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The AAP Asthma Guideline for Pediatric Primary Care

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In partial fulfilment of the requirements for the Doctor of Nursing Practice

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Table of Contents

Abstract	1
Introduction	2
Background	3
Problem Statement	4
Purpose Statement	4
Project Question	5
Project Objectives	5
Search Terms	6
Literature Review	6
Theoretical Framework	11
Project Design	16
Intervention/Project Timeline	21
Ethics/Human Subjects Protection	23
Plan for Analysis/Evaluation.	23
Analysis of Results	25
Discussion of Findings	26
Significance/Implications for Nursing	28
Limitations	29
Dissemination	29

Project Sustainability	30
References	31
Tables	
Table 1	32
Table 2	44
Table 3	45
Table 4	47
Table 5	48
Appendices	
Appendix A : The Plan-Do-Study-Act Model	49
Appendix B : The AAP Asthma Guideline (2013)	50
Appendix C : Chart Review Tools	52
Appendix D : Inhaler Spacer Factsheet	55
Appendix E : Asthma Action Plan	56
Appendix F : Presentation for the AAP Asthma Guideline Implementation	58
Appendix G : Permission Letter for Project	59

Abstract

Introduction: The purpose of this quality improvement project was to implement the AAP Asthma Guideline in a primary care setting. This project aimed to improve the quality of asthma care and decrease ER/urgent care transfer for pediatric asthma exacerbation.

Method: This project used a pre-post chart review design by educating four primary care providers were educated on the use of the AAP Asthma Guideline to identify and treat the pediatric population between 2-17 years with asthma at a pediatric primary clinic.

Result: Provider compliance with the guideline was analyzed by two-tailed Wilcoxon Signed Rank Test and ER/Urgent care transfer rate due to asthma exacerbation was analyzed by Odds Ratio analysis. During the project, the providers' compliance rate with the AAP guideline significantly increased. The providers achieved patient-centered education by communicating with their caregivers through asthma action plan. Improving providers' documentation in the Electronic Medical Record facilitated collaborating and communicating among team members which are essential to gain successful outcomes in the delivery of health care. However, the assumption that the AAP asthma guideline implementation will lead to lower the ER/urgent care transfer for asthma exacerbation could not be objectively demonstrated in this project since only a small sample size in a short timeframe was examined.

Introduction

Asthma is a complex environmental and genetic illness that affects individuals across the globe. It is one of the most prevalent chronic illnesses affecting approximately 6 million children in the United States (Zahran et al., 2018). Acute asthma exacerbations are high in children and are one of the leading causes of frequent childhood hospitalizations. Approximately, 60% of children with asthma have one or more acute exacerbations annually, which cost the US government more than \$6.1 million (Pardue et al., 2016). In 2016, more than 6.1 million children in the US were diagnosed with asthma, which accounted for 135,000 hospitalizations annually and 383,000 deaths as of 2015 (Sawick et al., 2016). Evidence-based assessment tools can guide healthcare providers in measuring the asthma severity, starting the treatment process, and setting up a functional partnership with the children and their caregivers to minimize asthma exacerbation and to establish asthma control.

Asthma exacerbation assessment helps healthcare providers determine the priorities for all patients with asthma depending on the severity level. After the assessment, healthcare providers shift their concern to the classification of asthma severity to start and monitor control over time and ensure that the treatment process can be changed (Ban et al., 2018). Carrying out simple assessments improves the management of asthmatic patients both in the Emergency Room (ER) and primary care setting. The management of mild, moderate, and acute severe asthma exacerbations in primary care needs an effective assessment tool for measuring the severity of asthma that makes it possible to offer efficient treatment.

Background

Asthma is one of the predominant chronic disorders that affects over 6.1 million children in the US and leads to high numbers of preventable mortality and morbidity (CDC, 2018). The issue of acute asthma exacerbation assessment was first recognized in 1991 when the National Asthma Education and Prevention Program (NAEPP) published the strategies for diagnosis and asthma management (National Institutes of Health [NIH], 1991). This classification was divided into mild, moderate, and severe groupings based on several symptoms such as the frequency of exacerbations and school attendance for children. The severity assessment also included the level of exercise tolerance and pulmonary function tests before the beginning of asthma treatment processes (Yawn et al., 2019). Most importantly, however, the domains of severity and control were measured in terms of future risks and impairment levels. In 1997, NAEPP introduced moderate-persistent, mild-intermittent, mild-persistent, and severe-persistent asthma based on peak expiratory flow, frequency of symptoms, and peak-flow variability within 24 hours (NIH, 1997). During this time, children less than five years were only assessed using the symptom data alone. National Asthma Education and Prevention Program initiated the first classification of severe asthma assessment although how the severity was measured and modified remained unclear for some time.

Today, many health practitioners use different strategies to assess asthma exacerbations among children. The exacerbations categorized as mild, moderate, severe, or very severe are based on symptoms, lung function, and physical examination parameters among other factors (NIH, 2007). Children suffering from severe asthma exacerbation exhibit signs like inspiratory and expiratory wheezing, cyanosis, and retractions of the chest wall (Asthma and Allergy Foundation of America, 2017). However, one sign is not enough indication of asthma severity.

but an combination of symptoms can provide more accurate and valid data bearing in mind the connection between pathophysiology of asthma medical features (Magpuri, Dixon, McCorkle, & Crowley, 2018). This fact has triggered the development of different assessment methods to quantify the severity of asthma exacerbations. Assessment of asthma control begins with the history of past asthmatic exacerbations and challenges placed on the patient's daily activities such as sporting activities, use of medications, devotion to therapy, and other factors that may complicate care.

Problem Statement

Use of evidence-based assessment tools is a strategy that can guide healthcare providers in measuring the level of asthma severity, starting the treatment process, and deciding the nature of pediatric patients with severe asthma exacerbation (O'connor et al., 2015). Many patients who experience asthma exacerbation seek help from primary care facilities (Price et al., 2017) and, therefore, utilizing evidence-based tools that guide healthcare providers is a necessity (Patel et al., 2019). This project will take place at a primary care clinic that frequently treats asthma exacerbations. The clinic does not currently enforce specific asthma guidelines for providers to use, instead allowing providers to choose methods they prefer. The use of evidence-based assessment tools enhances patient wellbeing, quality of treatment, and intensifies the movement of asthmatic patients in the primary care setting. This project explores the AAP Asthma Guideline as an evidence-based guideline tool to assess the severity of asthma exacerbations among children in the primary care setting and to develop an effective treatment plan.

Purpose Statement

Primary care providers recognize the need to offer high quality care to all patients. The purpose of this OI project is to educate primary care providers on the use of the AAP Asthma

Guideline to identify and treat the pediatric population between 2-17 years with asthma and

evaluate if that improve the providers' compliance rate with the evidence-based asthma guideline

and reduces the number of ER/urgent care visits due to asthma attacks with a period of six

weeks.

Project Question

Among asthmatic patients between 2-17 years of age, does the use of the AAP Asthma

Guideline by healthcare providers to identify and treat the population increase the providers'

compliance with the guideline and reduce the number of ER/urgent care transfers due to asthma

exacerbations within the next six weeks compared to patients treated by healthcare providers

without the use of the AAP Asthma Guideline?

P=Population of patients: Healthcare providers in pediatric clinic.

I=Intervention: Use of the AAP Asthma Guideline to asthmatic patients between 2-17 years of

age.

C=Comparison: The compliance rate with the AAP Asthma Guideline and the number of

incidence of ER/urgent care transfers due to asthma exacerbations of patients treated by

providers prior to the implementation of the use of the AAP Asthma Guideline.

O=Outcome: Increased the providers' compliance rate with the guideline and reduced ER/urgent

care visits for asthma attacks.

T=Timeframe: Six weeks.

Project Objectives

In the timeframe of this DNP project, the project site will:

5

- Educate providers regarding implementation of the AAP Asthma Guideline prior to implementation of the DNP project.
- b) Implement the AAP Asthma Guideline to treat asthmatic patients between 2-17 years of age in the primary pediatric clinic.
- c) Increase the providers' compliance with the using the guideline.
- d) Reduce ED/urgent care transfers for asthma exacerbation within the next 6 weeks.

Search Terms

An electronic search using databases including CINAHL, PubMed, Medscape, TUN-Jay Sexter Library, Google Scholar, UpToDate, and other governmental regulatory agencies such as the NIH and CDC were used. Search terms included "asthma", "asthma exacerbation", "pediatric", "primary care", "asthma guideline", and "the AAP Asthma Guideline" were used. Inclusion criteria included literature published from 2014 to 2019, meta-analysis, systematic reviews, randomized control trials, and cohort studies. Studies published prior to 2014 were considered only if historically relevant. Exclusion criteria included studies not relevant to the primary care environment. There were 40 eligible articles identified through the search, and 30 were selected for appraisal and literature review.

Literature Review

This review of the literature was conducted to identify the following:

- 1. Significance and risk factors of asthma exacerbations.
- 2. Prevention and management of asthma exacerbations in children.
- 3. Asthma severity assessments in children with acute asthma exacerbation.

Impact and Significance

Asthma exacerbation rates among asthmatic children in the US increased from a rate of 48 percent among children 18 years old and under in 2014 to 54 percent in 2016 (CDC, 2016). In 2015, a multicenter study in the US showed that the rate of ER visits with first-listed diagnosis of asthma were most frequent among children aged one to four years (14.6 visits per 1,000 population) and five to nine years (12.3 visits per 1,000 population) (McDermott, Stocks, & Freeman, 2018). Among one year and older pediatric age groups, asthma was the second most common category of first-listed ER conditions in 2015 (McDermott, Stocks, & Freeman, 2018). These statistics suggest that the identification of asthma exacerbation and treatment of asthma are crucial in the primary care setting to prevent ER visits.

Risk Factors for Asthma Exacerbations.

Race, ethnicity, and socioeconomic status. In the United States, asthma exacerbations are more prevalent among children who are African American, Puerto Rican, or economically undeserved (Frieden, 2013). Non-Hispanic (NH) blacks and Hispanics have a high risk of ER visits for asthma attack, compared with NH whites (Mehta et al., 2013).

Viral Infections. 85% of asthma attack in children is derived from respiratory viral infections including human rhinovirus (HRV) (Saraya et al., 2014). However, other respiratory viruses including respiratory syncytial virus (RSV), parainfluenza, influenza, and others often cause repetitive wheezing and severe asthma exacerbations in young children (Stein et al., 1999). Deunasa et al. (2016) reported that HRV was most commonly detected (73%), followed by influenza A (27%) and RSV (7.7%) in 175 children age 2 to 15 years. Also, multi-infection with two or more viruses was identified in 15 % of the participants (Duenas et al., 2016).

Air Pollution. Trasande et al. (2005) reported severe asthma exacerbation in children is caused by traffic-related air pollution. Estimated nine million dollars spent on health care for pediatric asthma exacerbations at two districts in California, Riverside and Long Beach, in 2012 (Brandt et al., 2012).

