

## 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 control group received no intervention.

The primary outcome of the study was the number of participants who quit smoking by the end of the 12-week 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 increasing the number of participants who quit smoking. The number of participants who quit smoking in the experimental group was significantly higher than the number of participants who quit smoking in the control group.

The results also showed that the intervention was effective in increasing the number of participants who reduced their smoking and the number of participants who were motivated to quit smoking.

The study has several limitations. First, the study was a pilot study and therefore the results may not be generalizable to a larger population. Second, the study was conducted in a community setting and therefore the results may not be generalizable to a clinical setting.

Despite these limitations, the study provides valuable information about the feasibility of the intervention and the potential for increasing the number of participants who quit smoking.

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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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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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### Internal Cost Savings (ICS)

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☐ If selected ‘Yes’ to question 1, has the Participant ☐ died ☐ or ☐  
☐ returned? ☐ Yes ☐ No ☐ Did not respond

### Net Payment Reconciliation Amount (NPRA)

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**r** **r** **d** **d** [Performance Period 1: October 1, 2018 through June 30, 2019]

☐ If you are submitting a document to the amendment section of the [Participant Portal](#), please upload the document to the amendment section of the [Participant Portal](#).

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☐ If selected 'Yes' to question [Participant Profile Selection](#), please upload the document to the amendment section of the [Participant Portal](#).

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## Compliance Plan

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1. ☐ If you are submitting a document to the amendment section of the [Participant Portal](#), please upload the document to the amendment section of the [Participant Portal](#). *[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)

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☐ If you are submitting a document to the amendment section of the [Participant Portal](#), please upload the document to the amendment section of the [Participant Portal](#).

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☐ If selected 'Yes' to question ☐ **Do you have any other information that you would like to provide?**  
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## BPCI Advanced Impact



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☐ If selected 'Yes' to either question 1 or 2, ☐ **d** ☐ **r** ☐



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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 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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