**Clinical Study Reporting Diagram**(modified from the CONSORT 2010 Flow Diagram)

Please complete the diagram below with the most recent data available. Fill in the applicable number of intervention or cohort arms. If your study has more than 5 arms, please use a second copy of this form.

Number of trial sites:

Target # subjects enrolled:

Target # subjects complete (accounting for dropout):

Study duration:

Pre-screened (n= ) as of *[Current Date]*

If applicable, list number of:

* Phone inquiries (related to advertising)
* Referrals
* Identified from databases
* Other

Screened in clinic (n= ) as of *[Current Date]*

Excluded (n = )

List reasons for screen failures, (i.e. negative amyloid PET):

* Criteria 1 (n = )
* Criteria 2 (n = )
* Criteria 3 (n = )
* Criteria 4 (n = )

Enrolled (n= ) as of *[Current Date]*

*Intervention/cohort*:

Allocated to

Arm 5 (n = )

*Intervention/cohort*:

Allocated to

Arm 4 (n = )

*Intervention/cohort*:

Allocated to

Arm 3 (n = )

*Intervention/cohort*:

Allocated to

Arm 2 (n = )

*Intervention/cohort*:

Allocated to

Arm 1 (n = )

Completed

(n = )

Completed

(n = )

Completed

(n = )

Completed

(n = )

Completed

(n = )

Lost to follow-up

(n = )

Lost to follow-up

(n = )

Lost to follow-up

(n = )

Lost to follow-up

(n = )

Lost to follow-up

(n = )