Allergens and Indoor Pollutants. In the United States, compared to children living in suburban areas, those living in urban areas in the Northeast are commonly exposed to certain allergens such as dust mites, cockroach, and mouse. Sensitized children are prone to develop poor asthma control and severe asthma exacerbations by exposure to allergens either indoors or outdoors (Kanchongkittiphon et al., 2015).

Psychosocial Stress. Chronic psychosocial stress is a cause of severe asthma exacerbations (Rosenberg et al., 2014). The potential mechanism for the undesirable effects of stress related to severe asthma attack includes a lack of adherence with controller medications, lower therapeutic responses (Brehm et al., 2015).

Poor Adherence to Treatment. Poor adherence to controller medications was significantly related to severe asthma exacerbations in pediatric population (Engelkes et al., 2015).

Current Management for Asthma Exacerbation.

An exacerbation is identified as a worsening in symptoms characterized by a change from the patient's previous status; occasionally the patients with the worsening symptoms are diagnosed as asthma. Optimal management of asthma exacerbation is essential such as self-management with an asthma action plan and appropriate escalation of intervention in primary care (NAEPP, 2007).

All asthma treatment guidelines consider the prevention of asthma exacerbations as an essential part of asthma control. The recommendations are following the AAP Asthma Guideline (2013) that is from the National Asthma Education and Prevention Program's (NAEPP's) expert panel (2007), and the NAEPP is part of the National Heart, Lung, and Blood Institute (NHLBI). The AAP Asthma Guideline (2013) set classification and management of asthma based on age group. The classification of the guidelines is based on the level of the children's risk including occurrence of acute severe exacerbations within the last 12 months and impairment including daily symptoms within the last 4 weeks. Based on risk and impairment, asthma is classified into intermittent, mild persistent, moderate persistent, and severe persistent. Control should be assessed by evaluating similar assessments and is classified as well controlled, not well controlled, or very poorly controlled. Following the classification of asthma severity and level of control, the AAP Asthma Guideline (2013) state taking a stepwise approach to pharmacologic treatment, which indicates increasing medications until asthma is controlled and adjusting medications when possible to reduce side effects.

The principal goals for managing asthma exacerbations include correction of significant hypoxia, an immediate reversal of airflow obstruction, and reduction of future recurrence. Short-acting inhaled β -agonists (SABAs) are the base of the treatment of acute asthma exacerbations. Systemic glucocorticoids should be considered for deterioration in symptoms and/or lung function after the first treatment with inhaled beta-agonists (AAP, 2013). Ganaie et al. (2016) reported appropriate administration of oral glucocorticoids for severe asthma exacerbations is recommended as the most effective intervention for decreasing ER visits and hospitalizations for asthma attacks. Additionally, the AAP Asthma Guideline (2013) recommends ensuring adequate baseline controller therapy including inhaled corticosteroids (ICSs) might be indicated to reduce

the recurrence of asthma exacerbation. ICSs improve lung function, reduce the incidence of asthma-related hospitalizations, decrease ER visits, and restrict the need for oral corticosteroids (Chauhan et al., 2012). The AAP Asthma Guideline (2013) recommends the intervention with inhaled corticosteroid/long-acting β -agonists (ICS/LABAs) for uncontrolled asthma with ICSs and/or LTRAs (AAP, 2013). However, severe asthma exacerbations continue in spite of helpful controller medications. According to the AAP Asthma Guideline (2013), the subspecialist consultation is necessary if 1) 0 to 4 years and step 3 care or higher is required, 2) 5 years or older and step 4 care or higher is required, and 3) difficulty in achieving or maintaining asthma control.

The AAP Asthma Guideline update (2019) reported the two major goals in managing asthma in children. First, the asthma management is for reducing impairment including supporting a good quality of life, normal activities without limitation, and decreasing missed school days. Second, the management is for reducing risk including staying children off systemic corticosteroids, decreasing ER/hospitalization, and supporting healthy lung function to avoid airway remodeling that is connected to chronic inflammation.

Addressing the Problem with Current Evidence

Even though the evidence-based asthma guidelines exist, unfortunately, asthma is related to significant morbidity and the intolerable number of preventable deaths (Booker, 2014).

Asthma related medical care and missed school continue to burden our communities, costing more than \$80 billion each year (Nurmagambetov et al., 2018). Prevention of asthma exacerbations has been placed as a crucial component of establishing best possible asthma control. The treatment of asthma is based on an assessment of severity and asthma control.

Asthma disproportionately affects minorities and socioeconomically disadvantaged children.

Black children have the highest asthma morbidity and mortality rates of any US group, with twice the ER visit/hospitalization rate and 9 times the mortality rate as white children. To bridge these gaps, it is important to practice guideline-based management with accurate assessment of asthma severity and control.

Poor adherence to controller medications for asthma leads to severe asthma exacerbations, including ER visits and hospitalizations (Williams et al., 2011). Poor adherence to controller medications might be caused by provider underdiagnosis or undertreatment of asthma, or patients' poor compliance with medications (Fischer et al., 2010). Poor adherence to ICSs accounts for approximately 60% of asthma-related hospitalizations (Williams et al., 2007).

Evidence gaps.

Despite of evidence-based treatments and guideline-based care, asthma remains a significant public health problem and significant disparities in care. To reduce overall asthma morbidity and its racial and ethnic disparities, it is essential to support efforts to prevent asthma, build a capacity for integrated primary care, improve the identification of at-risk children, and remove barriers to guideline-based care.

Theoretical Framework

The theoretical framework to guide this DNP project was created by Dr. Deming who recognized organizations should focus on constant improvement and collaborate as a team. The Plan-Do-Study-Act (PDSA) cycle is utilized to assess Quality Improvement (QI) ideas (Byrne, Gang, & Carr, 2015) by developing, testing, and implementing new ideas (Gilliam & Siriwardena, 2013). The PDSA cycle is comprised of four cyclical, repeating phases: Plan, Do, Study, and Act (see Appendix A).

Historical Development of the Theory

The PDSA cycles introduced by Edward Deming which includes the four stages used for testing and implementing (Deming, 1919). The PDSA cycle is a powerful tool for achieving improvements in quality and safety and has been commonly used by healthcare organizations to accelerate healthcare processes and desirable results. The practical concepts of PDSA cycles recommend examining the selecting changes on a small scale if they produce QI and expand the tests to larger samples (IHI, 2016). These rapid assessments allow healthcare organizations to modify the change easily following analysis to support desirable results that are recognized (Damschroder et al., 2009).

Applicability of Theory to Current Practice

The concepts of QI have been implemented in healthcare to minimize errors and variation in results (Christoff, 2018). PDSA Methodology is commonly utilized for QI and PDSA cycles provide a frame for iterative testing of the changes to achieve the quality of system. Approaching to QI in healthcare is supported by modification on care systems. PDSA cycles aim at the crux of change, the conversion of goals and plans into actions (Taylor, 2014)

Frisina et al (2019) performed QI-interventions to fulfil the Healthy People (HP) 2020's influenza vaccination goal of 90% among people working in healthcare settings. The QI interventions were focused on staff education, increased accessibility to influenzas vaccines, and improving communication with staff during multiple PDSA cycles. The researchers described QI-interventions remarkably improved the overall vaccination rates from 70.2% (2011–14 influenza seasons) to 84.9% (2014–15 influenza season) in PDSA 1, and 91.1% (2015–16

influenza season) in PDSA cycle 2. This research reported that using PDSA could perform as a method to improve patient outcomes by QI-interventions.

Bansal et al (2016) also utilized PDSA cycles in their study on communication and cancer. Using the PDSA cycle, the researchers reported that educating in the communication of goal of chemotherapy and utilization of checkboxes in progress note template could achieve competency in communication of goal of therapy in cancer patients. Improved patient satisfaction rates were attained on obtaining adequate information and reason for which they were given chemotherapy, and readiness of oncologists to listen, answer questions, and explain things thoroughly to them.

Hendriksma (2017) performed a QI program based on PDSA cycles in the safety net clinic to improve the quality of care in volunteer healthcare professionals. The formalization of the QI program in the safety net clinic during this PDSA cycles provides a strong foundation from which to launch the next PDSA cycle focusing on greater volunteer involvement (Hendriksma, 2017).

Major Tenets

Plan. In the plan stage, a change designed for improvement is recognized. An organization planned to improve with a distinct and solid intention (IHI, 2016). The plan should be completed at a determined time for the specific population who will be affected (IHI, 2016). Complying with the plan is essential; so is distributing the people and resources required to achieve the plan (IHI, 2016). In 2001, the Institute of Medicine (IOM, 2001) published a second report, Crossing the Quality Chasm: A New Health System for the 21st Century, which indicates six comprehensive "Aims for Improvement" for health care. First, we should prevent patients

from injuries from the care for help (IOM, 2001). Second, we should coordinate care with evidence-based science to prevent redundant care and inappropriate care (IOM, 2001). Third, we should respect the individual and their choices as patient-centered service (IOM, 2001). Fourth, we need to avoid unnecessary waiting for patients and their caregivers (IOM, 2001). Fifth, it should be efficient (IOM, 2001). At last, we should eliminate racial and ethnic gaps in health outcomes (IOM, 2001). Once the aim is recognized, it is necessary to make projections about what will occur and why (IOM, 2001). Also, it is important to create a plan to assess the change including who, what, when, where, what information required to be obtained (IOM, 2001).

Do. In the do stage, tested the change is identified. Once a team has chosen a plan and goal, developed criterions to verify whether a change contributes to an improvement, the following step is to carry out a change in the real worksite (IHI, 2016). The PDSA cycle is a simple tool for evaluating a change by planning, carrying it out, discovering the results, and complying with what is learned (IHI, 2016). This is the scientific strategy that leads to taking practical action to deal with the issue (IHI, 2016). The purpose to test changes are to assess the improvement from the change, to evaluate whether the suggested change will succeed in the real situation, to examine which combinations of changes will have the favorable results on the significant measures of quality, to estimate costs, social effect, and side effects from a suggested change, and to reduce resistance from application (IHI, 2016).

Study. In the study stage, the success of the change is examined. Evaluating the analysis of the data concurrently could bring remarkable quality improvements (THA, 2019). Concurrent data enables a healthcare facility to recognize the chance to announce or adjust an issue in real-time (THA, 2019). After compared the data to the projections, the next step is to outline and review what was learned (IHI, 2016). This phase provides the researchers to decide if the

changes are required to achieve the outcome successfully (THA, 2019). The data could be reviewed by checklists, direct monitoring, or surveys (THA, 2019).

Act. In the act stage, adaptations are identified and introduced to the next steps to inform a new cycle. Depending upon what was discovered from the test, it is necessary to decide what adjustments should be considered (IHI, 2016). The process procedures might be modified to more acceptable steps or included more staff training or other strategies to increase the responsibility for the new plan (THA, 2019). At last, a modified plan for the future test will be prepared (IHI, 2016). As a result of the act phase, the PDSA cycle makes a new beginning until the result has been fulfilled (THA, 2019).

