

ISSN: 2230-9926

ORIGINAL RESEARCH ARTICLE

Available online at http://www.journalijdr.com



International Journal of Development Research Vol. 08, Issue, 10, pp.23327-23334, October, 2018

OPEN ACCESS

COMPARISON OF ASTHMA CONTROL TEST (ACT) WITH (GINA) GUIDELINES IN THE ASSESSMENT OF ASTHMA CONTROL AND DETERMINE IF CAN USE ACT AS ALTERNATIVE TO GINA GUIDELINES IN CONTROL OF ASTHMA

*Ahmed Hassan Jabber

Hilla Teaching Hospital, Babylon, Iraq

ARTICLE INFO

Article History: Received 17th July, 2018 Received in revised form 13th August, 2018 Accepted 20th September, 2018 Published online 29th October, 2018

Key Words:

Asthmatics, ACT, Spirometer.

ABSTRACT

The gold standard in assessing asthma control is the Global Initiative for Asthma (GINA) criteria. and because of the difficulties of access to pulmonary functions tests, The ACT has the added advantage that it does not require lung function assessment. The aim of this study is to assess asthma control through ACT score and GINA guideline, and to determine if the ACT can be as useful as the GINA-guidelines criteria in assessing asthma control in Iraq. Cross sectional study with comparing ACT vs. GINA guideline in control of asthma level. This study was conducted at Respiratory consultation unit of the Iraqi National center of early detection of Cancer, Baghdad-Iraq, The study was conducted during the period from 1stNovember 2012 to 1stJuly 2013. A total of 71 adult asthmatic patients who were attended to the respiratory consultant unit were asked to participate and were enrolled in this study regardless their age or gender. Their asthma diagnosed and proved clinically by a combination of history, clinical finding In addition objective measurements using spirometry (FEV1) measured by the reversibility test which is defined as (an increment of>12% or 200 ml of FEV1 after 20 minutes of administration of inhaled short acting B2-agonist). There were 71 patients enrolled in this study, of them 66 (92.96%) had an ACT score of \leq 19 and 5 patients (7.04%) had an ACT score of > 19, it had been found the number of male is (27) and (26) (39.4%) out of them had an ACT<19and only (1) (20.0%) had ACT>19 and number of female is (44), (40) (60.6%) out of them had ACT<19 and (4) (80.0%) had ACT>19. No significant differences had been found in between those patients with ≤ 19 ACT score vs. those with > 19, regarding the age and gender, in both comparison P>0.05 it had been found that good agreement present between ACT and GINA, 92.9%.ACT agreed the GINA in (37 patients with uncontrolled asthma, 24 patients with partially controlled and 5 patients with controlled). ACT can served as an alternative diagnostic tool in assessing asthma control even without an aid of a spirometer or a peak flow meter. An ACT score of more than 19 can classify patient as controlled asthmatic while an ACT score < 19 can classify the patient as uncontrolled and partially controlled asthmatics.

Copyright © 2018, *Ahmed Hassan Jabber*. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Citation: Ahmed Hassan jabber, 2018. "Comparison of Asthma Control Test (ACT) with (GINA) guidelines in the Assessment of Asthma Control and determine if can use ACT as alternative to Gina guidelines in control of asthma", *International Journal of Development Research*, 8, (10), 23327-23334.

INTRODUCTION

Asthma has defied as "a chronic inflammatory disorder of the airways" (National Asthma Education and Prevention Program, 2007) However, this description omits the characteristic waxing and waning character of airflow obstruction in asthma.

**Corresponding author:* Ahmed Hassan Jabber, Hilla Teaching Hospital, Babylon, Iraq. A more useful definition would combine the central roles of inflammation and bronchial hyper responsiveness with the characteristic clinical symptoms. As an example, asthma may be defined as "a common chronic disorder of the airways that is complex and characterized by variable and recurring symptoms, airflow obstruction, bronchial hyper responsiveness, and an underlying inflammation.

