513(g) Requests for Information

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I have a product and I am not sure if it is a medical device?

I have a product and I am not sure how it is classified?

What do I do?
Learning Objectives

1. Describe how FDA addresses device determination inquiries

2. Outline the 513(g) Request process

3. Explain when it is appropriate to send an informal device determination email or a formal 513(g) request for information
Definition of a Medical Device

Section 201(h) of the Food, Drug & Cosmetic Act (FD&C Act) defines a device as:

An instrument, apparatus, implement, machine, contrivance, implant, in vitro reagent, or other similar or related article, including a component part or accessory which is:
Definition of a Medical Device (Continued)

• Recognized in the **official National Formulary**, or the **United States Pharmacopoeia**, or any supplement to them,

• Intended for use in the **diagnosis of disease** or other **conditions**, or in the **cure, mitigation, treatment, or prevention of disease** in man or other animals, or

• Intended to **affect the structure or any function** of the body of man or other animals
Definition of a Medical Device
(Continued)

• And does **not** achieve its primary intended purposes through **chemical action** within or on the body of man or other animals and which is not dependent upon being **metabolized** for the achievement of its primary intended purposes

• The term "device" does **not** include **software functions excluded pursuant to section 520(o)"**
Regulatory Classes

Devices are classified based on level of risk

- Class III: High Risk (PMA)
- Class II: Moderate Risk (510(k))
- Class I: Low Risk majority are (510(k) Exempt)
Regulatory Classes

Devices are classified based on level of risk

Class III
High Risk
(PMA)

Class II
Moderate Risk
(510(k))

Class I
Low Risk
majority are (510(k) Exempt)

Pre- market Approval (PMA)

Pre-market Notification 510(k)

510(k) Exempt = No 510(k) submission required
De Novo Classification Request (De Novo)

- Automatically been placed in Class III due to lack of an appropriate predicate
- Alternate pathway to classify novel devices of low to moderate risk

Acceptance Review for De Novo Classification Requests
www.fda.gov/regulatory-information/search-fda-guidance-documents/acceptance-review-de-novo-classification-requests
Device Categories

There are different types of medical devices:

- General Wellness
- Aesthetic
- Diagnostic
- Therapeutic
- Software as a Medical Device (SAMD)
- Combination Products
- Mobile Medical Application
Is My Product A Medical Device?
Your Device Determination Steps

1. Know the intended use and technology of the device. What does it do? How does it do it?
2. Does it meet the definition of a medical device per section 201(h) of the FD&C Act?
3. Check FDA databases for existing or similar device

Still Unsure? Obtain Feedback from Us
Mechanism For Obtaining Feedback

Option 1 - Informal

Email Device Determination Mailbox

CDRH provides quick and informal response

Option 2 - Formal

Submit a 513(g) Request

FDA provides formal response with device classification if applicable
Option 1 - Informal Device Determination

- Email inquiry to Device Determination Team at: DeviceDetermination@fda.hhs.gov

- Include the following information in your email:
  - Brief device description
  - Clear intended use
  - List or picture of all labeling claims
Example Email Inquiry

• I have a heat pack made of fabric that contains oats
• The pack is heated in the microwave and can be reused
• The heat pack is intended to provide pain relief and increase blood circulation
• Is my product a medical device regulated by the FDA?
CDRH Informal Device Determination Response

• Yes/no response with additional relevant information (e.g. link to regulation or guidance document) when appropriate

• Generally provided within 7 days

• If inquiry is complex and cannot be adequately addressed via email, we may recommend submission of a 513(g) Request
Example Email CDRH Response

• Based on the information you provided, the heat pack product appears to be a medical device per Section 201(h) of the Federal Food Drug & Cosmetic (FD&C) Act

• This is our best judgment on how the product would be regulated. This response is not a classification decision and does not constitute FDA clearance or approval for commercial distribution of a product
Option 2 - Formal Device Determination 513(g) Request

• Request for classification information and regulatory requirements for a particular product

• Governs requests "for information respecting the class in which a device has been classified or the requirements applicable to the device."
Information to Include in a 513(g) Request

1. Cover Letter
2. Device Description
3. Intended Use
4. Labeling Claims

Guidance: FDA and Industry Procedures for Section 513(g) Requests for Information under the Federal Food, Drug, and Cosmetic Act

1. Cover Letter

- Date of the request
- Company name
- Contact person
- Contact information (email and phone number)
- Device Name
- Question(s) to FDA
2. Device Description

- Brief description
- Describe mechanism of action
- Describe disease or condition the device is intended for
- Include picture or schematic, if available
- Patient population
3. Intended Use

- Describe what the device is used for
- Physiological purpose

**Example:** The device is intended for removal of body fat
4. Labeling Claims

• Promotional claims beyond the intended use

Example: Device removes 10% of body fat
Types of 513(g) Questions

**APPROPRIATE**

- Is the product subject to FDA regulations?
- What is the appropriate regulatory pathway?
- Is the product exempt from 510(k)?

