



www.mearis.cms.gov/public/home

Register: To submit and manage a MEARIS™ application or request, you must have or create a registered account.

Login: To log in to MEARIS™, applicants and CMS users must be registered.

Resources: Click from a topic list to see information related to that application, request, or the MEARIS™ platform. Request application related questions can be submitted to CMS using the form available under “Contact” on the Resources screen.

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MEARIS™
Medicare Electronic Application
Request Information System™

Login Resources

One place to manage Medicare coding and payment related applications

Register to submit applications and requests

[Register](#)

Hospital Inpatient Applications and Requests

The Inpatient Prospective Payment System (IPPS) sets forth a system of payment for operating costs of acute care hospital inpatient stays under Medicare Part A. Each case is categorized into a diagnosis-related group (DRG) and a payment weight is assigned to it. Medicare makes payments for the costs of approved graduate medical education (GME) programs to hospitals, including hospitals that are not paid under the IPPS.

Click an application to learn more about it.

- New Technology Add-on Payments (NTAP)
Coming Soon!
- The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)
Request Deadline 2022: 12/2/2022
- New & Revised Medicare Severity Diagnosis Related Groups (MS-DRG)
Request Deadline 2024: 10/20/2022



Requestors are strongly encouraged to review each of the links provided on the “Important Information” screen before starting a request

The screenshot shows the MEARIS (Medicare Electronic Application Request Information System) interface. At the top, there is a dark blue navigation bar with the MEARIS logo and the text 'Medicare Electronic Application Request Information System™'. To the right of the logo are navigation links for 'Home', 'Tasks', 'Applications', and 'Teams', along with a user profile icon. Below the navigation bar is a white main content area. At the top center of this area is a green icon of a medical bag with a cross. Below the icon is the heading 'Welcome to the International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS) Request'. Underneath the heading is a section titled 'Important Information' which contains a list of six links, each preceded by a green information icon: 'Application Guidance', 'Application Process and Timeline', 'How the Online Application Works', 'Preview New Code Application PDF', 'Click to see the latest ICD-10-PCS codes', and 'PRA Disclosure Statement'. At the bottom of the page, there is a red circle highlighting a blue button labeled 'Go' and a white button labeled 'Cancel' below it.



Technical support is available under “Useful Links” at the bottom of the MEARIS™ site

mearis.cms.gov/secure/icd-10-pcs/introduction

- [Application Process and Timeline](#)
- [How the Online Application Works](#)
- [Preview New Code Application PDF](#)
- [Click to see the latest ICD-10-PCS codes](#)
- [PRA Disclosure Statement](#)



Ready to get Started?

[Go](#)

[Cancel](#)

Useful Links

- [CMS Web Policies](#)
- [Technical Support](#)
- [Resources](#)

A federal government website managed and paid for by the U.S. Centers for Medicare & Medicaid Services, 7500 Security Boulevard, Baltimore MD 21244
Medicare Electronic Application Request Information System™ (MEARIS™)
v2.12.4944



New ICD-10-PCS Request:

Requests to add new code(s)

Revise ICD-10-PCS Request:

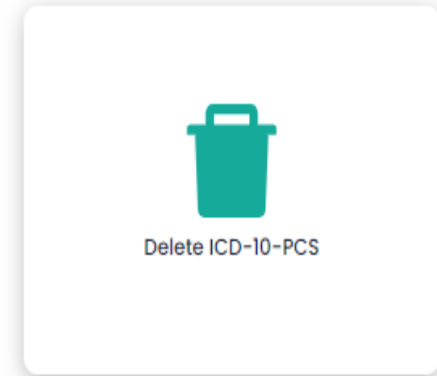
Requests to revise existing ICD-10-PCS code(s) (e.g. add additional approach)

Delete ICD-10-PCS Request:

Requests to delete existing ICD-10-PCS code(s)

Any request submitted should include a description of the new code or change being requested, and rationale for why the new code or change is needed.

What type of ICD-10-PCS request would you like to complete?



Back



Spartans | The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)

Contact Info

Drug or Technology Info

New Code

NTAP Info

FDA Info

Attachments

Summary

Who is the primary contact?

First name

First name is required.

Middle name (optional)

Last name

Last name is required.

Organization

Organization is required.

Occupation/Job Title

Occupation/Job Title is required.

US Phone Number

Phone Number is required.

Extension (optional)

Email address

Please enter a valid email address.

Country

United States

Mailing address line 1

Mailing address line 1 is required.

Mailing address line 2 (optional)

City

City is required.

State

State is required.

Zip code

Zip code is required.

Relationship

Relationship is required.

Next



Spartans | The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)

Contact Info

Drug or Technology Info

New Code

NTAP Info

FDA Info

Attachments

Summary

Who is the secondary contact?

