

DEPARTMENT OF HEALTH AND HUMAN SERVICES  
Food and Drug Administration  
**FY 2017 MDUFA SMALL BUSINESS  
QUALIFICATION AND CERTIFICATION**  
*For a Business Headquartered in the United States*

OMB Number 0910-0508  
Expiration Date: June 30, 2019  
PRA Statement: See page 3.

**Section I – Information about the Business Requesting Small Business Status**

1. Name of business claiming MDUFA Small Business status:		2. Federal Employer Identification Number:	
2a. Organization ID Number (Org ID)		3. Address where business is physically located:	
4. Name of person making this Certification:		5. Your telephone number ( <i>include area code</i> ): (     )	
6. Your mailing address: <input type="checkbox"/> Check if same as item 3.		7. Your e-mail address:	
8. What is your relation to the business claiming MDUFA Small Business status?			
9. Have you listed all of the business's affiliates in Section II of this form? Check <b>one</b> response: <input type="checkbox"/> Yes <input type="checkbox"/> This business has no affiliates.			

10. Complete, sign, and date the following Certification:

I certify that \_\_\_\_\_  
*Name of business (must be identical to response to item 1)*

Check **one** response:

has no affiliates and reported "gross receipts or sales" of no more than \$100,000,000 on its most recent Federal income tax return. I have attached a true and accurate copy of the business's most recent Federal income tax return.

has only the affiliates listed in this Certification, and together with those affiliates reported total "gross receipts or sales" of no more than \$100,000,000 for the most recent tax year. I have attached a true and accurate copy of the entity's most recent Federal income tax return, and a true and accurate copy of the most recent Federal income tax return, or an FY 2017 Foreign Small Business Qualification and Certification, for each of the entity's affiliates.

I further certify that, to the best of my knowledge, the information I have provided in this Certification is complete and accurate. I understand that submission of a false certification may subject me to criminal penalties under 18 U.S.C. § 1001 and other applicable federal statutes.

Signature of person making this Certification (*must be signed by the person identified in item 4*):  
\_\_\_\_\_

Date of this Certification: \_\_\_\_\_

**Section II – Information about You and Your Affiliates**

a. Name of Affiliate	b. Taxpayer ID Number	c. Gross Receipts or Sales
1.		\$
2.		\$
3.		\$
4.		\$
5.		\$
6.		\$
7.		\$
8.		\$
9.		\$
10.		\$
11.		\$
12.		\$
13.		\$
14.		\$
15.		\$
16.	Total Gross Receipts or Sales of All Affiliates <i>(sum of lines 1 through 15)</i>	\$
17.	Gross Receipts or Sales of the Business Making this Certification	\$
18.	<b>Total Gross Receipts or Sales Used to Determine Qualification as a Small Business</b> <i>(sum of lines 16 and 17)</i>	\$

**PRIVACY ACT NOTICE**

This notice is provided pursuant to the Privacy Act of 1974, 5 U.S.C. 552a. The collection of this information is authorized by 21 U.S.C. 379i and 379j. FDA will use the information to assess qualification as a small business, collect and process user fee payments, and facilitate debt collection under the Debt Collection Improvement Act. FDA may disclose information to courts and the Department of Justice in the context of litigation and requests for legal advice, to other Federal agencies in response to subpoenas issued by such agencies, to HHS and FDA employees and contractors to perform user fee services, to the National Archives and Records Administration and General Services Administration for records management inspections, to the Department of Homeland Security and other Federal agencies and contractors in order to detect or respond to system breaches, to banks in order to process payment made by credit card, to Dun and Bradstreet to validate submitter contact information, and to other entities as permitted under the Debt Collection Improvement Act. Furnishing the requested information is mandatory for a business requesting for qualification as a "small business." Failure to supply the information could prevent FDA from processing requests for small business determinations and user fee payments. Additional details regarding FDA's use of information is available online:  
<http://www.fda.gov/regulatoryinformation/foi/default.htm>

**FDA Use Only**

- Review:**  Information verified  
 Information not verified
- Decision:**  Qualifies for Small Business fee discounts  
 Qualifies for Small Business fee discounts and fee waiver for first premarket application
- SBD17 \_\_\_\_\_
- Does not qualify

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The information below applies only to requirements of the Paperwork Reduction Act of 1995.

