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1 **Diagnosis of Alzheimer’s disease patients with Rapid Cognitive**  
2 **Decline in clinical practice: interest of the Deco questionnaire.**

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1 **Abstract**

2

3 *Background:* Patients with Alzheimer’s disease (AD) who deteriorate rapidly are likely to  
4 have a poorer prognosis. There is a clear need for a clinical assessment tool to detect such a  
5 decline in newly diagnosed patients.

6 *Objective:* To identify the predictive factors of rapid cognitive decline (RCD) in a cohort of  
7 patients with mild to moderate AD ; and to validate a self-questionnaire for caregivers as a  
8 diagnostic tool for rapid decline.

9 *Design and analysis:* an open-label, observational, 12-month, multicenter, French study.

10 Physicians were asked to record data of three eligible rivastigmine naïve (or on rivastigmine  
11 for < 1 year) AD patients. Risk factors of RCD and the detection power of the Détérioration  
12 Cognitive Observée scale (Deco), a 19 item self-questionnaire for caregivers, were assessed at  
13 endpoint using regression analyses.

14 *Results:* Out of the 361 patients enrolled in the study, 91 (25.2%) were excluded due to loss of  
15 follow-up. Among subjects using cholinesterase inhibitors or memantine, 161 (59.6%)  
16 experienced a stabilization (29.2%) or an improvement (30.4%) in global functioning as  
17 measured by the CGI-C. Sixty of the remaining 204 patients retained for analysis (29.6%, CI  
18 95% [23.4; 35.8]) lost three or more points on the MMSE score between the inclusion and one  
19 of the follow-up visit. In the multivariate logistic regression analysis, institutionalization,  
20 higher level of education and the loss of 3 points or more on the MMSE were found to be  
21 significant predictors of a rapid cognitive loss in this population. The threshold which  
22 maximizes the predictive values of the Deco score as a diagnostic tool of rapid cognitive  
23 decline was significantly different according to the age of the patient (below or over 75 years  
24 old). A score below 16 for patients < 75 years old and below 14 for patients  $\geq$  75 years old  
25 consistently predicted a RCD within the next year.

26 *Conclusion:* The Deco test appears to be a simple tool to alert the physician to the possibility  
27 of an aggressive course of the disease which warrants particular management.

28

29 **Key words: Alzheimer’s disease, rapid cognitive decline, diagnostic tool**

1 **INTRODUCTION**

2 Neurodegenerative dementias such as Alzheimer Disease (AD) are characterized by a  
3 progressive cognitive decline. However, the rate of progression can be highly variable from  
4 one patient to another [1] and some patients could have a more rapid cognitive decline (RCD).  
5 These patients are likely to have a poorer prognosis, with a higher mortality rate or a lower  
6 survival rate without severe dementia, than those with slower progression rates [1-3].

7 A prospective observational study performed in a population of 455 AD patients  
8 followed for at least one year, demonstrated that the risk of deterioration was significantly  
9 decreased in patients taking cholinesterase inhibitors (ChEIs) for at least one year compared  
10 to untreated patients [4]. To date, rivastigmine is the only drug which has been specifically  
11 studied in patients with RCD. A meta-analysis of four 6-month, double-blind, placebo-  
12 controlled phase studies [5] showed that patients who experienced rapid decline, i.e. a decline  
13 of at least 4 points of the ADAS-cog score, during 26 weeks of placebo treatment, improved  
14 significantly more after 12 weeks of rivastigmine therapy than those who had declined by less  
15 than 4 points in the first 26 weeks. Thus, identifying AD patients at risk of RCD is important  
16 as these patients might benefit from specific primary or secondary care including more  
17 frequent clinical and neuropsychological evaluations, more frequent adaptation of drug  
18 treatment and more effective support for the caregiver.

19 Several factors may predict RCD in AD patients: cognitive status, presence of Extra-  
20 Pyramidal Signs, psychotic symptoms and hallucinations, prominent sub-cortical pathology  
21 with attention and executive deficits and poor nutrition [6; 7]. However many of these factors  
22 are of limited interest in a clinical setting for individual patients as it is difficult to obtain  
23 valid information about most of them during the time of a consultation. There is thus a clear  
24 need for a simple clinical assessment tool, drawing on caregiver input, to detect RCD in  
25 recently diagnosed patients. The “Détérioration Cognitive Observée” (Deco) scale is a 19-item  
26 questionnaire administered to a person with at least a monthly contact with the patient. Its aim  
27 is to evaluate the change in cognitive functioning of the patient over the previous year. This  
28 instrument has been shown to be highly sensitive to early changes in cognitive functioning in  
29 previous studies [8].

