

# **Lungs 4 Life (Preventing non-cystic fibrosis (CF) Bronchiectasis in children)**

**By Adele Cluett, Alison Howitt, Annie Nicholas  
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# The Lungs 4 Life Project Team

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# Background

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- Previous studies have documented that the rate of non-cystic fibrosis (CF) bronchiectasis is higher amongst Maaori and Pacific Island children than other ethnic groups.
- Key Risk Factors identified were:
  - Hospitalisation with pneumonia
  - Recurrent hospitalisation (> x3 admissions) with lower respiratory infection under the age of 2 years old.
  - Radiologic abnormalities
  - Chronic wet cough or wheeze
  - Pacific or Maaori ethnicity
  - Low socioeconomic areas

# What we found

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- There is no best practice approach to identifying, treating and managing these high risk children.
- There exists a health equity gap for Maaori & Pacific Island children
- As Nurses working at Kidz First we see a number of recurrent patients with multiple significant LRTI admissions
- No clear integration between primary and secondary care services

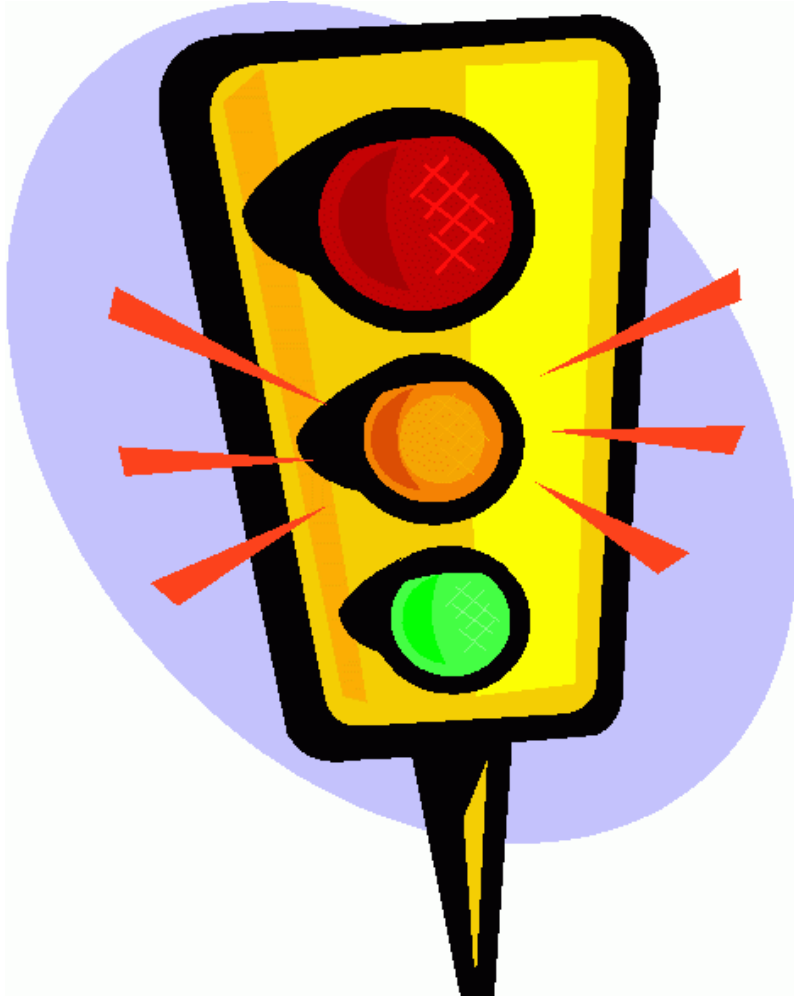
# Case review

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- 10 month old Baby A.
  - >10 Presentations to hospital
  - 2 x ICU admissions requiring CPAP
  - Multiple diagnoses via CXR of pneumonia and Bronchiolitis
  - > 3 days of admission to hospital.
- There was no follow up plan for Baby A.
- Baby A is one of the many children that present to Kidz First with a significant history, and at “high-risk” for developing Bronchiectasis.
- This demonstrated the need for preventative pathways.

# Present emerging ideas

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We use a Traffic light system to help identify the “at-risk” & “high risk” children in the hospital.