Theory Application to the DNP Plan

Many patients who experience asthma exacerbation seek help from primary care facilities (Price et al., 2017) and, therefore, utilizing evidence-based tools that guide healthcare providers is a necessity. Evidence-based assessment tools can guide healthcare providers in measuring the asthma severity, starting the treatment process, and setting up a functional partnership with the children and their caregivers to minimize asthma exacerbation and to establish asthma control. The PDSA model served a theoretical framework that guided the implementation and evaluation of the AAP Asthma Guideline in a pediatric primary clinic serving pediatric populations.

The PDSA model was utilized to implement this DNP Project at the host site. As part of the Plan phase of this PDSA cycle, the AAP Asthma Guideline was selected to collaborate with the pediatric clinic providers. The AAP Asthma Guideline establishes a standard process to assessment and management of pediatric patients with asthma exacerbation. During the Do phase, the procedure the AAP Asthma Guideline was educated to the providers in a pediatric

clinic. Throughout the Study phase, the documentation compliance data, as well as the outcomes from the chart review, were analyzed. Documentation compliance was compared for two sixweek periods: before this PDSA cycle and after the implementation of this PDSA cycle. The hope is that analysis of the outcome will reveal a reduction of ER/urgent care transfer for asthma exacerbation during the six weeks since the AAP Asthma Guideline implementation. As part of the Act phase, the outcomes reported during this PDSA cycle were reviewed. And the recommendations will be made for future PDSA cycles within the pediatric clinic. The ultimate goal is that the AAP Asthma Guideline will be integrated into the structure and process of the pediatric clinic to ensure protocol sustainability beyond this PDSA cycle.

Project Design

Description of project design

This project consists of QI methods to reduce asthma exacerbation by introducing the AAP Asthma Guideline to asthmatic patients between 2-17 years of age in a pediatric primary care setting. Many pediatric patients who experience asthma exacerbation seek help from primary care facilities (Price et al., 2017) and, therefore, utilizing evidence-based tools in measuring the level of asthma severity to guide healthcare providers is a necessity (Patel et al., 2019). The project lead met with the healthcare providers (N=4) to review the AAP Asthma Guideline for asthmatic children at the pediatric clinic. The lead educated the providers to implement the tool to evaluate asthma for the target population during a 6-week period. Data collected during the 6-week period were analyzed using a chart review tool developed exclusively for this project. The tool will determine the provider's compliance using the AAP Asthma Guideline and the efficacy of the guideline via the percent reduction of ER/urgent care transfer.

Setting

The project setting was a pediatric primary care clinic located in Houston, Texas. The clinic provides well-child checkups, immunizations, lab workup, asthma treatment, and other health services for the pediatric population from newborn to twenty-two years of age. The clinic serves approximately 60 individuals per day. The population served includes approximately 90% African Americans and Hispanics and 10% Caucasians, Middle Eastern, and Asians. At the time of the project, the clinic was supported by 4 providers including 1 board certified pediatricians, and 3 nurse practitioners. The site administrator has approved the project and staffs have been informed. Approval (Appendix G) have been granted from the project's clinic site.

Population of Interest

The direct population of interest are the four providers who implemented the AAP Asthma Guideline. The inclusive criteria are pediatricians and nurse practitioners who currently work for the pediatric clinic and agree to participate in the QI project. There are no exclusion criteria. The provider's patients are an indirect population of interest considering that this project intends to evaluate the percent reduction in ED/urgent care transfers. The inclusive criteria are all the patients with asthma/exercise-induced bronchospasm/asthma exacerbation age 2 to 17 years old at the pediatric clinic during the six weeks period. There are no exclusion criteria based upon sex/gender or race/ethnicity.

Stakeholders

The stakeholders are with an interest in the practice site and the services being offered, which would include the patients, pediatricians, nurse practitioners, medical assistants and managerial staffs. Those identified as key project stakeholders are essentially involved in the

healthcare organization and would be significantly affected by changes to the system. Due to the large number of stakeholders, it was important to encourage effective communication. To promote effective communication and rapport, the goals and plans were shared with all stakeholders before project initiation. Additionally, the stakeholder feedback was requested. Feedback may be used to modify or adapt project implementation methods to achieve the project outcomes more efficiently.

Recruitment methods

Following project approval from the clinic's administrator, the project lead did email/call to recruit the clinic's providers at the project site about the project and answer for the concerns and questions during the week 1. The project lead hosted the AAP Asthma Guideline training for the clinic's providers to the project. A lunch was provided during the training as an incentive for attendance as compensation. All information collected from the providers were kept confidential. Each chart was designated a number and a codebook was generated to record data in Excel worksheets. The data saved on the laptop were password protected. In addition, data collected were stored in Research Electronic Data Capture (REDCap), a secure web-based application to assist data capture for this project. Every effort was made to protect the data collected while accounting for privacy and confidentiality issues (AHRO, 2018).

Tools

Several tools were developed and/or utilized for this DNP Project. Tools include the AAP Asthma Guideline, a chart review tool, and educational materials. Each aspect of the tool development is described further below.

AAP Asthma Guideline. The guideline (Appendix B) is an evidence-based, pediatric asthma guideline that provide accurate assessment of asthma severity and control. The AAP Asthma Guideline is based on the National Asthma Education and Prevention Program's (NAEPP's) expert panel (2007), and the NAEPP is part of the National Heart, Lung, and Blood Institute (NHLBI). Multiple research reported that the benefits of decreased ED visit from the national asthma guideline-based care (NAEPP, 2007; Garro et al., 2011; Lovinsky et al., 2010; Sin et al., 2002; Blais et al., 2004; Ly et al., 2007; Ducharme et al., 2011; Rowe et al., 1999). The AAP Asthma Guideline includes how to assess pediatric patients with an acute asthma exacerbation by recognizing early signs and symptoms of worsening asthma (AAP, 2013). Also, the guideline includes optimal pharmacotherapy options with an asthma action plan (AAP, 2013). The treatment options include an inhaled short-acting beta2-agonist (SABA) bronchodilator, systemic oral corticosteroids (OCS), a daily inhaled corticosteroid (ICS), long-acting beta2-agonist (LABA), and montelukast.

Chart review tools. Chart review tools (Appendix C) were utilized to evaluate the providers' compliance using the AAP Asthma Guideline. The measures include: 1) performing assessment with spirometry only for patients older than 5 years, 2) providing optimal treatment options based on stepwise approach, 3) Inspecting medications, inhaler, and spacer and educating the administration skills and knowledge for self-management, 4) providing asthma action plan based on the asthma severity and level of asthma control. Also, the chart review tools were used to determine if the patients transferred to ER/urgent care during the six weeks period. The tools were reviewed by the project team and reviewed and approved at the project site by stakeholders.

PowerPoint and education materials. The project lead developed training materials for staff. The training materials covered the AAP Asthma Guideline (Appendix B), and chart review tools (Appendix C), and asthma action plan (Appendix E) with patient education (Appendix D). The PowerPoint (Appendix F) guided the training and encouraged participation and feedback from the providers. These tools were reviewed by the project team and reviewed and approved at the project site by stakeholders.

Data collection procedures

The clinic electronic medical record (EMR) databases were accessed with permission to extract charts including the ICD 10 codes for asthma. The specific ICD 10 codes related to asthma include J45.20, J45.21, J45.22, J45.30, J45.31, J45.32, J45.40, J45.41, J45.42, J45.50, J45.51, J45.52, J45.901, J45.902, J45.909, J45.990, J45.991, and J45.998 which are listed on the chart review tools (Appendix C). The project lead reviewed the selected charts to determine providers' compliance with the AAP Asthma Guideline and the number of patients transferred to ER/urgent care pre/post-implementation of the AAP Asthma Guideline. Data collected were stored in Research Electronic Data Capture (REDCap), a secure web-based application to assist data capture for this project.

Providers' compliance with the AAP Asthma Guideline. Each provider that attended training and agreed to implement the AAP Asthma Guideline for six weeks was tracked by the project lead. Chart review tools were utilized to evaluate the providers' compliance using the AAP Asthma Guideline. The total 100 charts were collected, 50 pre-implementation and 50 post-implementation of the guideline to look at the percent compliances per provider before and after.

ED/urgent care transfer. The project lead reviewed the provider's notes to determine if the patients transferred to ER/urgent care due to asthma exacerbation before/during/after the visit during the six weeks period. This allowed a comparison of the number of patients transferred to a higher level of care pre/post-implementation of the AAP Asthma Guideline.

Intervention/Project Timeline

The proposed timeline for the DNP project is as follows:

Week 1

Week/Date	Activity
Week 1	The project lead did email/call to recruit the clinic's providers at
February 21- February 27,	the project site about the project on February 21 and answer for
2020	the concerns and questions until February 26. The site
	administrator has approved the project. Approval (Appendix G)
	has already been granted from the project's clinic site. The
	educational training for the providers was performed by the
	project lead at 10 AM on February 26 th . It covers the AAP
	Asthma Guideline, chart review tools, asthma action plan, and
	inhaler spacer factsheet by using PowerPoint presentation. A
	total of 4 providers including 1 pediatrician and 3 nurse
	practitioners including project lead agreed to participate in the
	project. The project lead collected the providers' email address
	and contact number to communicate and answer any questions
	during the project implementation.

Week 2-6

Week/Date	Activity
Week 2	The project implementation phase continued at the practice
February 28 – March 3, 2020	site. The project lead communicated and received feedback
Week 3	with other providers by email or text message that the
March 4- March 10, 2020	implementation is going well without any change. The project
	lead completed 40 percent of the pre-implementation chart
	review.
Week 4	The project implementation phase continued at the practice
March 11- March 17, 2020	site. The project lead completed 100 percent of the pre-
	implementation chart review which is the total 50 charts.
Week 5	The project lead communicated and received feedback with
March 18- March 24, 2020	other providers by email or text message that the
	implementation is going well without any change.
Week 6	The project implementation phase continued at the practice
March 25- March 31, 2020	site. The project lead completed 100 percent of the post-
	implementation chart review which is the total 50 charts.

Week 7

Week/Date	Activity	
Week 7	The project lead analyzed project findings based on the chart	
April 1- April 7, 2020	review. The evaluation results will be disseminated to	
	stakeholders upon completion.	

Ethics/Human Subjects Protection

The project lead submitted Institutional Review Boards (IRB) determination forms per TUN policy. It is expected that the project will likely fall under the category of TUN quality improvement project which would not require IRB review. IRB is not required by the project site. Asthma management and prevention incorporated into this project are standard clinical procedures and consistent with established AAP Asthma Guideline. There are no direct risks for participants in this project. One potential benefit is that the AAP Asthma Guideline provide appropriate, evidence-based treatment of pediatric asthma exacerbation that gives the providers a blueprint of asthma exacerbation assessment and treatment. Patient's may benefit by learning their asthmatic status and taking the necessary precautions to reduce asthma exacerbation. A lunch was provided during the training as an incentive for attendance. No ethical issues or conflicts of interest have been identified for this project.

The clinic electronic medical record (EMR) databases were entered with permission to extract charts including the ICD codes for asthma. Data collected were stored in Research Electronic Data Capture (REDCap), a secure web-based application to assist data capture for this project. This is to protect the privacy and confidentiality of the participants while being engaged in the project.

