

Bundled Payments for Care Improvement (BPCI) Advanced Annual Check-in Questionnaire

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Purpose:

Participants' activities and barriers related to designing and implementing the BPCI Advanced Model

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Instructions:

The first part of the study was a pilot study. The purpose of the pilot study was to determine the feasibility of the study and to estimate the sample size required for the main study. The pilot study was conducted with 10 participants. The results of the pilot study showed that the study was feasible and that the sample size required for the main study was 100 participants.

The main study was conducted with 100 participants. The participants were recruited from a variety of sources, including social media, community organizations, and university databases. The participants were randomly assigned to two groups: the experimental group and the control group. The experimental group received the intervention, and the control group did not receive the intervention.

The intervention was a 12-week program that focused on improving the participants' understanding of the risks and benefits of the intervention. The intervention was delivered through a series of workshops and group discussions. The control group received no intervention.

The primary outcome of the study was the participants' understanding of the risks and benefits of the intervention. This was measured using a series of questions that were administered at baseline and at the end of the 12-week program. The secondary outcome was the participants' willingness to participate in the intervention. This was measured using a series of questions that were administered at baseline and at the end of the 12-week program.

The results of the study showed that the intervention significantly improved the participants' understanding of the risks and benefits of the intervention. The participants in the experimental group were more likely to understand the risks and benefits of the intervention than the participants in the control group.

The results also showed that the intervention significantly increased the participants' willingness to participate in the intervention. The participants in the experimental group were more likely to participate in the intervention than the participants in the control group.

The study has several limitations. First, the study was a pilot study, and the results may not be generalizable to a larger population. Second, the study was conducted with a convenience sample, and the results may be biased. Third, the study did not include a blinding procedure, and the results may be biased.

Despite these limitations, the study provides valuable information about the feasibility of the study and the effectiveness of the intervention. The results suggest that the intervention is a promising approach for improving the participants' understanding of the risks and benefits of the intervention and increasing their willingness to participate in the intervention.

Diagram illustrating the location of the *rDNA* and *rD* genes on the *E. coli* chromosome. The *rDNA* gene is located at approximately 100 minutes, and the *rD* gene is located at approximately 110 minutes. The *rDNA* gene is transcribed into rRNA, and the *rD* gene is transcribed into 16S and 23S rRNA.

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Participant Profile

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Care Redesign Plan

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CEHRT Use

1. *To meet this criterion, at least 75 percent of eligible clinicians in each participating APM Entity group, or each hospital if hospitals are the APM Entities, to use CEHRT to document and communicate clinical care.*



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Partner Agreements

1. **Definition:** A **Discrete-Time Signal** is a signal that is defined only at discrete intervals of time. It is represented by a sequence of values, often denoted as $x[n]$, where n is the discrete time index.

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4.

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Diagram illustrating the structure of a DNA sequence. The sequence is represented by a horizontal bar divided into segments. The segments are labeled with letters: D, r, d, r. The segments are color-coded: D is blue, r is green, d is red, and r is green. The segments are arranged in a sequence: D, r, d, r. The segments are connected by lines, indicating a continuous sequence.

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Compliance Plan

1. ☐ If your response is ‘yes’, please upload this document to the amendment section of the [Participant Portal](#). If your Compliance Plan includes multiple appendices, you can upload the main document only.]
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Waivers (Fraud & Abuse Waivers and Medicare Payment Policy Waivers)

Participant Profile Selection Does Not Limit Use of Waivers”

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If selected ‘Yes’ to question “Do you have any other information about the person?”

Respondent’s name

Relationship to respondent

Address

Phone number

E-mail address

Other information

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BPCI Advanced Impact



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Document Submission



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Self-disclosed Investigations or Sanctions

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2. I selected 'Yes' to the question "Do you agree to participate in the study?"
- I understand that my participation in the study is voluntary and that I may withdraw from the study at any time without penalty or loss of benefits to which I am entitled.
- Additional information is located under Article 13 of the [Participation Agreement](#).

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Other Issues Not Previously Listed

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