux disease GERD)

d. Family History

- History of asthma, allergy, sinusitis, rhinitis, eczema, or nasal polyps in close relatives

e. Disease Development and Treatment

- Age of onset and diagnosis
- History of early-life injury to the airway (e.g. parental smoking, bronchopulmonary dysplasia, pneumonia)
- Progression of the disease (better or worse)
- Current treatment and response, including plans for the management of exacerbations
- Frequency of short acting β 2-agonist use
- The need for oral corticosteroid and frequency of use

f. Social History

- Daycare, workplace and school characteristics that interfere with adherence to treatment
- Social factors that interfere with adherence, such as substance abuse
- Social support/social network
- Level of education
- Work / job

g. History of Exacerbations

- Usual prodromal signs and symptoms
- Rapidity of onset
- Duration
- Number and severity of exacerbations in the past years
- Life-threatening exacerbations (e.g. intubation, intensive care unit admission)
- Usual patterns and management

h. Perception of the Disease

- Patient's perception and beliefs regarding use and long term effects of medications
- Family knowledge and beliefs of asthma

i. Impact of Asthma on Patient and Family

- Episodes of unscheduled care (emergency department (ED), urgent care, hospitalization)
- Number of days missed from school/work
- Limitation of activity, especially sports and strenuous work
- History of nocturnal awakening
- Effect on growth, development, behavior, school or work performance, and lifestyle
- Impact on family routines, activities, or dynamics
- Economic impact

4. Examination

- Height and weight (assessment of growth), nose, throat and paranasal sinuses (nasal polyps may be associated with asthma)
- Features of atopy
- A good general examination and recording of the vital signs including the respiratory rate
- Auscultation of the chest can be normal or one can hear a prolonged expiratory phase or diffuse wheezing

5. Investigations:

The medical history and physical examination are not always reliable in excluding other diagnoses or in assessing the lung status.

a. Peak Expiratory Flow (PEF)

Peak flow measurement is effort dependant and requires patient understanding. In certain patients measuring PEF prior to and after a bronchodilator or an exercise challenge may help in confirming the diagnosis.

Measurement of PEF variability by comparing the morning and evening PEF over a period of 2 weeks is also helpful. Variability over 20% supports the diagnosis. It is calculated as follows:

Mean peak flow variability $\% = (highest - lowest) \times 100$ highest

b. Spirometry:

Spirometry is generally recommended, rather than measurements by a peak flow meter, due to wide variability in peak flow rate and reference values. The absence of airway obstruction does not rule out asthma, especially if the patient is not symptomatic at the time of testing, since airflow obstruction in asthma is, by definition, intermittent (Table1).

Table 1: Interpreting Spirometry

Diagnosis	FEV ₁	FVC	FEV ₁ /FVC%	
Obstructive disease	Decreased	Normal or Decreased	Decreased	
Restrictive disease	Decreased or normal	Decreased	Normal or increased	
Reversible airway obstructionIncrease of >12 % of the base line value after administration of bronchodilator				
FEV_1 = Forced Expiratory Volume in One Second ,FVC = Forced Vital Capacity , FEV ₁ /FVC% = FEV ₁ as Percentage of FVC				

c. Additional Tests:

Additional pulmonary function studies like diffusing capacity, lung volumes or bronchial challenges studies may be needed to complete the evaluation. Other tests, which are routinely not necessary but may be useful, are:

- Chest X Ray
- Absolute eosinophil count
- IgE
- Exhaled Nitric oxide
- Sputum eosinophils

6. Differential Diagnoses:

Not all patients who wheeze have asthma.

Diagnosis has to be reassessed if the patient is not responding to conventional treatment. Some of the alternate diagnoses to be considered are:

- Chronic obstructive pulmonary disease (COPD)
- Aspiration
- Mechanical obstruction of the airways (benign or malignant tumors)
- Pulmonary infiltration with eosinophils
- Congestive heart failure
- Pulmonary embolism
- Cough secondary to drugs (e.g. angiotensin-converting enzyme inhibitors, and beta blockers)
- Foreign body in trachea or bronchus
- Vocal cord dysfunction (VCD)
- Vascular rings or laryngeal webs

<u>1. ASSESSING SEVERITY</u>

Classification of asthma by severity based on the level of symptoms, airflow limitation, and lung function is useful when decisions are being made about management at the initial assessment of a patient

A. Intermittent Asthma:

a. Clinical Features:

- Intermittent symptoms less than once a week
- Nocturnal symptoms not more than 2 times a month
- Asymptomatic in between attacks

b. Peak flow measurement:

- FEV1 or PEF \geq 80% predicted
- FEV1 or PEF variability < 20%

B. Mild Persistent Asthma:

a. Clinical Features:

- Mild, brief symptoms, more than once per week, but less than once a day
- Nocturnal symptoms more than twice a month
- Asymptomatic between exacerbations
- Exacerbations may affect activity and sleep

b. Peak Flow Measurement:

- FEV1 or PEF \geq 80% predicted
- FEV1or PEF variability < 20-30%

C. Moderate Persistent Asthma:

a. Clinical Features:

- Symptoms daily
- Exacerbations may affect activity and sleep
- Nocturnal asthma symptoms more than once a week
- Daily use of inhaled β2-agonist

b. Peak Flow Measurement:

- FEV1or PEF 60-80% predicted
- FEV1or PEF variability > 30%

D. Severe Persistent Asthma

a. Clinical Features:

- Frequent exacerbations
- Daily symptoms
- Frequent nocturnal symptoms
- Limited physical activities

b. Peak Flow Measurement:

- FEV1or PEF $\leq 60\%$ predicted
- FEV1or PEF variability >30%

2. ASSESSING CONTROL

Asthma severity can change over time or with treatment. The classification of asthma severity based on the level of control achieved with treatment is the best way to take ongoing treatment decisions. Table 2 lists the characteristics of Controlled, Partly controlled and Uncontrolled asthma. Asthma control can also be assessed by using a validated questionnaire like Asthma Control Test (Appendix 1)

Characteristics	Characteristics Controlled		Uncontrolled
Day time symptomsNone or ≤ 2 times/week		> 2 times /week	\geq 3 features of partly controlled asthma present in any week
Activity limitation None		Any	
Night symptoms None		Any	
Salbutamol use	None or ≤ 2 times/week	> 2 times / week	
PEF or FEV1 Normal		< 80 %	
Exacerbations None		$\geq 1/$ year	One in any week

Table 2: Levels of Asthma Control

SECTION 3:

The aim of treatment is to achieve complete control of asthma symptoms and maintain it for prolonged periods. Over a period of time the lowest dose and the amount of medications have to be identified to keep the patient symptom free.

<u>1. TREATMENT STEPS</u>

The principle of step wise approach to treatment is to increase the dose, the frequency or the number of medications if needed or to decrease the same when possible to achieve and maintain control (Figure 1).

- During each visit the patient's present treatment, compliance, inhaler technique and the level of asthma control achieved with current treatment should be assessed
- If not controlled go to the next higher step as discussed below
- If partly controlled, weigh the risk and benefit of further treatment option and how far control can be achieved in a particular patient
- If adequate control is achieved and maintained for 3 months, bring down the treatment to the next level
- For very mild intermittent asthma one can begin with **Step 1**
- Patients with persistent symptoms may need **Step 2** care to begin with
- More severe symptoms may need **Step 3 or Step 4** medications initially

STEP 1: AS NEEDED RELIEVER MEDICATIONS

This treatment is useful for patients with very mild symptoms like cough or wheezing of short duration occurring twice or less per week. This step is also recommended as the initial treatment option for patients with bronchospasm on exercise

Recommended: A short-acting inhaled $\beta 2$ agonist (e.g. Salbutamol inhaler) as required

STEP 2: RELIEVER MEDICATION PLUS A SINGLE CONTROLLER

A reliever medication (e.g. Salbutamol) should be used as required

- **Recommended:** A low-dose inhaled corticosteroid is recommended as the initial controller treatment for asthma patients (dose schedule of inhaled corticosteroid is given in Table 3).
- Alternative: Oral leukotriene modifiers (e.g. Montelukast 10mg daily) can be advised especially in patients refusing or unable to use inhaled corticosteroid or when they experience local side effects.

STEP 3: RELIEVER MEDICATION PLUS ONE OR TWO CONTROLLERS

A reliever medication (e.g. Salbutamol) should be used as required

- **Recommended:** A low-dose inhaled corticosteroid with an inhaled long acting β 2agonist is recommended either in a combination inhaler device or as separate components (dose schedule of inhaled corticosteroid is given in Table 3).
- Alternative: Increase to medium dose inhaled corticosteroid (dose schedule of inhaled corticosteroid is given in (Table 3). If this dose is given as an inhaler, the use of a spacer device is recommended.
- Others: Combine low-dose inhaled corticosteroid with oral leukotriene modifiers (e.g. Montelukast 10mg daily) or a sustained release Theophylline.