30 The objectives of the prospective study we report here were to identify the predictive factors  
31 of RCD in a cohort of patients with mild to moderate AD ; and to analyze the predictive value  
32 of a self-questionnaire for caregivers (the Deco score) as a simple diagnostic tool for RCD.

33  
34

## 1 PATIENTS AND METHODS

### 2 *Study Design*

3 EXPLORE was an open label, observational cohort study, 12-month, multicenter,  
4 French study for which 350 neurologists, geriatricians and psychiatrists were contacted in  
5 order to finally include 250 of them. Each was asked to record data of three eligible patients.  
6 Eligibility criteria included: outpatients with AD over fifty years old, satisfying the criteria of  
7 the fourth edition of the Diagnostic and Statistical Manual of Mental Disorders (DSM-IV) for  
8 dementia and those of NINCDS-ADRDA Work Group; with Mini-Mental State Examination  
9 (MMSE) [9] scores between 10 and 26; initiating therapy for the first time with rivastigmine  
10 or having already received rivastigmine for less than one year; with a planned follow-up visit  
11 within 12 months of enrolment and a caregiver's burden assessment; with an available  
12 informant. Patients not treated according to the product labeling, patients involved in a  
13 clinical trial or patients treated by an experimental drug within the previous 4 weeks were  
14 excluded. Investigators were advised to follow product labeling.

15 The Steering Committee, the French National Council of Medical Practitioners Order (CNO)  
16 and the National Council of Informatics and Freedom (CNIL) approved the study. All the  
17 patients were given a study information sheet.

18 For the present analysis, only subjects followed during twelve months were considered  
19 (n=204). Figure 1 is a detailed flow chart of patients 'selection and exclusion.  
20

### 21 *Study assessments*

22 At study entry, a detailed medical history of each patient was taken and baseline and  
23 follow-up assessments of cognitive performances was performed.  
24 Rapid cognitive decline was defined as a decline of at least three points of MMSE during the  
25 one year of follow-up. The following potential predictive factors of RCD were collected at  
26 baseline: MMSE, presence of behavioral disorders reported by the patient or his family  
27 (delusions, hallucinations, agitation/aggression), presence of extrapyramidal signs, lack of  
28 autonomy for the key Instrumental Activities of Daily Living (IADL) items (budget, phone,  
29 drugs, transport) [10], lack of stabilization with previous IChE or memantine treatment, Body  
30 Mass Index (BMI) and level of education. The clock drawing test[11], the 5-word test[12] and  
31 a verbal fluency test were also taken into account.

32 The 19-item caregiver questionnaire, the Deco score was collected at the end of the 12-  
33 month follow-up.

1 Safety (vital signs, adverse events and a physical examination) regarding the use of  
2 IChE was assessed throughout the study as needed.

### 4 *Data analysis*

5 Characteristics of the patients who were seen at the 12-month visit were compared to  
6 those of all subjects enrolled in the study by appropriate univariate statistical tests.  
7 Using simple and multiple logistic regressions, we studied the risk of RCD according to  
8 baseline characteristics of the patients. The multivariate analysis included all variables  
9 significantly associated with the risk in the univariate analysis at a p-value < 0.10. Moreover,  
10 we compared mean values of the Deco score according to some baseline characteristics of the  
11 patients. We used student t-test for the comparison.  
12 In addition, we ran a Receiver Operating Characteristic (ROC) analysis to detect the best  
13 threshold of diagnosis of RCD for the Deco score. Sensitivity, specificity, positive and  
14 negative predictive values were calculated for the best thresholds found from the observed  
15 ROC curve. We systematically sought for interactions between age and Deco score on the risk  
16 of RCD. Indeed, the risk of rapid decline is higher among younger subjects so we  
17 hypothesized that the predictive value of Deco might have a differential association with RCD  
18 according to age. SAS statistical software version 9.2 was used to perform analyses.

## 21 **RESULTS**

### 22 *Patients*

23 Three hundred and sixty one patients were enrolled in the study by 152 neurologists,  
24 geriatricians and psychiatrists. Among subjects using ChEI or memantine, 161 patients  
25 (59.6%) experienced a stabilization (29.2%) or an improvement (30.4%) in global  
26 functioning, as measured by the CGI-C. One hundred and fifty seven patients (25.2%) were  
27 excluded from analysis because of loss of follow-up between baseline and the 12-month  
28 follow-up (n=151) or missing value on MMSE (n=6) (figure 1). Baseline characteristics of the  
29 analyzed and enrolled populations were comparable (Table 1). In the analyzed population, 89  
30 patients (43.6%) had received prior ChEI treatment and 115 (56.4%) were de novo patients.  
31 Of the 89 patients who had received prior ChEI treatment, 63 patients were already treated by  
32 rivastigmine and 26 discontinued from a previous treatment by another ChEI.

