

## Proposed Survey Questions

1. How much experience do you have working with FDA?
  - 0-3 years
  - 3-6 years
  - 7-10 years
  - 10+ years
  
2. How do you currently receive information from CDER SBAP? (select all that apply)
  - Email
  - Hard Copy documents
  - Website postings
  - CDERLearn courses
  - In-person meetings
  - Webinars
  - Text Messages
  - Other \_\_\_\_\_
  
3. What are your most preferred means for receiving FDA related information? (select three)
  - Email
  - Video Conference
  - CDERLearn
  - Conference Call
  - Webinar, Workshop
  - Community Forum
  - Twitter
  - Hard Copy (regular mail)
  - Text Messages
  - Voicemail
  
4. Other than the CDER outreach materials, have you accessed materials to educate you about the drug application process?
  - Websites, if so which ones \_\_\_\_\_
  - Webinars, if so which ones \_\_\_\_\_
  - Training or other educational materials from another organization. If so, which organization(s)? \_\_\_\_\_
  - I have not accessed any materials from a non FDA CDER source
  
5. How satisfied are you with CDER SBAP's outreach? (1-Very Satisfied: 5 – Very dissatisfied)
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6. Using a scale from 1- 5, with 5 being most effective, 1 being least effective, and N/A to indicate you haven't accessed the item, how would you rate the effectiveness of the current materials and information developed by CDER SBAP?
- Website
  - Webinar
  - Conference
  - Tweets
  - Other
7. What are the top 3 issues you have had getting your drug application reviewed and approved?
- \_\_\_\_\_
  - \_\_\_\_\_
  - \_\_\_\_\_
8. What are the top 3 information products you believe CDER should provide to improve communications with small pharmaceutical businesses? (Select from list below)
- FDA 101 Material: Introduction to the basic operations, authorities, processes and requirements of the FDA
  - A directory of official and unofficial industry guidance sources
  - Checklists for different stages and processes
  - A forum where small pharmaceutical businesses in need of specialized services and consultants offering such services can connect
  - Sample completed application/registration forms
  - A forum where small pharmaceutical businesses can share advice and best practices
  - NDC numbers for raw materials
  - Information about potential sources of grants, loans, or alternative financing
  - Step-by-step process guide for bringing a new drug to market
  - CDER SBAP 101: Basic information about the resources and assistance that CDER SBAP does and does not offer, as well as CDER administrative personnel and processes
  - Lists of currently applicable regulations for a product/process type, and identification of potential regulations that may be applied in the near term
  - Searchable Q&A database
  - List of specific steps needed to comply with FDA requirements for specific products/processes
  - Enhanced options for emailing CDER SBAP and obtaining a specific response to a question
  - Information about average or approximated cost ranges for specific steps/processes
  - Average timelines for specific steps/processes
  - Comparison of US and EU requirements
  - Contact information for subject matter experts inside FDA
  - Tips for Passing Inspection
  - Templates and mock versions of required documents

- Other \_\_\_\_\_

9. Please provide any other comments to help CDER SBAP improve their communication methods.

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10. Would you be open to providing follow-up information? If yes, please provide your name, email and phone number below.

- Name \_\_\_\_\_
- Name of Company \_\_\_\_\_
- Email Address \_\_\_\_\_
- Phone Number \_\_\_\_\_

11. What is your company/organization size?

- 1-9 employees
- 10-49 employees
- 50-499 employees

500+ employees