The Asthma Control Test (ACT): A simple 5-question test for asthma has been developed and validated in several studies



(Reddel, 2009; Liu, 2007; Thomas, 2009) The ACT was initially developed in a study which looked at 25 of the most common questions that doctors ask when talking to patients about asthma control, with 5 questions standing out as being the most accurate predictors (Reddel, 2009). The 5 questions take less than a minute to answer and can be asked by the health care professional or the patient can complete the test themselves. There is a score of 1-5corresponding to a high level of symptoms. Studies have shown that the ACT score effectively discriminates between patients who differ in asthma control, is responsive to changes in control, and can discriminate between groups of patients in different lung function ranges. A score of 20-25 means that a patient's asthma is controlled. A score of 15-19 is partially control means that it may be possible to increase the level of asthma control and a full review of the treatment plan, including education on inhaler technique and the important of compliance with treatment, is warranted.

A score of 14 or less indicates that asthma is poorly or not controlled and that an urgent review of and changes to the patient's management are needed. Although there are no randomized studies that demonstrate that use of the ACT translate into better asthma control, its use is highly likely to improve patient outcomes as asthma therapy can be confidently adjusted up if control is demonstrated to be poor. The (GINA) guidelines based on clinical symptoms including daytime symptom and limitation of activities and nocturnal shortness of breath, spirometric studies with FEV1, (GINA) guidelines classification of symptom control into control, partially control, uncontrol have promoted the progression and the improvement of asthma management (Hasegawa et al., 1998; Hasegawa, 2012). To use these guidelines appropriately, it is extremely important to evaluate strategy essentially depends on the level of asthma control (Bateman, 2008; Ohta, 2011). Studies of actual clinical care have indicated that there is poor use of pulmonary functiontests, including forced expiratory volume at 1 second, (PEF), which are require under most circumstances for proper evaluation of asthma control under these guidelines. The Asthma Control Test (ACT), (Nathan et al., 2004) developed in 2004, consisting of 5 questions. This tool is recognized as a superior for achieving asthma control. One of the greatest benefits of ACT is that no respiratory function tests are required to evaluate asthma control. The ACT is thus suitable for administration using questionnaire surveys for asthmatic patient easy to use in the actual clinical care setting.

Measurements of lung function: Although the diagnosis of asthma is usually based on the presence of characteristic symptoms, patients with asthma frequently have poor recognition of their symptoms and poor perception of symptom severity, especially if their asthma is long-standing ⁽¹⁴⁾, assessment of symptoms such as dyspnoea and wheezing by physicians may also be inaccurate. For patients >5 yrs of age, measurements of lung function to confirm airflow limitation, and particularly the demonstration of reversibility of lung function abnormalities, greatly enhance diagnostic confidence. Quality control and adequate instruction for patients on how to perform the forced expiratory maneuvers essential (Aaron et al., 1998). The degree of reversibility in forced expiratory volume in one second (FEV₁) that indicates a diagnosis of asthma is generally accepted as $\geq 12\%$ and ≥ 200 mL from the pre-bronchodilator value. Repeated testing at different visits is advised.

Because many lung diseases may result in reduced FEV₁, a useful assessment of airflow limitation is the ratio of FEV₁ to forced vital capacity (FVC), (PEF) measurements made using a peak flow meter can also be an important aid in both diagnosis and monitoring of asthma. However, measurements of PEF are not interchangeable with other measurements of lung function, such as FEV₁ in adults (Davies *et al.*, 2006) or children (Lock *et al.*, 1996), because values obtained with different peak flow meters vary and the range of predicted values is too wide. PEF measurements are also very effort dependent, and quality may be poor. Therefore, measurements should always be compared with the patient's own previous best measurements (Bernstein *et al.*, 1996) using his/her own peak flow meter. The previous best measurement is usually obtained when the patient is asymptomatic and controlled.

Asthma control: In general, the term control may indicate disease prevention or even cure. However, in asthma, where neither of these are realistic options at present, it refers to control of the manifestations of disease. There is evidence that reducing inflammation with controller therapy achieves clinical control, but because of the cost and/or general unavailability of tests to routinely assess airway inflammation (Green et al., 2002; Pizzichini et al., 1996; Smith, 2005), it is recommended that treatment is aimed at controlling the clinical features of disease, including lung function abnormalities. Complete control of asthma is commonly achieved with treatment, the aim of which should be to achieve and maintain control for prolonged periods with due regard for the safety of treatment, potential for adverse effects, and the cost of treatment required to achieve this goal. Validated measures for assessing the clinical control of asthma score goals as continuous variables and provide numerical values to distinguish different levels of control. Examples of validated instruments are: Childhood Asthma Control Test (Liu, 2007), Asthma Control Test (http://www.asthmacontrol. com/ Date last updated: January 8, 2001. Date last accessed: July 15, 2007), Not all of these instruments include a measure of lung function. Their value in clinical practice, as distinct from the research setting, although suggested in several reports, requires further evaluation.