1 There were more women than men (58.3% versus 41.7%) and the mean age was  
2 approximately 78 years (range 51-102 years). At baseline, patients included in this analysis  
3 had a mean MMSE score of  $19.7 \pm 4.4$ .  
4 For patients who had been previously treated by ChEIs, the mean duration of treatment before  
5 starting the study was  $14.0 \pm 14.3$  months with a mean MMSE decrease of  $2.1 \pm 3.0$  ( $-1.1 \pm$   
6  $2.6$  for rivastigmine;  $-4.1 \pm 2.8$  for donepezil and  $-3.6 \pm 3.0$  for galantamine).  
7 For the analyzed population, the MMSE decrease within the previous year was only known  
8 for 101 patients (i.e., data missing for 103).  
9 The mean follow-up period was  $377.6 (\pm 61.2)$  days.

10

### 11 *Predictive factors of cognitive decline*

12 Sixty of the 204 patients retained for analysis (29.4%, 95%CI [23.4; 35.8]) were  
13 qualified as rapid cognitive decliners as they lost three or more points on the MMSE score  
14 between the inclusion and the 12-month follow-up visit.

15 Table 2 shows that the demographic and neuropsychological characteristics  
16 significantly associated with RCD by univariate analysis are: living in an institution and a  
17 high level of education (p-value < 0.03). In addition, a weaker level of autonomy and the loss  
18 of 3 points or more on the MMSE during the year prior to inclusion tended to be associated  
19 with RCD (p-value < 0.07). As there was a high number of missing values for the variable  
20 “loss of 3 points or more on the MMSE during the year prior inclusion”, two multivariate  
21 models were performed to analyze the risk of RCD (table 3). Model 1 included residential  
22 status, level of education and loss of autonomy. In this model, only the level of education  
23 remained significantly associated with RCD (OR = 2.21, 95%I.C. (1.12 – 4.37)). The second  
24 model (model 2) was further adjusted for the loss of more than 3 points on MMSE prior  
25 inclusion. In this model, the level of education and the loss of more than 3 points on MMSE  
26 were significant predictors of rapid decline.

27

### 28 *The Deco questionnaire*

29 The mean Deco questionnaire score was  $19.0 \pm 8.8$ . It was significantly different  
30 between rapid decliners (15.5 [13.6; 17.5]) and slow decliners (20.3 [18.8; 21.8]) (p for  
31 difference = 0.001) (table 4).

32 Table 4 displays the mean of the Deco score according to baseline characteristics of  
33 the patients. No significant association was found for socio-demographic characteristics.

1 Significant lower values of Deco scores were observed for subjects with low level of (p-value  
2 = 0.07).

3 Figure 2 shows the ROC curve of the predictive value of RCD according to the Deco  
4 score. Visually, the threshold which maximizes the specificity and the sensitivity of the score  
5 seems to be between 16 and 18. Table 5 reports the risks of RCD as well as the results from a  
6 sensitivity/specificity analysis for three thresholds (a Deco score  $\leq 16$ ,  $\leq 17$  or  $\leq 18$ ). The  
7 best model in terms of risk was observed for the threshold of 16. The risk of RCD for a  
8 patient having a score  $\leq 16$  for the Deco questionnaire was three times higher than for a  
9 patient scoring over 16 (p=0.002). However, we found an interaction between age and the  
10 threshold of 16 on the risk of RCD. For this threshold, the risk of RCD was significantly  
11 increased among younger subjects (age  $< 75$ ) but not among older ones (respectively OR =  
12 6.7, 95% C.I. (1.8-25.1) and OR = 2.1, 95% C.I. (0.9-4.8)). As a consequence we ran new  
13 analyses to evaluate the best predictive thresholds of RCD according to age-group (below or  
14 over 75). Results are reported in table 6. We found that a threshold of 16 among subjects aged  
15 below 75 years old and a threshold of 14 for subjects older than 75 years old maximized the  
16 predictive values of the Deco score. Sensitivity, specificity and predictive values were  
17 consistently better than in the analysis pooling the two age-groups.

18

## 19 **DISCUSSION**

20 In this one year observational study we found that 29.4% of patients with AD lost 3  
21 points or more at the MMSE score and were thus classified as rapid decliners over the study  
22 period. Living in an institution, having a high educational level and being impaired for at least  
23 two activities in daily living were significantly associated with RCD. Finally, the Deco score  
24 was differently related to the occurrence of RCD according to the age of the patient. A  
25 threshold of 16 was found to be the best diagnostic tool for RCD among subjects aged  $< 75$   
26 years old while a threshold of 14 was found for subjects aged  $\geq 75$  years old.

