

Appointment Lag Time and Tracking Transition in Young Adults

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Acknowledgements: T1D Exchange



Background

- Transfer from pediatric to adult providers is a sensitive time for Type 1 diabetes care
- Transfer is part of a larger transition process that should start early in adolescence
- If transition is not appropriately discussed, transfer of care is often frustrating for both patients and providers. Delay in transfer can lead to gaps in care and sub-optimal outcomes

Aim Statement

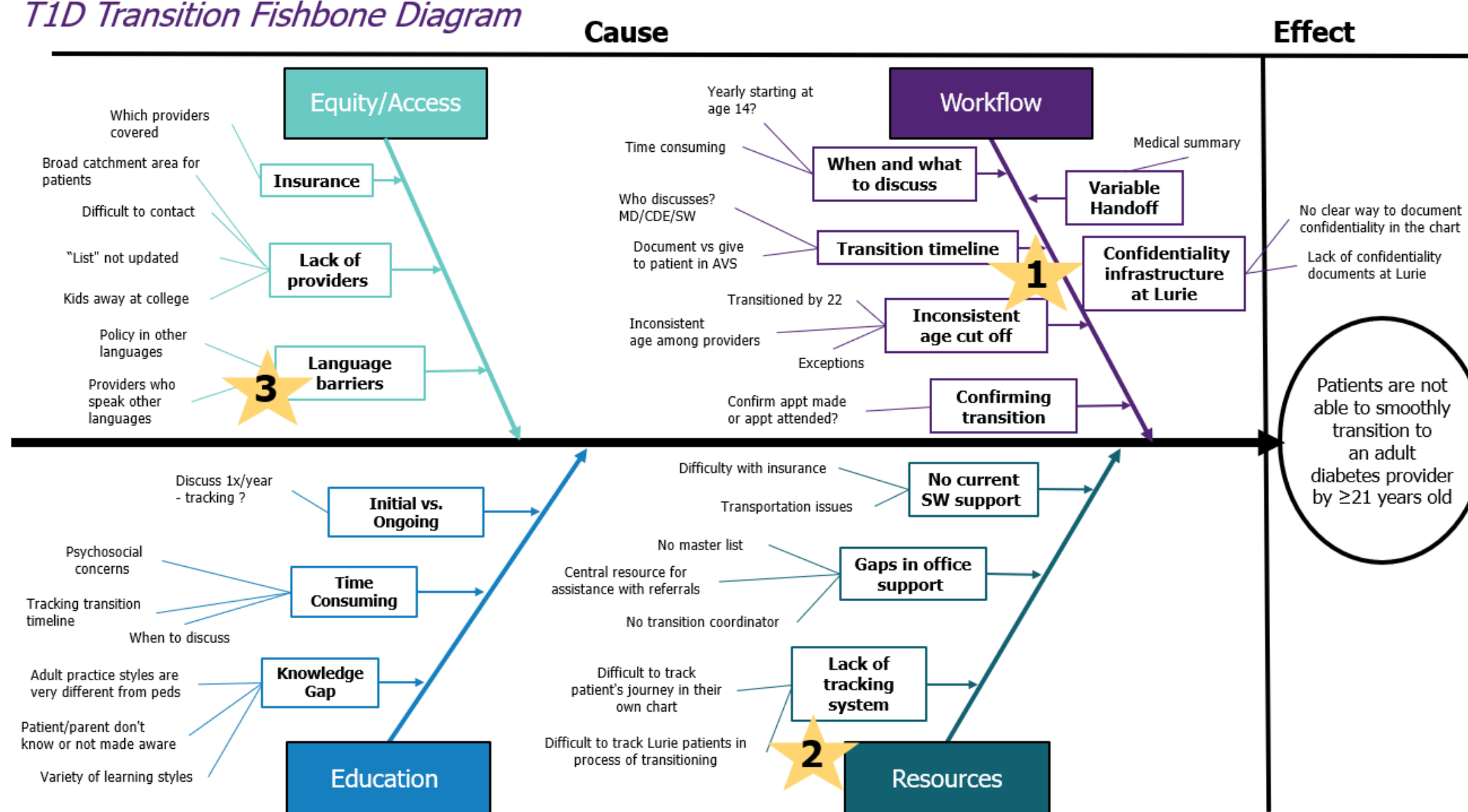
- Decrease the number of Type 1 diabetes patients ≥ 21 years old who are seen each month in pediatric endocrinology by 10% by 12/31/2023
- Decrease the time between the last pediatric visit and the first adult visit (new intervention; no baseline data) and increase the number of patients with a completed adult appointment each month by 10% by 12/31/2023
- Increase the discussion of transition during clinic appointment for patients ≥ 14 years old (new intervention; no baseline data) by 12/31/2023

