Regulamento de UDI do FDA dos EUA e Atualizações Recentes

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20 de janeiro de 2022

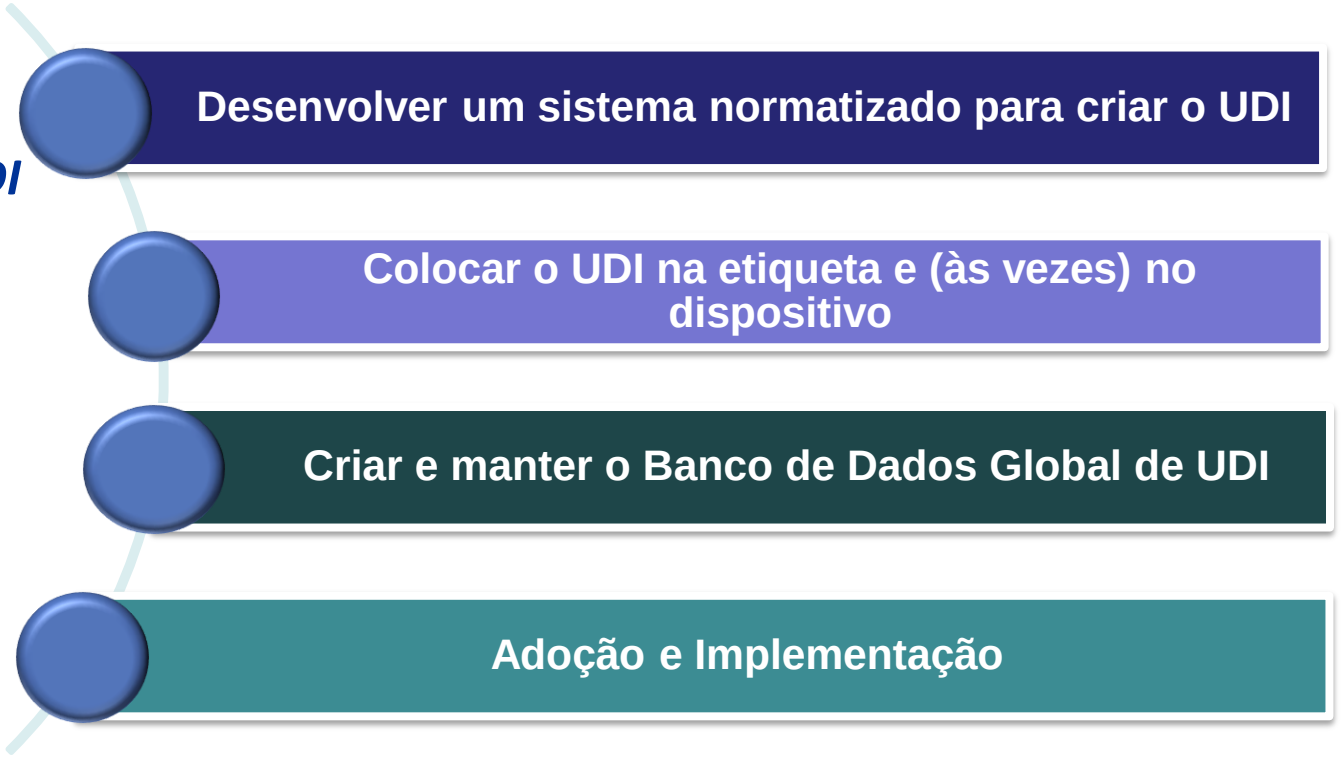
Tradução não oficial

Estabelecendo um Sistema UDI

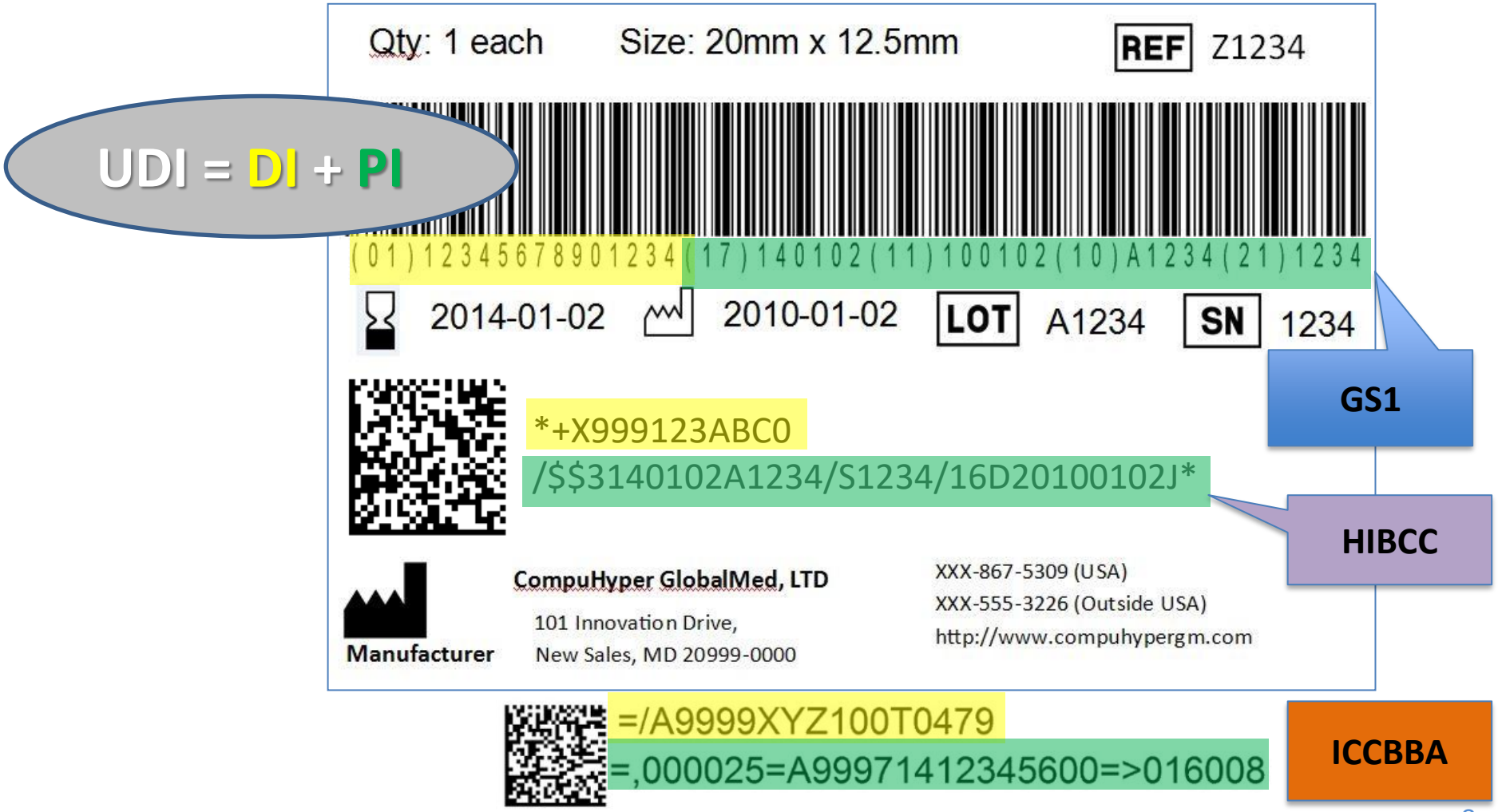
Regra final do UDI

[78 FR 58786]

24 set 2013



Identificador Único de Dispositivos (UDI)



Prazo de Implementação

	Data	Deve suportar um UDI e Enviar dados ao GUDID	Marcação Direta (para certos usos pretendidos)
✓	**24 set 2014**	**Dispositivos de classe III** Dispositivos licenciados pela Lei PHS	
✓	**24 set 2015**	Dispositivos implantáveis, de suporte de vida e de sustentação de vida (I/LS/LS)	Dispositivos LS/LS
✓	**24 set 2016**	**Dispositivos de classe II** (que não são I/LS/LS)	Dispositivos de classe III e dispositivos licenciados pela Lei PHS
✓	**24 set 2018**	Dispositivos de classe I Dispositivos não classificados (que não são I/LS/LS)	Dispositivos de classe II (que não são LS/LS)
✓	**24 set 2020**	**Dispositivos de classe I** Dispositivos não classificados (que não são I/LS/LS)	Dispositivos de classe I Dispositivos não classificados (que não são LS/LS)
	24 set 2022	**Dispositivos de classe I Dispositivos não classificados (que não são I/LS/LS)**	**Dispositivos de classe I Dispositivos não classificados (que não são LS/LS)**