27 In our study we found a relatively low proportion of rapid decliners compared with  
28 other studies. Although other definitions of cognitive decline have been proposed [3; 13], a  
29 decline of 3 points or more of the Mini-Mental State Examination (MMSE) over one year is a  
30 commonly used definition of RCD. Indeed, several clinical or population-based studies have  
31 shown that this threshold of decline was strongly linked to mortality or entry in an institution  
32 [2; 14]. Moreover, the use of MMSE compared to other neuropsychological tests is the only  
33 one to fit with clinical practice. In addition, all our patients were using ChEI so the low



1 proportion of rapid decliners found is consistent with the fact that rivastigmine, which is the  
2 only therapy studied in this population, has been shown to reduce the risk of rapid cognitive  
3 decline in patients using ChEI [4; 15].

4 As AD patients with RCD have a poorer prognosis, they warrant particular attention  
5 from the clinician. Previous efforts have been made to identify risk factors of rapid decline  
6 which could be targeted for timely introduction of preventive actions [6; 7; 16; 17] . In our  
7 study, demographic factors as living in institution and high level of education, and previous  
8 rapid decline appear as independent factors correlated with a rapid disease progression, while  
9 the usual clinical predictors as poor nutritional status, psychotic or extrapyramidal signs were  
10 not found to be correlated. While this lack of correlation could be attributed to the greater  
11 benefits of rivastigmine in this population [5; 7], the low proportion of patients in our study  
12 with such clinical symptoms at baseline should also be taken into account.

13 Even if certain factors may be good predictors of rapid decline, available data remain  
14 contradictory [18] and are not sufficient or relevant in routine practice for clinical-decision  
15 making, especially for newly diagnosed patients. In this respect, we aimed to assess the  
16 capacity of a new scale to evaluate the one-year change in cognitive functioning of patients  
17 according to their caregiver (or a person in contact with them). The Deco questionnaire used  
18 in this study appeared to be a valuable clinical tool for the detection of RCD. Indeed, when  
19 taking into account the age of the patients, thresholds of detection were strongly associated  
20 with RCD. A Deco score of less than 16 among patients under 75 years old, and of less than  
21 14 among patients over 75 years old, was found to be strongly associated with RCD. The risk  
22 of being a rapid decliner was 3 to 6 times higher among patients whose caregiver scored less  
23 than the threshold found for the Deco score. Moreover, the positive predictive value was close  
24 to 50% in the younger age-group which is a relevant value for clinical decision making.  
25 Indeed, it indicates that if an informant reports a score lower than 16 for a patient, there is  
26 50% chance that this subject has actually encountered a rapid decline within the previous  
27 year. Preventive care for these subjects appears necessary and will be useful for one patient  
28 out of 2. In addition, it is to be noted that the predictive values of the Deco score were better  
29 in the younger group of patients. This finding may be explained by the fact that older patients  
30 complain less than younger ones and they, as well as their caregivers, may attribute their  
31 changes in cognitive performances to age rather than to a pathological process. They may  
32 report less frequently little cognitive changes. Hence, the Deco questionnaire could represent  
33 a simple way of detecting patients with a high probability of rapidly progressing symptoms  
34 and thus to adapt disease management accordingly.

1           While our results are promising, this study does have some limitations. It remains  
2 merely a historical comparison and no definitive conclusion can be drawn. Moreover, only a  
3 low number of neurologists finally recruited patients. Together with the fact that a quarter of  
4 the patients were lost to follow-up, this could have introduced a bias. Our sample may not be  
5 fully representative of the AD population and thus our results should be interpreted with  
6 caution. Nonetheless, when enrolled patients' baseline characteristics were compared to those  
7 of the finally analyzed population, no difference was found, especially in the items of level of  
8 autonomy and cognitive impairment.

9           Overall this study suggests that AD patients at risk of RCD can be identified in the  
10 setting of a regular consultation by means of the Deco questionnaire. Once alerted to the  
11 possibility of an aggressive disease course, the physician could pay particular attention to  
12 these patients and adapt disease management accordingly.