First name First name is required.	Middle name (optional)	Last name Last name is required.
Organization Organization is required.	Occupation/Job Title Occupation/Job Title is required.	
US Phone Number Phone Number is required.	Extension (optional)	Country United States
Email address Please enter a valid email address.		
Mailing address line 1 Mailing address line 1 is required.		
Mailing address line 2 (optional)		
City City is required.	State State is required.	Zip code Zip code is required.
Relationship Relationship is required.		

Back

Next



If **“Yes”** is selected, another screen will appear to allow the requestor to provide the manufacturers’ demographic information.

If **“No”** is selected, the system will navigate to a screen where the requestor will briefly explain why they have indicated that there is no manufacturer associated with this application.

The screenshot shows the MEARIS web application interface. At the top left is the MEARIS logo with the text 'Medicare Electronic Application Request Information System™'. To the right of the logo are navigation links: 'Home', 'Tasks', 'Applications', and 'Teams', each with a corresponding icon. A user profile icon is also visible in the top right corner. Below the navigation bar, the page title is 'Spartans | The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)'. A horizontal menu contains several tabs: 'Contact Info', 'Drug or Technology Info', 'New Code', 'NTAP Info', 'FDA Info', 'Attachments', and 'Summary'. The 'Contact Info' tab is currently selected and highlighted with a teal underline. The main content area displays the question 'Is there a manufacturer associated with this application?' in a large, bold, black font. Below the question are two large, white, rounded square buttons. The left button features a teal checkmark icon and the text 'Yes'. The right button features a teal 'X' icon and the text 'No'. At the bottom left of the screen, there is a 'Back' button with a teal border and rounded corners.



Spartans | The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)

Contact Info

Drug or Technology Info

New Code

NTAP Info

FDA Info

Attachments

Summary

Select the appropriate category



Drug/Therapeutic Agent



Procedure/Technology

Back



Drug/Therapeutic Agent

Describe the drug/therapeutic agent:

- Describe the mechanism of action
 - What is it?
 - What does it do?
 - What are the procedural steps involved?
- What are the routes of administration for the drug?

MEARIS™
Medicare Electronic Application Request Information System™

Home Tasks Applications Teams

Spartans | The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)

Contact Info **Drug or Technology Info** New Code NTAP Info FDA Info Attachments Summary

Describe the drug/therapeutic agent

Briefly describe the drug/therapeutic agent for which you are requesting a new procedure code.

Soliris® (eculizumab), a first-in-class complement inhibitor, is the first and only FDA approved therapy for adults with anti-AQP4 antibody-positive NMOSD.

155 / 3000

Please describe the steps and route of administration of the drug/therapeutic agent.

Soliris® (eculizumab) is administered via an IV infusion by a healthcare professional according to the following schedule:

- 900 mg weekly for the first 4 weeks, followed by
- 1200 mg weekly for the fifth dose 1 week later, then
- 1200 mg every 2 weeks thereafter

263 / 3000

Back Next



- What diagnoses are associated with or indicated for use of the drug/therapeutic agent?
- How is the indication currently treated or managed?
- Have there been any associated complications/sequela/adverse events? If yes, how many and what did they consist of? (E.g. fever, shortness of breath, anaphylaxis, etc.)

Drug/Therapeutic Agent

MEARIS™
Medicare Electronic Application Request Information System™Home Tasks Applications Teams

Spartans | The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)

Contact infoDrug or Technology infoNew CodeNTAP InfoFDA InfoAttachmentsSummary

Provide the diagnostic details for this drug/therapeutic

Which diagnosis/diagnoses is the drug/therapeutic agent indicated to address? Please provide ICD-10-CM codes. (optional)

Neuromyelitis Optica Spectrum Disorder (NMOSD) - (ICD-10-CM G36.0) is a rare autoimmune disorder of the Central Nervous System (CNS). This disease affects approximately 1.96 per 100,000 individuals and is characterized by neuroinflammatory relapses that result in progressive and irreversible damage to the optic nerve and spine. Relapses in NMOSD are unpredictable and can cause blindness, paralysis and increased overall mortality. Approximately 90% of patients with NMOSD suffer relapses, approximately 50% of patients with NMOSD have experienced a relapse within one year and approximately 76% of patients do not recover completely from their first relapse. Therefore, a primary goal in the treatment of NMOSD is prompt initiation of relapse prevention upon a diagnosis of NMOSD.

786 / 3000

[Click to see the latest ICD-10-CM codes](#)

Have there been any adverse outcomes or complications associated with the administration of the drug/therapeutic agent? If yes, how many and what was the nature of the adverse outcome? (E.g. rash, fever, shortness of breath, etc.)

