

Supplementary Figure S1

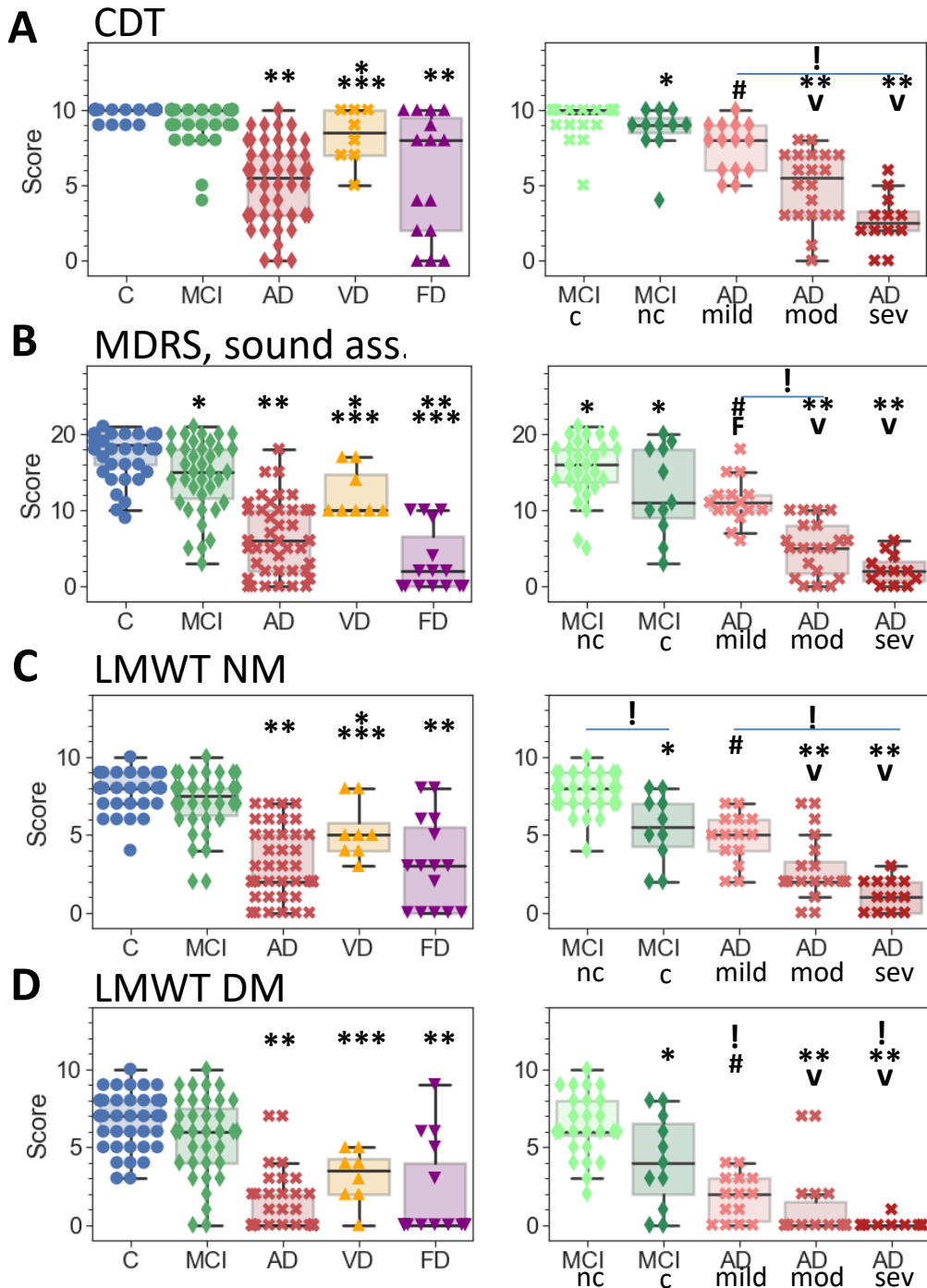


Figure S1. The results of study participants using psychometric tests and scales: (A) clock drawing test (CDT); (B) Mattis dementia rating scale (MDRS), **sound associations subtest; (C, D) Luria memory words test (LMWT, NM – non-intermediate memorization, DM – delayed memorization). MCI – mild cognitive impairment (nc – non-converter, c – converter); AD – Alzheimer’s disease (mild, moderate, and severe subgroups); VD – vascular dementia; FD – frontotemporal dementia. Lines inside boxes show medians; box flanges – 25-75 percentiles; whisker range \pm SD. Significantly different results, with p -values < 0.01 according to Mann-Whitney U-test, are shown with the following symbols: * - significantly different to the control group; ** - to control and MCI groups; *** - to control, MCI, and AD groups; **** - to control and AD groups; ***** - to control, MCI, AD, and VD groups; # - to control and MCI-nc; “F” – to FD; “V” – to VD; “!” – significant difference between subgroups.**