Current State Analysis

- The current state analysis information was gathered in a variety of ways including rounding with front line staff
- Additionally, providers spent time speaking with patients about their barriers to transitioning to adult care
- The core transition team analyzed the barriers and prioritized three areas to begin improvement work

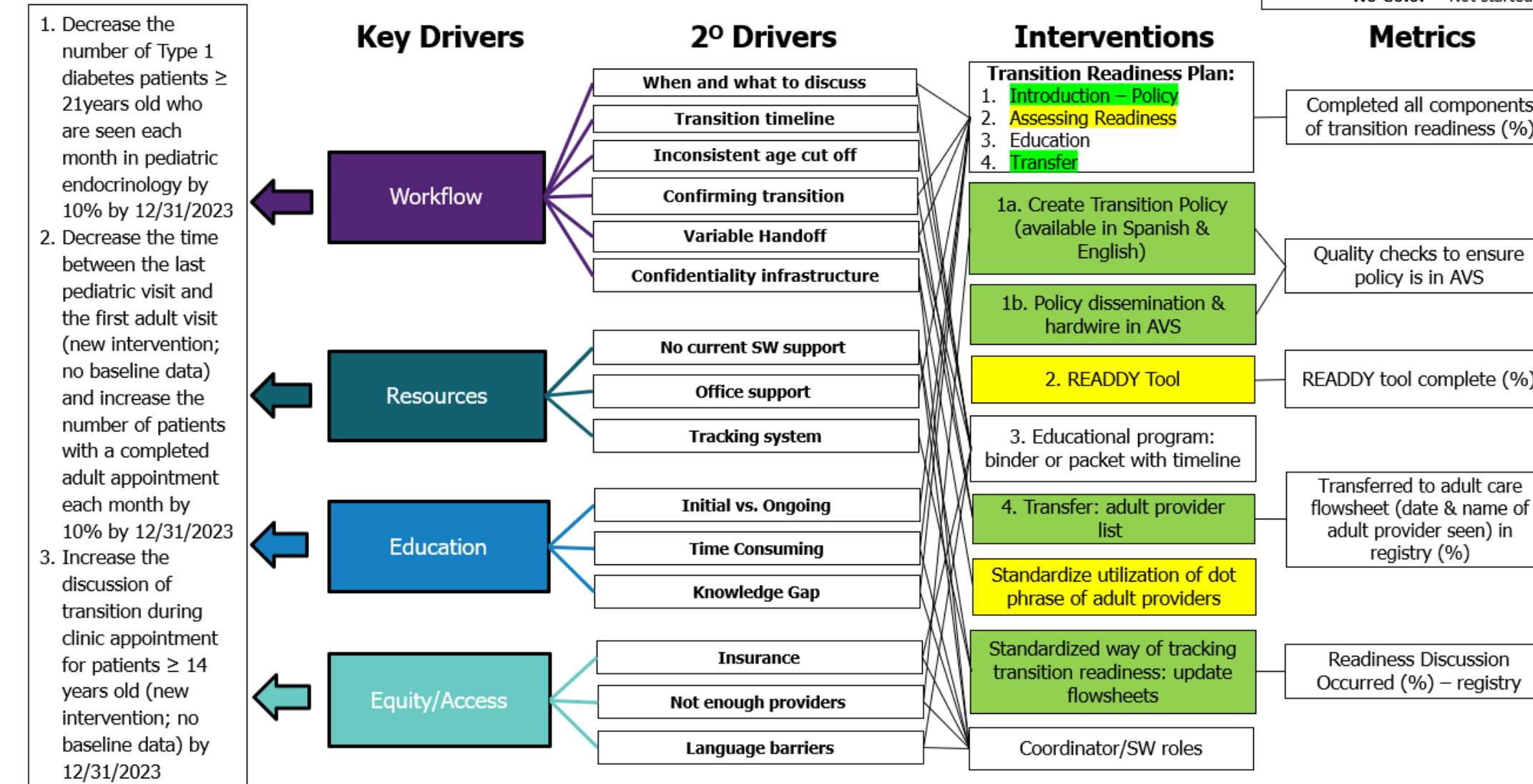
Fishbone Diagram

T1D Transition Fishbone Diagram



Key Driver Diagram

T1D Exchange: Transition Readiness Key Driver Diagram

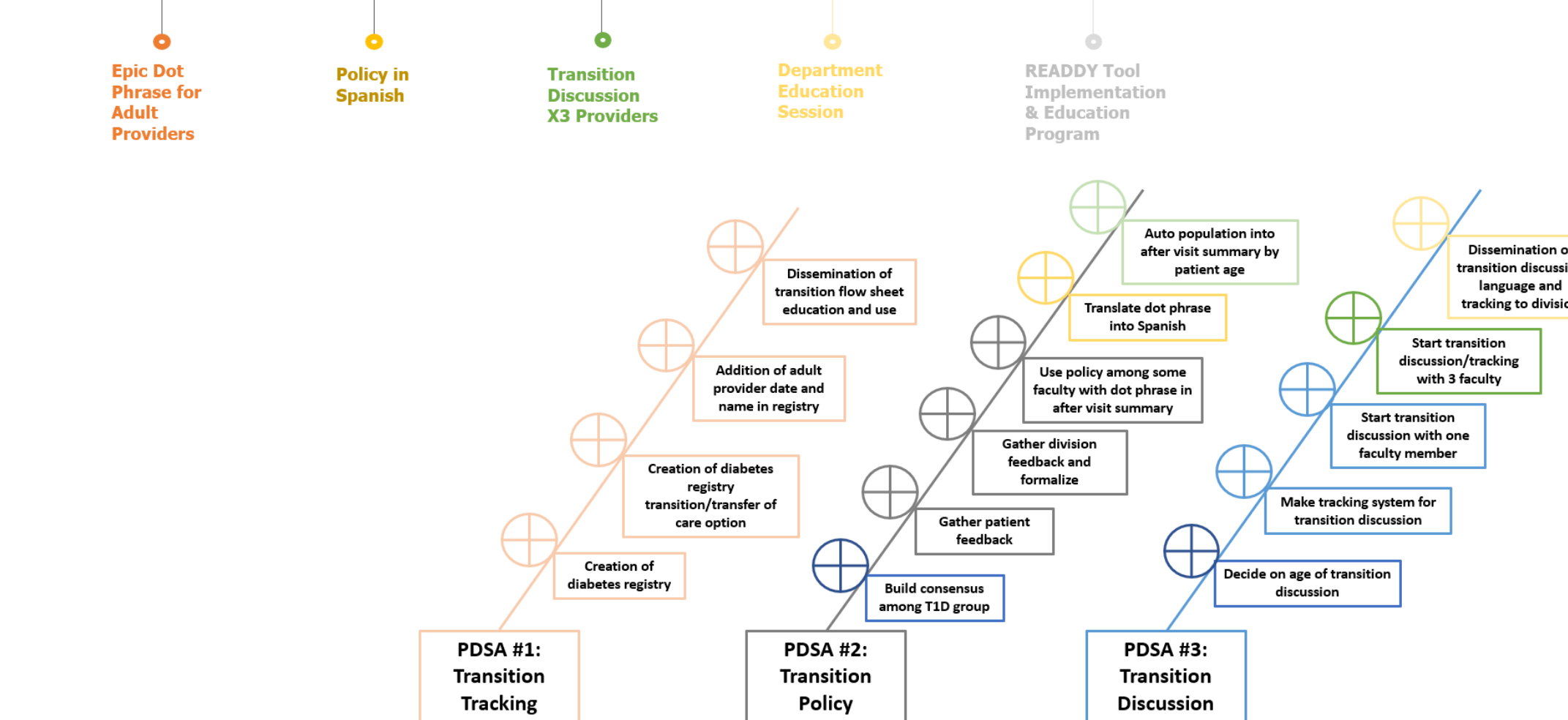
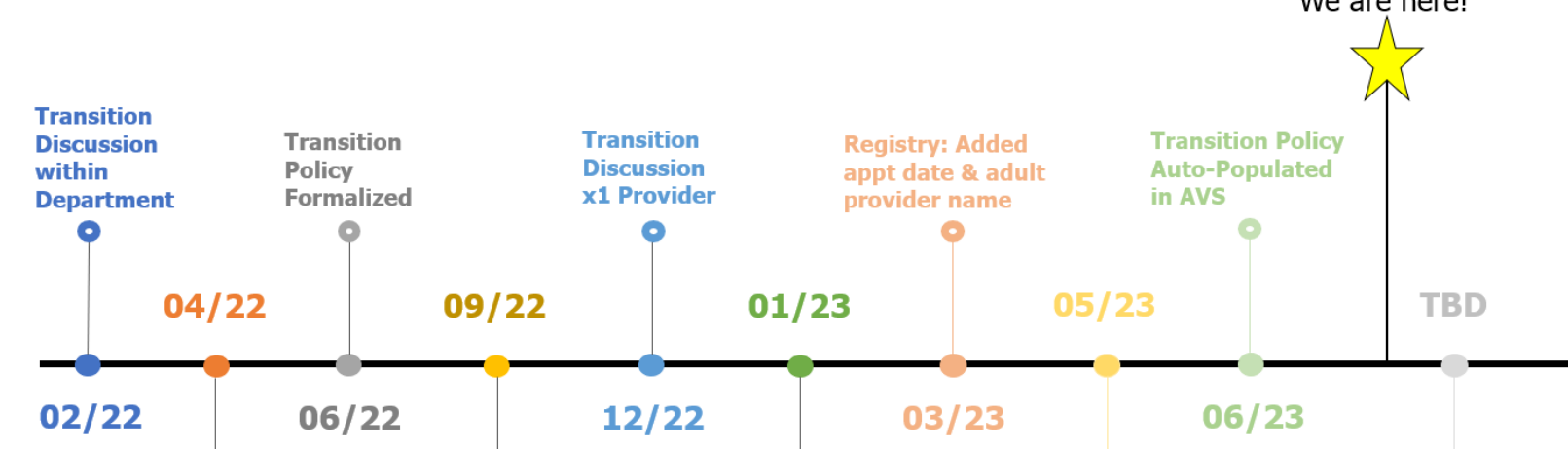


