Cohort – Seniors Quid

Title of the cohort
Cohort – Seniors Quid

Acronym for cohort
PAQUID

Name of Principal Investigator
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First name
Last name DARTIGUES

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Funding source
CNSA BEAUFOR IPSEN, NOVARTIS PHARMA

1. The cohort includes, or expects to include, incidence of the following conditions

• Alzheimer’s disease and other dementias
• Parkinson’s disease

When studies on the above condition(s) are expected to become possible

• Already possible

2a. Stated aim of the cohort

Studying brain and functional aging after 65 years, distinguish the normal and pathological conditions,
and identify people at high risk of physical or mental deterioration in which preventive action could be possible

2b. Features distinguishing this cohort from other population cohorts

Representative cohort of the French population aged 65 and over

3a. i) Number of publications that involve use of cohort to date

151

3a. ii) Up to three examples of studies to date (PI, Institution, Title of Study)

Risk factor for dementia and Alzheimer’s disease (social support, reactivation of herpes viruses, leisure)
Natural history of Alzheimer’s disease
Risk factor for Parkinson’s disease

3b. Publication list/link to where data or publications are accessible (if available)


Amieva, H., R. Stoykova, et al. (2010). “What aspects of social network are protective for dementia? Not the quantity but the quality of social interactions is protective up to 15 years later.” Psychosom Med 72(9): 905-911.


3c. Information (i.e. research findings) expected to be gained from the population cohort
4a. Study criteria: age range of participants at recruitment
   Age in years from: 65
   To (‘until death’ if applicable): until death

4b. Study criteria: inclusion criteria
   Persons aged 65 and over living at home and enrolled as voters in the Dordogne or Gironde

4c. Study criteria: exclusion criteria
   People not living at home

5. Size of the cohort (i.e. number of participants enrolled)
   - 1,000 – 5,000 participants

6a. Measures used to characterise participants
   age, sex, level of education, city of residence

6b. Additional measures for participants with a clinical disorder
6c. Are there defined primary and secondary endpoints (e.g. defined health parameters)
   Dementia and dependency, mortality

7. Study design
   - Prospective cohort

8. Cases matched by
   - Age
   - Sex
9a. Does the study include a specialised subset of control participants

- No

9b. If yes, description of specialised subset of control participants

10a. i) Data collection start date

01-01-1988

10a. ii) Data collection end date

01-01-2018

10a iii) Data collection for this study is

- Data collection ongoing
- Data analysis ongoing
- Closed to new patients

10b. Plans to continue the cohort study beyond the current projected end date

- No

11. Data collected

- Only through the study

12. System in place to enable re-contact with patients for future studies

- No

13a. Format and availability of data stored in a database

<table>
<thead>
<tr>
<th>Format</th>
<th>Available</th>
</tr>
</thead>
<tbody>
<tr>
<td>Data summarised in database</td>
<td>no</td>
</tr>
<tr>
<td>Database is web-based</td>
<td>no</td>
</tr>
<tr>
<td>Database on spreadsheet</td>
<td>no</td>
</tr>
<tr>
<td>Database is on paper</td>
<td>no</td>
</tr>
<tr>
<td>Other (specify)</td>
<td>no</td>
</tr>
</tbody>
</table>

13b. Format and availability of data held as individual records

Language used:

14a. Are data available to other groups
14b. Access policy/mechanisms for access if data are available to other groups
15. Data sharing policy specified as a condition of use
   - No policy exists

16a. Are tissues/samples/DNA available to other groups
No

16b. i) Description of available tissues/samples/DNA
16b. ii) Form available tissues/samples/DNA are supplied in
16b. iii) Is the access policy/mechanism for obtaining samples the same as that for obtaining data

17. Is information on biological characteristics available to other groups
   - No