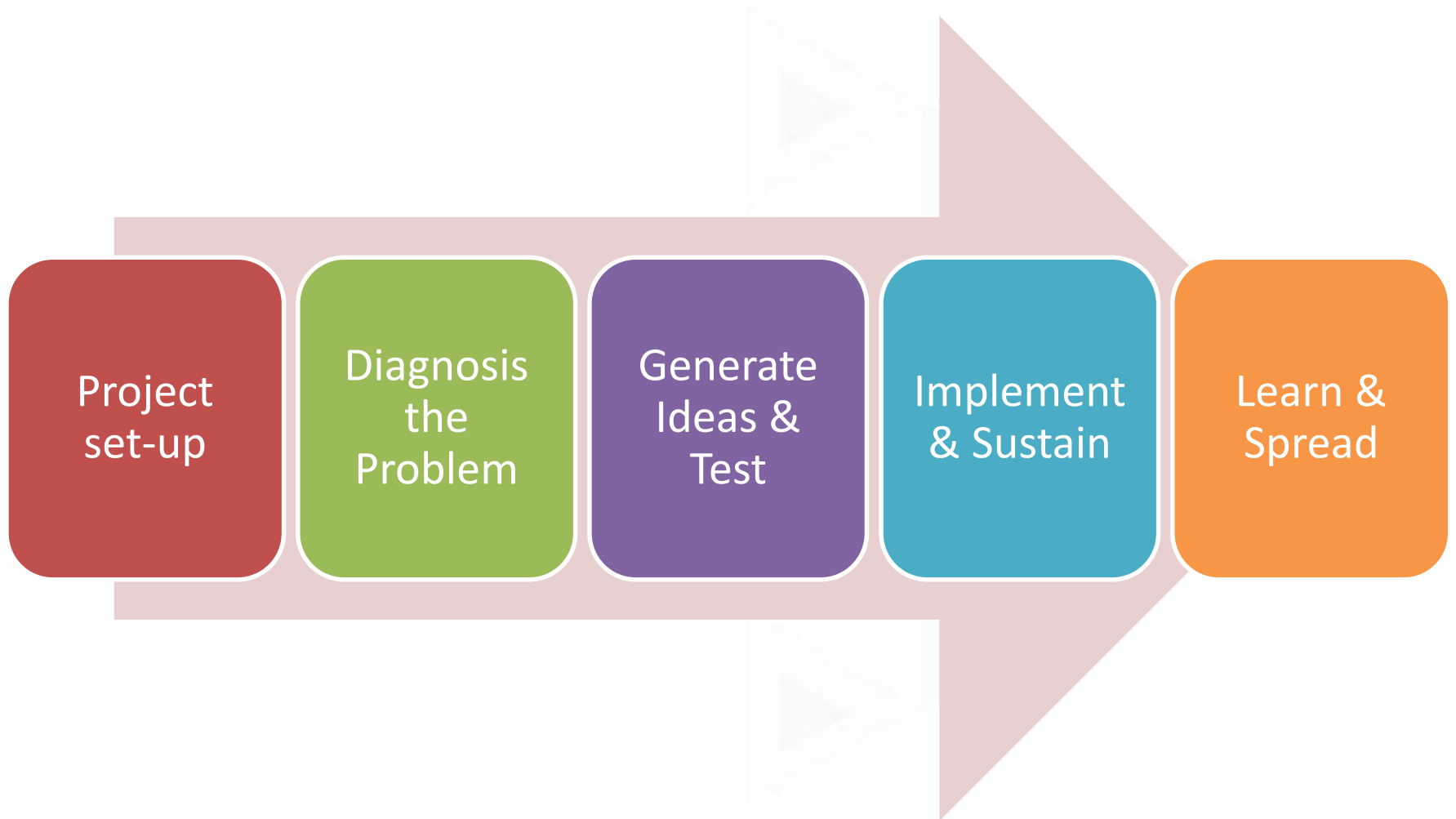
**Yellow** = First Admission with significant Lower respiratory tract infection (LRTI)

