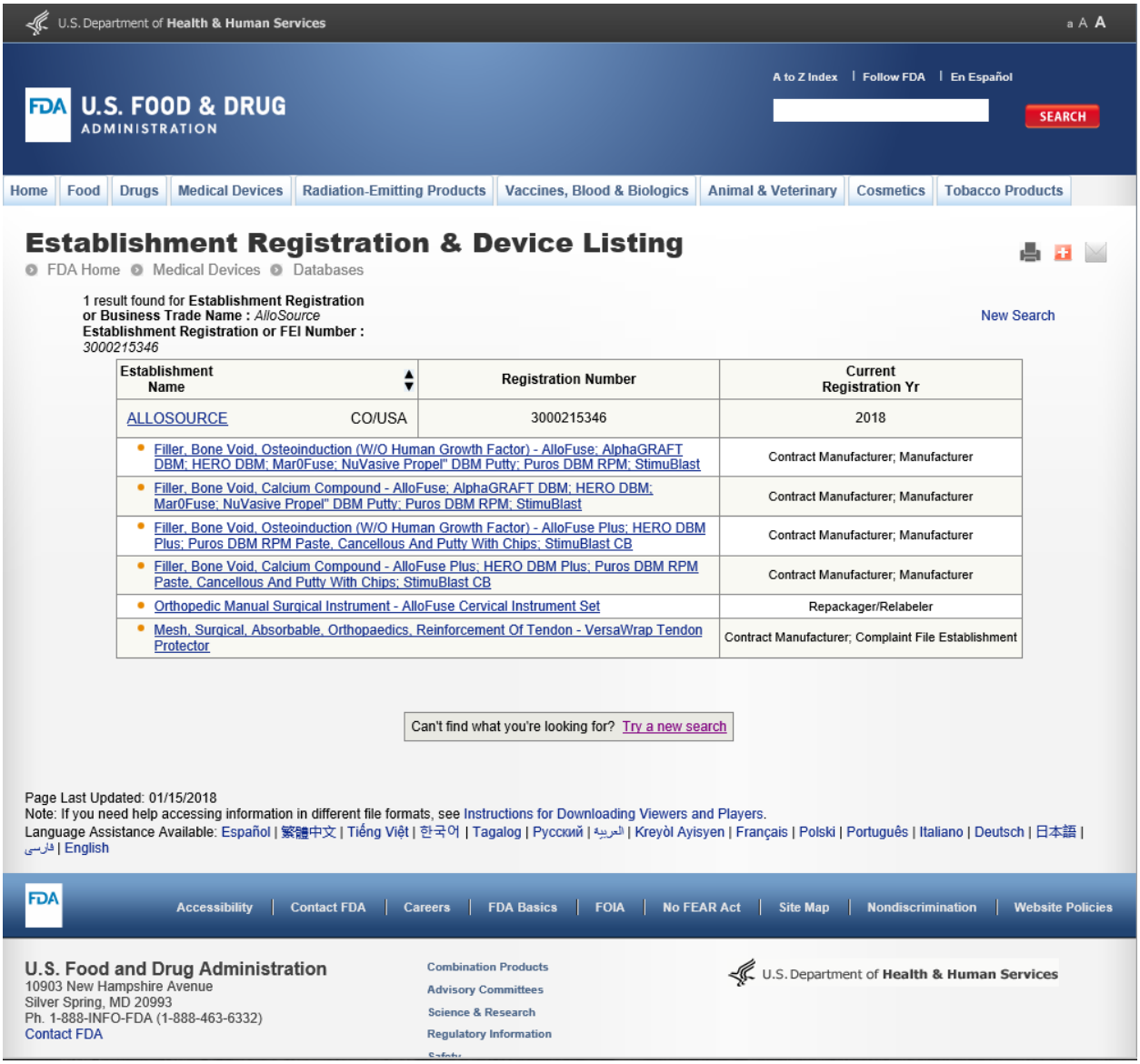
FDA 2018 Establishment & Device Listing for AlloSource Medical Devices

Medical Device Registrations are issued and maintained by the FDA Center for Devices and Radiological Health (CDRH). The following Establishment & Device Listing screenshot was taken on January 15, 2018, and reflects the annual establishment registration & device listing has been complete for fiscal year 2018.

Up-to-date registrations can be viewed at: http://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfRL/rl.cfm by entering 'AlloSource' in the *Establishment Name* field, then clicking *Search*.



U.S. Department of Health & Human Services

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Establishment Registration & Device Listing

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1 result found for **Establishment Registration or Business Trade Name :** *AlloSource*
Establishment Registration or FEI Number : *3000215346*

New Search

Establishment Name		Registration Number	Current Registration Yr
ALLOSOURCE	CO/USA	3000215346	2018
• Filler, Bone Void, Osteoinduction (W/O Human Growth Factor) - AlloFuse; AlphaGRAFT DBM; HERO DBM; Mar0Fuse; NuVasive Propel" DBM Putty; Puros DBM RPM; StimuBlast			Contract Manufacturer; Manufacturer
• Filler, Bone Void, Calcium Compound - AlloFuse; AlphaGRAFT DBM; HERO DBM; Mar0Fuse; NuVasive Propel" DBM Putty; Puros DBM RPM; StimuBlast			Contract Manufacturer; Manufacturer
• Filler, Bone Void, Osteoinduction (W/O Human Growth Factor) - AlloFuse Plus; HERO DBM Plus; Puros DBM RPM Paste, Cancellous And Putty With Chips; StimuBlast CB			Contract Manufacturer; Manufacturer
• Filler, Bone Void, Calcium Compound - AlloFuse Plus; HERO DBM Plus; Puros DBM RPM Paste, Cancellous And Putty With Chips; StimuBlast CB			Contract Manufacturer; Manufacturer
• Orthopedic Manual Surgical Instrument - AlloFuse Cervical Instrument Set			Repackager/Relabeler
• Mesh, Surgical, Absorbable, Orthopaedics, Reinforcement Of Tendon - VersaWrap Tendon Protector			Contract Manufacturer; Complaint File Establishment

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