13

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- 16  
17

1 Table 1 – Comparison of baseline characteristics of enrolled and analyzed patients

2

|   | <b>Enrolled Population</b><br>N = 361 | <b>Analyzed population</b><br>N = 204 |
|---|---------------------------------------|---------------------------------------|
| <b>Age</b> (years), m ± SD                      | 77.9 ± 7.55                           | 78.0 ± 7.5                            |
| <b>Sex</b> (female)                             | 215 (59.6%)                           | 119 (58.3%)                           |
| <b>Level of education</b>                       |                                       |                                       |
| * long secondary school                         | 92 (25.6%)                            | 53 (26.0%)                            |
| <b>Residential status</b> (out-patients)        | 302 (83.7%)                           | 169 (82.8%)                           |
| <b>Duration of the disease</b> (years) , m ± SD | 1.8 ± 1.5                             | 1.8 ± 1.4                             |
| <b>Psychotropic drugs</b>                       | 170 (47.1%)                           | 106 (52.0%)                           |
| * neuroleptics                                  | 31 (8.6%)                             | 20 (9.8%)                             |
| * antidepressive drugs                          | 98 (27.1%)                            | 60 (29.4%)                            |
| <b>Previous treatment by ChEI</b>               |                                       |                                       |
| * none  | 214 (59.3%)                           | 115 (56.4%)                           |
| * rivastigmine                                  | 97 (26.9%)                            | 63 (30.9%)                            |
| * other ChEI                                    | 50 (13.8%)                            | 26 (12.7%)                            |
| <b>Memantine</b>                                | 26 (7.2%)                             | 10 (4.9%)                             |
| <b>MMSE score</b> , m ± SD                      | 19.6 ± 4.6                            | 19.7 ± 4.4                            |
| <b>mini-Zarit score</b> , m ± SD                | 3.1 ± 1.7                             | 3.1 ± 1.7                             |
| <b>4-IADL score</b> (≥ 2 activities)            | 225 (62.4%)                           | 123 (60.3%)                           |
| <b>5 words test</b> , m ± SD                    | 5.6 ± 2.3                             | 5.7 ± 2.3                             |
| <b>Verbal fluency</b> , m ± SD                  | 10.7 ± 5.5                            | 10.5 ± 5.1                            |

3 Figures are given as n and % unless otherwise specified

4 SD: Standard Deviation

1 Table 2. Association between baseline characteristics of patients and the risk of rapid  
 2 cognitive decline. Univariate logistic regression.

|   | Rapid cognitive decline |             | OR (95%I.C.)            | p-value      |
|---|-------------------------|-------------|-------------------------|--------------|
|   | Yes<br>n=60             | No<br>n=144 |                         |              |
| <b>Sex</b>  |                         |             |                         |              |
| Male  | 25 (29.4)               | 60 (29.4)   |                         |              |
| Female  | 35 (70.6)               | 84 (70.6)   | n.a                     |              |
| <b>Age</b>  |                         |             |                         |              |
| <75 years old   | 16 (26.7)               | 43 (29.9)   | 1.00                    |              |
| >= 75 years old   | 44 (73.3)               | 101 (70.1)  | 1.17 (0.60-2.30)        | 0.647        |
| <b>Residential status</b>                               |                         |             |                         |              |
| Out-patients  | 44 (73.3)               | 125 (86.8)  | 1.00                    |              |
| Institution   | 16 (26.7)               | 19 (13.2)   | <b>2.39 (1.13-5.06)</b> | <b>0.022</b> |
| <b>Level of education</b>                               |                         |             |                         |              |
| primary or short secondary school                       | 22 (36.7)               | 31 (21.5)   | 1.00                    |              |
| long secondary school                                   | 38 (63.3)               | 113 (78.5)  | <b>2.11 (1.09-4.08)</b> | <b>0.026</b> |
| <b>BMI</b>  |                         |             |                         |              |
| < 18.5 kg/m <sup>2</sup>                                | 9 (23.1)                | 30 (20.8)   | 0.68 (0.26-1.78)        | 0.432        |
| 18 <= BMI < 25 kg/m <sup>2</sup>                        | 15 (30.6)               | 34 (23.6)   | 1.00                    |              |
| >= 25 kg/m <sup>2</sup>                                 | 36 (60.0)               | 80 (55.6)   | 1.02 (0.50-2.10)        | 0.957        |
| <b>MMSE</b>   |                         |             |                         |              |
| >=20  | 35 (58.3)               | 78 (34.2)   | 1.00                    |              |
| <20   | 25 (41.7)               | 66 (45.8)   | 0.84 (0.46-1.55)        | 0.586        |
| <b>Verbal fluency</b>                                   |                         |             |                         |              |
| >=15  | 12 (20.3)               | 27 (18.9)   | 1.00                    |              |
| <15   | 47 (79.7)               | 116 (81.1)  | 0.91 (0.43-1.95)        | 0.811        |
| <b>5 words</b>  |                         |             |                         |              |
| >=8   | 10 (16.7)               | 36 (25.2)   | 1.00                    |              |
| <8  | 50 (83.3)               | 107 (74.8)  | 1.68 (0.77-3.66)        | 0.190        |
| <b>Clock</b>  |                         |             |                         |              |
| >2  | 22 (37.9)               | 48 (35.6)   | 1.00                    |              |
| <=2   | 36 (62.1)               | 95 (66.4)   | 0.83 (0.44-1.56)        | 0.557        |
| <b>Symptoms</b>   |                         |             |                         |              |
| Agitation   | 13 (21.7)               | 22 (15.3)   | 1.72 (0.76-3.93)*       | 0.194        |
| Hallucination   | 9 (15.0)                | 16 (11.1)   | 1.64 (0.65-4.18)*       | 0.298        |
| Strolling   | 6 (10.0)                | 13 (9.0)    | 1.35 (0.46-3.92)*       | 0.584        |
| Depression  | 14 (23.3)               | 38 (26.4)   | 1.08 (0.50-2.31)*       | 0.851        |
| EPS   | 6 (10.0)                | 18 (12.5)   | 0.97 (0.35-2.73)*       | 0.959        |
| <b>Loss of autonomy for at least 2 activities</b>       |                         |             |                         |              |
| No  | 18 (30.0)               | 63 (43.8)   | 1.00                    |              |
| Yes   | 42 (70.0)               | 81 (56.2)   | <b>1.81 (0.95-3.45)</b> | <b>0.069</b> |
| <b>Previous ChEI treatment</b>                          |                         |             |                         |              |
| No  | 32 (53.3)               | 83 (57.6)   | 1.00                    |              |
| Yes   | 28 (46.7)               | 61 (42.4)   | 1.19 (0.65-2.18)        | 0.572        |
| <b>Loss &gt;= 3 points on MMSE in the previous year</b> |                         |             |                         |              |
| No  | 10 (29.4)               | 33 (49.3)   | 1.00                    |              |
| Yes   | 24 (70.6)               | 34 (50.8)   | <b>2.33 (0.97-5.61)</b> | <b>0.060</b> |