Plan for Analysis/Evaluation

This QI project used a pre-post chart review design to evaluate the effectiveness of the AAP Asthma Guideline aimed to improve the providers' compliance rate with the evidence-based asthma guideline and decrease ER/urgent care transfer for pediatric asthma exacerbation. The analysis plan is to assess: 1) percent compliance with the AAP Asthma Guideline per

provider, and 2) the number of patients transferred to ER/urgent care pre/post-implementation of the AAP Asthma Guideline.

To address the first aim of the analysis of the project, patient charts (N=100) were reviewed, 50 pre-implementation and 50 post-implementation of the AAP Asthma Guideline. The AAP Asthma Guideline includes the essential 4 items including assessment, optimal treatment plan, inspection and education, asthma action plan with follow up plan based on the stepwise approach to managing asthma. A pre-implementation chart review between February 27th, 2019, through March 31st, 2019 was conducted by the project lead. A pre- and postimplementation chart review evaluated each providers' charts including the ICD codes relating to asthma diagnosis by utilizing chart review tools. Each provider that attended training and agreed to implement the AAP Asthma Guideline for six weeks between February 21st, 2020, through March 31st, 2020, was tracked by the project lead. Following the data collection, the percent compliances with the AAP Asthma Guideline per provider were analyzed by Wilcoxon Signed Rank Test with 95% confidence intervals. The Wilcoxon test is the appropriate analysis to compare differences derived from the same population when the dependent variable is ordinal or continuous. It is used to assess differences from matched pair designs or repeated measures. The distribution of the differences between the two related groups would be symmetrical in shape.

To address the second aim of the project, the same charts were reviewed by the project lead to determine the number of ER/urgent care transfer pre/post-implementation of the AAP Asthma Guideline. Odds Ratio with a 95% confidence interval was utilized to determine the effects of the AAP Asthma Guideline interventions which may reduce ER/urgent care transfers for asthma exacerbation. The odds ratio is the appropriate analysis to quantify the relationship between an exposure and a result in a case-control study. The exposure would be the AAP

Asthma Guideline implementation and the result would be ER/urgent care transfer due to asthma exacerbation. It was used to assess that the AAP Asthma Guideline implementation might be a protective factor against the ER/urgent care transfer.

Analysis of Results

The implementation of the AAP Asthma Guideline for preventing asthma exacerbation increased the providers' compliance rate of the asthma guideline (Table 2). The providers' compliance rate on the AAP Asthma Guideline increased from an average of 58% during the pre-implementation period to 94% during the post-implementation period. The number of ER/urgent care transfers for asthma exacerbation was also tracked at the practice site during the six-week implementation of the AAP guideline, but insufficient data was found to conduct statistically significant analysis (Table 5).

During the six-week implementation, the clinical providers served 75 asthmatic patients of 1413 patients. The project lead extracted 50 asthmatic patients' charts between 2-17 years of age for each pre-implementation period and post-implementation period (Table 1). The pre-implementation phase was selected from Feb 27th, 2019 to Mar 31st, 2019. The pre-implementation period of the provider 2 and provider 4 are different than others since they did not practice at the project site during the timeline. The post-implementation phase was performed from Feb 28th, 2020 to Mar 31st, 2020.

A two-tailed Wilcoxon Signed Rank Test with 95% confidence intervals was performed. This test assumes dependent samples, assessing before and after observations, and independence of the paired observations. The data showed a significant difference between each providers' compliance rate on the AAP Asthma Guideline at pre-implementation and post-implementation

with p-value (p < .05) (Table 4). The results showed that the implementation of the AAP Asthma Guideline increased the providers' compliance with evidence-based asthma care to patients (Table 2).

For this study, the following are the 4 levels of compliance used: one item is 25%, two items are 50%, three items are 75%, and four items are 100%. The measures used are classifying asthma severity and level if asthma control, providing optimal treatment options based on the stepwise approach, inspecting medications, inhaler, and spacer and educating the administration skills and knowledge for self-management, and providing asthma action plan and follow up plan. This implies that if the participant is compliant with one item, it is a 25% compliance rate, two items is 50% compliance rate, three items is 75 % compliance rate, and all four items is 100% compliance rate. Each provider is categorized depending on the compliance rate, which is based on the number of transactions they process during the period. Also, details on how the researcher reached the percent compliances are included in Appendix C.

Odds Ratio with 95% confidence intervals was performed to examine ER/urgent care transfers for asthma exacerbation. The data failed to show statistically significant to the reduction of ER/urgent care transfers for asthma exacerbation during the six-week implementation period of the AAP guideline (Table 5).

Discussion of Findings

The AAP Asthma Guideline was implemented by four providers for six weeks to identify and treat asthmatic children between 2-17 years of age in the primary pediatric clinic. Before implementation of the DNP project, the educational meeting for the participants executed to cover the AAP Asthma Guideline (Appendix B), chart review tools (Appendix C), inhaler spacer

factsheet (Appendix D), and asthma action plan (Appendix E) by using PowerPoint presentation (Appendix F). The project lead evaluated providers' charts to assess the providers' compliance rate with the AAP Asthma Guideline and ER/urgent care transfer rate for asthma exacerbation to compare between pre- and post-implementation of the AAP guideline. During the project, providers' documentation improved significantly at the site regarding the description of asthma severity and the history of the level of asthma control (Table 3) for initial diagnosis and treatment to achieve asthma control which helped communication between providers about the patients.

Utilizing handouts, patient education regarding inhaler use, and spacer use (Appendix D) was provided for effective medication application. Without the proper technique, patients could not gain the benefit of the therapy or achieve control. Providing an asthma action plan (Appendix E) improved the evidence-based practice of assessing asthma severity level and control of asthma exacerbation. Patients learn self-management skills and maintain long-term care for quality asthma management with the asthma action plan. NAEPP (2007) reported asthma control focuses on both arenas, 1) reducing impairment of symptoms and functional limitations; 2) reducing the risk of future asthma exacerbations.

By implementing the AAP Asthma Guideline, the practitioners not only provided the evidence-based optimal asthma management for the patients but also introduced the skills and tools for the patients and their caregivers to continue with their self-care at home and school. During each visit, providers became more engaged and invested in practice changes because primary care health professionals are at the forefront of care to minimize asthma exacerbation.

Data analysis revealed that the adherence to the AAP resulted in a decrease of asthmarelated ER/urgent care transfers at the project site. However, the assumption that the AAP Asthma Guideline implementation will lead to lower the ER/urgent care transfer for asthma exacerbation could not be objectively demonstrated in this project (Table 5) since only a small sample size in a short timeframe was examined.

Significance/Implications for Nursing

This project is significant to the nursing profession because it addresses the care of asthma among children leading to high numbers of preventable mortality and morbidity. Among one year and older pediatric age groups, asthma was the second most common category of first-listed ER conditions in 2015 (McDermott, Stocks, & Freeman, 2018). The identification of asthma exacerbation and treatment of asthma are crucial in the primary care setting to prevent ER visits. In addition, nurses have key roles to play in improving health outcomes for their patients through education, implementing evidence-based practice, and communicating with others. Throughout the DNP project, providers performed their essential roles to the vulnerable population with asthma. First, the providers' compliance rate significantly increased the guideline-based management for the targeted patients with asthma (Table 4) who are at higher risk for ER visits. Second, providers achieved patient-centered education by communicating with their caregivers and school nurse through asthma action plan. Third, improving providers' documentation in the EMR facilitated collaborating and communicating among team members which are essential to gain successful outcomes in the delivery of health care.

The implications for future practice are to maintain the AAP Asthma Guideline to the project site. It is crucial to implement the evidence-based guideline for providing quality asthma care and reducing asthma exacerbation for the pediatric population. Uncontrolled asthma has major adverse effects on the quality of life, and overall healthcare costs (Holmes, 2017). Nurses in primary care settings play a crucial role to manage chronic disease by providing health

education (Holmes, 2017). As leaders, DNP nurses have the education and competency to play the roles of advocates, educators, and clinicians capable of implementing the evidence-based guideline that serves to improve healthcare outcomes.

Limitations

According to Olufowote (2017), examining limitations of a QI project demonstrates awareness of the problems being examined. Highlighting limitations is essential as it addresses specific issues in the QI project. In this project, the primary limitation is a short time frame as the project lead had six weeks period for the implementation of evidence-based guidelines for asthma care. Gamble et al. (2017) mention that sufficient time is needed for research to be completed in the desired manner as the results of the study depend on the interaction between the researchers and the respondents. During this QI project, there were several barriers in data recruitment and collection techniques such as short timeframe, language barrier, difficulty in measuring AAP Asthma Guideline, and small sample size. The language barrier of Spanishspeaking caregivers which was resolved by providing educational materials in English and Spanish. The comprehensive asthma care based on the AAP Asthma Guideline was challenging to measure, which was determined by creating chart review tools. During the data analysis, the small sample size in the project impacted the failure to verify the assumption that the AAP Asthma Guideline implementation led to a decrease in ER/urgent care transfer due to asthma exacerbation.

Dissemination

Dissemination of findings is an essential element of DNP projects. Marín-González et al. (2016) stated that dissemination of a QI project is necessary as it results in an increased awareness of the project as well as maximizes the impact that a QI project can have in enhancing

the healthcare outcomes of the clients that will benefit from it. In this case, the QI project will be prepared as a PowerPoint presentation and delivered to Touro University Nevada faculty through Zoom online meetings on June 24th, 2020. Correspondingly, the project will be prepared as a poster presentation and delivered to NCNP (National Conference of Nurse Practitioner) 2020 Fall Conference in Las Vegas, NV. The QI project will also be submitted to the DNP repository. Grey (2013) stated that DNP students should submit their final doctoral projects to the DNP repository to have the opportunity to share their final evidence-based projects with other DNP students and faculty. The DNP candidate also plans to present this QI project to stakeholders and providers in current healthcare organizations on 2020 Fall meeting.

Project Sustainability

According to Daneshpour (2015), project sustainability is critical as it allows projects to promote sustainable development goals that shape society. In this project, the implementation of evidence-based asthma guidelines for the pediatric population was essential to the reduction of asthma-related ER/Urgent care visits. The project site has adopted the protocol as part of their daily clinical practice, and it is integrated into the EMR as a simple, easy to use template for checking asthma diagnosis or status, and a template plan including patient education.

Sustainability is necessary for long term changes. Ideally, the project lead had to buy in from providers as their participation is crucial in planning, monitoring, and delivery of project delivery and support processes (Gemünden, 2016). A few supportive efforts encouraged providers to continue applying the guideline in the pediatric clinic, including the following: reminding providers of implementing the AAP Asthma Guideline by using the templates during every monthly meeting for the next 6 months and providing providers of the multiple copies of

the asthma action plan and inhaler spacer factsheet in English and Spanish to promote patients' education.

