Penumbra System Reperfusion Catheters

Select Device Based on Vessel Size



Prior to use, please refer to the Instructions for Use for complete product indications, contraindications, warnings, precautions, potential adverse events, and detailed instructions for use.

PENUMBRA SYSTEM — Intended Use

The PENUMBRA SYSTEM is intended for use in the revascularization of patients with acute ischemic stroke secondary to intracranial large vessel occlusive disease using continuous aspiration.

Potential Adverse Events

Possible complications include, but are not limited to, the following: allergic reaction and anaphylaxis from contrast media; acute occlusion; air embolism; arteriovenous fistula; death; device malfunction; distal embolization; emboli; false aneurysm formation; hematoma or hemorrhage at access site; inability to completely remove thrombus; infection; intracranial hemorrhage; ischemia; kidney damage from contrast media; neurological deficits including stroke; vessel spasm, thrombosis, dissection, or perforation; radiation exposure that may lead to cataracts, skin reddening or burns from x-ray exposure.

Product availability varies by country. Please contact your local Penumbra representative for more information.

Copyright ©2020 Penumbra, Inc. All rights reserved. The Penumbra P logo, Penumbra System, MAX, ACE, and Penumbra JET are registered trademarks or trademarks of Penumbra, Inc. in the USA and other countries. 17628, Rev. A 03/20 EU