Diretriz de 2018

Diretriz de 2020

Diretriz de 2018

Harmonização Internacional

Tradução não oficial

IMDRF/UDI WG/N7 Final: 2013
Diretriz sobre UDI

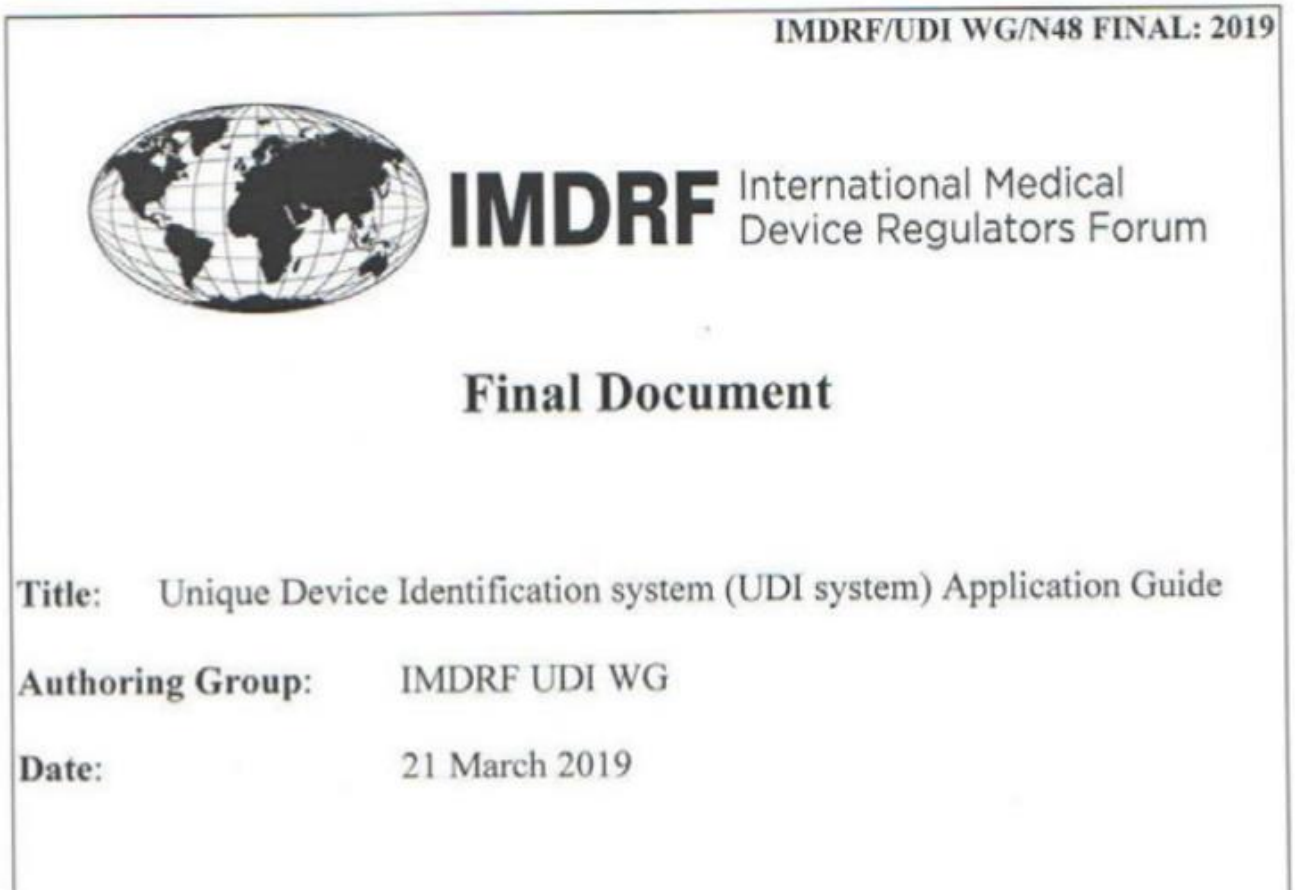
IMDRF International Medical Device Regulators Forum