Asthma control by using markers of inflammation: These include measurement of nonspecific airway hyper responsiveness, cells and mediators from induced sputum, exhaled nitric oxide and components of exhaled breath air. There is at present no good peripheral blood analysis to measure airway inflammation. treatment strategy directed at normalization of the nonspecific bronchial responsiveness was found to reduce exacerbations and normalize airway inflammation In another study, a strategy aimed at normalizing induced sputum eosinophil counts was found to reduce asthma exacerbations and admissions without the need for additional anti-inflammatory treatment (Green et al., 2002) Another study showed similar results and suggested that assessment of asthma medication needs from evaluation of sputum eosinophilia was mostly useful in preventing exacerbations of the eosinophilic type Among the ACSS (Boulet, 2002) was the first to suggest including sputum eosinophils as a potential additional parameter to assess control.

Loss of control

Passive and active smoking: Around 24% of the US population are current smokers (Centers for Disease Control and Prevention, 2004).

The prevalence of smoking in asthmatics is similar to that in the general population Moreover, many nonsmokers are exposed to environmental tobacco smoke (Eisner, 2002) Smoking asthmatics have poorer asthma control and increased acute care needs (Boulet, 2006) By comparison with nonsmokers, they have more respiratory symptoms, worse quality of life, and more emergency department visits and hospitalizations.

Respiratory infections: Viral respiratory infections represent the most common cause of asthma exacerbations and, hence, contribute to a loss of asthma control. The importance of viruses eliciting asthma exacerbations was suspected and confirmed with polymerase chain reaction methods. Rhinovirus is the most frequently identified causal agent (Rakes, 1999; Jacoby, 2002) However, other viruses such as the human metapneumovirus have recently been identified ⁽³¹⁾ The role of glucocorticosteroids in the prevention and treatment of virus-induced exacerbations is still a matter of discussion.

Acute and chronic allergen exposure: Allergens have been considered potential key contributors to the etiology and clinical course of asthma (Yssel, 1998) although in epidemiologic studies the relationship between asthma and allergy is not obvious (Pearce, 2000) The importance of environmental allergen exposure in the development of asthma exacerbations has not been fully defined except in some cases like thunderstorm-induced asthma or the Barcelona asthma epidemic Evidence that chronic allergen exposure caused by house dust mites or pollens can have the same effect is weaker possibly because chest symptoms in asthma may also be related to nasal symptoms. In experimental studies with asthmatic patients, a single, high-dose allergen challenge can easily reproduce most, if not all, features of an asthma exacerbation (Bentley, 1997).

Aim of study: The aim of this study is to assess asthma control through ACT score and GINA guideline, and to determine if the ACT can be as useful as the GINA-guidelines criteria in assessing asthma control in Iraq.

PATIENTS AND METHODS

This is a cross sectional study with comparing ACT vs. GINA guidelines in control of asthma level. This study was conducted at Respiratory consultation unit of the Iraqi National center of early detection of Cancer, Baghdad-Iraq. The study was conducted during the period from 1stnovember 2012 to 1stJuly 2013. A total of 71 adult asthmatic patients who were attended to the respiratory consultant unit were asked to participate and were enrolled in this study regardless their age or gender. Their asthma diagnosed and proved clinically by a combination of history and symptom: 1) cough which worsens at night, 2) wheeze, 3) difficulty of breathing, 4) chest tightness. In addition, objective measurements of airflow obstruction using spirometry (FEV1) measured by the reversibility test which is defined as (an increment of>12% or 200 ml of FEV1 after 20 minutes of administration of inhaled short acting B2-agonist) according to the British guidelines on the management of asthma.

Inclusion criteria

 Previously diagnosed asthmatic patients i.e. previously attend respiratory clinic and underwent spirometric test.

- Patients who were aged 15 years and more were included.
- Both genders were eligible able to underwent spirometer test.

