Audit Review Period:		
Issue of non-compliance:	Medication errors	
Scope:	• The scope of this Impact Analysis is limited to 50% of the participants enrolled during the audit review period who were not included in the provision of	
	services sample selection.	
	The auditor will select the participants to be reviewed and enter their identifying information on the Participant Impact tab.	
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Instructions:	• Review only the participant medical records selected by the auditor. The selected participants are identified in the Participant Impact tab.	
	Review the selected medical records to determine if any medication errors occurred.	
	a Respond to the questions in the participant impact tob	
	 Respond to the questions in the participant impact tab. 	
• The review timeframe is the audit review period. Errors noted before or after the audit review period should not be included.		
	 The review timetrame is the audit review period. Errors noted before or after the audit review period should not be included. 	
	After completing the Impact Analysis, if any changes need to be made to the Root Cause Analysis, please update the changes in the RCA tab.	
Impact Analysis Due Date:		

			Detailed Description of the Issue
(MM/DD/YY)	(Completed By The CMS Audit Lead)	(Completed By The CMS Audit Lead)	
(Completed By The CMS			(Explain what happened)
Audit Lead)			(Remaining fields to be Completed by PACE Organization)

Root Cause Analysis for the Issue	Methodology - Describe the process that	# of Individuals	Action Taken to Resolve System/	Date System/ Operational Remediation	Date System/ Operational Remediation
(Explain why it happened)	was undertaken to determine the # of	Impacted	Operational Issues	Initiated	Completed (MM/DD/YY)
	individuals (e.g. participants) impacted			(MM/DD/YY)	

Actions Taken to Resolve Negatively Impacted Individuals Including Outreach Description and Status	Initiated	Date Individual Outreach and Remediation Completed (MM/DD/YY)

For the purpose of this Impact Analysis, a medication error is defined as: any preventable event that may cause or lead to inappropriate medication use or participant harm while the medication is in the control of the PACE Organization or one of it's contracted providers. Such events may be related to professional practice, health care products, procedures, and systems, including prescribing, order communication, product labeling, packaging, and nomenclature, compounding, dispensing, distribution, administration, education, monitoring, and use.

Participant First Name	Participant Last Name	Participant ID	Date of Enrollment	Date of Disenrollment	Did the participant experience a medication error during the	Medication Name
					audit review period?	
			MM/DD/YYYY	MM/DD/YYYY		List each medication that was associated with a medication error
					(Yes/No)	in a new row.
					If NO, the PO may enter NA in all remaining fields.	

Medication Dosage	Medication Route	Medication Frequency	Medication Start Date	Medication Discontinue Date
				Enter NA if the medication has not been discontinued.

Describe the type of Medication Error	In what setting was or should the medication have been administered? (PACE Center, SNF, ALF, Home)
(Examples: Did not give medication, Gave wrong medication, etc.)	

Date the Medication Error Began (First occurrence of the Date the Medication Error Ended (Last Occurrence of medication error) MM/DD/YYYY MM/DD/YYYY	omitted in error between the first and last date?	If the participant experienced negative outcomes, did they occur, in some part, as a result of the failure to provide the item or service? (Yes/No)
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If yes, describe the negative outcomes.	Optional: Please note, you do not have to complete this column.
	If there are any mitigating factors that you would like CMS to consider related to a specific participant, please enter the information in this column.