STEP 4: RELIEVER MEDICATION PLUS TWO OR MORE CONTROLLERS

A reliever medication (e.g. Salbutamol) should be used as required

- **Recommended:** A medium or high dose inhaled corticosteroid with an inhaled long acting $\beta 2$ agonist (LABA) is recommended either in a combination inhaler device or as separate components (dose schedule of inhaled corticosteroid is given in Table 3).
- Alternative: Add a third controller, oral leukotriene modifiers (e.g. Montelukast 10mg daily) or a sustained release Theophylline to the medium or high dose inhaled corticosteroid with the inhaled long acting $\beta 2$ agonist.

STEP 5: RELIEVER MEDICATION PLUS ADDITIONAL CONTROLLERS

This is recommended only when all the measures like environmental control, treatment of precipitating problems like post nasal drip, gastro esophageal reflux ...etc fail and the diagnosis of asthma is properly established.

A reliever medication (e.g. Salbutamol) should be used as required

- Recommended:Addition of oral corticosteroid to the combination of the high dose
inhaled corticosteroid, inhaled long acting β2-agonist and leukotriene
modifiers (e.g. Montelukast 10mg) or a sustained release Theophylline
is recommended when asthma remains totally uncontrolled.
- Alternative:Addition of anti-IgE treatment to other controller medications has been
found to be of use in allergic asthma when oral corticosteroid fails.
Anti-IgE treatment is better reserved for specialists in the referral
centers and should not be considered by practitioners in general clinics.

Table 3: Equipotent Daily Doses of Inhaled Corticosteroid for Adults

Drug	Drug Low Daily Dose		High daily Dose
Beclomethasone	200 - 500µg	>500 -1000µg	> 1000 - 2000µg
Budesonide	200 - 400µg	>400 - 800µg	> 800 - 1600µg
Fluticasone	100 - 250µg	>250 - 500µg	> 500 - 1000µg

2. STEPPING DOWN

- Improvement may be noticed within days of starting controller medications. But the full benefit may be seen only after 3 to 4 months
- Once the physician is satisfied on the control obtained, stepping down the treatment can be planned after discussing with the patient. The preferred options are as follows:

CURRENT TREATMENT	STEPPING DOWN PLAN
Medium- high dose inhaled steroids	50% reduction in dose
Inhaled steroids + LABA	50% reduction in dose of inhaled steroids while continuing LABA. Further reduction to reach low dose and then to discontinue LABA OR Discontinue LABA while continuing high dose inhaled steroids
Low dose inhaled steroids	Switch to once daily dosing
Once daily low dose inhaled steroids	Controller treatment may be stopped if asthma remains controlled on a low dose of controller and there were no recurrences for one year

Table 4: Stepping Down Options

FIGURE 1: STEP UP CARE OF ASTHMA IN ADULTS

				STEP 5
			STEP 4	Inhaled Salbutamol prn
		STEP 3	Inhaled Salbutamol prn	RECOMMENDED THERAPY
	STEP 2	Inhaled Salbutamol prn	RECOMMENDED THERAPY	Add oral steroids to
	Inholod Solbutomol pro	RECOMMENDED THERAPY	Medium to high dose ICS + High dose ICS + L_{A}	High dose ICS + LABA + Montelukast
STEP 1	Inhaled Salbutamol prn	Low dose ICS + Long Acting B2 Agosists (LABA)	- Beclomethasone 1000 - 2000µg/day	/ Theophylline
	RECOMMENDED THERAPY	- Salmeterol 50µg BD	Or Budesonide 800 -1600µg/day Or Fluticasone >500 -1000µg/day Alternative options Add a third controller + Montelukast OR oral Theophylline	Alternate options
		Or Formeterol 4.5 - 12µg BD		Addition of IgE treatment
RECOMMENDED THERAPY	Low dose inhaled Corticosteroids (ICS)	Alternative options		to other controllers
Inhaled Salbutamol prn	- Beclomethasone 200 - 500μg/day Or Budesonide 200 - 400μg/day Or Fluticasone 100 - 250μg/day	y - Beclomethasone 500 -1000μg/day Or Budesonide 400 - 800μg/day		
	Alternate options	OTHERS		
	Montelukast 10µg daily	Low dose ICS + Montelukast OR oral Theophylline		
			1	

1. ASSESSMENT

Exacerbations of asthma are episodes of progressive increase in shortness of breath, coughing, wheezing, chest tightness or a combination of these symptoms. Severe exacerbations are potentially life threatening and so have to be identified early and closely observed. The aim of treatment is to relieve the airway obstruction, correct the hypoxia and prevent further worsening.

- Treatment is to be initiated as soon as an asthma attack is recognised, even before completing a full history or physical examination
- The history should include disease severity, duration of symptoms, current medications, the doses taken after the exacerbations, the response to the treatment and the risk factors for asthma related deaths (Table 5)
- The focus of evaluation should be on determining the severity of the attack (Table 6), identifying the precipitating factors and considering an alternate diagnosis
- Physical examination should start with a rapid evaluation of the patient's general appearance including anxiety level, mental status and the level of respiratory distress. Altered mental status, severely laboured breathing and accessory muscle use are signs of an impending respiratory failure
- PEF should be measured in all patients

Disease related	Psychosocial factors	
Previous near fatal asthma (ICU admissions)	Non-compliance with treatment	
Previous asthma admissions	Failure to attend appointments	
Requiring \geq 3 classes of asthma medications	Self-discharge from hospital	
Recent heavy use of Salbutamol	Psychosis, depression	
Repeated attendances at A&E	Alcohol or drug abuse	
	Social, employment, income problems	

	MILD	MODERATE	SEVERE
Breathlessness	While walking	While talking	While at rest
	Can lie down	Prefers sitting	Hunched forward
Talks in	Sentences	Phrases	Words
Alertness	May be agitated	Usually agitated	Usually agitated
Respiratory rate	Increased	Increased	Often >30/minute
Accessory muscles use	Usually not	Usually	Usually
Wheeze	Moderate often end expiratory only	Loud	Loud
Pulse rate (/ minute)	< 100	100 -120	>120
PEF	> 80%	60 - 80%	< 60%
SaO ₂ (room air)	> 95%	91-95%	< 91%

Table 6: Levels of Severity of Asthma Exacerbations

Life Threatening Asthma

Drowsiness or confusion, cyanosis feeble respiration, bradycardia, hypotension inaudiable breath sounds (silent chest), paradoxical thoracoabdominal movement and a PEF <33% all indicates a life threatening attack. PaCO₂ is usually low in acute severe asthma due to CO₂ washout, so a normal PaCO₂ or a raised PaCO₂ indicates severity.

2. TREATMENT

- Put the patient in a comfortable sitting position, legs down if possible, so that he/she can bend forward if needed and have support for the hands and legs
- The patient should not be left alone until the condition has clearly improved.
- a. Oxygen
 - Correct hypoxia urgently using high concentrations of oxygen to maintain SaO2 around 92% at least
 - Oxygen driven nebulizers preferred to air driven

b. Nebulized Salbutamol

- Repeated doses:
- 2.5 to 5 mg every 20 minutes x 3 doses
- Then 2.5 to 10 mg every 1 to 4 hours (depends on the response)

c. Nebulized Ipratropium Bromide

- 0.5mg every 20 minutes x 3 doses, then 0.5mg every 6 hours
- Combining nebulized Ipratropium with β2-agonist produce significantly greater bronchodilatation than β2-agonist alone

d. Corticosteroids

- IV Hydrocortisone 100mg immediately and then every 6 hours
 OR IV Methyl prednisolone 125mg immediately and 60mg every 8 hours
- Should be changed to oral prednisolone once the patient improves

e. Magnesium Sulphate

- Only in hospital where monitoring facilities are available
- Single dose of 1.2 to 2 grams as IV infusion over 20 minutes
- Contra indicated in renal insufficiency
- Acts by blocking calcium influx thus preventing bronchoconstriction

f. Others

- Antibiotics: not routinely required
- IV Fluids may be needed for some patients
- Cough syrup: No definite role in the management of an exacerbation
- Aminophylline is not recommended for regular use

SECTION 4:

3. ADMISSION TO INTENSIVE CARE UNIT (ICU)

In certain situations (see below), the patient should be admitted to the ICU for close observation and monitoring.