References

American Academy of Nursing. (2014). Raise the voice: Edge runners. Retrieved from www.aannet.org/edgerunners

us/Documents/medicalhome resources keypointsforasthma.pdf

- American Academy of Pediatrics (AAP). (2013). The AAP Asthma Guideline (2013). Retrieved from https://www.aap.org/en-
- Almutairi, N., Vlahu-Gjorgievska, E., & Win, K. T. (2019). Asthma management application for consumers: Nudging as a feature.
- Arnold, D. H., Gebretsadik, T., Abramo, T. J., Moons, K. G., Harrell, F. E., & Hartert, T. V.(2014). Development and internal validation of a pediatric acute asthma prediction rule for hospitalization. Journal of Allergy and Clinical Immunology in Practice, 3(2), 228-235.
- Arnold, D. H., O'Connor, M. G., & Hartert, T. V. (2015). Acute Asthma Intensity Research

 Score: updated performance characteristics for prediction of hospitalization and lung

 function. Annals of allergy, asthma & immunology: official publication of the American

 College of Allergy, Asthma, & Immunology, 115(1), 69
- Asthma and Allergy Foundation of America (2017). Your guide to managing asthma. Retrieved from https://www.aafa.org/media/1751/your-guide-to-managing-asthma.pdf

Asthma Community Network (2017). Collaborating for Better Care: Strategies for Successful

Partnerships Between Health Plans and Asthma Programs. Available at Asthma

Community Network. Retrieved from

http://www.asthmacommunitynetwork.org/system/files/webinar/pdf/CollaboratingForBet

terCare-WebinarSlides.pdf

- Ban, A. Y. L., Omar, A., Chong, L. Y., Lockman, H., Zaliza, I. Z., Ali, I. A. H., ... & Zim, M. A.M. (2018). Management of asthma in adults in primary care. Malaysian FamilyPhysician, 13(3), 20-26.
- Bansal, M., Abdallah, A., Pennisi, A., Arnaoutakis, K., Makhoul, I., Mehta, P. (2016). Improving Communication on Intent of Chemotherapy Using QOPI Scores and PDSA Cycles, J Canc Educ, 31, 736. https://doi.org/10.1007/s13187-015-0897-x
- Berg, K. T., O'Connor, M. G., Lescallette, R. D., Arnold, D. H., & Stack, L. B. (2015). AAIRS score overview: The acute asthma intensity research score. Academic Emergency Medicine, 22(10), E25–E26.
- Birken, C.S., Parkin, P.C., Macarthur, C. (2004). Asthma severity scores for preschoolers displayed weaknesses in reliability, validity, and responsiveness. J Clin Epidemiol, 57(11), 1177-81.
- Booker, Rachel. (2014). Asthma in children: diagnostic and management dilemmas. Practice Nurse, 44(11), 13-18.
- Brandt SJ, Perez L, Künzli N, Lurmann F, McConnell R. (2012). Costs of childhood asthma due

- to traffic-related pollution in two California communities. Eur Respir J, 40, 363–370.
- Brehm JM, Ramratnam SK, Tse SM, Croteau-Chonka DC, Pino-Yanes M, Rosas-Salazar C, Litonjua AA, Raby BA, Boutaoui N, Han YY, et al. (2015). Stress and bronchodilator response in children with asthma. Am J Respir Crit Care Med, 192, 47–56.
- Byrne, J., Gang, X., & Carr, S. (2015). Developing an intervention to prevent acute kidney injury: Using the plan, do, study, act (PDSA) service improvement approach. Journal of Renal Care, 41(1), 3-8. doi:10.1111/jorc.12090
- Cajigal, S., Wells, K. E., Peterson, E. L., Ahmedani, B. K., Yang, J. J., Kumar, R., ... & Williams, L. K. (2017). Predictive properties of the asthma control test and its component questions for severe asthma exacerbations. The Journal of Allergy and Clinical Immunology: In Practice, 5(1), 121-127.
- Carroll C.L., Sekaran A.K., Lerer T.J., et al. A modified pulmonary index score with predictive value for pediatric asthma exacerbations. Ann Allergy Asthma Immunol, 2005;94(3):355-9.
- Carpenter, D. M., Geryk, L. L., Sage, A., Arrindell, C., & Sleath, B. L. (2016). Exploring the theoretical pathways through which asthma app features can promote adolescent self-management. Translational Behavioral Medicine, 6(4), 509-518.
- Centers for Disease Control and Prevention (2016). Data from yearly National Health Interview Survey. National Center for Health Statistics. US Department of Health and Human Services. Retrieved from http://www.cdc.gov/asthma/most_recent_data.htm.

- Centers for Disease Control and Prevention. (2018). Asthma. Retrieved from https://www.cdc.gov/asthma/inhaler_video/default.htm
- Centers for Disease Control and Prevention. (2018). Morbidity and Mortality Weekly Report:

 Vital Signs: Asthma in Children United States, 2001–2016. Retrieved from

 https://www.cdc.gov/mmwr/volumes/67/wr/mm6705e1.htm.
- Chauhan BF, Ducharme FMAnti-leukotriene agents compared to inhaled corticosteroids in the management of recurrent and/or chronic asthma in adults and children. Cochrane D atabase Syst Rev [serial on the Internet]. 2012 [accessed January 2015];5:CD002314.

 Available from: http://summaries.cochrane.org/
- Chhabra, S. K. (2015). Clinical application of spirometry in asthma: Why, when and how often?

 Lung India: Official Organ of Indian Chest Society, 32(6), 635.
- Christoff, Patricia. (2018). Running PDSA cycles. Current Problems in Pediatric and Adolescent Health Care, 48(8), 198-201.
- Damschroder, L.J., Aron, D.C., Keith, R.E., Kirsh, S.R., Alexander, J.A., Lowery, J.C. (2009).

 Fostering implementation of health services research findings into practice: a

 consolidated framework for advancing implementation science. Implement Sci, 4, 50.
- Daneshpour, H. (2015). Integrating Sustainability into Management of Project. *International Journal Of Environmental Science And Development*, 6(4), 321-325. doi: 10.7763/ijesd.2015.v6.611.

- Dinakar, C., & Chipps, B. E. (2017). Clinical tools to assess asthma control in children.

 Pediatrics, 139(1), e20163438.
- Deming, W.E. (1986). Out of the crisis. Cambridge, MA: Massachusetts Institute of Technology Center for Advanced Engineering Study xiii, 1991;507.
- Duenas Meza E, Jaramillo CA, Correa E, Torres-Duque CA, García C, González M, Rojas D, Hernández A, Páez AM, del Pilar Delgado M. (2016). Virus and Mycoplasma pneumoniae prevalence in a selected pediatric population with acute asthma exacerbation. J Asthma, 53, 253–260.
- Engelkes M, Janssens HM, de Jongste JC, Sturkenboom MC, Verhamme KM. (2015).

 Medication adherence and the risk of severe asthma exacerbations: a systematic review.

 Eur Respir J, 45, 396–407.
- Fischer MA, Stedman MR, Lii J, Vogeli C, Shrank WH, Brookhart MA, Weissman JS. (2010).

 Primary medication non-adherence: analysis of 195,930 electronic prescriptions. J Gen Intern Med, 25, 284–290.
- Frieden TR; Centers for Disease Control and Prevention (CDC). (2013). CDC health disparities and inequalities report: United States, 2013. MMWR Suppl, 62:1–2.
- Frisina, P.G., Ingraffia, S.T., Brown, T.R., Munene, E.N., Pletcher, J.R., Kolligian, J. (2019).

 Increasing influenza immunization rates among healthcare providers in an ambulatory-

- based, University Healthcare Setting. International Journal for Quality in Health Care, 1–6. doi: 10.1093/intqhc/mzy247.
- Gamble, A., Chatfield, S., Cormack, M., & Hallam, J. (2016). Not Enough Time in the Day: A Qualitative Assessment of In-School Physical Activity Policy as Viewed by Administrators, Teachers, and Students. *Journal Of School Health*, 87(1), 21-28. doi: 10.1111/josh.12464.
- Gemünden, H. (2016). Project Governance and Sustainability Two Major Themes in Project Management Research and Practice. *Project Management Journal*, 47(6), 3-6. doi: 10.1177/875697281604700601.
- Gillam, S., & Siriwardena, A. N. (2013). Frameworks for improvement: Clinical audit, the Plan-do-study-act cycle and significant event audit. Quality in Primary Care, 21(2), 123-130
- The Global Initiative for Asthma (GINA). (2019). Resources include Pocket Guide for Asthma

 Management and Prevention, and GINA Report, Global Strategy for Asthma

 Management and Prevention. Retrieved from www.ginasthma.org.
- Gould, C. F., Perzanowski, M. S., Evans, D., & Bruzzese, J. M. (2018). Association of exercise-induced wheeze and other asthma symptoms with emergency department visits and hospitalizations in a large cohort of urban adolescents. Respiratory medicine, 135, 42-50.
- Grey, M. (2013). The Doctor of Nursing Practice: Defining the Next Steps. *Journal Of Nursing Education*, *52*(8), 462-465. doi: 10.3928/01484834-20130719-02.

- Hendriksma, K.J. (2017). A Quality Improvement Program in a Safety Net Clinic Serving

 Vulnerable Populations. Doctoral Projects. 18. Retrieved from

 http://scholarworks.gvsu.edu/kcon_doctoralprojects/18
- Herzog, R., & Cunningham-Rundles, S. (2012). Pediatric asthma: natural history, assessment, and treatment. Mount Sinai Journal of Medicine: A Journal of Translational and Personalized Medicine, 78(5), 645-660.
- Institute for Healthcare Improvement. (2016). Science of improvement: Testing changes. In

 Institute for Healthcare Improvement. Retrieved from

 http://www.ihi.org/resources/Pages/HowtoImprove/ScienceofImprovementTestingChanges.aspx
- Institute of Medicine. (2001). Crossing the Quality Chasm: A New Health System for the 21st Century. National Academies Press.
- Kanchongkittiphon W, Mendell MJ, Gaffin JM, Wang G, Phipatanakul W. (2015). Indoor environmental exposures and exacerbation of asthma: an update to the 2000 review by the Institute of Medicine. Environ Health Perspect, 123, 6–20.
- Langley, G.J. (1996). The improvement guide: A practical approach to enhancing organizational performance. 1st edn. San Francisco: Jossey-Bass Publishers.
- Lougheed MD, Lemiere C, Ducharme FM, et al. (2012). Canadian Thoracic Society 2012

- guideline update: diagnosis and management of asthma in preschoolers, children and adults. Can Respir J, 19, 127.
- Maekawa T, Ohya Y, Mikami M, Uematsu S, Ishiguro A (2018). Clinical Utility of the Modified Pulmonary Index Score as an Objective Assessment Tool for Acute Asthma Exacerbation in Children. JMA, 1(1):57-66.
- Magpuri, A. T., Dixon, J. K., McCorkle, R., & Crowley, A. A. (2018). Adapting an evidence-based pediatric acute asthma exacerbation severity assessment tool for pediatric primary care. Journal of Pediatric Health Care, 32(1), 10-20.
- Marín-González, E., Malmusi, D., Camprubí, L., & Borrell, C. (2016). The Role of Dissemination as a Fundamental Part of a Research Project. *International Journal Of Health Services*, 47(2), 258-276. doi: 10.1177/0020731416676227.
- Mason, D.J., Jones, D.A., Roy, C., Sullivan, C.G., & Wood, L.J. (2015). Commonalities of nurse-designed models of health care. Nursing Outlook, 63(5), 540-553.
- Mehta NK, Lee H, Ylitalo KR. Child health in the United States: recent trends in racial/ethnic disparities. Soc Sci Med 2013;95:6–15.
- National Asthma Education and Prevention Program. (1991). Expert Panel Report (EPR):

 Guidelines for the Diagnosis and Management of Asthma-Summary Report 1991. NIH publication, 91, 3642.
- National Asthma Education and Prevention Program. (1997). Expert Panel Report 2 (EPR-2):

- Guidelines for the Diagnosis and Management of Asthma-Summary Report 1997. NIH publication, 97, 4051.
- National Asthma Education and Prevention Program Expert Panel Report 3 (EPR-3) (2007).