**Orange** = Second Admission with significant LRTI. And/or frequent presentations without significant LRTI episode.

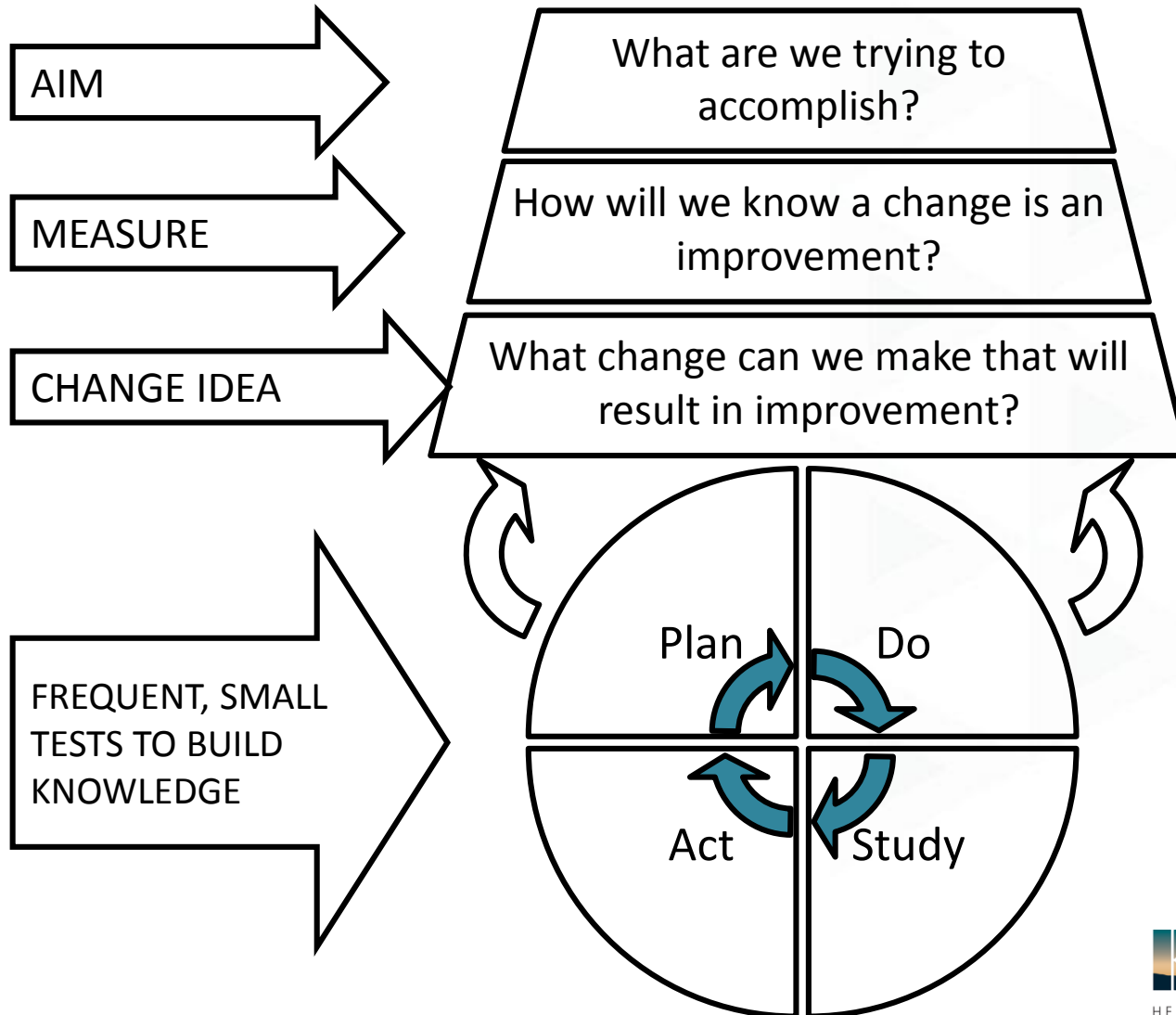
**Red** = Three or more admissions with significant LRTI. Any child who presents with a history of chronic wet cough or wheeze.