**\*DO NOT SEND YOUR COMPLETED FORM TO THE PRA STAFF EMAIL ADDRESS BELOW.\***

The burden time for this collection of information is estimated to average 1 hour per response, including the time to review instructions, search existing data sources, gather and maintain the data needed and complete and review the collection of information. Send comments regarding this burden estimate or any other aspect of this information collection, including suggestions for reducing this burden, to:

Department of Health and Human Services  
Food and Drug Administration  
Office of Operations  
Paperwork Reduction Act (PRA) Staff  
*PRAStaff@fda.hhs.gov*

*“An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB number.”*

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## **Instructions for Form FDA 3602 for FY 2017 (FY 2017 MDUFA Small Business Qualification and Certification, for a Business Headquartered in the United States)**

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You should complete and submit Form FDA 3602 for FY 2017 (FY 2017 MDUFA Small Business Qualification and Certification) if you wish to be eligible for reduced or waived fees for medical device submissions you make during FY 2017 (submissions received by FDA from October 1, 2016 through September 30, 2017).

You should also submit:

- a copy of your most recent Federal (U.S.) income tax return, *and*
- if you have any affiliates:
  - a copy of the most recent Federal income tax return of *each* of your domestic (U.S.) affiliates, *and*
  - a copy of an FY 2017 MDUFA Foreign Small Business Qualification and Certification for *each* of your foreign affiliates.

See Sections 738(d)(2) and 738(e)(2) of the Food, Drug, and Cosmetic (FD&C) Act.

FDA will use these materials to decide whether you qualify as a small business within the meaning of MDUFA.

You should mail your FY 2017 MDUFA Small Business Qualification and Certification, and copies of the Federal income tax returns that support your Certification, to FDA at this address:

FY 2017 MDUFA Small Business Qualification  
Division of Industry and Consumer Education  
10903 New Hampshire Avenue  
Building 32, Room 3215  
Silver Spring, MD 20993

If you need assistance, please contact the Division of Industry and Consumer Education at 800-638-2041 or 301-796-7100.

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### **Section I – Information about the Business Requesting Small Business Status**

**1. Name of business claiming MDUFA Small Business status.** Provide the full legal name of the business:

When completing the Form FDA 3602 please assure the business name is the same name as the business name on your U.S. Federal Tax Form

- If the business is a corporation, limited liability company, partnership, or other legal entity, provide the name used in its articles of incorporation, articles of organization, partnership registration, or other similar instrument filed with the State or other government under whose laws the firm was created.
- If the business is a sole proprietorship owned entirely by one individual, provide the name used when filing Federal, State, or other taxes.

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**2. Federal Employer Identification Number.** Your business's Federal Employer Identification Number (EIN) was assigned to you by the U.S. Internal Revenue Service and uniquely identifies your business.

When completing the Form FDA 3602 please assure the Federal Employer Identification Number matches the Federal Employer Identification Number (EIN) on your tax form.

**2a. The organization number (Org ID).** The Org ID is a system generated number assigned to a new organization during the FDA User Fee account creation process. See instructions for obtaining Org ID number on page 17 of the Guidance.

**3. Address where business is physically located.** This is the address where the business is physically located (the address you would give to a person who needed to travel directly to the business's primary establishment).

**4. Name of person making this Certification.** This is the person who is responsible for the accuracy and completeness of the information provided in the Certification and who must sign the Certification (see item 10). This is also a person whom FDA will contact for all communications regarding your FY 2017 MDUFA Small Business Qualification and Certification.

**5. Your telephone number.** This is the telephone number where FDA can reach you if we have a question concerning your FY 2017 MDUFA Small Business Qualification and Certification.

**6. Your mailing address.** This is the address to which you want FDA to send its decision letter informing you that you are, or are not, a small business. If your mailing address is the same as item 3, you can just check the box rather than repeating the information.

**7. Your e-mail address.** This is the e-mail address where FDA can reach you if we have a question concerning your FY 2017 MDUFA Small Business Qualification and Certification. We will be communicating with you via e-mail, please make sure your e-mail address is correct and functioning.

**8. What is your relation to the business claiming MDUFA Small Business status?** Briefly explain your position within the business (e.g., Chief Financial Officer; Vice President; Chief Counsel; or other relationship that gives you authority to provide an FY 2017 MDUFA Small Business Qualification and Certification on behalf of the business).

We will be communicating with you via e-mail, please make sure your e-mail address is correct and functioning.

**9. Have you listed all of the business's affiliates in Section II of this form?** If you have any affiliates, check the first box ("Yes") and list them in Section II of the form. If you do not have any affiliates, check the second box ("This business has no affiliates.").

- **What is an affiliate?** This term is defined by § 737(12) of the FD&C Act. *Affiliate* means a business entity that has a relationship with a second business entity where, directly or indirectly:
  - (a) one business entity controls, or has the power to control, the other business entity; or
  - (b) a third party controls, or has the power to control, both of the business entities.