Hypothesis/Future State Design

- Building in structure to talking about transition and transfer during visits will increase transfer to adult provider
- Creating a policy for the division that standardizes timing of transition discussions and transfer will increase timely and successful transfer
- Tracking transfer systematically will allow for improved follow up and completion of transfer to adult provider

Implementation

Intervention Timeline

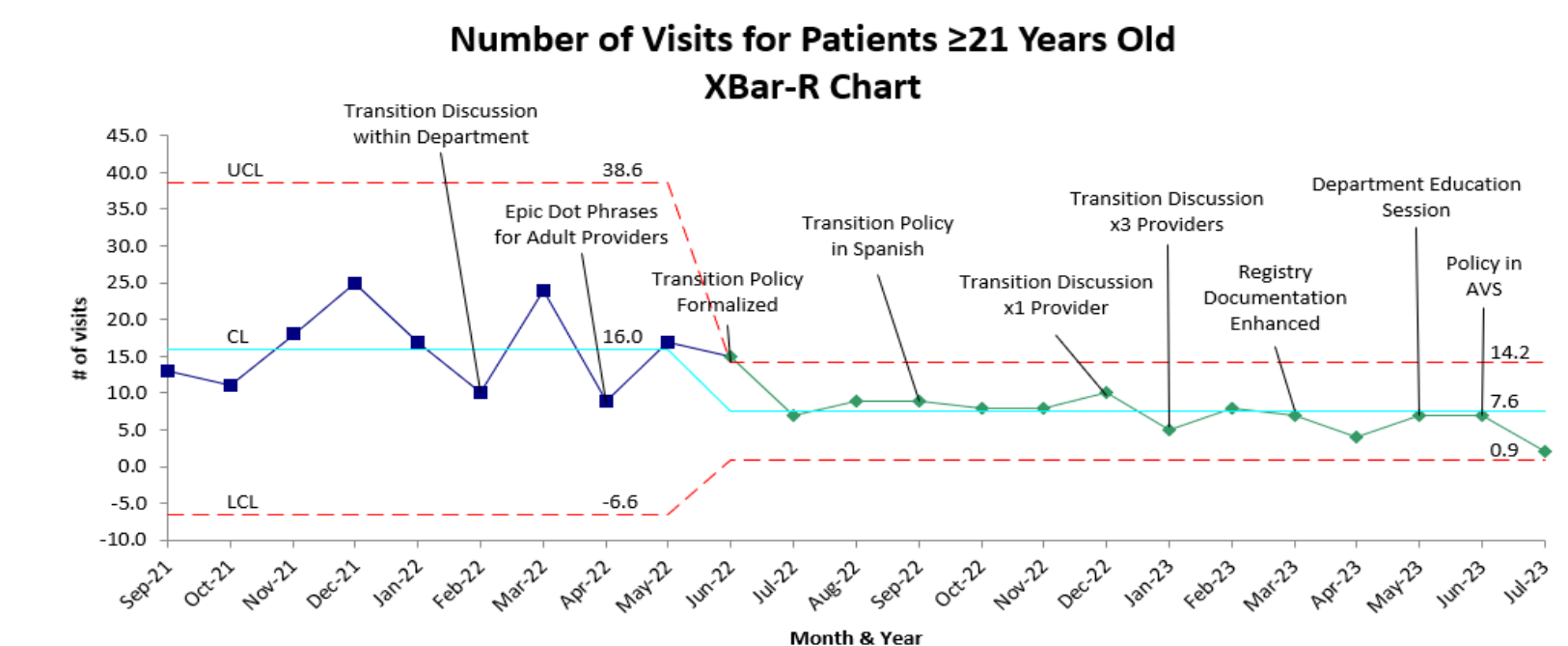