Exclusion criteria: Patient was excluded if he\she had one of the following criteria:

- Had been hospitalized for Asthma.
- Acute upper or lower respiratory tract infection within 4 weeks.
- A known respiratory disorder other than asthma.
- Smokers who were smoked more than 10 pack-year.
- pregnancy.

Data collection: Data were collected via full medical history and complete clinical examination and the data were recorded in a pre-constructed data sheet which was included:

Socio-demographic data; age, gender, and clinical examination data which included, Pulmonary Function Test Prebronchodilator FEV1 and FEV₁/FVC were measured using office spirometry in pulmonary function test outpatient clinic in Baghdad Teaching Hospital.

Assessment of Control of Asthma:- All patients were assessed for their control of asthma by using ACT scoring and GINA guideline as a golden standard, ACT scoring is a self administered 5 item questionnaire developed for assessing asthma control level. It evaluates the most recent 4 week time period. The Asthma Control Test (ACT) contains five items: the effect of asthma on daily activities, daytime and nocturnal symptoms, use of rescue inhaler medications and self assessment of asthma control, and dealing with asthma control during the previous 4weeks; each item is scored between 1 and 5, with the total-score ranging from 5 to 25. An ACT score of 25indicates that asthma is "controlled," whereas a score between15 and 19 shows partially controlled asthma and a score of <15 indicates "uncontrolled" asthma. And those with a score of <19 were re-grouped uncontrolled and partially control asthmatics. Then, the totally controlled patients were re-grouped as controlled patients (ACT>19). After the Asthma Control Test (ACT), patients had an interview wherein they were classified according to the GINA symptom severity, The GINA classification of symptom control into control, partially control, uncontrol. This is based on clinical symptoms including daytime symptom and limitation of activities and nocturnal shortness of breath, spirometric studies with FEV1.

Statistical analysis: Data of all patients were entered and analyzed by using the statistical package for social sciences (SPSS) software for windows version 18. Descriptive statistics were presented as mean \pm standard deviation (SD) for continuous variables and as frequencies and proportions (%) for categorical variables. Student's t test (independent 2 samples) was used to compare means of age, FEV1 in between two groups according to ACT level (\leq 19 or > 19). Chi square was used to assess the significance of association in between groups regarding the categorical variables. Agreement between ACT and GINA was calculated using percent agreement calculation and Kappa statistics.

RESULTS

There were 71 patients enrolled in this study, of them 66 (92.96%) had an ACT score of \leq 19 and 5 patients (7.04%)

1.11

Ahmed Hassan Jabber, Comparison of asthma control test (act) with (Gina) guidelines in the assessment of asthma control and determine if can use act as alternative to gina guidelines in control of Asthma

Table 1. Gina guideline classification asthma control

	Controlled	Partly controlled (any present in any week)	Uncontrolled
Daytime symptoms	None (2 or less\week)	More than twice \week	
Limitation of activities	None	Any	Three or more
Nocturnal symptoms/ awakening	None	Any	features of partly
Need for rescue/ "reliever" treatment	rescue/ "reliever" None (2 or less\week)		asthma
Lung function (PEF or FEV ₁)	Normal	< 80% predicted or personal best (if known) on any day) present in any

Table 2. Asthma control test score

 In <u>the past 4 weeks</u>, how much of the time did your <u>asthma</u> keep you from getting as much done as usual at work, school, or at home? 					
1	□2	□3	4		
All of the time	Most of the time	Some of the time	A little of the time	None of the time	Score
2. During	the past 4 wee	eks, how often hav	e you had shortr	ness of breath?	
1	2	□3	4	5	
More than once a day	Once a day	3 to 6 times a week	Once or twice a week	Not at all	Score
3. During the <u>past 4 weeks</u> , how often did your <u>asthma</u> symptoms (wheezing, coughing, shortness of breath, chest tightness, or pain) wake you up at night or earlier than usual in the morning?					
□1	□2	□3	□4	□5	
4 or more nights a week	2 to 3 nights a week	One night a week	One or two nights in the last 4 weeks	Not at all	Score
4. During the <u>past 4 weeks</u> , how often have you used your rescue inhaler or nebulizer medication (such as albuterol)?					
□1	□2	□3	□4	□5	
3 or more times per day	1 or 2 times per day	2 or 3 times per week	Once a week or less	Not at all	Score
5. How would you rate your asthma control during the past 4 weeks?					
□1	□2	□3	□4	□.5	
Not controlled at all	Poorly controlled	i Somewhat controlle	d Well controlled	Completely controlled	Score
Total score:				e:	

