



STAGE 2 AND 3 REPORTS TRACKING

ID NUMBER:

FORM CODE:

S	R	T
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DATE: 11/08/2011
Version 1.0

ADMINISTRATIVE INFORMATION

0a. Completion Date: / /
Month Day Year

0b. Staff ID:

The purpose of this form is to document when the Retinal Results report and/or the MRI Results report are sent to the participant and whether an atypical cognitive results letter and/or B12 and TSH results report is sent to the participant.

ATYPICAL LETTER

1. Was an atypical cognitive results letter sent to the participant?

Yes

No → **GO TO QUESTION 3**

2. Date the letter was sent: / /
Month Day Year

B12 AND TSH RESULTS REPORT SENT TO PARTICIPANT

3. Was the Vitamin B12 and TSH results report sent to the participant?

Yes

No → **GO TO QUESTION 5**

4. Date the letter was sent: / /
Month Day Year

RETINAL RESULTS REPORT SENT TO PARTICIPANT

5. Was a copy of the Retinal Results report sent to the participant?

Yes

No → **GO TO QUESTION 7**

6. Date the report was sent: / /
Month Day Year

MRI RESULTS REPORT SENT TO PARTICIPANT

7. Was a copy of the MRI Results report sent to the participant?

Yes

No → **END OF FORM**

8. Date the report was sent: / /
Month Day Year



INSTRUCTIONS FOR THE STAGE 2 AND 3 REPORTS TRACKING (SRT) FORM

I. General Instructions

The Stage 2 and 3 Reports Tracking Form (SRT) is used to document when the Retinal Results report, the MRI Results report, an “atypical” neurocognitive results letter, or the Vitamin B12/Thyroid Stimulating Hormone report is sent to the participant. It is important that the relevant sections of the SRT are completed when results are sent to the participant so that the DCC is aware that reports have been sent to the participant.

II. Detailed Instructions for Each Item

- 1 – 2. Record whether an “atypical” neurocognitive results letter was sent or given to the participant and, if so, when the letter was sent.
- 3 – 4. Record whether the B12/TSH results report was sent to the participant (together with or separately from the “atypical” letter, Retinal Results report, and/or MRI Results report) and, if so, when the report was sent.
- 5 – 6. Record whether the Retinal Results report was sent to the participant and, if so, when it was sent.
- 7 – 8. Record whether the MRI Results report was sent to the participant and, if so, when the report was sent.