- Rapidly worsening asthma or a lack of response to the initial therapy in the emergency department
- If patients have confusion, drowsiness, signs of impeding respiratory arrest, or loss of consciousness
- Impending respiratory arrest, as indicated by hypoxemia (PO₂ < 60mmHg) despite supplemental oxygen and/or hypercarbia (PCO₂ > 45mmHg)
- If intubation is required because of the continued deterioration of the patient's condition despite optimal treatment

4. DISCHARGE

Acute asthma relapse after discharge needs to be prevented. Many patients with asthma have contact with the health care only during exacerbations. So on discharge the attending doctor has to ensure the following:

- Review within 48 hours in the primary health care centre for patients discharged from Emergency department
- Monitor symptoms and PEF
- Check inhaler technique
- Provide written asthma action plan
- Modify treatment according to guidelines for chronic persistent asthma (ICS, inhaled LABA)
- Address factors that could have contributed to the present admission

5. MANAGEMENT OF EXACERBATIONS

MILD SYMPTOMS

- Oxygen + Nebulized Salbutamol every 20 minutes+ Prednisolone 40mg stat
- REASSESS AFTER ONE HOUR (Physical examination, PEF, SaO₂)
- Improved: Discharge on a five to seven day course of oral prednisolone, and their regular preventer and controller inhalers

MODERATE SYMPTOMS

- Oxygen + Nebulized Salbutamol + Nebulized Ipratropium bromide every 20 minutes + Hydrocortisone 100mg IV
- REASSESS AFTER ONE HOUR (Physical examination, PEF, SaO₂)
 - Improved: Continue nebulised Salbutamol every 30 minutes or one hour and reassess after three hours
 - If not improving: Give IV Magnesium Sulphate and continue as above
- Improved- Reassess and consider discharge
- Not improving- admit for continuation of treatment

SEVERE SYMPTOMS

- Oxygen + Nebulized Salbutamol + Nebulized Ipratropium bromide every 20 minutes + Hydrocortisone 100mg IV
- If not improving give IV Magnesium Sulphate
- CONTINUOUS ASSESSMENT
 - Improved: Admit for continuation of treatment
 - o If not improving: Admit to ICU for further treatment

SECTION 5:

1. INTRODUCTION

Proper management of asthma starts from the primary health care (PHC). PHC physicians should manage intermittent, mild persistent and moderate persistent asthmatics requiring PRN short acting $\beta 2$ agonist, inhaled steroids and long acting $\beta 2$ agonist.

2. DIAGNOSIS:

- Detailed history should include allergy, drugs, occupational, environmental and family history
- Full physical examination should include sinuses, nose, throat, skin and respiratory system

3. INVESTIGATIONS:

- Peak expiratory flow measurement before and after inhaled short acting $\beta 2$ agonist should be performed
- Complete Blood Count
- Chest X Ray
- Spirometry

4. MANAGEMENT:

- Assess severity
- Assess control
- Manage as outlined in Section 4
- Check inhaler technique
- Educate the patient and the family

5. FOLLOW UP:

- Periodic follow up, at least 4-6 times a year should be arranged at the Primary Care Centre
- Inhaler technique (Appendix 6), Peak expiratory flow (Appendix 8) and Compliance should be checked at each visit

6. WHEN TO REFER TO SPECIALIST CLINIC

Consider referral to specialist if:

- Signs and symptoms are atypical
- Diagnosis is uncertain
- The patient is not responding to therapy
- The patient developed side effects from medications
- The patient requires frequent courses of oral corticosteroid
- The patient has occupational asthma
- Abnormal lung function persists when the symptoms are apparently controlled
- Achieving control is difficult
- Necessary asthma medications or spacer devices are unavailable
- Additional testing is indicated

7. WHEN TO REFER ASTHMA EXACERBATION TO SECONDARY CARE

- Patients with severe or life threatening asthma exacerbation
- Persistent severe dyspnoea despite short acting β2 agonist given repeatedly 2-3 times at 20-30 minutes intervals
- The symptoms worsen while on management
- Frequent or persistent exacerbations
- Lack of family monitoring and social support at home
- Patients from far away places, who have minimum access to health centers

<u>1. EXERCISE-INDUCED ASTHMA (EIA):</u>

- EIA is a problem among active children, adolescents, and young adults
- A history of wheezing, fatigue or poor performance on exercise may be the presentation. Physical examination at the time of symptoms may be helpful. Pulmonary function or exercise test may be needed often
- Patient should be advised to avoid exercise in cold weather, in places when pollen or air pollution levels are high and to do proper warm-ups before vigorous exercise
- Taking short acting $\beta 2$ agonist 20-30 minutes before exercise is recommended
- Leukotriene modifiers should be considered in the management of EIA

2. PREGNANCY

- In approximately one-third of women asthma becomes worse; in one-third asthma becomes less severe; and in the remaining one-third it remains unchanged during pregnancy
- Poorly controlled asthma can have an adverse effect on the fetus, resulting in increased perinatal mortality, increased prematurity, and low birth weight
- There is no evidence to suggest an increased risk to the fetus for most medications used to treatment of asthma
- Inhaled corticosteroids, $\beta 2$ agonists, leukotriene modifiers (e.g. Montelukast) and appropriately monitored use of Theophylline are completely safe in pregnancy
- As in other situations, the focus of asthma treatment must remain on control of symptoms and maintenance of normal lung function
- Acute exacerbations should be treated aggressively in order to avoid fetal hypoxia. Treatment should include nebulized β2 agonists and oxygen. Systemic corticosteroid should be instituted when necessary
- Pregnant patients with asthma should be advised that the greater risk to their baby lies with poorly controlled asthma, and all the present drugs are safe in pregnancy

3. RESPIRATORY INFECTIONS

- Respiratory infections have an important relationship to asthma as they provoke wheezing and increased symptoms in many patients. Microorganisms associated with increased asthma symptoms are often respiratory viruses
- Treatment of an infectious exacerbation follows the same principles as treatment of other asthma exacerbations. Because increased asthma symptoms can often persist for weeks after the infection is cleared, anti-inflammatory treatment should be continued

4. SURGERY

- Airway hyperresponsiveness, airflow limitation, and mucus hypersecretion predispose patients with asthma to intraoperative and postoperative respiratory complications
- The likelihood of these complications depends on the severity of asthma at the time of surgery and the type of surgery
- Pulmonary function should be measured. If the patients FEV1 is less than 80% of personal best, a brief course of oral corticosteroid should be considered
- Patients who have received systemic corticosteroid within the past 6 months should have systemic coverage during the surgical period (100mg hydrocortisone every 8 hours intravenously). This should be rapidly reduced 24 hours following surgery, as prolonged systemic corticosteroid therapy may inhibit wound healing

5. RHINITIS/SINUSITIS/NASAL POLYPS

- Many patients with asthma have a history of rhinitis and up to 30% of patients with persistent rhinitis have or develop asthma
- Treatment of Rhinitis may improve asthma symptoms. Intra-nasal corticosteroid as well as leukotriene modifiers, anticholinergics, allergen-specific immunotherapy, and anti-IgE therapy can be effective in both conditions
- Sinusitis is a complication of upper respiratory infections, allergic rhinitis, nasal polyps, and other forms of nasal obstruction. Both acute and chronic sinusitis can worsen asthma. Treatment should include topical nasal decongestants or topical nasal or even systemic corticosteroid and a 10 day course of antibiotics
- Nasal polyps associated with asthma and rhinitis, and sometimes with aspirin hypersensitivity, are seen primarily in patients over 40 years old. Nasal polyps are quite responsive to topical corticosteroid. A limited number of patients with corticosteroid-refractory polyps may benefit from surgery

6. OCCUPATIONAL ASTHMA

- Once a diagnosis of occupational asthma is established, complete avoidance of the relevant exposure is advisable
- Occupational asthma may persist for several years after removal from exposure to the causative agent, especially when the patient has had symptoms for a long time before cessation of exposure
- Pharmacologic therapy for occupational asthma is identical to therapy for other forms of asthma, but it is not a substitute for adequate avoidance

7. GASTROESOPHAGEAL REFLUX

- The relationship of increased asthma symptoms, particularly at night, to gastroesophageal reflux remains uncertain
- A diagnosis of gastroesophageal reflux in patients with asthma can best be made by simultaneously monitoring esophageal pH and lung function
- Medical management should be given for the relief of reflux symptoms as it is often effective. Patients may be advised to eat smaller, more frequent meals; avoid food or drink between meals and especially at bedtime; avoid fatty meals, alcohol and Theophylline

8. ASPIRIN-INDUCED ASTHMA (AIA)

- Up to 28% of adults with asthma, but rarely children, suffer from asthma exacerbations in response to aspirin and other non steroidal anti-inflammatory drugs (NSAIDs)
- The majority of patients first experience symptoms, which may include vasomotor rhinitis and profuse rhinorrhea, during the third to fourth decade of life. Asthma and hypersensitivity to aspirin often develop subsequently
- Patients with AIA should avoid aspirin, products containing it, other analgesics that inhibit cyclooxygenase-1 (COX-1). Where an NSAID is indicated, a COX-2 inhibitor may be considered with appropriate physician supervision and observation for at least one hour after administration

<u>1. MANAGING ASTHMA DURING TRAVEL AND HOLIDAYS</u></u>

- Advise the patient to consult their doctor before they travel
- Give a list of medications and explain the role of each
- Provide them with an Asthma Action Plan (Appendix 3 & Appendix 4)
- Advise them to take enough medications to last the whole trip
- Remind them to continue all medications especially preventers

2. PEAK EXPIRATORY FLOW METER (PEFM)

PEF is the fastest rate at which air can move through the airways during a forced expiration starting with fully inflated lungs, measured by a device called Peak Flow Meter.

