FDA DOCUMENTATION FOR THE GENERIC CLEARANCE OF COLLECTION OF QUALITATIVE DATA ON TOBACCO PRODUCTS AND COMMUNICATIONS (0910-NEW)

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TITLE OF INFORMATION COLLECTION: [insert]

DESCRIPTION OF THIS SPECIFIC COLLECTION

1. **Statement of need:**
   [insert]

2. **Intended use of information:**
   [insert]

3. **Description of respondents:**
   [insert]

4. **Date(s) to be Conducted:**
   [insert]

5. **How the Information is being collected:**
   [insert]

6. **Confidentiality of Respondents:**
   [insert]

7. **Amount and justification for any proposed incentive**
   [insert]

8. **Questions of a Sensitive Nature**
   [insert]

9. **Description of Statistical Methods**
   [insert]

BURDEN HOUR COMPUTATION *(Number of responses (X) estimated response or participation time in minutes (/60) = annual burden hours):*

Type/Category of Respondent	No. of Respondents	Participation Time (minutes)	Burden (hours)

REQUESTED APPROVAL DATE: [insert]

NAME OF PRA ANALYST & PROGRAM CONTACT: [insert]
FDA CENTER: Center for Tobacco Products