**NOT APPROPRIATE**

- Type of performance data required?
- Adequacy of performance data?
- Adequacy of predicate?
- Request for reclassification?
Overview of 513(g) Review Process

513 Request is received by FDA via the document control center and checked for user fee and eCopy

Lead Reviewer reviews submission & provides recommendation

Management concurrence before letter issuance

FDA User Fee Programs
www.fda.gov/industry/fda-user-fee-programs

eCopy Program for Medical device Submissions
www.fda.gov/medical-devices/how-study-and-market-your-device/ecopy-program-medical-device-submissions
FDA Review 513(g) Request

- All information provided is reviewed by a lead reviewer, who may work with other subject matter experts.
- Regulation and precedent set by existing medical devices is considered.
- A medical device determination is made for the product based on technology, intended use, and medical claims.
- If needed, additional information is requested.
- FDA provides a recommendation and issues a letter via email within 60 calendar days.
FDA’s Response

• Response via formal letter
• Letter includes:
  o Device determination
  o Device class if applicable (i.e. Class I, II, or III)
  o Indicate whether device is under enforcement discretion, when appropriate
  o Recommended regulatory pathway (e.g. 510(k), exempt, De Novo, etc.)
  o Other applicable information when appropriate (e.g. guidance document)
EXAMPLE

Pear Company
5c Johnathan Doe
Consultant
Medical Device Company, LLC
121 Apple Bay Court, NE
Miami, Florida 33704

Re: C191234
   Product Name: Device A
   Dated: May 5, 2019
   Received: May 7, 2019

Dear Johnathan Doe:

We have reviewed the above referenced request for information, submitted in accordance with Section 513(g) of the Federal Food, Drug, and Cosmetic Act (Act), regarding the regulatory requirements applicable to the Device A. Based on the information provided in your submission, we believe that the Device A is an a Heat And Moisture Condenser (Artificial Nose). The Food and Drug Administration (FDA) has classified a Heat And Moisture Condenser (Artificial Nose) as a Class I type device under Title 21 of the Code of Federal Regulations (CFR) 868.3375. A Heat And Moisture Condenser (Artificial Nose) is a Class I type device, exempt from the premarket notification [510(k)] requirements of the Act, subject to the limitations to the exemption found in 21 CFR 868.9.
Responses and Information

• Responses provided via email or 513(g) are recommendations and not binding feedback and does not constitute FDA clearance or approval for commercial distribution of a product

• Stakeholder information provided via the device determination mailbox and 513(g) Request for Information is confidential
References


• User Fees for 513(g) Requests for Information Guidance (2017)
  www.fda.gov/regulatory-information/search-fda-guidance-documents/user-fees-513g-requests-information

• Is a product a medical device? FDA website
  www.fda.gov/medical-devices/classify-your-medical-device/product-medical-device
References

• General Wellness: Policy for Low Risk Devices Guidance (2016)

• Medical Mobile Application FDA website
  www.fda.gov/medical-devices/digital-health/mobile-medical-applications

• Software as a medical device FDA website
  www.fda.gov/medical-devices/digital-health/software-medical-device-samd
Providing Industry Education

1. CDRH Learn – Multi-Media Industry Education
   • Videos, audio recordings, power point presentations, software-based “how to” modules describing aspects of medical device and radiation emitting product regulations: [www.fda.gov/CDRHLearn](http://www.fda.gov/CDRHLearn)

2. Device Advice – Text-Based Education
   • Text-based resource that explains many aspects of medical device laws, regulations, guidances, and policies: [www.fda.gov/MedicalDevices/DeviceRegulationandGuidance](http://www.fda.gov/MedicalDevices/DeviceRegulationandGuidance)

3. Division of Industry and Consumer Education (DICE)
   • If you have a question - Email: [DICE@fda.hhs.gov](mailto:DICE@fda.hhs.gov)
   • Phone: 1(800) 638-2041 or (301) 796-7100 (Live Agents 9a –12:30 pm; 1-4:30 pm EST)
Summary

• A product is considered to be a medical device if it meets the definition in Section 201(h) of the FD&C Act

• There are two mechanisms to receive feedback, email us at device determination or submit a 513(g) request
Your Call To Action

- Search for similar devices in FDA databases
- Email DeviceDetermination@fda.hhs.gov for a simple inquiry
- Submit a 513(g) Request for a complex inquiry and/or if you are seeking device classification
- Refer to the FDA 513(g) Request For Information Guidance for instructions when submitting a 513(g)