- Can be used not only in hospital and clinic settings but also in home and office to help assess asthma severity, and evaluate response to therapy
- Can serve as early warning systems because a change in measurement from a patient's normal can signal trouble sufficiently early to take action to prevent an attack
- Healthy persons have good PEF but it gets reduced in presence of asthma and the reduction is proportional to the severity of asthma
- PEF is assessed by comparing it with the patients' personal best or with the predicted PEF value (Appendix 8)
- The predicted values are provided with all peak flow meters and are adjusted for height, race, sex, and age
- Personal Best Value: is the highest peak flow rate measured during a 2 week period when asthma is under control, or the highest reading of PEF in a year period (Appendix 9)
- Asthmatic patient should be trained on how to use PEFM, instructed how to find personal best and how to use the PEF reading (Appendix 7)
- PEF measurement should be performed at each visit and recorded for comparison during follows up visits
3. ASTHMA ACTION PLAN

An asthma action plan is a written plan developed by doctors, health educator or specialized nurses to help patients in the management of asthma episodes. It is a customized plan that tells the patient what to do based on changes of symptoms and peak flow readings.

A simple, easy-to-use asthma action plan uses the traffic light analogy as it relates to symptoms and daily peak flow monitoring. The green zone is "go" the yellow zone is "caution" and the red zone is "danger" (Appendices 3 & 4). Any asthma action plan should include at least the following information:

- A list of peak flow meter readings and zones based on personal best
- A list of routine symptoms such as coughing, wheezing, tightness in the chest, shortness of breath, and excess mucus production, and what they should do if these symptoms occur
- The name and dose of the preventer medication, the name and dose of the reliever medication
- Emergency telephone numbers and locations of emergency care
- Instructions about when to contact the doctor, and when to go directly to the hospital or health center for emergency

4. SPECIAL INSTRUCTIONS

- To select the best inhaler device for the patient, first evaluate his/her cognitive, inhalation, physical and visual abilities. Check selection criteria for inhaler devices (Appendix 5)
- When carer availability is essential in the management, make sure that they understand their role fully, and provide them with all necessary information
- To prevent inhaled corticosteroids and propellant side effects, advise the patient to gargle with tap water immediately after taking the inhaled medication and to use a large volume spacer
- Instruct patient how to clean and take care of their spacer devices
- Advise patient to keep all inhaled medications at room temperature
- Provide patients with a list of the names, dose and frequency for all prescribed asthma medications

5. QUALITY MANAGEMENT IN ASTHMA:

Centers which manage asthma patients in various departments such as PHC, Emergency department, In-Patient or Out- Patients specialty clinics, should have a system to maintain and assess asthma management according to the standards of care. This would include assessment, management and follow up patients in a structured process. Oman Respiratory Society Asthma Management Protocol (pages 63-66) is developed to guide health care providers in managing asthma and to improve the quality of care provided to patients. The protocol can also be use to evaluate and audit the management.

PART TWO

ASTHMA MANAGEMENT IN CHILDREN

<u>1. DIAGNOSIS</u>

Diagnosis of asthma in children especially those younger than four years old can be difficult and has important implications. Many are under-diagnosed and, therefore, do not receive adequate therapy. Similar to adults, assessment and monitoring of asthma in children should include history of symptoms, physical examination and assessment of quality of life. Other alternative diagnoses must be excluded.

A. Asthma Diagnosis in Children 0-4 Years of Age

Consider asthma in infants and young children who have four or more episodes of wheezing in the past year that lasted more than 1 day and affected sleep, <u>AND</u> who has either;

a. One of the following:

- A physician's diagnosis of atopic dermatitis
- Evidence of sensitization to aeroallergens
- A parental history of asthma

<u>OR</u>

- b. Two of the following:
 - Evidence of sensitization to foods
 - $\geq 4\%$ peripheral blood eosinophilia
 - Wheezing apart from colds

B. Asthma Diagnosis in Children 5-12 Years of Age

Consider asthma if any of the following signs or symptoms is present:

a. History of any of the following:

- Cough, worse particularly at night
- Recurrent wheeze
- Recurrent difficulty in breathing
- Recurrent chest tightness

b. Symptoms occur or worse at night

c. Symptoms occur or worse in the presence of:

- Exercise Animals with fur
- Pollens Aerosol chemicals
- Smoke Changes in temperature
- House dust mites
- Drugs (aspirin, beta blockers)
- Strong emotions
- Respiratory (viral) infections

d. Reversible and variable airflow limitation as measured by a spirometer or a peak expiratory flow meter

2. INVESTIGATIONS:

- Spirometry and PEF can be done in child above 5 years of age as in adults
- Chest X-Ray may be done to exclude any other causes in atypical presentation
- Review also asthma investigations in adult (page 15 -16)

3. DIFFERENTIAL DIAGNOSIS:

Other causes of episodic or chronic wheeze, cough and breathlessness in children include:

- Cystic Fibrosis
- Primary immune deficiency
- Tracheomalacia
- Congenital heart disease
- Vascular rings
- Foreign bodies
- Chronic rhino-sinusitis
- Gastro-oesophageal reflux
- Recurrent lower respiratory tract infections
- Congenital malformations causing narrowing of the intrathoracic airways

1. CLASSIFICATION OF ASTHMA SEVERITY

- a. Children age 0 4 years (Table 7)
- b. Children age 5 12 years (Table 8)

Table 7: Classification of Asthma Severity and Therapy in Children 0-4 Years of Age

INTERMITTENT ASTHMA		MILE) PERSISTENT	MODERA	TE PERSISTENT	SEVERE PERSISTENT		
COMPONENTS OF SEVERITY	FREQUENCY	COMPONENTS OF SEVERITY	FREQUENCY	COMPONENTS OF SEVERITY	FREQUENCY	COMPONENTS OF SEVERITY	FREQUENCY	
Symptoms	\leq 2 days/week	Symptoms	> 2 days / week but not daily	Symptoms	Daily	Symptoms	Throughout the day	
Night-time awakenings	\leq 2 days/month	Night-time awakenings	1-2 / month	Nigh time awakenings	3-4 / month	Nigh time awakenings	> 1 time/week	
SABA used for symptom control (not EIA)	\leq 2 days/week	SABA used for symptom control (not EIA)	> 2 days/week but not daily	SABA used for symptom control (not EIA)	Daily	SABA used for symptom control (not EIA)	Several times / day	
Interference with normal activity	None	Interference with normal activity	Minor	Interference with normal activity	Some limitation	Interference with normal activity	Extremely limited	
Exacerbations	0 -1 year	Exacerbations	≥ 2 in 6 months requiring systemic corticosteroids, <u>or</u> ≥ 4 wheezing episodes/1 year lasting >1 day AND risk factors for persistent asthma	Exacerbations	 ≥2 in 6 months requiring systemic corticosteroids, or ≥4 wheezing episodes/1 year lasting >1 day AND risk factors for persistent asthma 	Exacerbations	≥ 2 in 6 months requiring systemic corticosteroids, <u>or</u> ≥ 4 wheezing episodes/1 year lasting >1 day AND risk factors for persistent asthma	
<u>RECOMMEND</u> Salbutamol MDI wit prn		RECOMM Salbutamol MDI with <u>Give one of the follow</u> - Beclomethasone MI -Fluticasone MDI 100 -Budesonide (neb) 2 <u>or</u> Montelukast oraly	<u>ving medications;</u> DI 100 - 200 μg/day,) - 200 μg/day, 50-500 μg/ day,	RECOMMEN Salbutamol MDI with space Give one of the following - Beclomethasone MDI 200 - Fluticasone MDI 200 - 5 or Budesonide (neb) 500 -	<u>medications;</u>)0-400 μg/day, 00 μg/day	<u>RECOMME</u> Consult asthma specia Salbutamol MDI with a <u>Give one of the followi</u> - Beclomethasone MDI - Fluticasone MDI 200 or Budesonide (neb) And add Montelukast	spacer; 2 puffs prn. ng medications; I 200 - 400 μg/day,) - 500 μg/day 500-1000 μg/ day,	

Table 8: Classification of Asthma Severity and Therapy in Children 5-12 Years of Age

