

Medtronic

Surgical heart valves





2024 coding and reimbursement




This information is provided for your consideration. It is the provider's responsibility to determine and submit appropriate codes, modifiers, and charges for the services rendered.

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Surgical valves physician coding (select codes related to Medtronic surgical valves)

Valve product	Composition	Site	CPT ^{®†} code (physician)
Aortic and mitral positions			
Avalus™ Bioprosthesis 	Bioprosthesis (stented tissue valve)	Aortic only	33405 (aortic)
Mosaic™, Mosaic Ultra™ 	Bioprosthesis (stented tissue valve)	Aortic or mitral	33405 (aortic) 33430 (mitral)
Hancock™ II, Hancock II Ultra™ 	Bioprosthesis (stented tissue valve)	Aortic or mitral	33405 (aortic) 33430 (mitral)
Medtronic Open Pivot™ 	Mechanical	Aortic or mitral	33405 (aortic) 33430 (mitral)
Valve product	Composition	Site	CPT code (physician)

Aortic root – for replacement of malfunctioning native or prosthetic aortic valves with the option of aortic root replacement

Freestyle™ Aortic Root Bioprosthesis 	Bioprosthesis (stentless tissue valve)	Aortic valve only (with aortic root)	33410 if coronary arteries are not involved and aortic root is not replaced or 33410-22 if coronary artery reimplantation is performed or 33863 if aortic root replaced, including coronary artery reimplantation in addition to an ascending aorta graft.
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Modifier -22: increased procedural service.

Medtronic CardioVascular Reimbursement Hotline: 1-866-616-8400

[†]CPT[®] is a registered trademark of the American Medical Association.

Surgical valves payment

CPT code	CPT description	2024 Work RVUs ¹	2024 Total Facility RVUs ¹	2024 Medicare national unadjusted amount ¹
33405	Replacement, aortic valve, with cardiopulmonary bypass; with prosthetic valve other than homograft or stentless valve	41.32	66.76	\$2,186
33410	Replacement, aortic valve, with cardiopulmonary bypass; with stentless tissue valve	46.41	74.68	\$2,445
33430	Replacement, mitral valve, with cardiopulmonary bypass	50.93	82.37	\$2,697
33863	Ascending aorta graft, with cardiopulmonary bypass, with or without valve suspension; with aortic root replacement using composite prosthesis and coronary reconstruction	58.79	92.18	\$3,018

¹ CY 2024 payment was calculated with the Conversion Factor (CF) of \$ 32.7442. CMS CY 2024 Medicare Physician Fee Schedule Final Rule. Medicare Fee Service Payment. CMS.gov.: CY 2024 payment was calculated with the Conversion Factor (CF) of \$ 32.7442. CMS CY 2024 Medicare Physician Fee Schedule Final Rule. Medicare Fee Service Payment. CMS.gov. Available at: <https://www.cms.gov/medicare/medicare-fee-service-payment/physicianfeesched/pfs-federal-regulation-notice/cms-1784-f>. Accessed on January 10, 2024.

Reimbursement disclaimer

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This document provides assistance for FDA approved or cleared indications. Where reimbursement is sought for use of a product that may be inconsistent with, or not expressly specified in, the FDA cleared or approved labeling (e.g., instructions for use, operator's manual, or package insert), consult with your billing advisors or payers on handling such billing issues. Some payers may have policies that make it inappropriate to submit claims for such items or related service.

For questions or more information, contact Medtronic Cardiovascular Health Economics, Policy & Payment at Cardiovascular Reimbursement Hotline 1-866-616-8400 or mail to: rs.cardiovasculartheconomics@medtronic.com

Brief Statements **AVALUS™ Bioprosthesis**

Indications:

The AVALUS bioprosthesis is indicated for the replacement of diseased, damaged, or malfunctioning native or prosthetic aortic valves.

Contraindications

None:

Warnings/precautions/adverse events:

Only physicians who have received proper training in valve replacement should use this device. Accelerated structural deterioration due to calcific degeneration of bioprosthesis may occur in: children, adolescents, young adults, and patients with altered calcium metabolism (e.g., chronic renal failure, or hyperparathyroidism). Adverse events can include: angina, cardiac dysrhythmias, endocarditis, heart failure, hemolysis, hemolytic anemia, hemorrhage, infection other than endocarditis, transvalvular or paravalvular leak, myocardial infarction, nonstructural valve dysfunction (leaflet entrapment/impingement, obstructive pannus ingrowth, suture dehiscence, inappropriate sizing or positioning, or other), pericardial effusion or tamponade, prosthesis regurgitation, prosthesis stenosis, prosthesis thrombosis, stroke, structural valve deterioration (calcification, leaflet tear or perforation, or other), thromboembolism, tissue dehiscence, and transient ischemic attack. These complications could lead to reoperation, explant of the bioprosthesis, permanent disability, or death.

