

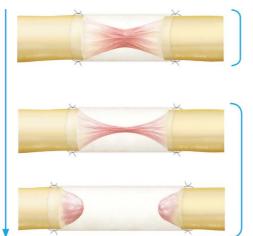
Peripheral Nerve Bridging Materials: Using Clinical Evidence to Optimize Outcomes



conduits were designed as a coaptation aid, not as a bridging material

Conduits are hollow tubes, relying on the body to form a fibrin cable.

Fibrin cable thins at longer gaps

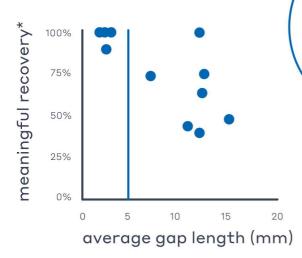


Originally designed to alleviate tension and reduce misalignment in short gaps less than 5 mm

What is the reliability with longer gaps?

conduits have a reliability threshold

Each datapoint represents one clinical publication in sensory nerve gaps.¹⁻¹⁰



*Defined as S3 or higher.

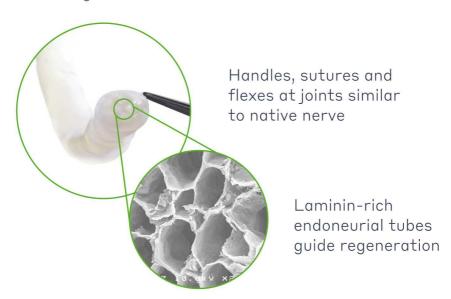
In gaps above 5 mm, more than 1 in 3 conduits fail to achieve meaningful recovery.¹⁻¹⁰

failures may result in:

- Lack of functional recovery
- Absence of protective sensation
- Revision procedure
- Chronic pain

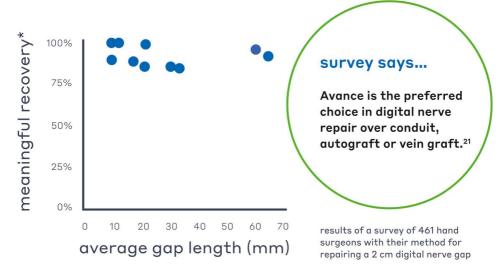
avance® nerve graft was developed to bridge nerve gaps

It is a biologically active processed human nerve allograft that eliminates the comorbidities and operative time associated with a second surgical site.



avance has demonstrated consistency and reliability

Each datapoint represents one clinical publication in sensory nerve gaps. 11-20



^{*}Defined as S3 or higher.

summary of clinical outcomes

Evidence based algorithm avoids potential complications of autograft harvest, without sacrificing patient outcomes.

% meaningful recovery st



^{*}Defined as S3 or higher. Data points derived from a weighted average of publications reporting meaningful recovery in sensory nerve gaps.^{1-20,22-24}



references

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axogen's portfolio of products

options for no transection



options for 0 mm to 5 mm



options for 5 mm to 70 mm



options for 70 mm+



this is the new normal

At least 66 portfolio peer-reviewed publications

Over 2/3 of US hand and microsurgery fellows were trained on this algorithm in 2018²⁵

Indications and Trademark Disclaimers

Avance Nerve Graft

REGULATORY CLASSIFICATION: Avance Nerve Graft is processed and distributed in accordance with US FDA requirements for Human Cellular and Tissue-based Products (HCT/P) under 21 CFR Part 1271 regulations, US State regulations and the guidelines of the American Association of Tissue Banks (AATB). Additionally, international regulations are followed as appropriate. Avance Nerve Graft is to be dispensed only by or on the order of a licensed physician.

INDICATIONS FOR USE: Avance Nerve Graft is processed nerve allograft (human) intended for the surgical repair of peripheral nerve discontinuities to support regeneration across the defect.

CONTRAINDICATIONS: Avance Nerve Graft is contraindicated for use in any patient in whom soft tissue implants are contraindicated. This includes any pathology that would limit the blood supply and compromise healing or evidence of a current infection.

Axoguard Nerve Connector

INDICATIONS FOR USE: The Axoguard Nerve Connector is indicated for the repair of peripheral nerve discontinuities with gaps up to 5 mm. The Axoguard Nerve Connector is supplied sterile and is intended for single use.

CONTRAINDICATIONS: The Axoguard Nerve Connector is derived from a porcine source and should not be used for patients with known sensitivity to porcine material.

Axoguard Nerve Protector

INDICATIONS FOR USE: The Axoguard Nerve Protector is indicated for the repair of peripheral nerve injuries where there is no gap. The Axoguard Nerve Protector is supplied sterile and is intended for one-time use.

CONTRAINDICATIONS: The Axoguard Nerve Protector is derived from a porcine source and should not be used for patients with known sensitivity to porcine material.

Avive Soft Tissue Membrane

REGULATORY CLASSIFICATION: Avive Soft Tissue Membrane is processed and distributed in accordance with US FDA requirements for Human Cellular and Tissue-based Products (HCT/P) under 21 CFR Part 1271 regulations, US State regulations and the guidelines of the American Association of Tissue Banks (AATB). Additionally, international regulations are followed as appropriate. Avive Soft Tissue Membrane is to be dispensed only by or on the order of a licensed physician.

INDICATIONS FOR USE: Avive Soft Tissue Membrane is processed umbilical cord intended for homologous use as a soft tissue covering.

CONTRAINDICATIONS: Avive Soft Tissue Membrane is contraindicated for use in any patient in whom soft tissue implants are contraindicated.

Disclaimer: Not all products are available internationally.

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