Patient Characteristics

Summary of patient characteristics by transition status, population includes all patients ≥ 20 years old in the diabetes registry as of 7/31/23

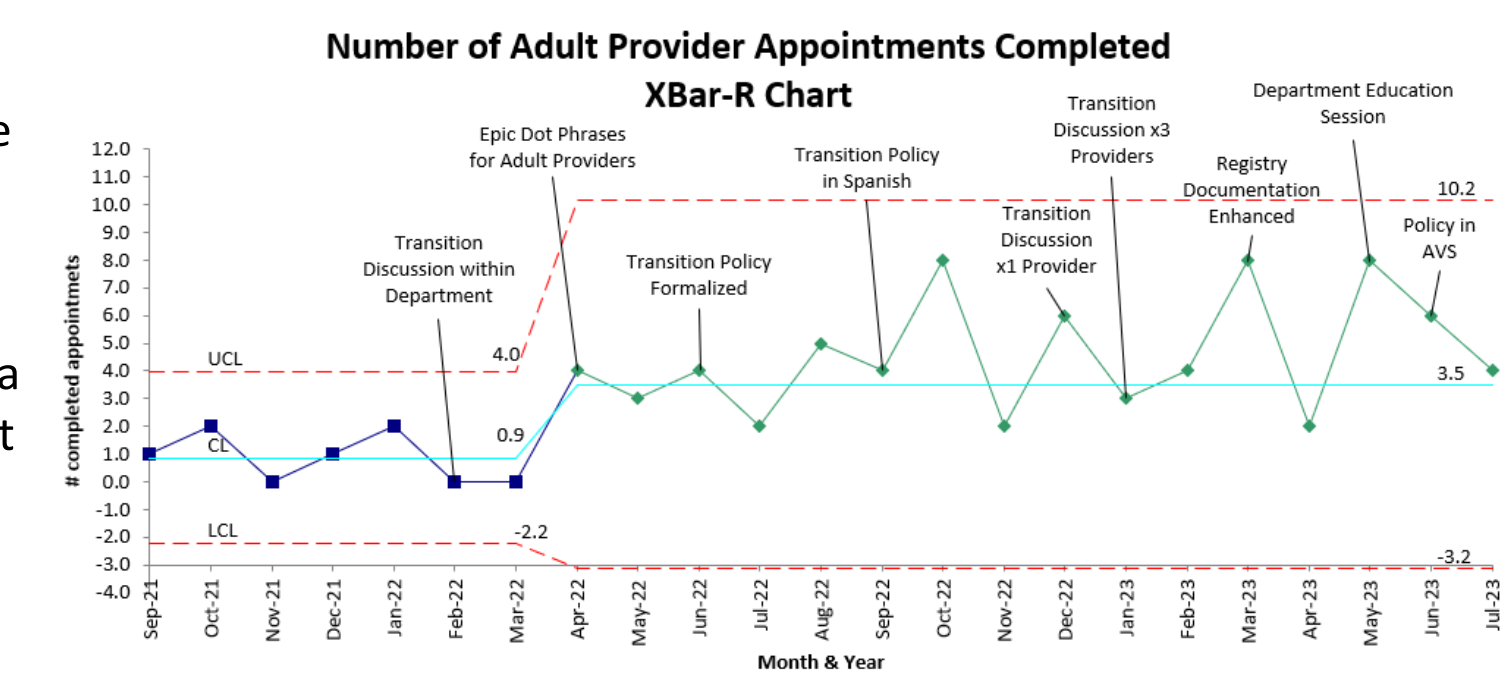
	Transitioned (n=83)	Not Transitioned (n=109)
Age (median, IQR)	21 years (20-21)	20 years (20-21)
Females (n, %)	41 (49.3%)	54 (49.5%)
Last HgA1C (median, IQR)	7.7% (6.9-9.7)	7.8% (7.0-9.1)
Lag Time to Adult Appt in months (mean and standard deviation)	4 (2-8)	NA