Caution: Federal law (USA) restricts this device to sale by or on the order of a physician.

For a listing of indications, contraindications, precautions, warnings, and potential adverse events, please refer to the Instructions for Use. For countries that use eIFUs, consult instructions for use at medtronic.com/manuals (opens new window). Note: Manuals can be viewed using a current version of any major internet browser.

Mosaic™ Bioprosthesis

Indications:

For the replacement of malfunctioning native or prosthetic aortic and/or mitral heart valves.

Contraindications:

None

Warnings/precautions/adverse events:

Accelerated deterioration due to calcific degeneration of bioprosthesis may occur in: children, adolescents, young adults, and patients with altered calcium metabolism (e.g., chronic renal failure, hyperparathyroidism). Adverse events can include: angina, cardiac arrhythmia, cardiac dysrhythmias, death, endocarditis, heart failure, hemolysis, hemolytic anemia, hemorrhage, transvalvular or paravalvular leak, myocardial infarction, nonstructural dysfunction, stroke, structural deterioration, thromboembolism, or valve thrombosis.

Caution: Federal law (USA) restricts these devices to sale by or on the order of a physician.

For a listing of indications, contraindications, precautions, warnings, and potential adverse events, please refer to the Instructions for Use. For countries that use eIFUs, consult instructions for use at this website medtronic.com/manuals (opens new window). Note: Manuals can be viewed using a current version of any major internet browser.

Open Pivot™ mechanical heart valve

Indications:

The Medtronic Open Pivot™ Heart Valve is indicated for the replacement of diseased, damaged, or malfunctioning native or prosthetic aortic or mitral valves.

Contraindications:

The Medtronic Open Pivot Heart Valve is contraindicated in patients unable to tolerate anticoagulation therapy.

Potential adverse events:

Adverse events potentially associated with the use of prosthetic heart valves include: cardiac arrhythmias, death, leaflet entrapment (impingement), endocarditis, hemolysis, anticoagulant-related hemorrhage, transvalvular or paravalvular leak, prosthesis thrombosis, structural deterioration, valve thromboembolism.

Caution: Federal Law restricts this device to sale by or on the order of a physician or properly licensed practitioner. Refer to the Instructions For Use packaged with each valve for a complete listing of warnings and precautions.

Freestyle™ Aortic Root Bioprosthesis

Indications:

For the replacement of malfunctioning native or prosthetic aortic valves with the option of aortic root replacement.

Contraindications:

None known.

Warnings/precautions/adverse events:

Accelerated deterioration due to calcific degeneration of bioprosthesis may occur in: children, adolescents, young adults, and patients with altered calcium metabolism (e.g., chronic renal failure, hyperparathyroidism). Adverse events can include: cardiac dysrhythmias, death, endocarditis, hemolysis, hemorrhage, transvalvular or paravalvular leak, nonstructural dysfunction, structural deterioration, thromboembolism, valve thrombosis, or intracardial hematoma. For additional information, please refer to the Instructions For Use provided with the product.

Caution: Federal law (USA) restricts this device to sale by or on the order of a physician.

Hancock™ II and Hancock II Ultra™ Bioprosthesis

Indications:

For patients who require replacement of their native or prosthetic aortic and/or mitral valves.

Contraindications:

None known.

Warnings/precautions/adverse events:

Accelerated deterioration due to calcific degeneration of bioprosthesis may occur in: children, adolescents, young adults, and patients with altered calcium metabolism (e.g., chronic renal failure, hyperparathyroidism). Adverse events can include: angina, cardiac arrhythmia, cardiac dysrhythmias, death, endocarditis, heart failure, hemolysis, hemolytic anemia, hemorrhage, transvalvular or paravalvular leak, myocardial infarction, nonstructural dysfunction, stroke, structural deterioration, thromboembolism, or valve thrombosis. For additional information, please refer to the Instructions for Use provided with the product.

Caution: Federal law (USA) restricts this device to sale by or on the order of a physician.

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