 Guidelines for the Diagnosis and Management of Asthma-Summary Report 2007. J

 Allergy Clin Immunol, 120, S94-138. Retrieved from:

 http://www.nhlbi.nih.gov/guidelines/asthma/asthgdln.htm.
- O'Connor, M. G., Berg, K. Stack, L. B., & Arnold, D. H. (2015). Variability of the acute asthma intensity research score in the pediatric emergency department. Annals of Allergy, Asthma & Immunology, 115(3), 244-245.
- Olufowote, J. (2017). Limitations of Research. *The SAGE Encyclopedia Of Communication Research Methods*, 1-5. doi: 10.4135/9781483381411.n297.
- Pardue J., B., Fleming, G. M., Otillio, J. K., Asokan, I., & Arnold, D. H. (2016). Pediatric acute asthma exacerbations: Evaluation and management from emergency department to intensive care unit. Journal of Asthma, 53(6), 607–617.
- Rosenberg SL, Miller GE, Brehm JM, Celedón JC. (2014). Stress and asthma: novel insights on genetic, epigenetic, and immunologic mechanisms. J Allergy Clin Immunol, 134, 1009–1015.
- Patel, S. J., Teach S. J. (2019). Asthma. Pediatrics in Review, 40, 549-567.
- Saraya T, Kurai D, Ishii H, Ito A, Sasaki Y, Niwa S, Kiyota N, Tsukagoshi H, Kozawa K, Goto

- H, et al. (2014). Epidemiology of virus-induced asthma exacerbations: with special reference to the role of human rhinovirus. Front Microbiol, 5, 226.
- Sawicki, G., Haver, K., Wood, R. A., Redding, G., & TePas, E. (2016). Asthma in children younger than 12 years: Initial evaluation and diagnosis.
- Scarfone RJ, Fuchs SM, Nager AL, Shane SA.(1993). Controlled trial of oral prednisone in the emergency department treatment of children with acute asthma. Pediatrics, 92, 513.
- Sonney, J. T., & Insel, K. C. (2016). Reformulating the Common Sense Model of Self-Regulation. Nursing Science Quarterly, 29(2), 154–159.
- Stein RT, Sherrill D, Morgan WJ, Holberg CJ, Halonen M, Taussig LM, Wright AL, Martinez FD. (1999). Respiratory syncytial virus in early life and risk of wheeze and allergy by age 13 years. Lancet, 354:541–545.
- Taylor, M.J., McNicholas, C., Nicolay, C., Darzi, A., Bell, D., Reed, J.E. (2014).

 Systematic review of the application of the plan-do-study-act method to improve quality in healthcare. BMJ Qual Saf, 23(4), 290-8. doi: 10.1136/bmjqs-2013-001862. Epub 2013

 Sep 11.
- Texas Hospital Association (THA) (2019). PDSA cycle. Retrieved from

 https://www.tha.org/THA-Foundation/Clinical-Initiatives-and-Quality/Initiatives/CAH-QI/CAHQI-Toolkits/PDSA-Cycle
- Trasande L, Thurston GD. (2005). The role of air pollution in asthma and other pediatric

- THE AAP ASTHMA GUIDELINE FOR PEDIATRIC PRIMARY CARE morbidities. J Allergy Clin Immunol, 115:689–699.
- Williams LK, Peterson EL, Wells K, Ahmedani BK, Kumar R, Burchard EG, Chowdhry VK, Favro D, Lanfear DE, Pladevall M. (2011). Quantifying the proportion of severe asthma exacerbations attributable to inhaled corticosteroid nonadherence. J Allergy Clin Immunol,128,1185–1191.e2.
- Wu, AC., Butler, MG., et al. (2015). Primary Adherence to Controller Medications for Asthma Is

 Poor. Annals of the American Thoracic Society, 12(2). Page numbers?
- Yawn, B. P., Brenneman, S. K., Allen-Ramey, F. C., Cabana, M. D., & Markson, L. E. (2019).

 Assessment of asthma severity and asthma control in children. Pediatrics, 118(1), 322-329.
- Zahran, H.S., Bailey, C.M., Damon, S.A., Garbe, P.L., and Breysse P.N. (2018). Vital signs:

 Asthma in children United States, 2001–2016. MMWR Morb Mortal Wkly Rep, 67, 149–155.

Table 1

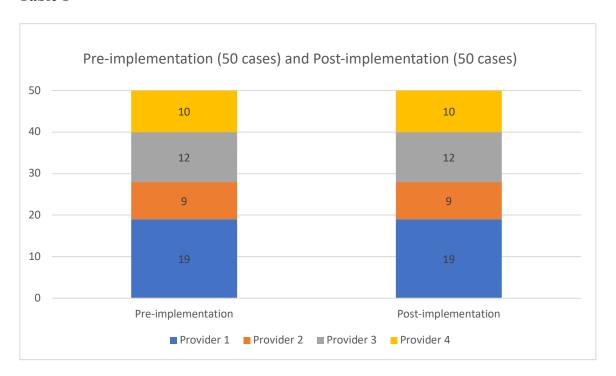


Table 2

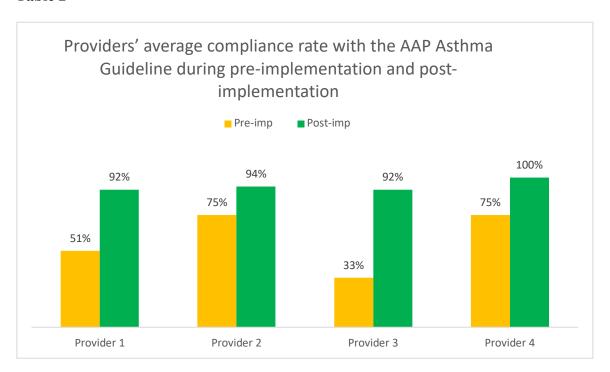
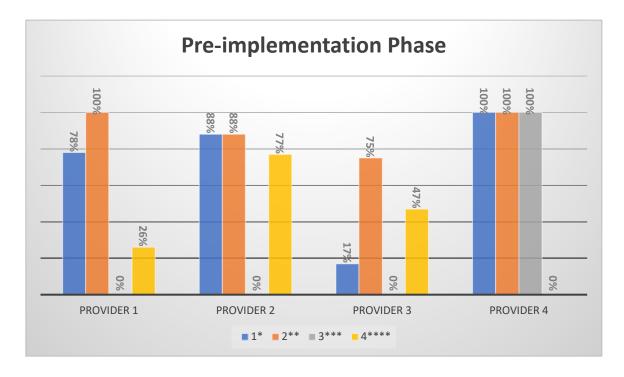
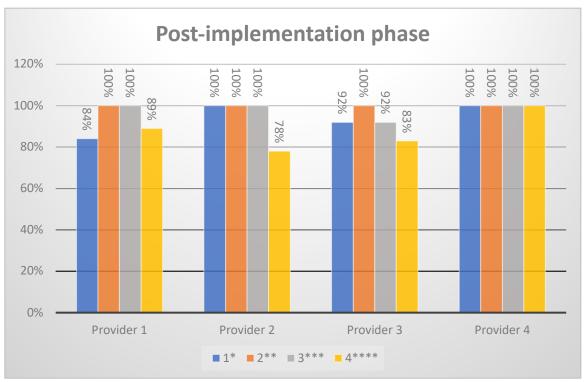


Table 3

Providers' average compliance rate with the four items of the AAP Asthma Guideline





1* is compliance rate with the classifying asthma severity and level of asthma control by using the AAP Asthma Guideline (Appendix B).

2** is compliance rate with the providing optimal treatment options based on the stepwise approach by using the AAP Asthma Guideline (Appendix B).

3*** is compliance rate with the inspecting medications, inhaler, and spacer and educating the administration skills and knowledge for self-management by using the Inhaler Spacer Factsheet (Appendix D).

4**** is compliance rate with the providing asthma action plan and follow up plan by using the Asthma Action Plan (Appendix E).

Table 4

Wilcoxon Signed Rank Test with 95% confidence intervals result						
Providers	A	В	С	D		
W-Value	0	0	0	0		
Mean Difference	-25	-72.73	-32.14	-25		
Sum of pos. ranks	0	0	0	0		
Sum of neg. ranks	171	66	28	55		
Significance of the result $(p < .05)$	Yes	Yes	Yes	Yes		

Table 5

	Transferred	Not transferred	Total
Pre-implementation	1	49	50
Post-implementation	1	49	50
Total	2	98	100
Odds Ratio with a 95% confide	ence interval result	<u> </u>	
Odds ratio	0.4898		
95 % CI	0.0430 to 5.5818		
z statistic	0.575		
Significance level	P = 0.5653		

Appendix A

The Plan-Do-Study-Act Model

What are we trying to accomplish? How will we know that a change is an improvement? What change can we make that will result in improvement? Act Plan Study Do

Figure A. The Plan-Do-Study-Act Model. Adapted from "Model for improvement: Plan-Do-Study-Act (PDSA) Cycles," by The Institute for Healthcare Improvement. Retrieved from http://www.ihi.org/resources/Pages/HowtoImprove/ScienceofImprovementTestingChanges.aspx.

Langley GL, Moen R, Nolan KM, Nolan TW, Norman CL, Provost LP. (2009). The Improvement Guide: A Practical Approach to Enhancing Organizational Performance (2nd edition). San Francisco: Jossey-Bass Publishers.

