

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, while 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 of smoking and the benefits of quitting. The intervention was delivered through a combination of group sessions and individual counseling. The group sessions were led by a trained facilitator, and the individual counseling was provided by a research assistant.

The control group received no intervention. The participants in the control group were asked to continue with their current smoking habits. The participants in the experimental group were asked to quit smoking during the 12-week intervention period.

The primary outcome of the study was the number of participants who quit smoking during the 12-week intervention period. The secondary outcomes were the number of participants who reduced their smoking and the number of participants who were motivated to quit smoking.

The results of the study showed that the intervention was effective in helping participants quit smoking. The number of participants who quit smoking during the 12-week intervention period was significantly higher than the number of participants who quit smoking in the control group.

The study has several limitations. First, the study was a pilot study, and the sample size was small. Second, the study was conducted in a community setting, and the results may not be generalizable to other populations. Third, the study did not measure the long-term outcomes of the intervention.

Despite these limitations, the study provides promising results for the use of a community-based intervention to help people quit smoking. Further research is needed to evaluate the effectiveness of the intervention in a larger, more diverse population.

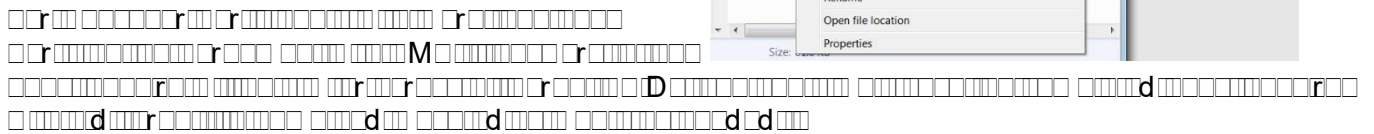
Diagram illustrating the location of the *rDNA* and *rD* genes on the *E. coli* chromosome. The *rDNA* gene is located near the origin of replication (*oriC*), and the *rD* gene is located near the *rDNA* gene. The *rDNA* gene is transcribed into rRNA, and the *rD* gene is transcribed into 16S and 23S 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 **Decision Tree** is a model for classification and regression tasks. It consists of a root node, internal nodes, and leaf nodes. The root node splits the data into two or more subsets based on a feature and a threshold. This process is repeated recursively for each subset, creating a tree structure. The leaf nodes represent the final classification or regression result.

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The diagram consists of two horizontal rows of square boxes. The top row contains 10 boxes; the 7th box from the left contains the letter 'r', and the 8th box contains the letter 'd'. The bottom row contains 16 boxes; the 9th box from the left contains the letter 'r', and the 14th box contains the letter 'r'. All other boxes are empty.

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Compliance Plan

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Participant Profile Selection Does Not Limit Use of Waivers”

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If selected ‘Yes’ to question “Do you have any children?”

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Enter respondent’s name.

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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?' and I understand that my participation is voluntary. I understand that I may withdraw from the study at any time without penalty or loss of benefits to which I am entitled. I understand that my participation in the study is confidential. I understand that my participation in the study is for my own benefit and the benefit of the community. I understand that my participation in the study is for the purpose of the study. I understand that my participation in the study is for the purpose of the study. Additional information is located under Article 13 of the [Participation Agreement](#).

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Other Issues Not Previously Listed

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