INTERMITTENT ASTHMA		MILC) PERSISTENT	MODER	ATE PERSISTENT	SEVERE PERSISTENT		
COMPONENTS OF SEVERITY	FREQUENCY	COMPONENTS OF SEVERITY	FREQUENCY	COMPONENTS OF SEVERITY	FREQUENCY	COMPONENTS OF SEVERITY	FREQUENCY	
Symptoms	\leq 2 days / week	Symptoms	> 2 days / week but not daily	Symptoms	>2 days/week but not daily	Symptoms	Throughout the day	
Night-time awakenings	\leq 2 days / month	Night-time awakenings	3-4/month	Nigh time awakenings	>1 /week but not nightly	Nigh time awakenings	7/week	
SABA used for symptom control (not EIA)	\leq 2 days / week	SABA used for symptom control (not EIA)	> 2 days/week but not daily	SABA used for symptom control (not EIA)	Daily	SABA used for symptom control (not EIA)	Several times / day	
Interference with normal activity	None	Interference with normal activity	Minor Limitation	Interference with normal activity	Some limitation	Interference with normal activity	Extremely limited	
Exacerbations	0-1 / year	Exacerbations	\geq 2 / year	Exacerbations	≥2 exacerbations/ year	Exacerbations	\geq 2 exacerbations/ year	
Lung Function	FEV1 > 80% predicted FEV1/FVC > 85%	Lung Function	FEV1 ≥ 80% predicted FEV1/FVC > 85%	Lung Function	FEV1 = 60- 80% predicted FEV1/FVC =75-85%	Lung Function	FEV1< 60% predicted FEV1/FVC < 75%	
RECOMMEN Salbutamol MDI w prn	DED THERAPY	RECOMM Salbutamol MDI with <u>Give one of the follow</u> Beclomethasone MD - Fluticasone MDI 10 - Budesonide (neb) 2 or Montelukast oraly	ving medications; I 100-200 μg/day, 00-200 μg/day, 50-500 μg/ day		cer; 2 puffs prn <u>he followings;</u>)0-400 μg/day,)0 μg/day, 1000 μg/ day followings;)0-200 μg/day,) μg/day,	Salbutamol MDI with Consult asthma spec Give one of the follor - Beclomethasone M or Fluticasone MDI or Budesonide (nel	cialist <u>wings:</u> IDI 200-400 μg/day 200 -500 μg/day	

2. GOALS OF ASTHMA THERAPY

The goal of asthma therapy is to maintain long-term control of asthma with the least amount of medication. The Initiation and adjustment of asthma therapies is to achieve the followings:

- Maintain normal activity levels and exercise
- Maintain near-normal pulmonary function
- Prevent acute episodes of asthma
- Minimize emergency department visits and hospitalizations
- Avoid adverse effects of asthma medications

<u>3. PRINCIPLES OF STEPWISE THERAPY IN CHILDREN (Tables 7&8)</u></u>

- Use stepwise approach to therapy, in which the dose and number of medications and frequency of administration are increased as necessary and decreased when possible to achieve and maintain control
- The level of impairment generally is judged on the most severe measure
- Regular follow up 1- 6 months interval is essential, depending on the level of control.
- Once well-controlled asthma is achieved and maintained for 3 months, a step-down on pharmacological therapy is recommended
- Treatment of young children is often in the form of a therapeutic trial; therefore, it is essential to monitor the child's response to therapy. If there is no clear response within 4–6 weeks, the therapy should be discontinued and alternative therapies or alternative diagnoses considered

4. INHALED CORTICOSTEROIDS (ICS) IN CHILDREN

ICS are the preferred therapy for initiating long-term control therapy in children of all ages.

- ICS, especially at low doses and even for extended periods of time, are generally safe
- The potential for the adverse effect of low- to medium-dose ICS on linear growth is usually limited to a small reduction in growth velocity, approximately 1cm in the first year of treatment that is generally not progressive over time. Children receiving ICS should be monitored, by using a stadiometer, for changes in growth
- The potential risks of ICS are well balanced by their benefits

Table 9: Equipotent Daily Doses of ICS for Children

Drug	Low Daily Dose	Medium Daily Dose	High daily Dose
Beclomethasone	$100-200\mu g$	$>200 - 400 \mu g$	> 400µg
Budesonide	$100-200\mu g$	$>200 - 400 \mu g$	$>400\mu g$
Fluticasone	$100-200\mu g$	>200-500µg	> 500µg
Budesonide -Neb	250 – 500µg	$>500 - 1000 \mu g$	> 1000µg

5. REDUCING RISKS WITH ICS THERAPY

- Use a spacer with metered dose inhalers to reduce oral deposition
- Rinse mouth/gargle after inhalation (with any form of device delivery)
- Wean to lower dose with sustained control
- Consider other add-on therapy rather than higher doses of ICS for incomplete control

6. MONITORING ASTHMA PROGRESSION

The following measures should be monitored over the course of follow up visits:

- Course of medications including frequency of use of Salbutamol
- Episodes of severe exacerbations requiring systemic corticosteroids
- Unscheduled clinic or emergency department visits
- Hospitalizations
- Pulmonary function test for 5 years and above
- Height and growth velocity of patients using ICS

7. LEVELS OF ASTHMA CONTROL

Table 2: Levels of Asthma Control

Characteristic	Controlled	Partly controlled	Uncontrolled
Day time symptoms	None or ≤ 2 times/week	> 2 times /week	≥ 3 features of partly controlled asthma present in any week
Activity limitation	None	Any	
Night symptoms	None	Any	
Salbutamol use	None or ≤ 2 times/week	> 2 times / week	
PEF or FEV1	Normal	< 80%	
Exacerbations	None	$\geq 1/$ year	One in any week

8. INDICATORS OF POOR ASTHMA CONTROL

- Awakens at night with symptoms
- Has unscheduled clinic or emergency department visits
- Has an increased need for short-acting inhaled β2 agonists (e.g. Salbutamol)
- Uses more than one canister of short acting $\beta 2$ agonist/month

9. MAIN REASONS FOR POOR CONTROL OF ASTHMA

- Poor inhaler technique
- Non- adherence to asthma therapy
- Wrong diagnosis
- Under treatment with anti inflammatory medications (ICS)
- Over reliance on short acting β2 agonists
- Presence of other coexisting conditions
- GERD
- Sinusitis
- Allergic rhinitis
- Continuous exposure to allergens

10. WHAT TO DO IF CONTROL IS NOT ACHIEVED

- Assess patient adherence and technique in using medications correctly and address as appropriate
- Address other factors that diminish control of asthma such as coexisting conditions, a new or increased exposure to allergens or irritants, or psychosocial problems
- In some cases, alternative diagnosis should be considered

<u>11. WHEN TO REFER TO SPECIALIST:</u>

Consider referral to specialist if:

- There are difficulties in achieving or maintaining control of asthma
- When moderate or higher doses of inhaled corticosteroids are required to achieve and maintain control
- Additional education is indicated to improve the patients' management skills or adherence
- Immunotherapy or other immunomodulators are considered, or additional tests are indicated to determine the role of allergy

These guidelines are *not* intended for the management of children with such associated conditions as cystic fibrosis, cardiovascular disease, chronic lung disease, Bronchopulmonary dysplasia (BPD) and immune deficiency syndromes.

1. ASSESSMENT OF SEVERE EXACERBATION

Categorize according to the most severe symptom or signs

	MILD	MODERATE	SEVERE
Breathlessness	While walkingCan lie down	While talkingPrefers sitting	 While at rest Infant stops feeding Sits upright
Talks in:	Sentences	Phrases	Words
Alertness	May be agitated	Usually agitated	Usually agitated
Respiratory rate (per minute)Expected normal< 2 mo	Increased	Increased	Very increased
and retractions	Usually not	Commonly	Usually
Wheeze	Moderate end expiratory	Loud throughout exhalation	Loud inspiratory and expiratory
Pulse rate / min Expected normal 2-12mo < 160	Mild increase	Moderate increase	Marked increase
Pulse oximetry (room air)	> 95 %	91 – 95 %	< 91 %

• Features of life threatening asthma (any one of the following):

- Drowsy or confused
- Apnea
- Inaudible breath sounds
- Paradoxical thoraco-abdominal movement
- Respiratory muscle fatigue / shallow respiration
- Cyanosis
- Bradycardia
- Silent chest

2. MANAGEMENT OF MILD TO MODERATE EXACERBATIONS

- * Sit up child, if possible
- * Give humidified O_2 to maintain $SaO_2 > 93\%$
- * Nebulized Salbutamol up to 3 doses in one hour at 20 minutes interval
- * Refer to the doses in the table of drug doses
- * Give Prednisolone 1-2mg/kg (max 40mg) by mouth
 - Or
- * IV Hydrocortisone 10mg/kg (max 250mg) intravenously stat if vomiting or not tolerating/taking oral prednisolone



3. MANAGEMENT OF SEVERE EXACERBATION

* Sit the child up / propped up position, if possible

* Give humidified O_2 to maintain $SaO_2 > 93\%$

* **Nebulized Salbutamol** upto 3 doses in one hour at 20 minutes interval or give continuous nebulized Salbutamol. Refer to table for doses and appropriate method of administration

* Add **nebulized Ipratropium** bromide 0.25 - 0.5mg with each dose of Salbutamol for 3 doses in one hour Start **IV fluids** at 2/3rd maintenance requirement

doses in one nour start **IV nuids** at 2/3 maintenance requireme

* Give Methylprednisolone / Hydrocortisone IV stat

* Consider giving MgSO₄ intravenously in a dose of 25-75mg/kg (max 2g) over 20 minute period in children with more severe exacerbations. MgSO₄ can be administered under monitoring even in Emergency Room if there is a delay in shifting the patient to HD/PICU Although this is a safe dose, it is recommended that BP must be monitored while administering MgSO₄ intravenously

Is there any improvement on re-assessment?

