DME Form: Letter of Medical Necessity

Patient Name:___

SSN:___

Diagnosis Code:

□ Adult acquired flatfoot – 734	□ Tendon rupture; ankle & foot – 727.68
□ Congenital flatfoot – 754.61	□ Chronic tibialis posterior tendonitis – 726.72
□ Pronation, acquired – 736.79	\Box Osteoarthrosis; ankle & foot – 715.17
□ Joint pain; ankle & foot – 719.47	\Box Traumatic arthropathy; ankle & foot – 716.17
\Box Tarsal coalition – 755.67	□ Instability of joint; ankle & foot – 718.87
□ Dropfoot – 736.79	□ Hemiplegia – 438.20
□ Calcaneofibular ligament sprain – 845.02	□ Deltoid ligament sprain – 845.01
\Box Charcôt's arthropathy – 713.5	□ Chronic Achilles tendonitis – 726.71
□ Other:	

Description of Orthosis:

The following Ankle-Foot Orthosis (AFO) & Component Parts have been prescribed for the above patient:

- $\hfill\square$ L1970: AFO, plastic, with ankle joint, custom fabricated
- L2210: Addition to lower extremity, dorsiflexion assist (plantar flexion resist), each joint
- L2270: Addition to lower extremity, varus/valgus correction strap, padded/lined or malleolus pad
- L2820: Addition to lower extremity, soft tissue interface for molded plastic, below knee section
- □ L1940: AFO, plastic or other material, custom fabricated
- L2275: Addition to lower extremity, varus/valgus correction, plastic modification, padded/lined
- □ L2280: Addition to lower extremity, molded inner boot
- □ L1960: AFO, posterior solid ankle, plastic, custom fabricated
- L2330: Addition to lower extremity, lacer molded to patient model, for custom fabricated orthosis only
- □ L2340: Addition to lower extremity, pretibial shell, molded to patient model

□ L3000: Foot insert, removable, molded to patient model. "UCBL" Type, Berkeley Shell, each. Plastic device, molded over model of patient's foot to provide control of the foot.

 \Box L3020: Foot insert, molded to patient model, longitudinal and metatarsal support, each. A device molded over a model of the patient's foot and placed in the shoe to provide support under the ball of the foot.

□ L1971: AFO, plastic or other material with ankle joint, prefabricated, includes fitting and adjustment □ Other:

Duration of Treatment/Medical Necessity:

The patient designated above qualifies for and will benefit from an ankle-foot orthosis based on the following criteria:

 \Box The condition necessitating the orthosis is expected to be permanent or of longstanding duration (more than 6 months)

 $\hfill\square$ The patient has had an orthopedic injury that requires bracing

 $\hfill\square$ There is a need to control the ankle or foot in more than one plane

 \Box The patient has a documented neurological, circulatory, or orthopedic condition that requires custom fabrication over a model of the patient's extremity to prevent tissue injury

□ The patient has a healing fracture which lacks normal anatomical integrity or anthropometric proportions

I hereby certify that the ankle-foot orthosis described above is a rigid or semi-rigid device which is used for the purpose of improving mobility, improving lower extremity stability, decreasing pain, and/ or facilitating soft tissue healing. The prognosis for this patient is excellent with the use of the above devices. A custom or pre-fabricated ankle-foot orthosis has been prescribed, based on the above diagnosis codes, in order to improve the patient's condition and ability to ambulate.