Variable		ACT s	core	P. value
		≤ 19	> 19	
		(n=66)	(n=5)	
Age (years)	Mean \pm SD	41.6 ± 14.5	51.2 ± 15.4	0.137 [NS]
Gender n (%)	Male	26 (39.4)	1 (20.0)	0.39 [NS]
	Female	40 (60.6)	4 (80.0)	
GINA classification	Uncontrolled	37 (56.1)	0	
	Partially controlled	29 (43.9)	0	
	Controlled	0	5 (100.0)	
FEV1 n (%)	> 80	0	2 (40.0)	
	60 - 80	35 (53.0)	3 (60.0)	
	< 60	31 (47.0)	0	
	Mean \pm SD	55.3 ± 15	74 ± 5.5	0.007 [sig]
	Mean ± SD	51(47.0) 55.3 ± 15	74 ± 5.5	0.007 [sig]

Table 3. Patients characteristics and GINA classification distributed by ACT score

Table 4. Distribution of ACT categories by GINA classification

	ACT			Total
GINA classification n (%)	Uncontrolled	Partially controlled	Controlled	
Uncontrolled	37 (88.1)	0 (0.0)	0 (0.0)	37 (52.1)
Partially controlled	5 (11.9)	24 (100.0)	0 (0.0)	29 (40.9)
Controlled Total	0 (0.0) 42 (100.0)	0 (0.0) 24 (100.0)	5 (100.0) 5 (100.0)	5 (7.0) 71 (100.0)
Percent agreement = 92.9%				

Table 5. Correlation between FEV1 and AC	ween FEV1 and ACT
--	-------------------

	ACT score level			P.value
FEV1	Uncontrolled	Partially controlled	Controlled	
> 80	0 (0.0)	0	2 (40.0)	
60 - 80	14 (33.3)	21 (87.5)	3 (60.0)	
< 60	28 (66.7)	3 (12.5)	0 (0.0)	
Mean \pm SD	49.7 ± 15.1	65.1 ± 8.4	74 ± 5.47	< 0.001 [sig]

had an ACT score of > 19, figure 1. Table 3. summarizes the patients characteristics distributed by ACT score level, No significant differences had been found in between those patients with ≤ 19 ACT score vs. those with > 19, regarding the age and gender, in both comparison P>0.05. Regarding the distribution of ACT score vs. GINA classification. It had been found that out of the 66 patients with ACT <19, 37 patients (56.1%) were labeled as uncontrolled on GINA, 29 (43.9%) labeled as partially controlled and none labeled as controlled, in contrast none of those with ACT score > 19 were labeled as uncontrolled or partially controlled on GINA, and Only the 5 patients with ACT score > 19 were labeled as controlled. On the other hand asthmatic patients with > 19 ACT score were significantly had higher FEV1 level 2 0f them (40%) had FEV1 of > 80 and 3 (60) of them had FEV1 of (60-80) while none of them had FEV1 < 60. Out of those patients with \leq 19 Act level, none had FEV1 >80, 35 (53%) had 60-80 and 31 (47%) had FEV1 < 60. On comparison of mean FEV1 in between groups, those with ACT \leq 19 had lower mean FEV1 as compared to those with >19 ACT, the mean FEV1 was 55.3 \pm 15 and 74 \pm 5.5 respectively, P=0.007.These finding indicating that FEV1 and ACT were directly correlated.

The distribution of ACT categories by the GINA classes is shown in table 4, it had been found that good agreement present between ACT and GINA, 92.9% and this percent is high due to small sample size. ACT agreed the GINA in 37 patients with uncontrolled, 24 patients with partially controlled and 5 patients with controlled) and had been found the uncontrolled patients by Gina is (37) while uncontrolled patients by ACT is (42) and this indicate small difference between them. In table 5, the distribution of FEV1 according to the ACT categories of the patients shows a direct correlation between FEV1 and ACT, and compared as a means, found P<0.001.