Supplementary Figure S2

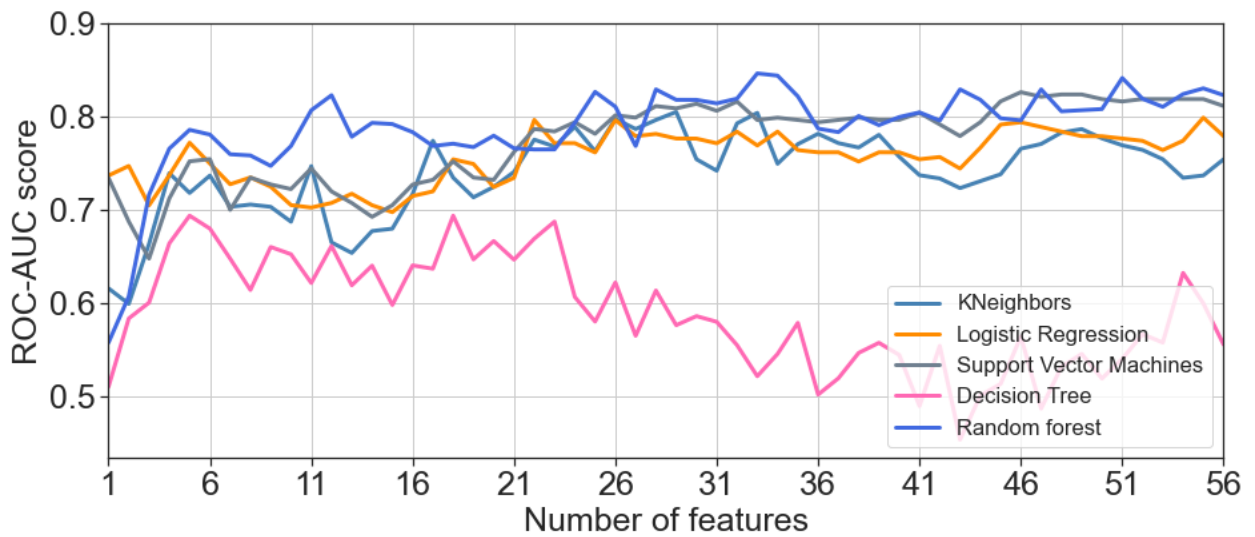


Figure S2. Preliminary evaluation of the effectiveness of different algorithms in the binary classification of AD versus control.