A total of 75 adverse events (AEs) were reported by participants in the phase 3 trial. Minor itching was the most common (24.3%)

128 / 3000

BackNext



Drug/Therapeutic Agent



Spartans | The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)

Contact Info

Drug or Technology Info

New Code

NTAP Info

FDA Info

Attachments

Summary

Provide the utilization details for this drug/therapeutic

Identify the number of times the drug/therapeutic agent has been (will be) administered.

In clinical trials, this therapeutic was administered to 300 patients

69 / 3000

What is the percentage of time the drug/therapeutic has been (will be) used across the following care settings? (optional)

Hospital Inpatient Facilities:

Number of Anticipated Cases

Percentage of Medicare beneficiaries %

Percentage of use Inpatient %

Outpatient Facilities/Physician Office:

Number of Anticipated Cases

Percentage of Medicare beneficiaries %

Percentage of use Outpatient %

Back

Next



Drug/Therapeutic Agent



The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)

Contact Info

Drug or Technology Info

New Code

NTAP Info

FDA Info

Attachments

Summary

How is the drug/therapeutic agent documented?

How and where (e.g. O.R. Report, Notes, etc.) will the drug/therapeutic agent be documented in the medical record?

Documentation would be found in the progress notes and medication administration record (MAR)

93 / 3000

Are there various terms that are used to describe the drug/therapeutic agent? (Please list)

Terms used to describe the drug/therapeutic are Soliris[®] or eculizumab

93 / 3000

Back

Next



Spartans | The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)

Contact Info

Drug or Technology Info

New Code

NTAP Info

FDA Info

Attachments

Summary

Select the appropriate category



Drug/Therapeutic Agent



Procedure/Technology

Back



Describe the device/technology/service or procedure.

- What is it?
- What does it do?
- How is it used?
- What are the procedural steps involved?
- If the technology is a device or implant, is only one device/implant routinely inserted or can multiple devices/implants be utilized?

Procedure/Technology

MEARIS™
Medicare Electronic Application Request Information System™

Home Tasks Applications Teams

Spartans | The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)

Contact Info **Drug or Technology Info** New Code NTAP Info FDA Info Attachments Summary

Describe the procedure/technology

Briefly describe the procedure/technology for which you are requesting a new procedure code.

Provide Response

0 / 3000

Please describe the procedure/technology and the steps involved in the performance of the procedure/technology.

Provide Response

0 / 3000

If the technology involves a device or implant, is only one device or implant routinely inserted/implanted or is it possible for multiple devices/implants to be utilized in one operative episode? (E.g. valves, stents, etc.)

Provide Response

0 / 3000



Describe the device/technology/service or procedure.

- If the technology involves a device or implant, is the device considered permanent?
- If the procedure involves vessels or specific body parts, is it beneficial or necessary to identify a range of the specific site? (E.g. 2-3 vertebrae, 4+ vessels or stents, etc.)
- Is the procedure/technology performed in conjunction with another procedure/technology or is it considered a standalone procedure/technology?

Procedure/Technology

MEARIS™
Medicare Electronic Application Request Information System™

Home Tasks Applications Teams

Spartans | The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)

Contact Info **Drug or Technology Info** New Code NTAP Info FDA Info Attachments Summary

Describe the procedure/technology (cont.)

If the technology involves a device or implant, is the device considered permanent? Would there ever be an occasion when a code for removal or revision would be needed?

Provide Response

0 / 3000

If the procedure involves vessels or specific body parts, is it beneficial or necessary to identify a range of the specific site? (E.g. 2-3 vertebrae, 4+ vessels or stents, etc.)

Provide Response

0 / 3000

Is the procedure/technology performed in conjunction with another procedure/technology or is it considered a standalone procedure/technology?

Provide Response

0 / 3000



Provide information regarding the clinical indication for this device/technology/service or procedure.

- What diagnoses are associated with or indicated for use of the device/technology/service or procedure?
- How is the indication currently treated or managed?
- Have there been any associated complications/sequela/adverse events? If yes, how many and what did they consist of? (E.g. dislodgement, failure, loosening, etc.)

Procedure/Technology

MEARIS™
Medicare Electronic Application Request Information System™Home Tasks Applications Teams

Spartans | The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)

Contact InfoDrug or Technology InfoNew CodeNTAP InfoFDA InfoAttachmentsSummary

Provide the diagnostic details for this procedure/technology

Which diagnosis/diagnoses is the procedure/technology indicated to address? Please provide ICD-10-CM codes. (optional)

Provide Response

0 / 3000

[Click to see the latest ICD-10-CM codes](#)

Have there been any adverse outcomes or complications? If yes, how many and what did they consist of? (E.g. dislodgement, failure, loosening, etc.)

