Race, Ethnicity, and Language Data Collection via Providers



Project Goals



Goal 1: Determine the feasibility and practicality of collecting patient REL data from providers



Goal 3: Gather qualitative provider feedback regarding their potential role in providing patient REL data to health carriers



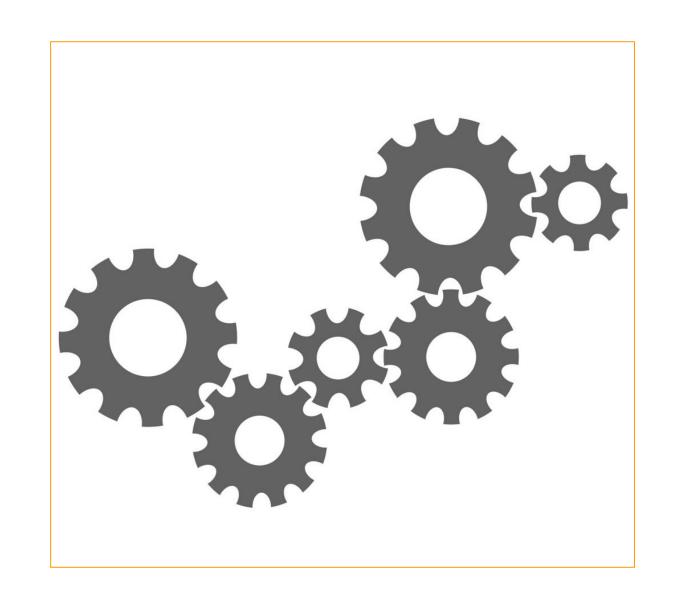
Goal 2: Determine what methods are the most efficient in collecting patient REL data from providers



Goal 4: Evaluate what incentives exist for providers to submit patient REL data to health carriers

Process Undertaken During DLP

- 1. Conducted provider focus groups with providers of varying size practices
- 2. Explored opportunities to conduct pilot test with identified UHC contracted providers
- 3. Explored opportunity to conduct pilot test with academic institution
- 4. Exploring opportunity to conduct pilot test with American Medical Association and other stakeholders



Challenges/Barriers

- Competing Internal business priorities
- Partnership approval process The process of approval to partner with the AMA took longer than expected, and identifying providers for the pilot continues to be an ongoing process.
- Stakeholder alignment In identifying additional stakeholders to partner with on this pilot, there has been the challenge of confirming mutually agreeable roles and project priorities.
- Provider participation Providers have a natural skepticism of projects done with insurance carriers because of potential future requirements that may be placed on providers.

Lessons Learned & Next Steps

- 1. Partnerships are key to implementation. A health insurance company alone may not be the best entity to conduct the pilot because of provider skepticism.
- 2. Providers need to be educated on the "WIIFM" to providing patient REL data to health insurance carriers.
- 3. Continue exploring opportunity with AMA and other stakeholders.