10. **The applicant's signature on the Form FDA 3602 in box 10 may be a wet (i.e., ink) signature or a valid digital signature. Complete, sign, and date the following Certification.** In this Certification, you should provide the following information:

- The name of the business that is claiming MDUFA Small Business status. This should be identical to your response to item 1.
- Check *one* response to indicate whether the business has any affiliates. Please make sure this agrees with the response in box 9.
  - Check the first box if the business has no affiliates and you have completed box 9 as “This business has no affiliates”.
  - Check the second box if the business has only the affiliates you listed in Section II of the form and you have completed box 9 as “Yes”.

- Check *one* response to indicate how the business determined it met the requirement that it have “gross receipts or sales” of no more than \$100 million:

- Check the first box if the entity reported “gross receipts or sales” of no more than \$100 million on its most recent Federal income tax return. Attach a true and accurate copy (a complete and unaltered copy) of the business’s most recent Federal (U.S.) income tax return. *FDA cannot accept a foreign tax return instead of a Federal (U.S.) income tax return.*

IRS Form	See Line Number
Schedule C (Form 1040)	1
Schedule C-EZ (Form 1040)	1
Form 1065	1a
Form 1065-B	1a
Form 1120	1a
Form 1120-F	Section II, 1a
Form 1120S	1a
Form 990	12
Any other form	<i>Please contact FDA.</i>

- **Where do I find my gross receipts or sales?** You reported your gross receipts or sales on your most recent Federal income tax return. Please note that the following list is not an all-inclusive list for IRS Forms that may contain information on your gross receipts or sales. You should provide all IRS Forms that contain information on your gross receipts or sales.

- Check the second box if the business *and* all of its affiliates *together* reported “gross receipts or sales” of no more than \$100 million on their most recent Federal income tax returns. You should attach a true and accurate copy (a complete and unaltered copy) of the entity’s most recent Federal income tax return *and* a true and accurate copy of each affiliate’s most recent Federal income tax return.
- **What is the most recent tax year?** The most recent tax year will be **2016**, except:
  - If you submit your FY 2017 MDUFA Small Business Qualification and Certification *before* April 15, 2017, *and* you have not yet filed your return for 2016, you may use tax year 2015.
  - If you submit your FY 2017 MDUFA Small Business Qualification and Certification *on or after* April 15, 2017, *and* have not yet filed your 2016 return because you obtained an extension, you may submit your most recent return filed prior to the extension, provided that you include IRS Form 7004 in your application.

- The person identified in item 4 (“Name of person making this Certification”) must sign the Certification.
- Date the Certification (this is the date you signed the Certification).

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## Section II – Information about You and Your Affiliates

Section II of the form provides space for listing up to 15 affiliates; if you have more than 15 affiliates, you may provide the additional information on one or more additional copies of Section II.

### Lines 1 through 15:

List each affiliate on a separate line. For each, you should provide the following information:

a. **Name of Affiliate.** Provide the full legal name of the affiliate:

- If the affiliate is a corporation, limited liability company, partnership, or other legal entity, you should provide the name used in its articles of incorporation, articles of organization, partnership registration, or other similar instrument filed with the State or other government under whose laws the firm was created.
- If the affiliate is a sole proprietorship (that is, it is owned entirely by an individual), you should provide the name used when filing Federal, State, or other taxes.

b. **Taxpayer ID Number.** This number uniquely identifies each business:

- If the affiliate is headquartered in the United States, you should provide the Employer Identification Number (EIN) assigned to the affiliate by the U.S. Internal Revenue Service.
- If the affiliate is headquartered outside the United States, you should provide the Taxpayer Identification Number provided by the National Taxing Authority where the affiliate has its headquarters.

c. **Gross Receipts or Sales.** For each affiliate headquartered in the United States, you should copy this number from the most recent Federal income tax return for the affiliate. See the instruction for item 9 to learn where you will find this information on a Federal income tax return. For each affiliate headquartered outside the United States, you should copy the information from item 3.b. of the National Taxing Authority Certification for the affiliate.

16. **Total Gross Receipts or Sales of All Affiliates.** This is the sum of the Gross Receipts or Sales shown in column c. of lines 1 through 15. If you have no affiliates please enter “0”.

17. **Gross Receipts or Sales of the Business Making this Certification.** This is the gross receipts or sales of the business identified in Section I, item 1.

18. **Total Gross Receipts or Sales Used to Determine Qualification as a Small Business.** This is the sum of lines 16 and 17. To qualify as a MDUFA small business for fee discounts, this sum must be **no more than** \$100 million. See Sections 738(d)(2)(A) and 738(e)(2)(A) of the FD&C Act.

