

**FDA/CBER  
DOCUMENTATION FOR THE GENERIC CLEARANCE  
on “Testing Communications on Biological Products”**

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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) and location(s):**  
[insert]
- 5. Collection procedures:**  
[insert]
- 6. Amount and justification for any proposed incentive:**  
[insert]

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

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

**REQUESTED APPROVAL DATE:** [insert]

**NAME OF CONTACT PERSON:** [insert]

**FDA/CDER OFFICE:** [insert]