



2 week Complications form - single

Participant Trial Number:

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Hospital site code:

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Please ask the participant the following questions about their trial tendon:

1. Have you/Are you currently experiencing any of the following at the injection site?

For office use Yes No

1	<input type="checkbox"/>	<input type="checkbox"/>
2	<input type="checkbox"/>	<input type="checkbox"/>
3	<input type="checkbox"/>	<input type="checkbox"/>
4	<input type="checkbox"/>	<input type="checkbox"/>
5	<input type="checkbox"/>	<input type="checkbox"/>
6	<input type="checkbox"/>	<input type="checkbox"/>
7	<input type="checkbox"/>	<input type="checkbox"/>
8	<input type="checkbox"/>	<input type="checkbox"/>

bruising at the injection site

fainting episode associated with tendon injection treatment

an infection diagnosed by a doctor

mild discomfort associated with tendon injection treatment

bleeding associated with tendon injection treatment

swelling

skin discolouration

allergic reaction

Other *please specify*

2. If any of those listed above have been ticked as **Yes**, have you received any treatment?

☐ Yes ☐ No

2.a. If **Yes**, what treatment

3. Had this/ Have these (that you ticked **Yes** to in Q1) now resolved?

☐ Yes ☐ No

3.a. If **No**, please detail

4. Are there any other details:

5. Please rate your pain on a scale of 0-100.....

number

0 being "No pain" and 100 being "Worst pain imaginable"

6. Have any contact details changed since the last appointment? ☐ Yes ☐ No

If **Yes**, please complete the **Change of Contact details form**

FOR OFFICE USE ONLY

Do any of the events detailed above meet the definition of an SAE?

☐ Yes ☐ No

If **Yes**, please notify the Trial Coordinator to initiate the completion of an SAE form.

Date Chief Investigator made aware of AEs/ SAEs :

d	d	-	m	m	m	-	y	y	y	y
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FORM COMPLETED BY (PRINT NAME):

SIGNATURE: DATE SIGNED:

d	d	-	m	m	m	-	y	y	y	y
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