YES ▼

FURTHER TREATMENT

* Monitor closely, including pulse oximetry

* Nebulized Salbutamol hourly and then gradually decrease the frequency to 1-4 hourly intervals when there is further improvement

* Nebulized Ipratropium bromide q 6 h may be considered

* Continue Methylprednisolone / Hydrocortisone IV

NO IMPROVEMENT

* Alert / admit in PICU

* Check ABG & UE, do chest X-ray

* Give loading dose A**minophylline** 6mg/kg over 30 minute period, followed by

continuous infusion at the following rate (max 50mg/h) depending on the age of child:

2-6mo 0.4mg/kg/h

6-11mo 0.7mg/kg/h

1-9 years 1mg/kg/h

10-12 years 0.8mg/kg/h

Cardiac /Liver failure 0.2mg/kg/h

* Continue nebulized Salbutamol h/ continuously

* Continue to monitor closely in PICU. Wean off infusions if there is improvement. If there is no improvement in spite of all above measures, consider intubation and mechanical ventilation

Patients with life threatening asthma require admission to ICU/PICU

4. DRUG DOSES FOR ASTHMA IN CHILDREN

MEDICATION	DOSE	COMMENTS
Salbutamol nebulizing solution	 0 – 1 month 1.25 to 2.5mg > 1 months 2.5 to 5mg To be given every 20 minutes x 3 doses, then every 1-4 h or 5mg/kg/h by continuous nebulization 	 Dilute up to 4ml with normal saline O₂ flow at 6-10 L/min The face mask should be tightly fitting
Ipratropium bromide nebulizing solution	 0-1 year 25mcg/kg >1-5 years 125-250mcg >5 -12 years 250-500mcg >12 years 500mcg To be given every 20 min in severe attack x 3 doses, then every 4-6 h 	 Can be mixed with Salbutamol solution Titrate to response
Aminophylline Only to be used where monitoring facility is available	Loading dose 6mg/kg IV infusion over 30 min, followed by maintenance infusion	 Use a loading dose of 3mg/kg in children on maintenance oral Theophylline Monitor serum levels
Hydrocortisone IV	10mg/kg stat and then 4-5mg/kg q6h (max 250mg)	Convert to oral Prednisolone when stabilized
Prednisolone tablets	1-2mg/kg/day orally in 2 divided doses for 3-5 days (max 40mg/day)	There is no need to taper if patients are started on inhaled corticosteroid
Methylprednisolone IV	0.5 to 1mg/kg/dose q6h then reducing to minimum effective dose	
Magnesium sulfate IV	25-75mg/kg (max 2g) IV infusion over 20 minutes	 Monitor BP during and for up to 90 min after infusion Monitor level if frequent doses or infusion is required

PART THREE

REFERENCES & APPENDICES

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APPENDIX 1: ASTHMA CONTROL TEST



APPENDIX 2: ASTHMA CONTROL TEST



APPENDIX 3: ASTHMA ACTION PLAN



APPENDIX 4: ASTHMA ACTION PLAN

Con the second			sthma Daily Manag be used with symptoms a			. 🛞
Name:	Date:	Personal Best:	Institution:	Specialis	t Name:	Emergency Tel:
Green Zone: A	Asthma is under control					
or wheeze, at n - You can prac	nest tightness, breathlessness ight or day time tice your usual daily activity piratory flow rate (PEFR) r personal best	2. Continue (Ibutamol MDI (,,,,,,,,) 1-2 get asthma symptoms e taking your daily preven), (Ibutamol MDI (,,,,,,) 1-2 fore exerciser or heavy effe	ter) puffs 30	week, inform a. Identify you b. Check your	albutamol > 2 times per your Doctor to: In triggers inhaler technique. asthma medication
Yellow Zone (O	Caution) : In this stage cond	lition may get bette	er or worse			
breathless or wi	cough, chest tightness heeze while taking Your daily cation tween 79-50% of your	y available ev 2. Continue (If your syn Green Zou 1. Continue puffs every 2. Continue	butamol MDI () 1-2 lized Salbutamol () i very 20 minutes for one ho e taking your daily prevent), (f machine is ur er) goes back to) 1-2 ore days er	or your PEFR Zone: 1. Take puffs or use ne every 20 minu 2. Continue tak (3. Take an app soon as possibl a. Identify you b. Check your	king your daily preventer), () ointment with your doctor as le to:
Red Zone: Me	dical Warning					
cough - Having trouble - Taking Salbuta symptoms	st tightness, breathlessness or e talking or walking umol have not reduced your 50% of your personal best	or nebuliz minutes 2. Continu	albutamol MDI () zed Salbutamol () eve ne taking your daily preve), (ry 20 nter	c at the same tin	pital/ health center immediately OR Call the ambulance & me continue taking Salbutamol 2 puffs every 20minutes

APPENDIX 5: SELECTING INHALER DEVICES



APPENDIX 6: INHALER TECHNICQUE STEPS

ОЪ	serve patient's technique & for each	step put a√or X in	appropriate c	olumn
Go	od technique (G) = all essential steps	(E) performed accu	arately	
	or technique (P) = one or more essen			ely * Common problem step
	A	1		
	R B			a ora
	17 19			
1		00		
E	6.		CITERIE D	
	METERED DOSE INHAL	ER (MDI) •		TURBUHALER®
	(Ventolin [®] , Becotide [®] or Becl	omethasone®)		(Salbutamol, Budesonide, Formoterol & Symbicort [®])
IE	Remove MDI Cap		1E	Unscrew cover
2E* 3	Shake vigorously		2 3E*	Exhale slowly & completely away from mouthpiece
4E	Exhale slowly & completely Insert MDI upright between teeth & c	loced line around it	JE≭ 4E	Hold turbukaler upright Turn the coloured grap as far as it will go
5E*	Inhale steadily and deeply through m		5E*	then turn back till click sound is heard
6E*	immediately depress canister for		6E	Insert mouthpiece between teeth & close lips around it
7E	one depression	211	7E	Inhale forcefully & deeply through mouth
8	Remove inhaler keeping lips closed		8	Remove turbuhaler [®] keeping lips closed
9	Hold breath as much as possible	VETZ	9	Hold breath for 5-10 seconds
10	Breath normally For 2 nd dose wait 20-30 seconds.		10	Breathe normally
11 12E	For 2 ^{an} dose wait 20-30 seconds.		11 12E	For a 2 nd dose wait 20-30 seconds then repeat steps 2-10
13	Wipe MDI mouthpiece with clean tiss	:11e	12E	Wipe mouthpiece with clean dry tissue
14	Replace cap of MDI		14	Replace cover
	MDI+ MOUTHPIECE S	PACER		MDI+FACE MASK SPACER
1E	Remove cap from MDI		1E	Remove MDI cap
2E	Hold inhaler t & shake vigorously		2E*	Shake vigorously
3E*	Insert MDI upright into spacer, oppos		3E	Insert upright MDI into spacer
4E	Exhale slowly & completely away fro		4E	Hold MDI & spacer together keeping MDI in upright position
5	Insert spacer mouthpiece between tee	th, close lips	5	Place mask over mouth & nose
6E	Depress canister once	\geq	6E	and press gently to seal mask
7E	Inhale slowly & deeply through mout	h 🔊	7E	Depress canister once 🏼 🍳
8E	Remove spacer keeping lips closed	17	8E	Inhale
9	Hold breath for as much as you can	/ /	9	through open mouth
10E	Breathe normally	EV.	10E	for 5 breaths
11 12	Remove MDI from spacer	B	11 12	For a 2 nd dose wait 20-30 seconds
12 13E	For a 2 nd dose wait 20-30 seconds then repeat steps 2-11	0	12 13E	then repeat steps 3-8
136	Remove MDI from spacer and replac		136	Remove MDI from spacer
14	Wipe spacer mouthpiece with clean t		14	Keep spacer in clean place
16	Keep spacer in clean place	155 UC	16	Wipe face with clean wet cloth
10	Keep spacer in clean place		10	wipe lace with clean wet cloth
	AEROLIZER® (Forme	oterol)		DISKUS [®] (Fluticasone, Salmeterol, & Seretide [®])
1E	Remove the aerolizer [®] blue cover		1	Hold the outer case in one hand and
2	Hold the base of the aerolizer firmly		2*	place your other thumb on the thumbgrip
3E	Open aerolizer [®] by turning the mouth	piece clockwise	3E	Push thumbgrip away as far as it will go until it clicks
4E	Remove one capsule from foil strip	-	4	Exhale slowly & completely away from mouthpiece
SE	Place the capsule in the inhaler slot	M	5	Hold diskus [®] with mouthpiece facing you
6E	Close aerolizer [®] following the arrow of	on mouthpiece	6E*	slide lever away as far as it will go until it clicks
7E*	Hold aerolizer [®] upright & press blue b	outtons at base	7E*	Insert mouthpiece between teeth & close lips around it
_	(listen for the clicking sounds of caps	ule piercing)		
3	then release the blue buttons	100	8E*	Inhale steadily & deeply through mouth
9E	Insert mouthpiece between teeth & cla		9	Remove diskus [®] keeping lips closed
10E	Inhale steadily & deeply through mou		10	Hold breath for 5-10 seconds
11	Remove aerolizer [®] keeping lips close	1	11	Breathe normally
12	Hold breath for 5-10 seconds		12	To close diskus [®] put your thumb in the thumbgrip
13	Breathe normally Open aerolizer [®] & check capsule is er		13 14	& slide it backwards till you hear a click For a 2 nd dose wait 20-30 seconds
14E 15E			14 15E	
15E 16E	If necessary repeat steps 9-13 till caps Remove empty capsule & close & rep	sue is empty	15E 16	then repeat steps 1-12 (* load 2 nd dose!) Wine mouthniese with clean dry tissue
16E 17		orace cab	10	Wipe mouthpiece with clean dry tissue To close diskus [®] repeat steps 12 & 13
	Replace blue cover Wine mouthniece with clean dry ticcy	10	1/	TO CLOSE QUEAUE - TEPERI SIEPS 12 60 15
10 :	Wipe mouthpiece with clean dry tissu	10		<u> </u>
18			** ****** -	nsistence approach & facilitate compliance