DISCUSSION

In our study There were 71 patients enrolled in this study, of them 66 (92.96%) had an ACT score of \leq 19 and 5 patients (7.04%) had an ACT score of > 19,it had been found the number of male is (27) and (26) out of them had an ACT<19and only (1) had ACT>19 and number of female is (44), (40) out of them had ACT<19 and (4) had ACT>19.



Figure 1. Distribution of Asthmatic patients according to ACT score



Figure 2. Comparison of mean FEV1 according to ACT level

In general characteristic of this study both male and female distributed according to ACT score level, found no significant differences between those patients with ≤ 19 ACT score vs. those with > 19, regarding the age and gender, in both comparison found P>0.05 in table (3) and this have similar report in Kurdistan-Iran by sigarin et al. and consistent with results of USA study (2006) and Philippines study (2007). This finding confirms the usefulness of ACT as a valid test in different populations. The present study revealed that 92.96% of studied asthmatic patients had ACT scores \leq 19 and 7.04% of them had ACT scores > 19. This finding regarding proportion of uncontrolled asthmatic patients measured by ACT is higher than that reported by Philippine study in 2007 (72%) and close to results of USA study (2004) in which, most of asthmatic patients were uncontrolled. This high proportion of uncontrolled asthma showed that in our country, still asthma is not totally contained. And a full review of the treatment plan, including education on inhaler technique and the important of compliance with treatment is warranted. Our study revealed that 56.1% of uncontrolled asthmatic patients assessed by ACT were categorized as uncontrolled by GINA classification and 43.9% of them were categorized by GINA classification as partially controlled. This finding is consistent with that reported by Spanish study (2006) that found 57% of uncontrolled asthmatic patients were labeled as uncontrolled with GINA. All the asthmatic patients categorized as controlled by ACT were categorized by GINA classification as controlled (p < 0.001). This finding is consistent with results of USA study (2005) on 522 subjects that showed ACT may serve as a useful screening tool in the community to determine

whether patients have controlled or uncontrolled asthma. On the other hand asthmatic patients were The distribution of ACT categories according to the GINA classes is shown in table (4), In the present study percent agreement between ACT and GINA was 92.9%. This finding is consistent with results of Cross-sectional survey (2008) comparing ACT score and GINA classification of asthma control among 2949 patients attending primary care physicians and specialists in France, Germany, Italy, UK, Spain and USA (Thomas et al., 2009). In this study we observed a stronger correlation between the ACT scores and mean FEV 1 (p < 0.001), and is consistent with the findings observed in other studies (Moy, 2001; Juniper, 19933) this have similar result with Kurdistan-Iran by sigarin et al. These results confirm that asthma control cannot be inferred from the clinical measure of airway function alone. In this study, the ACT was useful in predicting GINA-defined asthma control categories and was particularly useful in confirming patients whose asthma was not controlled according to the GINA classification. We found that an ACT score of < 19correctly predicted GINA 'partly controlled' or 'uncontrolled' asthma 100%. Stempel, et al (2005), also in other study showed that ACT may serve as a useful screening tool in the community to determine whether patients have controlled or uncontrolled asthma Consequently, this makes it an excellent diagnostic tool for screening asthma severity.

Conclusion

- Asthma control test (ACT) can served as an alternative diagnostic tool in assessing asthma control even without an aid of a spirometer or a peak flow meter in an out-patient basis or as home based easily and quickly completed by patients.
- It can serve as a guide in the case management of asthmatic patients by step up and step down treatment when the asthmatic patient is control or un control according to ACT score to guide adjustments in asthma therapy.
- The ACT may promote communication and partnership between patients and physicians, which helps the patients to establish confidence in asthma management, and improves physicians' performance and treatment outcome.

Recommendation

- Asthma control questionnaire test should be applied routinely in our daily practice (especially respiratory clinics) to assist in future studying and planning for proper asthma management.
- Encouraging patients for using ACT score questionnaire in home and in work which is easily and quickly completed by patients to guide treatment and, follow the patient conditions when attend respiratory clinic.
- ACT score is a simple, inexpensive tool that can be used especially in our country Iraq where financial resources are limited and disabling our patient to do the standard diagnostic test such as spirometry.

REFERENCES

Aaron SD., Dales RE., Pham B. 1998. Management of steroiddependent asthma with methotrexate: a meta-analysis of randomized clinical trials. *Respir Med.*, 92:1059–1065.

