

Site Number:
Date of Visit:
Person Completing Form:

Participant ID:
Participant Letters:

A. STUDY DRUG ADMINISTRATION

1. Was subcutaneous injection given? Y N
a. If NO, specify why:

2. Did the subject experience any problems following the drug administration? Y N

Site evaluation	1) Time Post Injection	2) Duration	3) Grade*
a. Redness	_____ min	_____ min	<input type="radio"/> 1 <input type="radio"/> 2 <input type="radio"/> 3 <input type="radio"/> 4
b. Swelling	_____ min	_____ min	<input type="radio"/> 1 <input type="radio"/> 2 <input type="radio"/> 3 <input type="radio"/> 4
c. Itching	_____ min	_____ min	<input type="radio"/> 1 <input type="radio"/> 2 <input type="radio"/> 3 <input type="radio"/> 4
d. Pain	_____ min	_____ min	<input type="radio"/> 1 <input type="radio"/> 2 <input type="radio"/> 3 <input type="radio"/> 4

3. Did the subject experience any other problems during study drug administration? Y N N/A
a. If YES, Specify

If any problems were encountered, complete an Adverse Event Report Form if \geq Grade 2 severity.
If the Adverse Event is Grade 1 record on source document.