\* Compared with none trouble

Table 3. Risk of rapid decline in two multivariate logistic regression analyses

|   | <b>Model 1 (n = 204)</b>  |                | <b>Model 2 (n = 101)</b>   |                |
|---|---------------------------|----------------|----------------------------|----------------|
|   | <b>OR (95%I.C.)</b>       | <b>p-value</b> | <b>OR (95%I.C.)</b>        | <b>p-value</b> |
| <b>Residential status</b>   |                           |                |                            |                |
| Out-patients  | 1.00                      |                | 1.00                       |                |
| Institution   | 1.95 (0.89 - 4.31)        | 0.097          | 2.88 (0.97 - 8.59)         | 0.057          |
| <b>Level of education</b>   |                           |                |                            |                |
| Primary or short secondary school                                   | <b>1.00</b>               |                | <b>1.00</b>                |                |
| long secondary school   | <b>2.21 (1.12 - 4.37)</b> | <b>0.022</b>   | <b>4.94 (1.83 - 13.32)</b> | <b>0.002</b>   |
| <b>Loss of autonomy for at least 2 activities</b>                   |                           |                |                            |                |
| No  | 1.00                      |                | 1.00                       |                |
| yes   | 1.71 (0.86 - 3.40)        | 0.124          | 0.72 (0.26 - 2.01)         | 0.529          |
| <b>Loss <math>\geq</math> 3 points on MMSE in the previous year</b> |                           |                |                            |                |
| No  |                           |                | <b>1.00</b>                |                |
| yes   |                           |                | <b>2.81 (1.03 - 7.65)</b>  | <b>0.043</b>   |

Table 4. Association between baseline characteristics of patients and the Deco score

|   | <b>m</b> | <b>sd</b> | <b>p-value</b> |
|---|----------|-----------|----------------|
| <b>Sex</b>  |          |           |                |
| Male  | 18,5     | 9,1       | 0,618          |
| Female  | 19,2     | 8,7       |                |
| <b>Age</b>  |          |           |                |
| <75 years old   | 19,1     | 8,8       | 0,803          |
| >= 75 years old   | 18,8     | 8,9       |                |
| <b>Residential status</b>                                 |          |           |                |
| Out-patients  | 18,8     | 8,8       | 0,727          |
| Institution   | 19,6     | 9,6       |                |
| <b>Level of education</b>                                 |          |           |                |
| Primary or short secondary school                         | 19,1     | 9,1       | 0,388          |
| long secondary school                                     | 18,3     | 8,1       |                |
| <b>Loss of autonomy for at least 2 activities</b>         |          |           |                |
| No  | 20,3     | 8,6       | <b>0,071</b>   |
| yes   | 17,9     | 8,9       |                |
| <b>Rapid decline between inclusion and 12-month visit</b> |          |           |                |
| No  | 20,2     | 8,9       | <b>0,0008</b>  |
| yes   | 15,1     | 7,4       |                |



Table 5. Predictive power of 3 thresholds of the Deco score. Univariate logistic regression.

