Appendix 1 Data collection elements

The following data will be collected at baseline for all participants:

- 1. Demographic data: date of birth, gender, ethnicity, medical insurance type, and kindred relationship of caregiver; education status of caregiver, family economic status, occupation of caregiver and residency.
- 2. Disease inducement.
- 3. asthma-like symptom: if there any wheezing, shortness of breath, cough, chest tightness/chest pain, nasal obstruction, nose itch, runny nose, dyspnea, attack or aggravation at night and/or early morning within the four weeks before enrollment and their frequency.
- 4. Response to bronchodilators: such as if asthma can be relieved by aerosol inhalation controller or reliever drugs.
- 5. Family history of asthma.
- 6. History of asthma.
- 7. Other history of illness: history of respiratory infection, sinusitis, obstructive sleep apnea, gastroesophageal reflux.
- 8. History of lung surgery.
- 9. History of allergy: eczema, allergic rhinitis, etc.
- 10. Acute attack information: such as frequency of hospitalization or emergency room admission due to acute asthma-like symptom within previous 12 weeks.
- 11. Physical examination and signs: height, weight, pulse rate, respiratory rate, body temperature, cyanosis, wet rales, three concave signs, wheezing sound, anxious/fidgety, shortness of breath.
- 12. Spirometry examination.
- 13. Protocol specified laboratory tests.
- 14. Protocol specified imaging examination.
- 15. Diagnosis.
- 16. Asthma-related drugs in the previous 12 weeks.
- 17. Other concomitant medication.

The following data will be collected for participants diagnosed as stable asthma:

- 1. Asthma control: defined according to Bronchial Asthma in Children Guideline for Its Diagnosis and Treatment (2016).
- 2. Asthma symptom score: children aged 4–11 years old will be assessed by the C-ACT score, while the older children (12–14 years of age) will be assessed by ACT score.

The following data will be collected for 1st follow-up (12±1 weeks):

- 1. Asthma-like symptom: if there any wheezing, shortness of breath, cough, chest tightness/chest pain, nasal obstruction, nose itch, runny nose, dyspnea, attack or aggravation at night and/or early morning within the four weeks before enrollment and their frequency.
- 2. Asthma control: stage of asthma (acute attack stage/chronic duration stage/clinical remission stage), control level classification [defined according to *Bronchial Asthma in Children Guideline for Its Diagnosis and Treatment (2016)*].
- 3. Acute attack information: such as frequency of hospitalization or emergency room admission due to acute asthma-like symptom within previous 12 weeks.
- 4. Asthma-related drugs in the previous 12 weeks.*
- 5. Asthma symptom score: children aged 4–11 years old will be assessed by the C-ACT score, while the older children (12–14 years of age) will be assessed by ACT score.
- 6. Medical cost for the diagnosis and treatment of asthma.
- 7. Study discontinuation status.

The following data will be collected for 2nd follow-up (24±1 weeks):

- 1. Asthma-like symptom: if there any wheezing, shortness of breath, cough, chest tightness/chest pain, nasal obstruction, nose itch, runny nose, dyspnea, attack or aggravation at night and/or early morning within the four weeks before enrollment and their frequency.
- 2. Asthma control: stage of asthma (acute attack stage/chronic duration stage/clinical remission stage), control level classification [defined according to *Bronchial Asthma in Children Guideline for Its Diagnosis and Treatment (2016)*].
- 3. Acute attack information: such as frequency of hospitalization or emergency room admission due to acute asthma-like symptom within previous 12 weeks.
- 4. Physical examination and signs: height, weight, pulse rate, respiratory rate, body temperature, cyanosis, wet rales, three concave sign, wheezing sound, anxious/fidgety, shortness of breath.
- 5. Spirometry examination.
- 6. Protocol specified laboratory tests.
- 7. Protocol specified imaging examination.
- 8. asthma-related drugs in the previous 12 weeks.*
- 9. Asthma symptom score: children aged 4–11 years old will be assessed by the C-ACT score, while the older children (12–14 years of age) will be assessed by ACT score.
- 10. Medical cost for the diagnosis and treatment of asthma.
- 11. Study discontinuation status.

*, treatment change include but not limit within the following situations: drug dosing change, drug administration schedule change, change drug (drug brand change is not included), add new treatment.

In addition to the data elements mentioned above, any AE/SAE reported at any time will be recorded.

Table S1 Data collection plan

Data collection	Baseline ^ª	1 st Visit	2 nd Visit
Window of visits (week ± week)	0±1	12±1	24±1
Informed consent	Х		
Demographics	Х		
Disease inducement	Х		
Asthma-like symptom	Х	Х	х
Acute attack information	Х	Х	х
Physical examination and signs	Х		х
Asthma-related drugs in the previous 12 weeks	х	Х	Х
Response to bronchodilators	Х		
Family history of asthma	Х		
listory of asthma	Х		
listory of lung surgery	Х		
listory of allergy	Х		
Other past history of illness	Х		
Spirometry examination	Х		Х
Protocol specified laboratory tests	Х		Х
Protocol specified imaging examination	Х		Х
Diagnosis	Х		
Concomitant medication	Х	Х	х
Asthma control ^b	Х	Х	Х
Asthma symptom score ^b	Х	Х	Х
AEs and SAEs		Х	х
Cost of diagnosis and treatment of asthma (if applicable)		х	Х
Study discontinuation status		Х	х

^a, to help assess the rates of participation, sites should maintain a log of the number of eligible participants who decline to participate in the study. ^b, the information of participants with diagnosed stable asthma in Phase I need to be collected. AE, adverse event; SAE, severe adverse events.