APPENDIX 7: HOW TO USE PEAK FLOW METER



APPENDIX 8: PEAK FLOW RATE FOLLOW UP CHART

Age:	C xpira 35 yr 67cm	tory I	Flow S	Sectio	n (S	ample)		Sex: F	
Da	ite	Pred	icted	Pre Meas		% of Predicted	Post RX	% of Change	
2\1\/	\2008 430		30	250		58%	360	360-250 =110 110/250 = 44 %	
				easuremen Male& Fer			Peak Expiratory Flow in N		
Height 109	Score 147	Height	Score 254	Height	Score 360	660 650 640 630 620 810	MEN 75 190 72 183 63 175 66 167 67 167 175	680 640 620 620 620	

109	147	130	254	150	360
112	160	132	267	152	373
114	173	135	280	155	387
117	187	137	293	157	400
119	200	140	307	160	413
122	214	142	320	163	427
124	227	145	334	165	440
127	240	147	347	168	454



References: Global Strategy for Asthma Management and Prevention, National Institute of Health, National Heart Lang g and Blood Institute, Publication No. 953-659, 1995, Stempel DA et al, Ann Allergy Asthma Immunol (U.S) Feb. 1996 76 (2)p 153 ISSUN:1081-1206.

Peak Expiratory Flow rate Reading Guidelines

Asthma Classification	Predicted	Variability **
Intermittent	80%	< 20%
Mild	80%	20% -30%
Moderate	60% - 80%	30%
Severe	<60%	> 30%

**Variability is between the night & morning reading REFERENCE: BADDAR, WORTHING, AL RAWAS, OSMAN, AL RIYAMI, REESPIRATORY CARE. DECEMBER 2006 VOL 51 NO 12

APPENDIX 9: HOW TO FIND PERSONAL BEST VALUE

How to Find PEF Personal Best

Personal best is the highest number that can be achieved on a peak

flow meter over a 2-3 week period when your asthma is under control, and should:

1. Take daily asthma preventer in the morning, wait at least 6 hours to take the reading

Peak Expiratory Flow Daily Recording Chart (Sample)

- 2. Take the reading between noon and 2:00 PM for 1-2 weeks period
- 3. Blow 3 consecutive reading and write down the highest number
- 4. Record the highest number on the daily recording chart for each day
- 5. The highest number during this period is called personal best
- 6. This number can change over time so the test must be repeated yearly

650 600 550 500 450 400 350	1/10/0 AM)8 PM	2/10/ AM	08 PM	3/10/ AM	08 PM	4/10/ AM	08 PM	5/10/ AM	08 PM	6/10/ AM	08 PM	7/10/ AM	08 PM
650 600 550 500 450 400			AM		AM	PM								
 600 550 500 450 400 	•		-	•										
550 500 450 400	•		•	•										
500 450 400	•		•	•						1				
450 40 400	•	0	•	•										
40														
> 400													9	
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							Q				_0			
7 <mark>5</mark> 300								P						
250														
200														
150														
100														
50														
Copyright, (Oman Resp	piratory Soc	dety											

Asthma Management Protoco	ol		
l. Clinical History Section:			
Name:	Fa	amily Hi	story of Asth
MRN:	D	uration o	f Symptoms:
Sex:			
Hospital / Health Center:			
Fill in A, B, C, D & E at First Visit Only.			
Please tick ($$) or (X) when appropriate.			
A. Smoker () EX Smoker () Passive Smoker ()			
3. History of: a. Allergic Rhinitis () b. Nasal Polyps () b. Ecze	ma/Atopio	Derma	titis ()
C. What asthma medication is the patient currently taking? a. Nothing () b. Inhaled short acting B2 agonist () c. Long d. Inhaled steroids () e. Oral steroids () f. Theophylline () g. Oral Salbutamol () h. Other ()	acting B2	agonist ()
D. Has the patient used nebulised Salbutamol in the last 12 months? a. Home () b. Hospital () c. Health center ()	d. Othe	ers ()	
E. Has the patient been admitted for asthma in the last 12 months? a.	General wa	ard () l	D. ICU ()
Please tick ($$) or (X) when appropriate	General wa	ard () t	D. ICU ()
	General wa	ard () t	D. ICU ()
Please tick (√) or (X) when appropriate Date: 1. Does the patient have:	General wa	ard () t	5. ICU ()
Please tick (√) or (X) when appropriate Date:	General wa	ard () t	o. ICU ()
Please tick (√) or (X) when appropriate Date: 1. Does the patient have:	General wa	ard () 1	o. ICU ()
Please tick ($$) or (X) when appropriate Date: 1. Does the patient have: a. Cough (when patient does not have common cold)	General wa	ard () t	D. ICU ()
Please tick () or (X) when appropriate Date: Date: I. Does the patient have: a. Cough (when patient does not have common cold) b. Phlegm production c. Breathlessness	General wa	ard () t). ICU ()
Please tick () or (X) when appropriate Date: Date: I. Does the patient have: a. Cough (when patient does not have common cold)	General wa	ard () 1	p. ICU ()
Please tick () or (X) when appropriate Date: Date: I. Does the patient have: a. Cough (when patient does not have common cold) b. Phlegm production c. Breathlessness	General wa	ard () t). ICU ()
Please tick () or (X) when appropriate Date:	General wa		D. ICU ()
Please tick () or (X) when appropriate Date: Date: I. Does the patient have: a. Cough (when patient does not have common cold) Image: Cough (when patient does not have common cold) b. Phlegm production Image: Cough (when patient does not have common cold) c. Breathlessness Image: Cough (when patient does not have common cold)	General wa		p. ICU ()
Please tick () or (X) when appropriate Date:	General wa		D. ICU ()
Please tick () or (X) when appropriate Date:	General wa		D. ICU ()
Please tick (\sqrt{s}) or (X) when appropriate Date: Date:	General wa		D. ICU ()
Please tick (\sqrt{s}) or (X) when appropriate Date:	General wa		p. ICU ()
Please tick (\sqrt{s}) or (X) when appropriate Date:	General wa		D. ICU ()
Please tick (\sqrt{s}) or (X) when appropriate Date:	General wa		D. ICU ()
Please tick (\sqrt{s}) or (X) when appropriate Date:	General wa		p. ICU ()

 I. Beta blocker
 2. ACE inhibitors (Captopril, Cilazapril, Lisinopril)
 3. Additives in inhalers
 4. NSAIDs

 REFERENCE: BADDAR, WORTHING, AL RAWAS, OSMAN, AL RIYAMI, REESPIRATORY CARE. DECEMBER 2006 VOL 51 NO 12
 3. Additives in inhalers
 4. NSAIDs