Final Document

Title:	UDI Guidance Unique Device Identification (UDI) of Medical Devices
Authoring Group:	IMDRF UDI Working Group
Date:	9 December 2013

Tradução não oficial

IMDRF/UDI WG/N48 Final: 2019
Guia de Aplicação do Sistema UDI



IMDRF/UDI WG/N48 FINAL: 2019

IMDRF International Medical Device Regulators Forum

Final Document

Title: Unique Device Identification system (UDI system) Application Guide

Authoring Group: IMDRF UDI WG

Date: 21 March 2019

Tradução não oficial

IMDRF/UDI WG/N53 FINAL: 2019

	A	B	C	D	E	F	G	H
1	Jurisdiction	IMDRF			European Union		United States	
2	Source	IMDRF/UDI WG/N7FINAL:2013 Section 9.2			http://ec.europa.eu/growth/sectors/medical-devices/new-regulations/guidance_en		https://www.fda.gov/MedicalDevices/DeviceRegulationandGuidance/UniqueDeviceIdentification/GlobalUDIDatabaseGUDID/ucm416106.htm	
3	Date revised				18-Feb-2019		18-Feb-2019	
4		Data Element #	Data Element	UDID Requirement	Data Element	EUDAMED Requirement	Data Element	GUDID Requirement
5	Device Identification & Description	--	--	--	Basic UDI-DI	R	--	--
6		1	UDI-DI (UDI type, e.g. GS1 GTIN, HIBC-LIC, ISBT-128 PPIC)	R	UDI-DI	R	Primary DI Number	R
7					Issuing Entity (UDI-DI)	R	Issuing Agency	R
8					Isuing Entity (Basic UDI-DI)	R	--	--
9		--	--	--	Single Registration Number	R	--	--
10		9	Brand Name	C	Name or Trade name (name of product)	R	Brand Name	R
11		11	Device model or version	R	Name or, if applicable, device model that identifies the BASIC UDI-DI Group in the technical documentation and/or certificate and declaration of conformity	R	Version or Model	R
12		12	Reference and/or catalogue number	C	Reference, article or catalogue number	R	Catalog Number	O
13		15	Additional product Description (optional) – Additional clinically relevant information, e.g. radio-opaque	O	Additional product Description	O	Device Description	O
14		--	--	--	Direct Marking (DM) DI	R	DM DI Different from Primary DI	C
15		--	--	--	--	--	DM DI Number	C
16		--	--	--	Issuing Entity (Secondary DI)	C	Issuing Agency (Secondary DI)	O
17		1	Additional device identifier(s) (if applicable) e.g. GS1, HIBC, or ISBT-128	C	Secondary UDI-DI	C	Secondary DI Number	O
18		--	--	--	--	--	Issuing Agency (Previous DI)	O
19		--	--	--	--	--	Previous DI Number	O
20		2	Unit of Use UDI-DI	R	Unit of Use (UoU) UDI-DI	C	Unit of Use DI Number	C
21		--	--	--	Quantity of devices	R	Device Count	R
22		--	--	--	Additional Trade Names	C	--	--

Tradução não oficial

IMDRF/UDI WG/N54 Final: 2019
Requisitos do sistema relacionados ao uso de UDI na saúde incluindo casos de uso selecionados

IMDRF/UDI WG/N54 FINAL:2019

IMDRF International Medical Device Regulators Forum

FINAL DOCUMENT

International Medical Device Regulators Forum

Title: System requirements related to use of UDI in healthcare including selected use cases

Authoring Group: IMDRF UDI WG

Date: 21 March 2019

Elena M. Astapenko, IMDRF Chair

Tradução não oficial

Página da Web sobre UDI do FDA dos EUA

UDI Rule and Guidances, Training, Resources, and Dockets

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This page provides a comprehensive list of links to:

- UDI rule and guidances
- UDI training for industry
- UDI and GUDID technical documents
- UDI dockets

www.fda.gov/udi

Diretriz Final sobre Formato e Conteúdo do UDI

Figure 1 shows an example of a UDI in both easily readable plain-text and AIDC forms.