- American Lung Association. Take the asthma control test and share the results with your doctor. http://www.asthma control.com/ Date last updated: January 8, 2001. Date last accessed: July 15, 2007.
- Bateman ED, Hurd SS, Barnes PJ *et al.* 2008. Global strategy for asthma management and prevention: GINA executive summary. *Eur Respir J.*,31:143-78.
- Bentley AM., Kay AB., Durham SR. 1997. Human late asthmatic reactions. ClinExp Allergy, 31:71–86.
- Bernstein IL., Bernstein DI., Dubb JW., et al. 1996. Placebocontrolled multicenter study of auranofin in the treatment of patients with corticosteroid-dependent asthma. Auranofin Multicenter Drug Trial. J Allergy Clin Immunol., 98:317–324.
- Boulet LP., Boulet V., Milot J. 2002. How should we quantify asthma control? A proposal. *Chest.*, 122:2217–2223
- Boulet LP., Lemiere C., Archambault F. *et al.* 2006. Smoking and asthma: clinical and radiologic features, lung function, and airway inflammation. Chest129:661–668.
- Centers for Disease Control and Prevention (CDC). Cigarette smoking among adults–United States, 1998. MMWR Morb Mortal Wkly Rep 2000;49:881–884.
- Davies H., Olson L., Gibson P. 2000. Methotrexate as a steroid sparing agent for asthma in adults. *Cochrane Database Syst Rev.*, CD000391
- Eastell R., Reid DM., Compston J. *et al.* 1998. A UK Consensus Group on management of glucocorticoid-induced osteoporosis: an update. *J Intern Med.*, 244:271–292.
- Eisner MD. 2002. Environmental tobacco smoke and adult asthma. *Clin Chest Med.*,23:749–761.
- Green RH., Brightling CE., McKenna S. *et al.* 2002. Asthma exacerbations and sputum eosinophil counts: a randomised controlled trial. Lancet, 360:1715–1721.
- Green, RH., Brightling CE., McKenna S., *et al.* 2002. Asthma exacerbations and sputum eosinophil counts: a randomised controlled trial. *Lancet.*, 360:1715–1721
- Hasegawa T., Koya T., Sakagami T. *et al.* 2012. Asthma control and management changes in Japan surveyed using questionnaire *Intern Med.*, 51:567-74.
- Hasegawa T., Suzuki E., Terada M. *et al.* 2005. Improvement of asthma management in actual practice consistent with prevalence of anti-inflammatory agents. -Based on questionnaire surveys in Niigata Prefecture, Japan from 1998to 2002 -. *Allergol Int.*, 54:555-63.
- http://www.ginasthma.com/guidelineitem.asp?? (Accessed 16 December 2006).
- Jacoby DB. 2002. Virus-induced asthma attacks. JAMA., 287:755–761.
- Juniper EF., Guyatt GH., Ferrie PJ., et al. 1993. Measuring quality of life in asthma. Am J Respir Dis., 147:832–838.
- Liu AH., Zeiger R., Sorkness C. *et al.* 2007. Development and cross-sectional validation of the Childhood Asthma Control Test. *J Allergy ClinImmunol.*, 119:817–825.
- Liu AH., Zeiger R., Sorkness C. *et al.* 2007. Development and cross-sectional validation of the childhood Asthma Control Test. *J Allergy ClinImmunol*.119 (4) :817-25.
- Lock SH., Kay AB., Barnes NC. 1996. Double-blind, placebocontrolled study of cyclosporin A as a corticosteroidsparing agent in corticosteroid-dependent asthma. *Am J Respir Crit Care Med.*, 153:509–514.
- Mendoza MMR., Cruz BO., Guzman-Banzon AV., Ayuyao FG., De Guia TS. 2007. Comparative Assessment of Asthma Control Test (ACT) and GINA Classification including FEV1 in predicting asthma severity. *Phil Heart Center J.*, 13 (2) :149-154