APPENDIX 10: ASTHMA MANAGEMENT PROTOCOL

	PEFR		ight:			Dat	e		D	ate				Date				Dat
Predicted value		Test				Dat	c			ate		+		Jate	,			Dat
Actual value																		
% of (Predicat	ed/Chan	ge)																
Good (G)	Comp		Patient	N														
	haler T			(1)														
Good (G)	, Poor (P), New I	Patient	(N)				_										
MDI, Spacer w Turbuhaler®	nth (MP	or Masl	k)		_							+						
Aerolizer®																		
Diskus®																		
Asthma Contr Date:	ol Test ((ACT)																
Date: Total Score:								+				+						
1 Dunie -	the next i	marken	U.	ft an		there a			. 6		inc.c.		hda				ahar	l or h -
1. During All the time	the past 4 1	Most of					preve	-	1	-	f the			ne at Non				
	_						ac tillie		A	ane o	и ше	ame	7	. NOI	or th	ie uii	ue J	
2. How oft	en have y 1	Once a		2		times	a week	3	1-2	time	saw	eek 4		Non	at al	1	5	
3. How off	-		-															night
earlier tha	in usual i	n the mo	orning?	лошз (т	, , , , , , , , , , , , , , , , , , , ,	tough	, enes	ugut		orca	une ss	ness,	pain	,		/u uj	p at	
\geq 4 times a we	eek 1	2 -3 nig	tts a we	ek 2	O	nce a v	veek	3	1-2	time	s / mo	nth	4	No	ot at	a11		5
4. How of													T.					
\geq 3 times a da	y 1	1- 2 tim	ies a day	7 2	2-3	times	a weel	2	3 ≤	1 tin	ie a w	eek	4	Non	at a	11	5	
5. How we Not controlled							past 4 1 t contro			Wall	contro	11.0.4	4	Com	alata		tral	5
Not controlled	• 1	Poorly	control	ed 2	Son	iewna	t contro	onea	5	well	contro	ollea	4	Com	piete	e con	trol	2
ACT I	Reference	: < 20 =	Not Co	ntrolled	20 -	24 = 1	Well C	ontro	lled	25 =	= Full	Cont	rol	Scor	e:		•	
	Peak Fr	iratory Flo	w Rate M	0.0				B	Dent		long to 2	law i	Park					
No	male)	68	<u></u>	Peak	CREGG A J.	NUNN. B	ow in N	iJournal 3	3 u bjec 73. 3. 282.	us I	-	690						
109		Height 130	Score 254	Height	360	04 630 630	0	K	-12	-163	K	1	-			4.59 640 630 629		
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		137	293	157	400	54 54 33 32	: //	38	landari diri landari dori	dian, two vitien, woe	en = 18	Mesmin.		1	1	510 510 539 529		
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122		142	320	163	427	480 471 460 451				152		1			•	(m) (m) (m) (m) (m) (m) (m) (m) (m) (m))	
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123		145	347	165	454	410		is it div	an Iban proti an IS litres's this protoal	etrot, and i non, heres th lisenita.	te 100 Bizera n warnen ber an predictet	R		1	11	489 409 390 380		
1.27	240		1		d lesting of	38 PEI Lie	A 1994	0 25	30	35 4	0 45		55	0 65				
Referen	es: Giobal Strategy Intional Heart Lung Ann Allergy Asth	for Asthena Manu	agerment and Pre-	ventore, ventore			4.0	03			OF IN YEARS	24	34		249			

APPENDIX 10: ASTHMA MANAGEMENT PROTOCOL

3. Medication Section: Dose & Frequency

For Asthma Severity; Review Q	Quick Referen	Date	Date	Date	Date	
Asthma Severity:						
Medication	Device	Strength				
Short acting B2 agonist						
Salbutamol	MDI	100µg/inh				
Salbutamol	Rotahaler®	200µg/cap				
Salbutamol	Rotahaler®	400µg/cap				
Salbutamol	Neb Solution	5mg/mL				
Salbutamol	Syrup	2mg/5mL				
Salbutamol	Tablets	2mg				
ICS						
Beclomethasone	MDI	50µg/inh				
Beclomethasone	MDI	250µg/inh				
Budesonide	Turbuhaler®	100µg/inh				
Budesonide	Turbuhaler®	200µg/inh				
Budesonide	Turbuhaler®	400µg/inh				
Fluticasone	MDI	50µg/inh				
Fluticasone	MDI	125/inh				
Fluticasone	MDI	250µg/inh				
Fluticasone	Diskus®	100µg/inh				
Fluticasone	Diskus®	250µg/inh				
LABA	DISKUS	2000,000				
Salmeterol	MDI	25µg/inh				
Salmeterol	Diskus®	50µg/inh				
Formoterol	Aerolizer®	12µg/ cap				
Formoterol	Turbuhaler®	9µg/ inh				
ICS+LABA (Combined)	Turbullater	Dose/inh				
Symbicort [®]	Turbuhaler®	80µg/4.5µg				
Budesonide+Formoterol)	Turbullator	per inh				
Symbicort®	Turbuhaler®	160µg /4.5µg				
Seretide [®] Fluticasone+Salmeterol)	Diskus®	100µg/50µg				
Seretide®	Diskus®	250µg/50µg				
Seretide®	Diskus®	500µg/50µg				
Other						
Sod.cromoglycate	MDI	5mg/inh				
Motelukast Sodium	Tablets	5mg /tab				
Motelukast Sodium	Tablets	10mg /tab				
Theophylline	SR Tablets	200mg /tab				
Theophylline	SR Tablets	300mg/tab				
Oral Steroids						
Prednisolone	Tablets	1mg /tab				
Prednisolone	Tablets	5mg /tab				
Prednisolone	Tablets	25mg/tab				
Other (non asthma medication)						
Referral to Specialist:	1	1				
Institution						
Reason for referral REFERENCE: BADDAR, WORTHING, AL RAWAS, OSMAL	N AL RIVAMI RECEDIDAT	ORY CARE DECEMBER 3005	01518012			L

APPENDIX 10: ASTHMA MANAGEMENT PROTOCOL

4. Quick Reference Guidelines

Assess severity to initiate therapy

During a patient's initial presentation, if the patient is not currently taking long-term control medication, asthma severity is assessed to guide clinical decisions for initiating the appropriate medication and other therapeutic interventions.

Step 1	Mild Intermittent Symptoms < 1 time / week Night symptoms ≤ 2 / month	Inhaled short acting B2 agonist when needed; Salbutamol (100-200µg)
	PEFR or FEV1 ≥ 80% of predicted or personal best • Variability < 20%	
Step 2	Mild Persistent Symptoms ≥ 1 time / week but not daily Night symptoms > 2 / month	Inhaled short acting B2 agonist PRN + Low dose ICS (Beclomethasone 200 – 500µg or Budesonide 200 - 400µg or Fluticasone 100 -250µg) or Leukotriene modifiers (Monteleukast 10 mg daily)
	Attacks may affect activity PEFR or FEV1 ≥ 80% of predicted or personal best • Variability between 20-30%	
Step 3	Moderate Persistent Daily symptoms Night symptoms >1 / week Attacks may affect activities PEFR or FEV1 60-80% of predicted or personal best • Variability > 30%	Inhaled short acting β2 agonist PRN + Low dose ICS +Inhaled long acting β2-agonist (LABA; Salmeterol 50µg BD or Formeterol 4.5-12µg BD) or Medium Daily Dose ICS (Beclomethasone >500 -1000µg or Budesonide > 400 - 800µg or Fluticasone >250 - 500µg) or Low dose ICS + Leukotriene modifiers or Low dose ICS +Sustained Release Theophylline
Step 4	Severe Persistent Continuous symptoms Frequent night symptoms Limited physical activity PEFR or FEV1 ≤ 60% of predicted or personal best • Variability > 30%	Inhaled short acting B2 agonist PRN + Medium or high dose ICS + LABA or Medium or high dose ICS + Leukotriene modifiers or Medium or high dose ICS + Sustained release Theophylline or High Daily Dose ICS Beclomethasone >1000 - 2000µg or Budesonide >800 - 1600µg or Fluticasone >500 - 1000µg
Step 5	Referral to Specialist Asthma is not Controlled with medication in Step 4	Inhaled short acting B2 agonist PRN + High dose ICS + LABA + Leukotriene modifiers +Oral steroids or High dose ICS + LABA + Sustained release Theophylline +Oral steroids

Classification of Asthma & Therapy

Step Up

If control is not achieved, consider step up, But first review:

a. Inhaler technique b. Compliance c. Trigger factors

Patients persisting in step 4, should be referred to a chest specialist

Step Down

Review treatment every 3 to 6 months.

If control is sustained for at least 3 months, a gradual stepwise reduction in treatment should be considered

Assess Control to Adjust Therapy

	Assess Control to	D Adjust Therapy	
Table2: Levels of asthma	control		
Characteristic	Controlled	Partly controlled	Uncontrolled
Day time symptoms	None (≤2 times/week)	> 2 times/week	\geq 3 features of partly
Activity limitation	None	Any	controlled asthma present in
Night symptoms	None	Any	any week
Salbutamol use	None (≤2 times/week)	> 2 times/week	
PEF or FEV1	Normal	< 80%	
Exacerbations	None	≥1 / year	One in any week

Exact Dations involte ≤ 1 / year REFERENCE: BADDAR, WORTHING, AL RAWAS, OSMAN, AL RIYAMI, REESPIRATORY CARE. DECEMBER 2006 VOL 51 NO 12

To Submit any Comments, Please Contact:

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