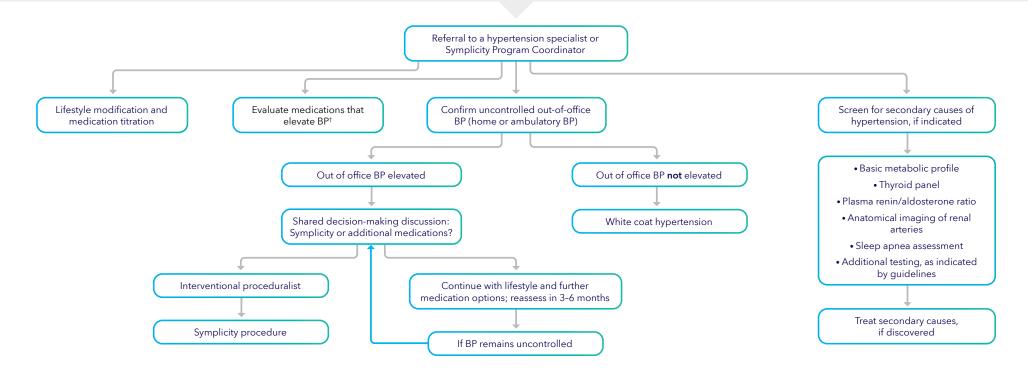
Medtronic

Symplicity™ blood pressure procedure

Patient screening flowchart¹⁻³



Consider the following for patients with hypertension (HTN):





Physician discretion should be used to make a final decision on the screening process and intervention. Refer back to AHA/ACC guidelines for additional information.

[†]Amphetamines, antidepressants, atypical antipsychotics, decongestants, herbal supplements, immunosuppressants, oral contraceptives, NSAIDs, recreational drugs, systemic corticosteroids, angiogenesis inhibitor (list is not exhaustive).

[†]AHA/ACC Hypertension Care Guidelines. 2017.

²Roger KL, et al. *Endovascular Today*. 2023;22:73-76.

³Kandzari DE, et al. Circ Cardiovasc Qual Outcomes. 2023;16:e008997.

Indications

The Symplicity Spyral[™] renal denervation system is indicated to reduce blood pressure as an adjunctive treatment in patients with hypertension in whom lifestyle modifications and antihypertensive medications do not adequately control blood pressure.

Contraindications

The Symplicity Spyral system is contraindicated in patients with any of the following conditions: \bullet Renal artery diameter < 3 mm or > 8 mm \bullet Renal artery fibromuscular dysplasia (FMD) \bullet Stented renal artery (< 3 months prior to RDN procedure) \bullet Renal artery aneurysm \bullet Renal artery diameter stenosis > 50% \bullet Pregnancy \bullet Presence of abnormal kidney (or secreting adrenal) tumor \bullet Iliac/femoral artery stenosis precluding insertion of the catheter.

Warnings and Precautions

A thorough understanding of the technical principles, clinical applications, and risks associated with vascular access techniques and percutaneous transluminal catheterization in renal arteries is necessary before using this device.

The safety and efficacy of the Symplicity Spyral system has not been established in patients with isolated systolic hypertension or in patients with prior renal artery interventions including renal stents, renal angioplasty, or prior renal denervation. The Symplicity Spyral system has not yet been studied in patients who are breastfeeding, under the age of 18, or with secondary hypertension. • Avoid treatment with the Symplicity Spyral at theter within 5 mm of any diseased area or stent. • Implantable pacemakers (IPGs) and implantable cardioverter defibrillators (ICDs) or other active implants may be adversely affected by RF ablation. Refer to the implantable device's Instructions for Use. • The patient's heart rate may drop during the ablation procedure. • Proper pain medication should be administered at least 10 min before ablating renal nerves.

Potential Adverse Events

Potential adverse events associated with use of the renal denervation device or the interventional procedures include, but are not limited to, the following conditions: • Allergic reaction to contrast • Arterial damage, including injury from energy application, dissection, or perforation • Arterial spasm or stenosis • Arterio-enteric fistula • AV fistula • Bleeding • Blood clots or embolism • Bruising • Cardiopulmonary arrest • Complications associated with medications commonly utilized during the procedure, such as narcotics, anxiolytics, or other pain or anti-vasospasm medications • Death • Deep vein thrombosis • Edema • Electrolyte imbalance • Heart rhythm disturbances, including bradycardia • Hematoma

 Hematoma – retroperitoneal • Hematuria • Hypertension • Hypotension (may cause end organ hypoperfusion) • Infection • Kidney damage including renal failure or perforation • Myocardial infarction • Nausea or vomiting • Pain or discomfort • Peripheral ischemia

 \bullet Pulmonary embolism \bullet Proteinuria \bullet Pseudoaneurysm \bullet Radiocontrast nephropathy \bullet Renal artery aneurysm \bullet Skin burns from failure of the dispersive electrode pad \bullet Stroke \bullet Other potential adverse events that are unforeseen at this time.

Please reference appropriate product *Instructions for Use* and *User Manual* for more information regarding indications, contraindications, warnings, precautions, and potential adverse events.

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

For further information, please call and/or consult Medtronic at 800-633-8766 or the Medtronic website at medtronic.com.

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medtronic.com/SymplicityProcedure