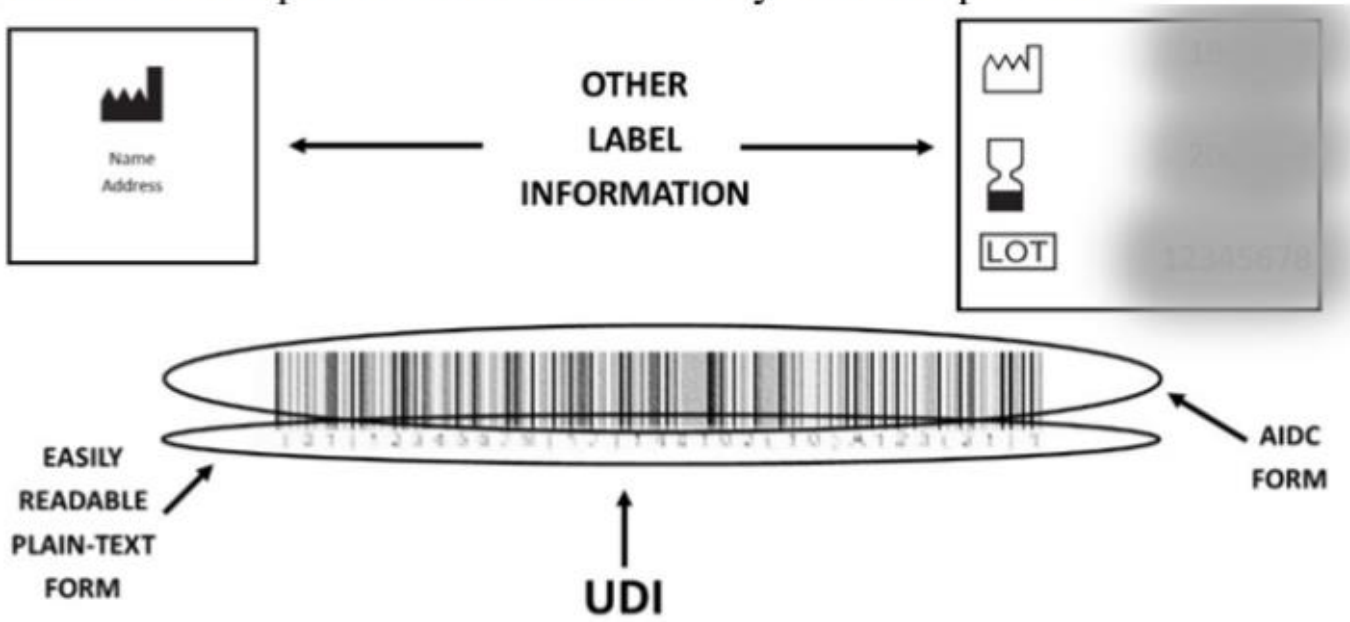

Figure 1. UDI in Easily Readable Plain-Text and AIDC Forms

Contains Nonbinding Recommendations

Draft – Not for Implementation

Select Updates for Unique Device Identification: Policy Regarding Global Unique Device Identification Database Requirements for Certain Devices

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Draft Guidance for Industry and Food and Drug Administration Staff

DRAFT GUIDANCE

This draft guidance document is being distributed for comment purposes only.

Document issued on October 14, 2021

You should submit comments and suggestions regarding this draft document within 60 days of publication in the *Federal Register* of the notice announcing the availability of the draft guidance. Submit electronic comments to https://www.regulations.gov. Submit written comments to the Dockets Management Staff, Food and Drug Administration, 5630 Fishers Lane, Room 1061, (HFA-305), Rockville, MD 20852. Identify all comments with the docket number listed in the notice of availability that publishes in the *Federal Register*.

For questions about this document regarding CDRH-regulated devices, contact UDI Regulatory Policy Support, 301-796-5995; email: GUDIDSupport@fda.hhs.gov. For questions about this document regarding CBER-regulated devices, contact the Office of Communication, Outreach, and Development (OCOD) at 1-800-835-4709 or 240-402-8010, or by email at ocod@fda.hhs.gov.