Supplementary Figure S3

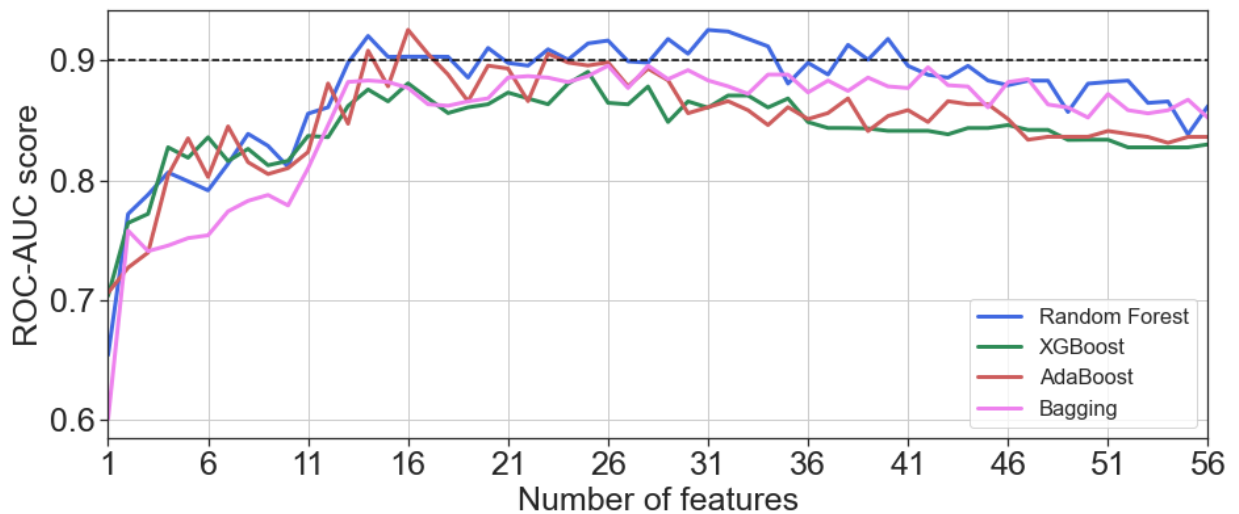


Figure S3. The optimization of models for binary classification of AD and control samples based on RF, AdaBoost, XGBoost, and Bagging algorithms.

Supplementary Figure S4

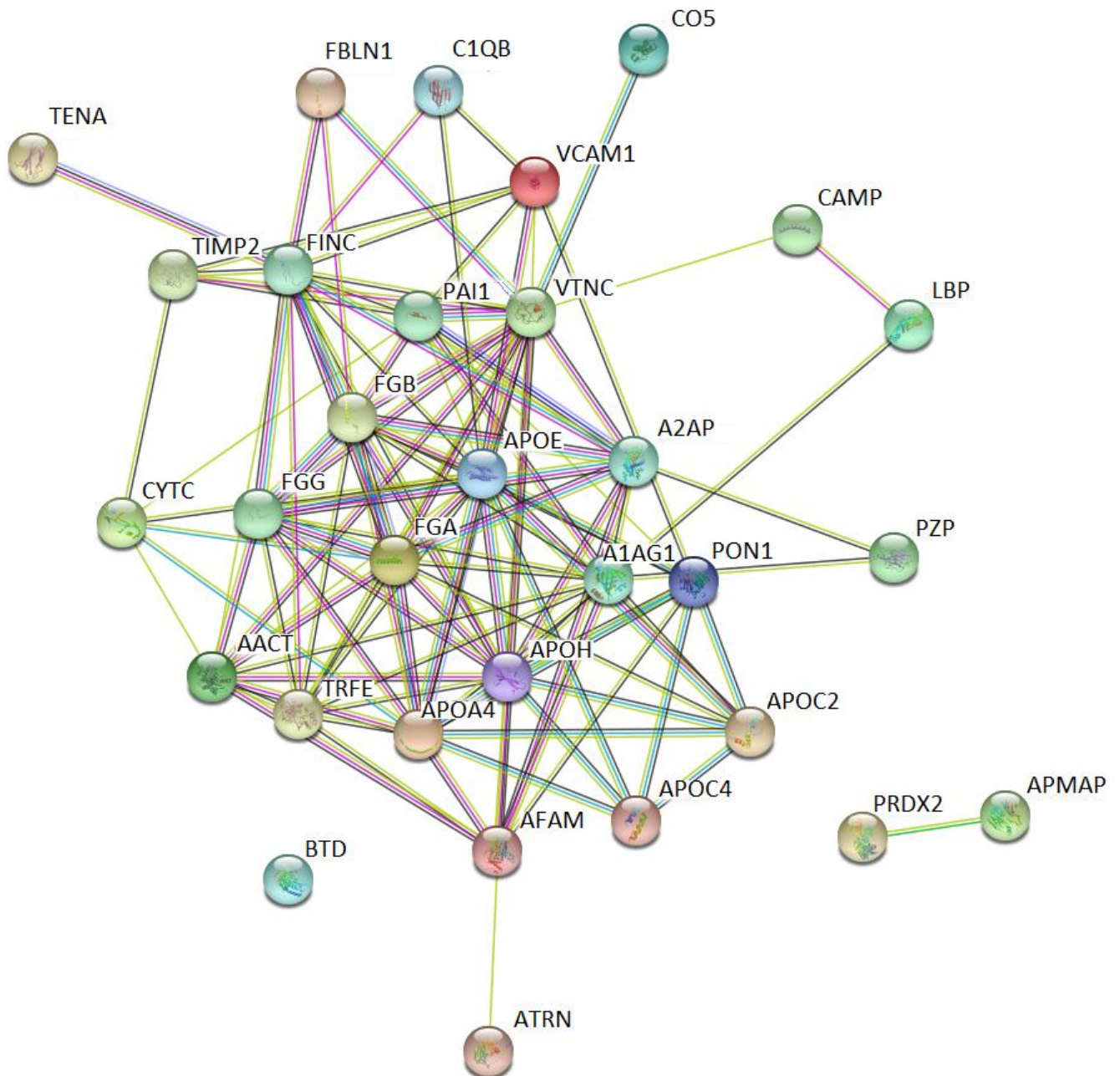


Figure S4. The STRING protein interaction network for 31 proteins important for AD distinction in this study.

