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Secondary Prevention of Alzheimer's Disease: the EPAD and AMYPAD studies

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Background

Alzheimer's disease (AD) is the main cause of dementia and affects more than 40 million people worldwide [1]. In order to develop a treatment to prevent or delay the onset of AD, the identification of subjects at a preclinical or prodromal stage is required [2]. Beyond studies in familial/genetic populations, opportunities exist to identify subjects at risk for sporadic AD based on APOE- ϵ 4, CSF biomarkers, imaging and other risk factors. Secondary prevention is the focus of two intimately related IMI-funded projects, EPAD and AMYPAD.



of β-amyloid imaging Quantifying patient-specific efficacy + Proof of Concept studies

European Prevention of AD (EPAD) study

6000 subject enrolled (existing natural history cohorts + studies of subjects with subjective memory complaints or mild cognitive impairment);
Aims to obtain ~80% positivity for amyloid (CSF or PET);

 The cohort will be followed longitudinally and constitute a trial-readiness cohort for early proof-ofconcept (PoC) studies targeting preclinical and prodromal AD. Amyloid Imaging to Prevent AD (AMYPAD) study
Aims to identify how ß-amyloid PET can be used, either alone or in combination with other assessments, for

- Early diagnosis of AD;
- Prognostic stratification of subjects enrolled in clinical trials;
- Therapy monitoring.
- Tracers:

• www.ep-ad.org

- 18F-Florbetaben (Piramal Imaging);
- 18F-Flutemetamol (GE Healthcare).

www.amypad.eu

Amyloid PET [3] Control AD

EPAD: European Prevention of	AMYPAD: A
Alzheimer's Disease	Alzhei
IMI 1 Project (start 01/2015)	IMI 2 Proj

/YPAD: Amyloid Prevention of Alzheimer's Disease IMI 2 Project (start 10/2016

*LCS: Longitudinal Cohort Study (EPAD) ; **PoC: Proof of Concept study

References

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[3] Tolboom N. *et al.* Detection of Alzheimer Pathology in vivo using both 11C-PIB and 18F-FDDNP-PET. *The Journal of Medicine* 2009; 50(2):191-197