Results

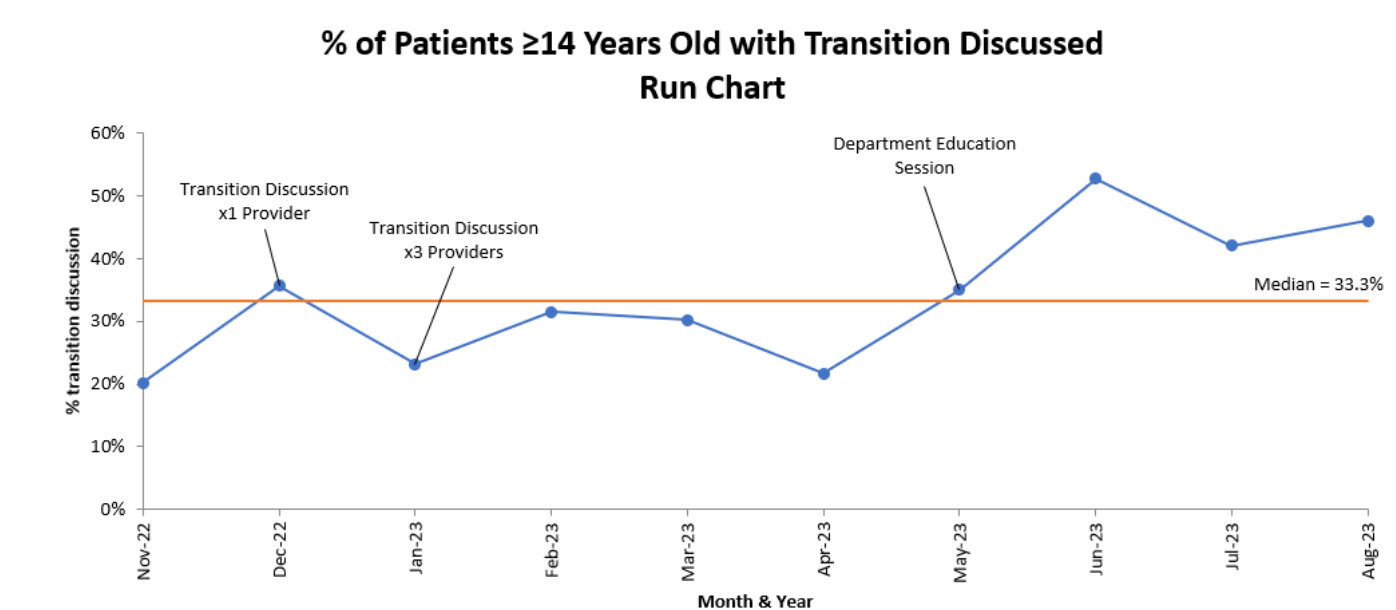


Aim #1: The number of patients ≥21 years old seen each month decreased from 16 to 7.6 visits per month (47.5% decrease)

Aim #2: Those who transitioned to adult providers did so at an average of 4 months after their last pediatric visit



The number of patients with a completed adult appointment increased from 0.9 to 3.5 completed appointments per month (25.7% increase)



Aim #3: The number of times that transition discussion occurred in clinic increased over time, though there is no baseline data available

Lessons Learned/Next Steps

- Relatively simple interventions of consensus building, creation of formalized policy and tracking/documentation led to substantive change in rates of transfer of care.
- Next steps will include integration of transition readiness assessment and curriculum building around transition.
- No data on race/ethnic group and equity, attempting add into registry so able to better track.



Contact Us!
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