Appendix B

The AAP Asthma Guideline (2013)

https://www.aap.org/en-us/Documents/medicalhome resources keypointsforasthma.pdf

Key Points for Asthma Guideline Implementation

GOALS OF THERAPY

Reduce Impairment

- · Prevent chronic and troublesome symptoms
- Minimize the need to use SABA for relief of asthma symptoms to ≤2 days/week
- . Maintain (near) normal pulmonary function
- · Maintain normal activity levels

Reduce Risk

- Prevent recurrent exacerbations
- · Provide optimal pharmacotherapy with minimal or no adverse effects
- · Minimize the need for ED visits or hospitalizations

Optimize Health and Function

- Provide initial and ongoing education to patient and family
- · Educate patient and family to recognize and avoid triggers
- Partner with patient and family to identify treatment goals and achieve well-controlled asthma that allows
 patient to fully and safely participate in activities (eg, physical education, recess, sports, etc)
- . Maintain patient's and family's satisfaction with asthma care

ASSESSMENT

- · Classify asthma severity and level of asthma control
- Identify precipitating and exacerbating factors (ie, asthma triggers, including those in the home, school, and child care settings)
- Identify comorbid medical conditions that may adversely affect asthma management
- · Periodically inspect medications, inhaler, and spacer to verify appropriate type
- Regularly assess the patient's and family's knowledge and skills for self-management, including medication administration and inhaler and spacer technique

VISIT Frequency

If asthma is not well controlled: Visits at 2- to 6-week intervals are recommended

If asthma is well controlled: Visits at 3- to 6-month intervals are recommended to monitor how well asthma control is maintained and to adjust medications as necessary

PATIENT AND FAMILY EDUCATION

Incorporate the following into every clinical encounter:

Use a written asthma action plan to share when and how to:

- . Take daily actions to control asthma
- · Adjust medication in response to signs of worsening asthma

Knowledge

- Basic facts about asthma
- · Role of medications

Skills

- Take medications correctly, use appropriate type of inhaler and spacer with proper technique
- Identify and avoid asthma triggers
- Self-monitor level of asthma control
- Recognize early signs and symptoms of worsening asthma
- Seek medical care as appropriate
- Communicate asthma information to school, child care center, and other caregivers

OBTAIN SUBSPECIALIST CONSULTATION IF

(see Table 1 on the following page)

- 0-4 years and Step 3 care or higher is required (may consider consultation at Step 2)
- CONSULTATION IF: 5 years or older and Step 4 care or higher is required (may consider consultation at Step 3)
 - Difficulty in achieving or maintaining asthma control

Information adapted from Texas Children's Health Plan's "Key Points for Asthma Guideline Implementation"



SABA = Short acting beta agonist LABA = Long acting beta agonist ICS = Inhaled corticosteroid OCS = Oral corticosteroid



Table 1: Stepwise approach to managing asthma

Steps	Preferred treatment
Step 1	SABA prn
Step 2	Low dose ICS
Step 3	0.4 years: Medium dose ICS + subspecialist referral ≥5 years: Low dose ICS + LABA or medium dose ICS
Step 4	Medium dose ICS + LABA or montelukast + subspecialist referral
Step 5	High dose ICS + LABA or montelukast + subspecialist referral
Step 6	High dose ICS + LABA or montelukast + OCS + subspecialist referral

Notes

- The stepwise approach is meant to assist—not replace—clinical decision making.
- Before step up, review adherence, inhaler technique, environmental control and comorbid conditions.
- If clear benefit is not observed within 4-6 weeks and/or technique and adherence is not satisfactory, consider adjusting therapy and/or alternative diagnoses.

Acronyms
SABA = Short acting beta agonist
LABA = Long acting beta agonist
ICS = Inhaled corticosteroid
OCS = Oral corticosteroid
ED = emergency department

Table 2: Classifying asthma severity and initiating therapy

Components of severity		Internalities.	Persistent			
		Intermittent	Mild	Moderate	Severe	
	Symptoms	≤2 days/week	>2 days/week	Daily	Throughout the day	
	Nighttime awakenings	0 (≤4 years) ≤2x/month (≥5 years)	1-2x/month (≤4 years) 3-4x/month (≥5 years)	3-4x/month (≤4 years) >1x/week (≥5 years)	>1x/week (≤4 years) Often 7x/week (≥5 years)	
	SABA use for symptoms	≤2 days/week	>2 days/week	Daily	Several times per day	
Impairment	Limitation of normal activity	None	Minor	Some	Extreme	
	Lung function *	FEV1>80% FEV1/FVC>85% (5-11 years) FEV1/FVC normal (≥12 years)	FEV1>80% FEV1/FVC>85% (5-11 years) FEV1/FVC normal (≥12 years)	FEV1>60% FEV1/FVC>75% (5-11 years) FEV1/FVC reduced by 5% (≥12 years)	FEV1<60% FEV1/FVC<75% (5-11 years) FEV1/FVC reduced >5% (≥12 years)	
Risk Exacerbations requiring OCS 0-1/year		0-1/year	≥2/6 months (0-4 years) ≥2/year (≥5 years)	s) **		
Recommended step for initiating therapy ***		Step 1	Step 2	Step 3	Step 3 (≤4 years) Step 3 or 4 (5-11 years) Step 4 or 5 (≥12 years)	

Table 3: Assessing asthma control and adjusting therapy

Co	mponents of control	Well controlled	Not well controlled	Very poorly controlled	
	Symptoms	≤2 days/week	>2 days/week or (if ≤11 years) multiple times ≤2 days/week	Throughout the day	
	Nighttime awakenings	≤1x/month (if ≤12 years) ≤2x/month (if >12 years)	≥2x/month (if ≤12 years) 1-3x/week (if >12 years)	≥2x/week (if ≤12 years) ≥4x/week (if >12 years)	
Impairment	Interference with normal activity	None	Some limitation	Extremely limited	
	SABA use for symptoms	≤2 days/week	>2 days/week	Several times per day	
	Lung function *	FEV1>80% FEV1/FVC>80%	FEV1 60-80% FEV1/FVC 75-80%	FEV1<60% FEV1/FVC<75%	
	Exacerbations requiring OCS	0-1x/year	2-3x/year (if 0-4 years) ≥2x/year (if ≥5 years)	≥3x/year (if 0-4 years) ≥2x/year (if ≥5 years)	
Risk	Reduction in lung growth	Requires long-term follow-up			
Treatment related to adverse effects		Medication side effects do not correlate with specific levels of control, but should be considered in overall assessment of risk.			
Recommended action for treatment ****		Consider step down if well controlled for ≥3 months.	Step up 1 step. Re-evaluate in 2-6 weeks.	Consider short course oral corticosteroid. Step up 1-2 steps. Re-evaluate in 2 weeks.	

- " Some individuals with smaller lungs in relation to their height (such as a thin individual with narrow A-F diameter to their chest) may normally have FEV1-80% and/or FEV1/WC-85%. Lung function measures should be correlated with clinical assessment of asthma severity.
- ** For 0.4 years, 54 wheezing episodes per year each lasting 5.1 day and risk factors for persistent asthma meets risk criteria for persistent asthma.
- of moderate or severe persistent asthma that is poorly controlled, consider a short course of OCS.
- **** Recommended guidelines

Appendix C

Chart Review Tools for 4 providers (1 pediatrician and 3 nurse practitioners)

			Number of char	rts
Evaluation	Measures		Pre-	Post-
			implementation	implementation
Patients betwee	en 2-17 years of age with the	following		
ICD codes*				
Providers'	Classifying asthma	The percent	Pre-	Post-
compliance	severity and level of	compliances	implementation	implementation
with the	asthma control**	4 items		
AAP	Providing optimal	(100%)		
Asthma	treatment options	3 items		
Guideline	based on the stepwise	(75%)		
	approach***	2 items		
	Inspecting	(50%)		
	medications, inhaler,	1 item		
	and spacer and	(25%)		
	educating the	0 item		
	administration skills	(0%)		
	and knowledge for	Total		
	self-management****			

	Providing asthma action plan and follow up plan****			
ED/urgent		Transferred	Not transferred	Total
care	Pre-implementation			
transfer	Post-implementation			
	Total			

^{*} ICD codes include below:

1) J45.20 Mild intermittent asthma, uncomplicated. J45.21 Mild intermittent asthma, with (acute) exacerbation. J45.22 Mild intermittent asthma, with status asthmaticus. 2) J45.30 Mild persistent asthma, uncomplicated. J45.31 Mild persistent asthma, with (acute) exacerbation. J45.32 Mild persistent asthma, with status asthmaticus. 3) J45.40 Moderate persistent asthma, uncomplicated. J45.41 Moderate persistent asthma, with (acute) exacerbation. J45.42 Moderate persistent asthma, with status asthmaticus. 4) J45.50 Severe persistent asthma, uncomplicated. J45.51 Severe persistent asthma, with (acute) exacerbation. J45.52 Severe persistent asthma, with status asthmaticus. 5) J45.901 Other and unspecified asthma, with (acute) exacerbation. J45.902 Other and unspecified asthma, with status asthmaticus. J45.909 Other and unspecified asthma, uncomplicated. 6) J45.990 Exercise induced bronchospasm. J45.991 Cough variant asthma. J45.998 Other asthma.

** The 2019 AAP Asthma Guideline assessment includes 1) impairment including symptoms, nighttime awakenings, SABA use for symptoms, limitation of normal activity, and lung function(spirometry), 2) risk for exacerbations requiring oral corticosteroid.

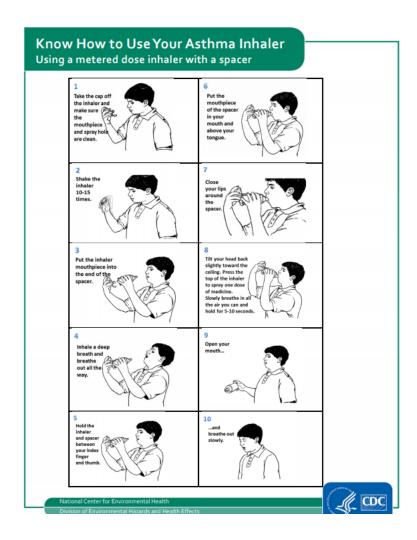
*** Step 1 includes SABA prn. Step 2 includes low dose ICS. Step 3 includes medium dose ICS + subspecialist referral for age 0 to 4 years, low dose ICS + LABA or medium dose ICS for age 5 years and older. Step 4 includes medium dose ICS + LABA or montelukast + subspecialist referral. Step 5 includes high dose ICS + LABA or montelukast + subspecialist referral. Step 6 includes high dose ICS + LABA or montelukast + OCS + subspecialist referral.