FDA U.S. FOOD & DRUG ADMINISTRATION

U.S. Department of Health and Human Services
Food and Drug Administration
Center for Devices and Radiological Health
Center for Biologics Evaluation and Research

cial

Envio de Dados

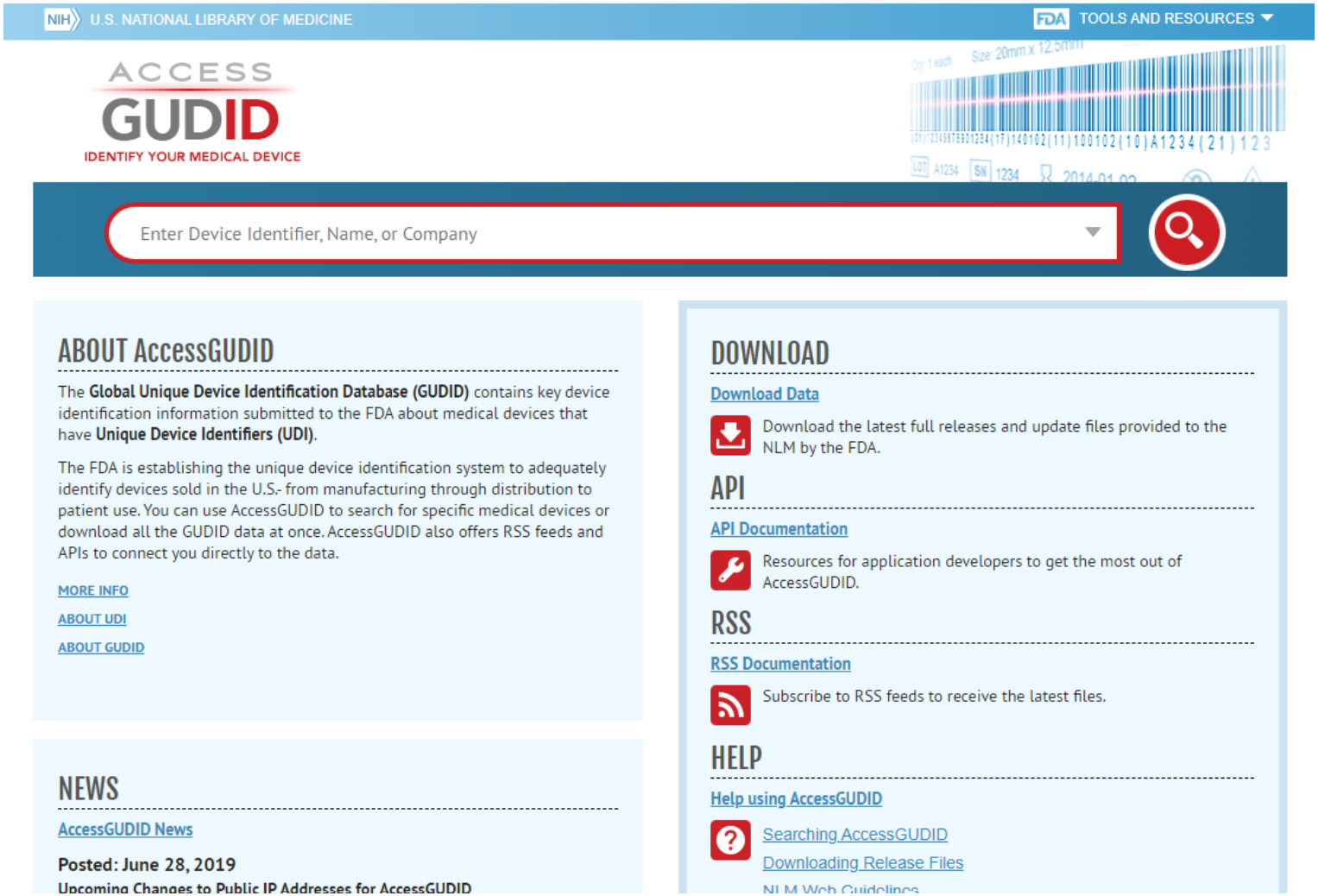
GUDID Global Unique Device Identification Database

ACCESS GUDID
IDENTIFY YOUR MEDICAL DEVICE

Uso de Dados

Informações de identificação de dispositivos de referência
Enviar uma vez – reutilizar em sistemas downstream
Otimizar qualidade de dados com base no uso

AccessGUDID



https://accessgudid.nlm.nih.gov/

Central de Ajuda para UDI do FDA

GUDIDSupport@fda.hhs.gov