# Ko Awatea Improvement Methodology

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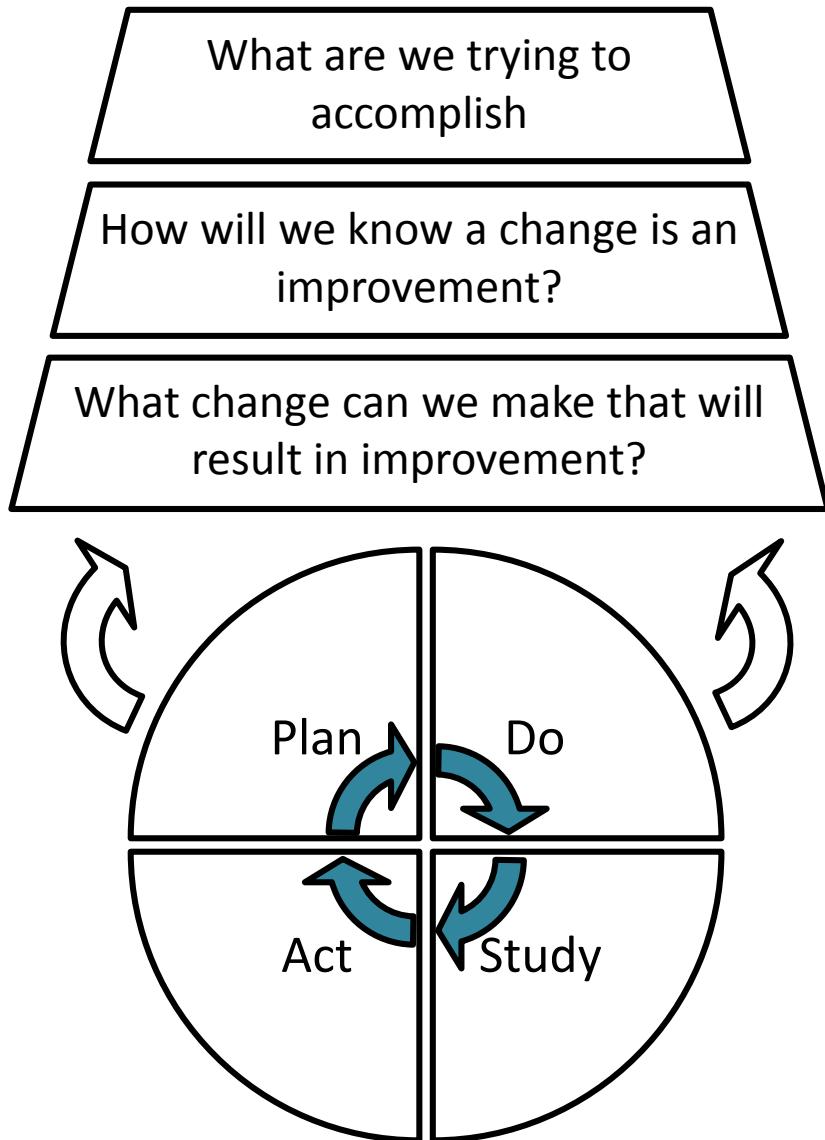


# IHI's Model for Improvement





# Model for Improvement



- Identification of “High-Risk” children and test change ideas to prevent the development of bronchiectasis.
- We developed measures for the project and each change idea we trialled e.g. Screening tool
- We reviewed previous studies and current practice to develop a set of change ideas to be tested in primary and secondary care.
- We’re currently testing these change ideas with the P-D-S-A Cycle to assess whether we Adopt, Adapt or Abandon.

# Success & Challenges

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- Having the screening tool helps to group these high risk children so we can implement preventative measures in their care.
- Developing consistent discharge planning for “high-risk” & “at-risk” children.
- Co-ordinating care between primary and secondary services. To enable better treatment and management of high risk children.
- Positive feedback from families about education and discharge planning.
- As clinicians, testing ideas at times was a challenge due to working schedules.

# Strategies & Reflections

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- Identifying these high risk children has highlighted the need to provide a service that will establish links between primary and secondary care.
- Utilising the tools being developed in other aspects of our nursing to support our families.
- Time management has been a key aspect in this campaign, working with different areas of the health care team.
- Creating Networks within various health care teams.

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# Any Questions?