****see Appendix E

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Appendix D

Inhaler Spacer Factsheet



Medications, inhaler, and spacer, and administration skills

https://www.cdc.gov/asthma/inhaler_video/default.htm

Appendix E

Asthma Action Plan

$\underline{http://www.pacnj.org/pdfs/atpfillablestudentengsp.pdf}$

Δethr	ma Treatn	nent	Plan – Stude	ent DAG	Sportsoned by	- Car
This asthma ac	tion plan meets NJ Law N.J.S	A. 18A:40-12.8	(Physician's Orders)	Pediatric/Adult Asth	ASSOCIATION.	Health (
Please P	rint)					
lame				Date of Birth	Effective Date	
octor			Parent/Guardian (if app	licable)	Emergency Contact	
hone			Phone		Phone	
EALTHY	(Green Zone)		ike daily control me ore effective with a			Triggers Check all items
	You have all of the	se: MED	ICINE	HOW MUCH to take an	d HOW OFTEN to take it	that trigger patient's asthma:
(201)	Breathing is good		tvair® HFA 🗌 45, 🗌 115, 🗌 23		vice a day	□ Colds/flu
	 No cough or wheeze Sleep through 		rospan™ vesco® 80, 160		2 puffs twice a day 2 puffs twice a day	□ Exercise
	the night	l i i	ilera® 🔲 100, 🔲 200	2 puffs to	vice a day	☐ Allergens
الأجر	Can work, exercise,		ulera® 🗆 100, 🗀 200 ovent® 🗀 44, 🗀 110, 🗀 220	2 puffs to	vice a day	 Dust Mites, dust, stuffed
	and play		rar®		nuffe twice a day	animals, carpet
		L A	Mair Diskus" 🔲 100, 🗀 250, 🗀	J JUUI IIIIIalali	on twice a day	 Pollen - trees, grass, weeds
		□ As	manex® Twisthaler® 🔲 110, 🗍	220	inhalations once or twice a day	o Mold
		님	ovent = DISKUS = [] 50 [] 100 [Ilmicort Flexhaler® [] 90 [] 19] 2501 inhalati 80	on twice a day inhalations once or twice a day	o Pets - animal
		□ Pı	lmicort Respules® (Budesonide) 🔲 🛭	.25, 🗌 0.5, 🗌 1.01 unit nel	bulized 🗌 once or 🗌 twice a day	o Pests - rodent
		☐ Si	ngulair® (Montelukast) 🗌 4, 🔲 5,			cockroaches
dior Dool	k flow above	□ Ot				Odors (Irritants)
u/or Pea/	k flow above			to ringo your mouth o	fter taking inhaled medicine	 Cigarette smoi & second hand
	If exercise trigge	s vour set			fter taking inhaled medicine minutes before exercise	smoke
AUTION	(Yellow Zone)	,	ontinue daily control me	edicine(s) and ADD q	uick-relief medicine(s).	cleaning products, scented products
	You have <u>any</u> of the Cough	ese: MED	ICINE	HOW MUCH to take an	d HOW OFTEN to take it	. Smoke from
(23.7	Mild wheeze	□ Al	buterol MDI (Pro-air® or Prove	ntil [®] or Ventolin [®]) _2 puffs	every 4 hours as needed	burning wood, inside or outsi
7	Tight chest		penex®		every 4 hours as needed	☐ Weather
3	· Coughing at night				nebulized every 4 hours as needed	 Sudden
2	Other:		ionebo		nebulized every 4 hours as needed	temperature change
V 6					nebulized every 4 hours as needed	o Extreme weath
	nedicine does not help wit	nin 🗀 In	ombivent Respimato	1 inhal	ation 4 times a day	- hot and cold
	or has been used more th	an Unin	crease the dose of, or add:			 Ozone alert da □ Foods:
	mptoms persist, call your the emergency room.		quick-relief medici	ne is needed mo	re than 2 times a	O
	flow from to		eek, except before			0
			•			0
MERGE	NCY (Red Zone)				and CALL 911.	Other:
9000	Your asthma is		Asthma can be a life	e-threatening illn	ess. Do not wait!	0
100	getting worse fast • Quick-relief medicin	did N	IEDICINE		ake and HOW OFTEN to take it	0
1	not help within 15-20	minutes] Albuterol MDI (Pro-air® or Pr	oventil® or Ventolin®)	4 puffs every 20 minutes	
600	 Breathing is hard or 	fast	Xopenex®		4 puffs every 20 minutes	This asthma treatmer
HH	 Nose opens wide • R Trouble walking and 	talking] Albuterol [_] 1.25, [_] 2.5 mg .] Duoneb®		1 unit nebulized every 20 minutes 1 unit nebulized every 20 minutes	plan is meant to assist not replace, the clinic
nd/or	Lips blue • Fingerna	ils blue	Xopenex® (Levalbuterol) 0.31	I, □ 0.63, □ 1.25 mg	1 unit nebulized every 20 minutes	decision-making
eak flow	Other:		Combivent Respimat®		1 inhalation 4 times a day	required to meet
elow			Other .			individual patient nee
Company Terrori Ste Design To- tor or les il reach the Design To- el Stee Design et al elitare disco- tive impletate de a reministrativo salar or spermiolitare a servicial della contra contra servicialità del della contra contra della contra di della contra contra discontra di della contra contra di contra di della della contra di della della contra di contra di della della della della della della della della della della della della de	and frames, made continues, comit, a looking of the	This student	Self-administer Medication: is capable and has been instructed method of self-administering of the	PHYSICIAN/APN/PA SIGNAT	Physician's Orders	DATE
	control of passing road seage. A lift Carllo dilaterar control translation fraction from the plants. In presenting the lateral conference from the principles.	non-nebulize	d inhaled medications named above	PARENT/GUARDIAN SIGNAT	URE	_
er Carrie and Premier and Capes or Carrie and Premier and Capes or and Carrie and Capes (Inc.)	and a second of the second and position of the second of t		e with NJ Law.	DUVCICIAN CTAND		
mental Protection Agreement and Administration Agreement and Agreement a	COUNTY To be to be an analysis of the later of the colors	This studen	t is <u>not</u> approved to self-medicate.	PHYSICIAN STAMP		

Plan de Tratamiento del Asma - Para Estudiantes The Pediatric/Adult Asthma Coalition Asthma Coalition of New Jersey **NJ/Healt** AMERICAN LUNG ASSOCIATIO (Este plan de acción para el asma cumple los requisitos de la ley de NJ N.J.S.A. 18A-40-12.8) (Ordenes Medicas) (en letra de imprenta) Nombre Fecha de nacimiento Fecha de vigencia Médico Padre, madre o tutor (si corresponde) Contacto de emergencia Teléfono Teléfono Teléfono Tome este medicamento(s) todos los días. Algunos inhaladores podrían ser Disparadores SALUDABLE (Verde Zona) ||||| mas eficaces si se usan con una cámara inhaladota - úsela como fue indicada. Marcar todos los factores que disparan Usted presenta MEDICAMENTO CUÁNTO tomar y CUÁNDO tomarlo el asma del naciente: todo esto: Advair® HFA 45, 115, 230 2 soplos dos veces al día Restriados · La respiración es buena 1 2 soplos dos veces al día 1 2 soplos dos veces al día Aerospan™ la influenza Ausencia de tos Ejercicios o silbido en su pecho Alergias 2 soolos dos veces al día Acaros, polvo, peluches, Duerme toda la noche 2 soplos dos veces al día 1 2 soplos dos veces al día 1 2 soplos dos veces al día · Puede trabajar, hacer alfombras Polen – árboles, ejercicio y jugar 1 inhalación dos veces al día césped, monte 1 2 inhalaciones una vez dos veces al día Moho 1 inhalación dos veces al día Mascotas - caspa Pulmicort Flexhaler® 90, 180 1 2 inhalaciones una vez dos veces al día Pulmicort Respules® _{Bultomolo} 0.25, 0.5, 1.0_1 unidad nebulizada una vez dos veces al día Singulair (Montelukast) 4, 5, 10 mg_1 pastilla diaria Indos veces al día de animales Pestes - ratones cucarachas Y/o fluio máximo mas de Olores (irritantes) Otro Humo de cigarrillo Ninguno y humo de segunda mano Recuerde enjuagarse la boca después de tomar medicamentos inhalados. o Perfumes Si el ejercicio desencadena el asma, tome soplo(s) minutos antes de hacer ejercicio. productos de Continúe tomando su(s) medicamento(s) de control diario y AGREGUE limpieza. ADVERTENCIA (Amarillo Zona) ||||| medicamento(s) de alivio rápida. perfumados Usted tiene alguno de Humo de la MEDICAMENTO CUÁNTO tomar y CUÁNDO tomarlo Madera estos síntomas: Albuterol MDI (Pro-air® or Proventil® or Ventolin®) __2 soplos cada 4 horas según necesite Ji Clima • Tos. • Silhido leve Cambios bruscos Xopenex® 2 soplos cada 4 horas según necesite · Pecho apretado Albuterol 1.25, 2.5 mg 1 unidad nebulizada cada 4 horas según lo necesite de temperatura Tos nocturna 1 unidad nebulizada cada 4 horas según lo necesite Temperaturas Duoneb Xopenex®(Levalbuterol) 0.31, 0.63, 1.25 mg 1 unidad nebulizada cada 4 horas según lo necesite extremas -Otro: caliente o fría Combivent Respimat® 1 inhalación quatro veces al día Y/o flujo máximo Olías de alerta Aumente la dosis o agregue: ozono de_ a Otro Alimentos Llame a su doctor o vava a la sala de emergencia si la medicina Llame a su medico si necesita la medicina de alivio rápido mas de alivio rápido no lo avuda en 15 a 20 minutos, o ha usado la de dos veces por semana, excepto antes de hacer ejercicio. medicina mas de dos veces y los síntomas persisten. EMERGENCIA (Rojo Zona) ||||| Tome estos medicamentos AHORA y LLAME al 911. Su asma empeora rápidamente: • El medicamento de alivio rápid El asma puede ser una enfermedad potencialmente mortal. ¡No espere! no le ayudó en un lapso de 15 a 20 minutos CUÁNTO tomar y CUÁNDO tomarlo MEDICAMENTO Albuterol MDI (Pro-air® or Proventil® or Ventolin®)_4 soplos cada 20 minutos La respiración es difícil o rápida Las alétas de la nariz se abren Xopenex® 4 soplos cada 20 minutos Este plan de tratamiento para el Se le ven las costillas. Albuterol 1.25, 2.5 mg 1 unidad nebulizada cada 4 horas según lo necesite esma ha sido diseñado para lene problemas para caminar Duoneb 1 unidad nebulizada cada 4 horas según lo necesite vudar no a reemplazar la y para hablar • Tiene los labios azules Xopenex®(Levalbuterol) 0.31, 0.63, 1.25 mg 1 unidad nebulizada cada 4 horas según lo necesite equeridas para satisfacer las Combivent Respirat® Tiene las ufias azules 1 inhalación quatro veces al día ecesidades individuales de Otro Otro cada paciente Y/o un flujo máximo por debajo de SOLO PARA MENORES: Este estudiante es capaz y se le ha enseñado MÉDICO/APN/PA FIRMA FECHA el método correcto, para que se administre (Ordenes Medicas) a sí mismo los medicamentos inhalados no nebulizados nombrados arriba, según la PADRE MADRE O TUTOR FIRMA Ley de NJ. Este estudiante no tiene la aprobación para SELLO DEL MÉDICO automedicarse. Print Save **Print Only Selected Medicines**

Appendix F

Presentation for the AAP Asthma Guideline implementation

Presentation for the AAP Asthma Guideline implementation.pptx

Appendix G

Permission Letter for Project

To,

Minjung Hong-DeCapio

Family Nurse Practitioner,

Touro University Nevada, Las Vegas, NV

February 10, 2020

Subject: Permission for conducting project

Dear Ms. Hong-DeCapio,

I am pleased to inform that MD Medical Group permit you in respect of your project request for implementing the AAP asthma guideline in MD Kids Pediatrics. Your initiative is appreciable, and we are ready to support the project at my best.

We wish you all the best in your project.

Yours truly,

Ana Veronica Rodriguez, MD

CMO MD Medical Group