- Moore PL. 2007. Practice management and chronic obstructive pulmonary disease in primary care. *Am J Med.*, 120:S23-7.
- Moy ML., Israel E., Weiss ST., *et al.* 2001. Clinical predictors of health-related quality of life depend on asthma severity. *Am J RespirCrit Care Med.*, 163:924–929.
- Naseh Sigari, Nader Sigari, HoomanGhasri, Ezzat Rahimi,*et al.* 2011. National Research Institute of Tuberculosis and Lung Disease, Iran. *Tanaffos.*, 10 (4) : 49-53
- Nathan RA., Sorkness CA., Kosinski M. 2004. Development of the asthma control test: a survey for assessing asthma control. *J Allergy Clin.mmunol.*, 113 (1) :59-65.
- Nathan RA., Sorkness CA., Kosinski M. *et al.* 2004. Development of the asthma control test: a survey for assessing asthmacontrol. *J Allergy ClinImmunol.*,113:59-65.
- National Asthma Education and Prevention Program, 2007. Expert panel report III: Guidelines for the diagnosis and management of asthma. Bethesda, MD: National Heart, Lung, and Blood Institute,. (NIH publicationno.08-4051) www.nhlbi.nih.Gov/guidelines/asthma/asthgdln.htm (Accessed on September 01, 2007).
- Nguyen VN., Chavannes N., Le LT., Price D. 2012. The Asthma Control Test (ACT) as an alternative tool to Global Initiative for Asthma (GINA) guideline criteria for assessing asthma control in Vietnamese outpatients. *Prim Care Respir J.*, 21:85-9. (pubmed,ivsl)
- Ohta K., Yamaguchi M., Akiyama K. *et al.* 2011. Japanese guideline for adult asthma. *Allergol Int.*, 60:115-45.
- Pearce N., Douwes J., Beasley R. 2000. Is allergen exposure the major primary cause of asthma? Thorax55:424–431.
- Pizzichini MM., Popov TA., Efthimiadis A. et al. 1996. Spontaneous and induced sputum to measure indices of airway inflammation in asthma. Am J RespirCrit Care Med., 154:866–869
- Rakes GP., Arruda E., Ingram JM. *et al.* 1999. Rhinovirus and respiratory syncytial virus in wheezing children requiringemergency care. IgE and eosinophil analyses. *Am J RespirCrit Care Med.*, 159:785–790.
- Reddel HK., Taylor DR., Bateman ED. *et al.* 2009. An official American Thoracic Society/European Respiratory Society statement: asthma control and exacerbations: standardizing endpoints for clinical asthma trials and clinical practice. *Am J RespirCrit Care Med.*, 180:59-99
- Roberts NJ., Smith SF., Partridge MR. 2011. Why is spirometry underused in the diagnosis of the breathless patient: a qualitative study. *BMC Pulm Med.*, 11:37.
- Schatz M., Mosen DM., Kosinski, M. 2007. Validity of the Asthma Control Test completed at home. *Am J Manag Care.*, 13 (12):661-7.
- Smith AD., Taylor DR. 2005. Is exhaled nitric oxide measurement a useful clinical test in asthma?. Curr Opin Allergy ClinImmunol., 5:49–56.
- Stemple D., Williams on A., Stanford R. 2005. Comparative assessment of asthma control with both ACT and Spirometry in subjects attending community events. *J Allergy ClinImmunol.*, 115:S216
- Thomas M., Kay S., Pike J. *et al.* 2009. The Asthma Control TestTM (ACT) as a predictor of GINAguideline-defined asthma control: analysis of a multinational cross-sectional survey. *Prim Care Respir J.*, 18 (1):41-9.
- Thomas M., Kay S., Pike J., *et al.* 2009. The Asthma Control Test TM (ACT) as a predictor of GINA guideline-defined asthma control: analysis of a multinational cross-sectional survey. *Primary Care Respiratory Journal.*, 18 (1): 41-49.

- Vega JM, Badia X, Badiola C. 2007. Validation of the Spanish version of the Asthma Control Test (ACT). J Asthma., 44 (10) :867-72
- Williams JV., Harris PA., Tollefson SJ. *et al.* 2004. Human metapneumovirus and lower respiratory tract disease in otherwise healthy infants and children. *N Engl J Med.*, 350:443–450.
- Yssel H., Abbal C., Pene J., Bousquet J. 1998. The role of IgE in asthma. *Clin Exp Allergy*, 28:104–109; discussion 117– 118.
- Zhou X., Ding FM., Lin JT., Yin KS. 2009. Validity of asthma control test for asthma control assessment in Chinese primary care settings. *Chest.*, 135:904-10.