Provide Response

0 / 3000



Procedure/Technology

Spartans | The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)

[Contact Info](#) [Drug or Technology Info](#) [New Code](#) [NTAP Info](#) [FDA Info](#) [Attachments](#) [Summary](#)

Provide the Utilization details for this procedure/technology

Identify the number of times the procedure has been (will be) performed using this technology.

Provide Response

0 / 2000

What is the percentage of time the procedure/technology has been (will be) performed/used across the following care settings? (optional)

Hospital Inpatient Facilities:

Number of Anticipated Cases

Percentage of Medicare beneficiaries %

Percentage of use Inpatient %

Outpatient Facilities/Physician Office:

Number of Anticipated Cases

Percentage of Medicare beneficiaries %

Percentage of use Outpatient %



Procedure/Technology

MEARIS™
Medicare Electronic Application
Request Information System™

[Home](#) [Tasks](#) [Applications](#) [Teams](#)

Spartans | The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)

Contact Info

Drug or Technology Info

New Code

NTAP Info

FDA Info

Attachments

Summary

How is the procedure/technology documented?

How and where (e.g. O.R. Report, Notes, etc.) will the procedure/technology be documented in the medical record?

Provide Response

0 / 3000

Are there various terms that are used to describe the procedure/technology? (Please list)

Provide Response

0 / 3000



Are there existing codes currently being reported by facilities to describe this drug or technology?



Yes



No



Other/Don't know

Back

Select:

- **Yes** – Requestor will enter the ICD-10-PCS code(s) currently used, if known, and indicate why they believe that existing codes do not adequately capture the drug or technology
- **No**
- **Other/Don't know** – Requestor will provide an explanation

Requestors will have the opportunity to provide a recommendation for possible new ICD-10-PCS code titles (e.g. approach, body part, device, qualifier)

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Contact Info

Drug or Technology Info

New Code

NTAP Info

FDA Info

Attachments

Summary

Have you applied or are you applying for New Technology Add-on Payments(NTAP)
for consideration?



Yes



No

 [Click here to learn more about starting an NTAP application](#) 

Back



Spartans | The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)

Contact Info

Drug or Technology Info

New Code

NTAP Info

FDA Info

Attachments

Summary

Provide some details about your NTAP application

What is the name of the technology?

Name

For which Fiscal year (FY) was the/will the NTAP application be submitted?

Year

What is the NTAP application confirmation number? (optional)

NTAP Application number

Back

Next



Approved – Requestor will provide the FDA approval date and, if applicable, any additional FDA details

Pending Approval – Requestor will provide the anticipated FDA approval date and, if applicable, any additional FDA details

Not Approved - Requestor will provide the FDA submission date and, if applicable, any additional FDA details

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Request Information System™

Home Tasks Applications Teams

Spartans | The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)

Contact Info Drug or Technology Info New Code NTAP Info **FDA Info** Attachments Summary

Is the drug/procedure/technology FDA approved?

Approved Pending Approval Not Approved

Back



To be considered complete, new ICD-10-PCS procedure code request submissions through MEARIS™ **must** include:

- A background paper which also indicates if the code request is for consideration for an October 1 or an April 1 implementation date;
- Section 508 Compliant PPT and PDF slide decks for presentation.

Examples of procedure code background papers and slide presentations presented at ICD-10 C&M Committee meetings can be found in agenda and meeting materials of this and previous meetings.

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Home Tasks Applications Teams

Spartans | The International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS)

Contact Info Drug or Technology Info New Code NTAP Info FDA Info **Attachments** Summary

Upload all supporting documentation, presentations, or other reference files (optional)

Uploaded Files
It looks like there is nothing here.

Supported formats include PDF, word, excel, powerpoint, JPEG, PNG, and plain text files

Drag and drop files to upload or [Browse Files](#)

[Back](#) [Next](#)



The time required for request application submission, including the time needed to gather relevant information as well as to complete the form may be extensive depending on the nature of the code request.

Requestors are encouraged to start in advance of the due date to ensure adequate time for submission.

ICD-10-PCS code request application submissions are due **December 2, 2022** to be considered for the March 7-8, 2023 ICD-10 Coordination and Maintenance Committee Meeting.

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Request Information System

Home Tasks Applications Teams

You have successfully submitted a International Classification of Diseases, 10th Revision, Procedure Coding System (ICD-10-PCS) request

Submission Confirmation Code: **ICD220114PV9LU**
Confirmation details have been sent to:
yewande.odukoya@ | mssmith@abc.com | jsmith@abc.com

[Download as PDF](#)

So, what's next?

- An email notification has been sent to the contact(s) you provided on this request.
- The contacts that you have identified will have access to the team and associated applications.
- This request can be viewed under the [applications list](#).
- While reviewing your request, CMS may request additional information or supporting documentation.

[Back To Home](#)

Useful Links

- [CMS Web Policies](#)
- [Technical Support](#)
- [Resources](#)

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