Supplementary Figure S5

AD classification score

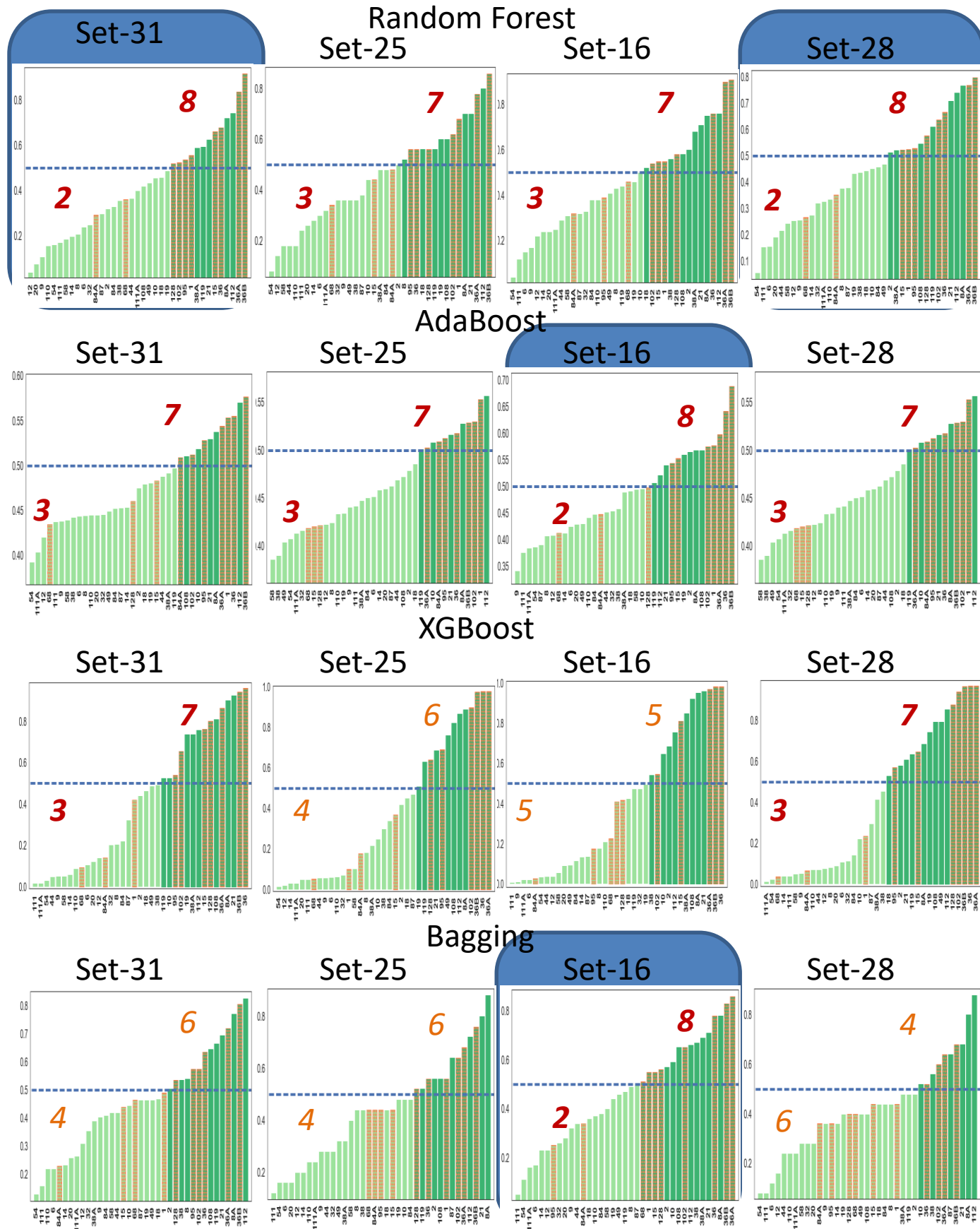


Figure S4. The differentiation of MCI patients using 16 classifiers. The most effective classifiers are marked with a blue background.