|            |           | <b>RCD</b> |                  | <b>OR (95%I.C.)</b> | <b>p-value</b> | <b>Se</b> | <b>Sp</b> | <b>PPV</b> | <b>NPV</b> |
|------------|-----------|------------|------------------|---------------------|----------------|-----------|-----------|------------|------------|
|            | Yes       | No         |                  |                     |                |           |           |            |            |
| DECO > 16  | 17 (37.8) | 85 (64.4)  | 1.00             |                     |                |           |           |            |            |
| DECO <= 16 | 28 (62.2) | 47 (35.6)  | 2.98 (1.46-6.00) | 0.002               | 62.2           | 64.4      | 37.3      | 83.3       |            |
| DECO > 17  | 16 (35.6) | 79 (64.4)  | 1.00             |                     |                |           |           |            |            |
| DECO <= 17 | 29 (59.9) | 53 (40.2)  | 2.70 (1.34-5.46) | 0.006               | 64.4           | 59.9      | 35.4      | 83.2       |            |
| DECO > 18  | 15 (33.3) | 72 (54.6)  | 1.00             |                     |                |           |           |            |            |
| DECO <= 18 | 30 (66.7) | 60 (45.5)  | 2.40 (1.18-4.87) | 0.015               | 66.7           | 54.6      | 33.3      | 82.8       |            |

RCD states for rapid cognitive decline, Se for sensitivity, Sp for specificity, PPV for positive predictive value, NPV for negative predictive value

Table 6. Best predictive thresholds of the Deco score according to age-groups. Univariate logistic regression.

|                             | <b>RCD</b> |           | <b>OR (95%I.C.)</b> | <b>p-value</b> | <b>Se</b> | <b>Sp</b> | <b>VPP</b> | <b>VPN</b> |
|-----------------------------|------------|-----------|---------------------|----------------|-----------|-----------|------------|------------|
|                             | <b>Yes</b> | <b>No</b> |                     |                |           |           |            |            |
| <b>Age &lt; 75 yrs old</b>  |            |           |                     |                |           |           |            |            |
| DECO > 16                   | 4 (26.7)   | 29 (70.7) | 1.00                |                |           |           |            |            |
| DECO <= 16                  | 11 (73.3)  | 12 (29.3) | 6.65 (1.76-25.1)    | 0.005          | 73.3      | 70.7      | 47.8       | 87.9       |
| <b>Age &gt;= 75 yrs old</b> |            |           |                     |                |           |           |            |            |
| DECO > 14                   | 14 (17.5)  | 66 (82.5) | 1.00                |                |           |           |            |            |
| DECO <= 14                  | 16 (39.0)  | 25 (61.0) | 3.02 (1.29-7.08)    | 0.011          | 57.8      | 74.2      | 43.3       | 83.8       |

RCD states for rapid cognitive decline, Se for sensitivity, Sp for specificity, PPV for positive predictive value, NPV for negative predictive value

