#### SUPPLEMENTARY MATERIALS

## **Supplementary Methods**

Personal history of smoking behavior was calculated by "pack/years" quantifying the packs smoked per day multiplied by years as a smoker, with the threshold was set to 15. Drinking was defined as an average of alcoholic drink ( $\geq$  50 ml) at least once per week lasting more than half a year. Body-mass index (BMI) was calculated as body weight (kg) divided by heights squared (m<sup>2</sup>).

The following variables were categorized according to the normal reference range [1-11] and we considered the following levels of biochemical indicators as the commonly used thresholds in this study: the low LDL level as less than 2.00 mmol/L [1], the high level of D-Dimer as more than 0.50mg/L [2], the high level of fibringen as more than 4.00g/L [3], the high level of hypersensitive C-reactive protein (hs-CRP) as more than 3.00mg/L [4], the high level of aspartate transaminase (AST) as more than 40IU/L [5], the low level of albumin as less than 35.00g/L [6], the low level of calcium as less than 2.10mmol/L [7], the high level of mean corpuscular volume (MCV), as more than 90.00fL [8], the high level of HCY as more than 15 mmol/L (the cutoff for hyperhomocysteinemia) [9], the high level of Cys C as more than 0.95mg/L [10] and the low level of fasting blood glucose (FBG) as less than 5.00mmol/L [11].

#### **Supplementary Results**

# Multiple linear regression analysis for the association of risk factors of dementia and MMSE score in PD, DM and PD-DM patients

Using multiple linear regression analysis, we found that a lower FBG (<5.00 mmol/L, B= -1.052, SE=0.433,  $\beta$ =-0.126, p=0.016), higher HCY (>15.00  $\mu$ mol/L, B= -1.336, SE=0.595,  $\beta$ = -0.115, p=0.026), and Cys C  $(>0.95 \text{ mg/L}, B= -0.908, SE=0.452, \beta= -0.103,$ p=0.046), and age (B=-0.140, SE=0.019,  $\beta$ = -0.376, p<0.001) were associated with lower MMSE score. No significant association was observed between MMSE score and hyperlipidemia (B= -1.232, SE=0.641,  $\beta$ =-0.098, p=0.056) in PD patients (adjusted  $R^2$ =0.173, F=14.469, p<0.001). In addition, higher fibringen  $(>4.00 \text{ g/L}, B=-1.857, SE=0.663, \beta=-0.174, p=0.006),$ lower LDL-C (<2.00 mmol/L, B= -1.876, SE=0.674,  $\beta$ = -0.173, p=0.006), age (B= -0.171, SE=0.032,  $\beta$ = -0.342, p<0.001) and SAE (B= -2.196, SE=0.640,  $\beta$ = -0.216, p<0.001) were associated with lower MMSE score in PD-DM patients (adjusted  $R^2=0.275$ , F=19.383, p<0.001). A higher AST (>40 IU/L, B= -3.278,

SE=1.023,  $\beta$ = -0.141, p=0.001) and Cys C (>0.95 mg/L, B= -1.878, SE=0.690,  $\beta$ = -0.119, p=0.007), anxiety or depression (B= -1.579, SE=0.531,  $\beta$ = -0.135, p=0.003), male gender (B= -1.182, SE=0.398,  $\beta$ = -0.131, p=0.003) and age (B= -0.163, SE=0.015,  $\beta$ = -0.481, p<0.001) were associated with lower MMSE score in DM patients (adjusted R²=0.310, F=34.277, p<0.001) (Supplementary Table 1). These observations are consistent with the results of multivariable logistic regression analysis for risk factors in dementia in PD, DM and PD-DM patients (Table 3A, Table 3B, Supplementary Table 4).

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