



(Non)/Therapeutic Foot/Insert Eval/Fit/Delivery Form

Name: _____ Date: _____ DOB: _____ Age: _____ Ht: _____ Wt: _____ Sex: ☐ M ☐ F

Date of evaluation: _____ Diagnosis: _____

Services referred for: _____

Describe condition/injury, including onset, surgeries, allergies, deformities, etc.: _____

Describe existing shoe/insert _____

New Shoes: ☐ N/A Manufacturer: _____ Style/Part #: _____ Size: _____

Style: ☐ Oxford ☐ Extra depth ☐ Other: _____ ☐ Prefabricated ☐ Custom fabricated

Mold taken: ☐ No ☐ Yes/method: ☐ Digital Scan ☐ Foam ☐ Plaster ☐ Other: _____

Modifications required: ☐ Velcro closures ☐ Steel Shank ☐ Rocker heel/sole ☐ Other: _____

Clinical rationale for shoe design: _____

New Inserts: ☐ N/A Manufacturer: _____ Style/Part #: _____ Size: _____

☐ Prefabricated Self molding ☐ Prefabricated heat molded ☐ Custom Fabricated

Mold taken: ☐ No ☐ Yes/method: ☐ Digital Scan ☐ Foam ☐ Plaster ☐ Other: _____

Describe materials/design to be utilized: _____

Clinical rationale for design: _____

Functional Goals for patient: (check all that are applicable)

☐ Protection of foot ☐ Reduction in pain ☐ Accommodation to deformity ☐ Correction of deformity

☐ Facilitate healing of injury ☐ Post surgical support/correction ☐ Other: _____

Shoe/Inserts to be ordered: ☐ N/A Estimated date of delivery: _____

X _____

(Only sign above if delivery is other than today's date. At time of final delivery, complete sections below and re-sign w/date of delivery as indicated at bottom)

Patient tolerated procedures without problem: ☐ Yes ☐ No (explain): _____

Functional Goals Met: ☐ Yes ☐ No (explain): _____

Device checked for structural safety/integrity and compliance with manufacturer guidelines: ☐ Yes ☐ No

The patient states satisfaction with the fit and function of shoes/inserts: ☐ Yes ☐ No (explain): _____

List additional supplies provided to patient: ☐ N/A _____

Written and or oral instructions provided to: ☐ Patient ☐ Parent ☐ Caregiver

☐ Donning/doffing of device ☐ Skin Inspection ☐ Care and cleaning ☐ Usage/break-in period ☐ Fitting issues

☐ Usage to ensure safety ☐ Patient is an experienced wearer

☐ How/whom to report problems related to device/change of physical condition

Patient tolerated the procedure without incident/problem: ☐ Yes ☐ No (explain): _____

Follow up scheduled: ☐ 1 week ☐ 1-3 weeks ☐ 1 month ☐ 3 months ☐ 6 months ☐ 1 year ☐ PRN

Practitioner Signature: _____ Date: _____