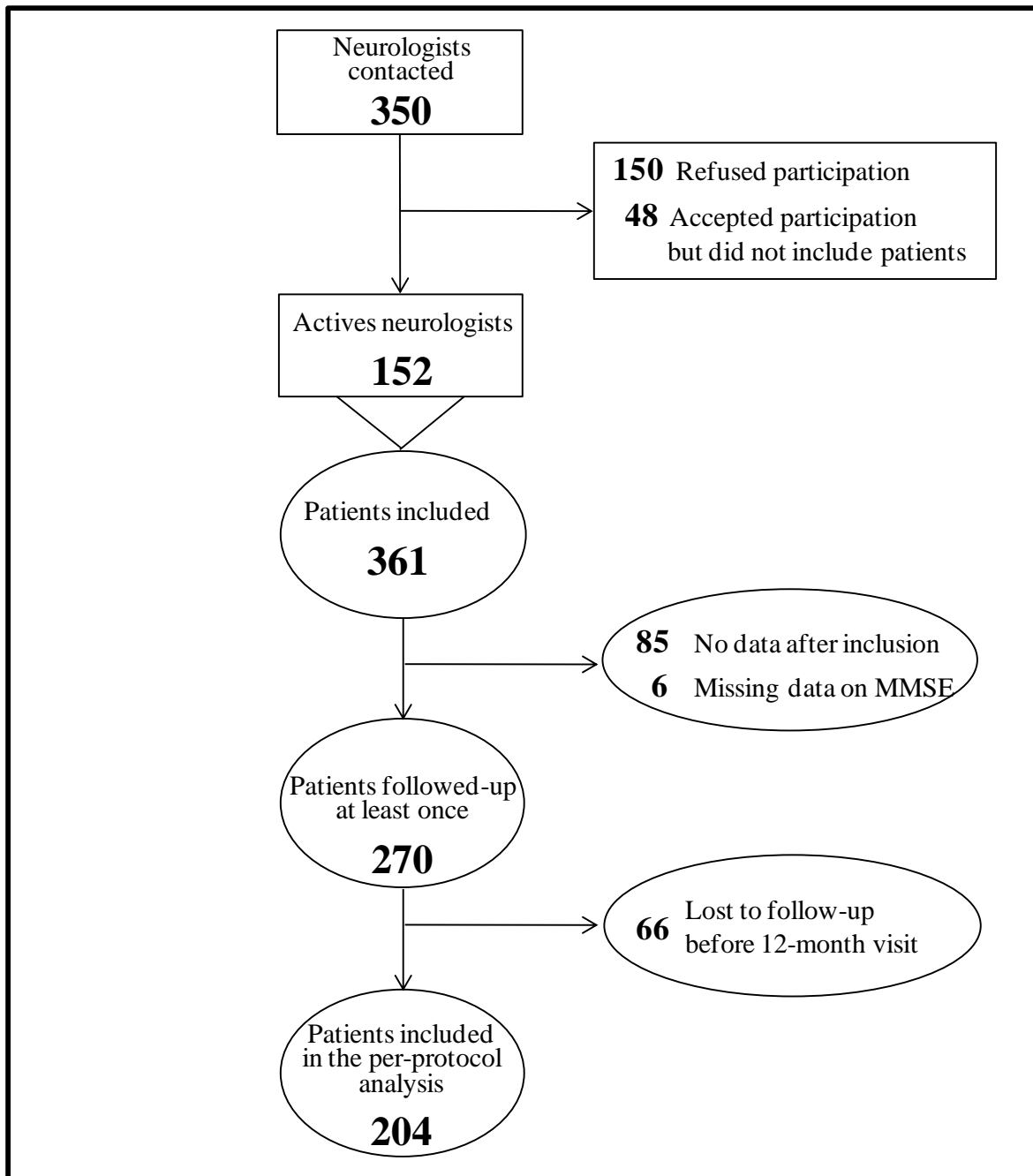


Figure 1: Flow chart of the patients' selection for the per-protocol analysis

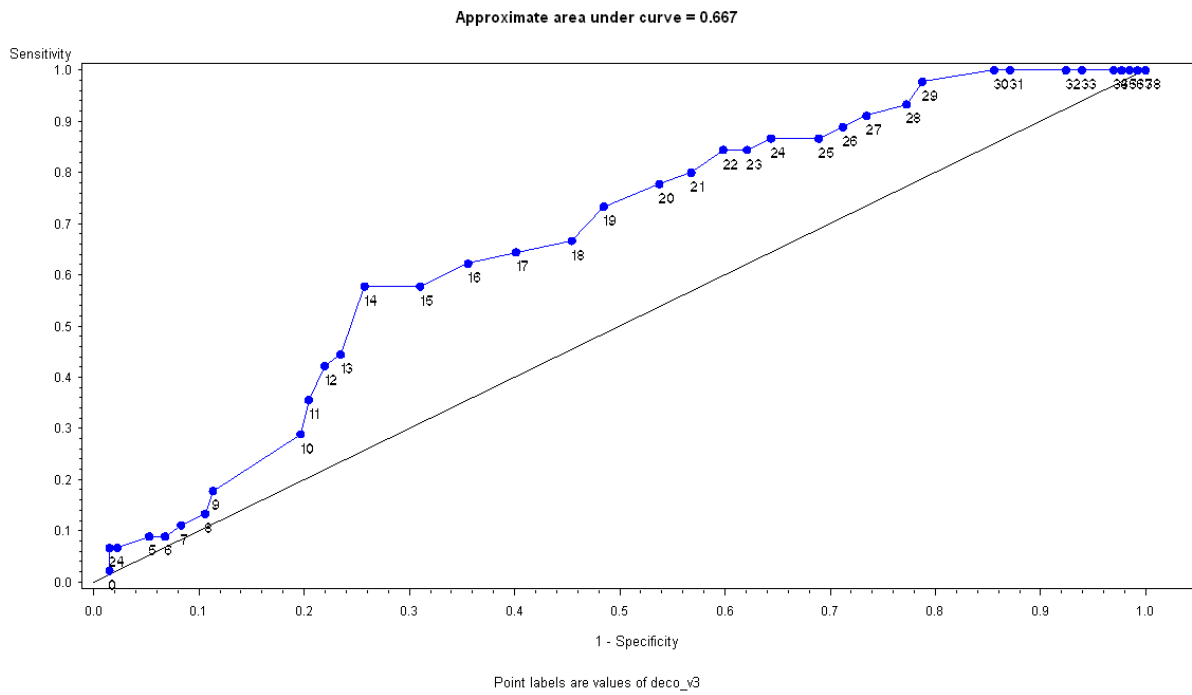


Figure 2: ROC curve for the prediction of rapid cognitive decline according to DECO scores