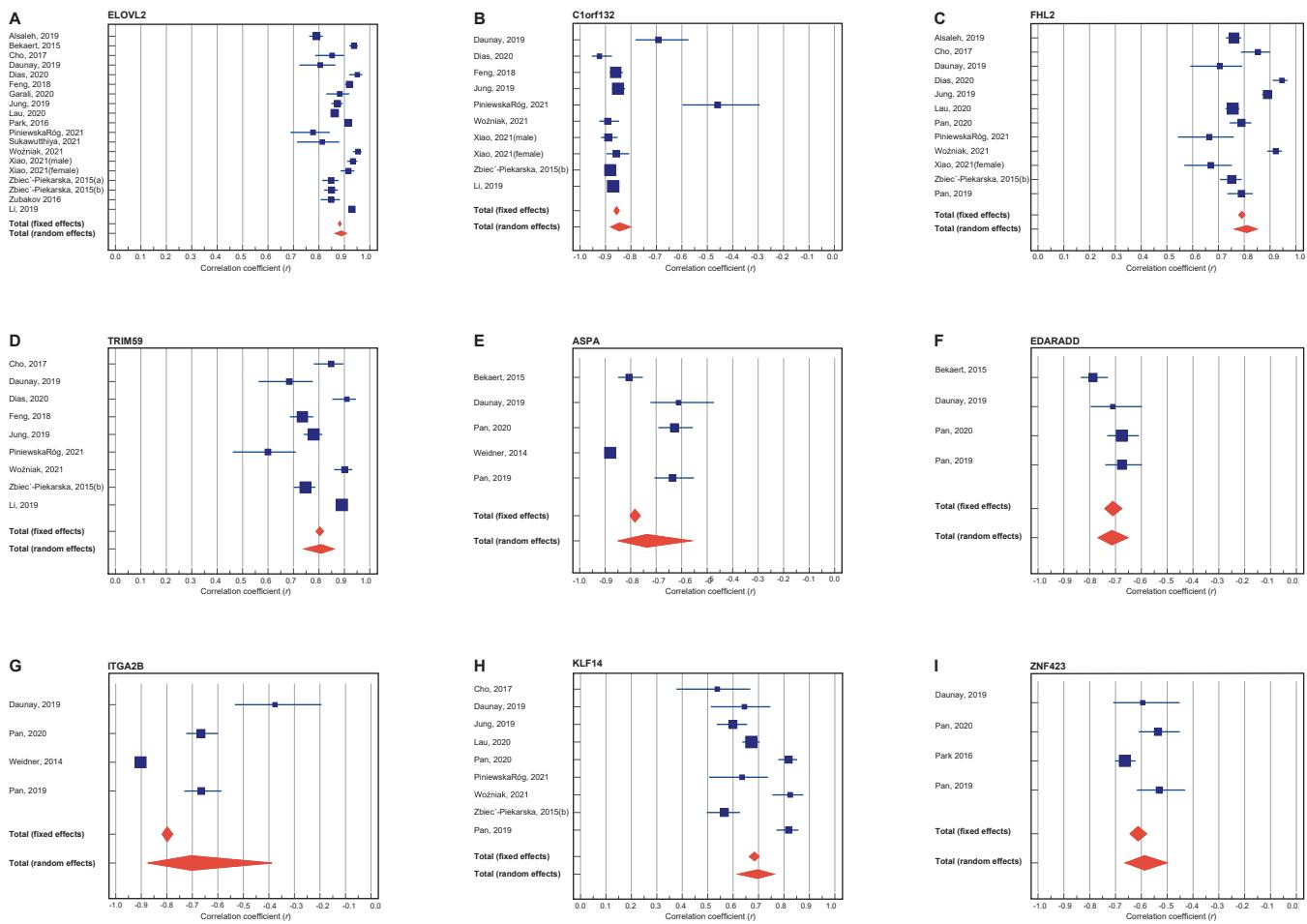


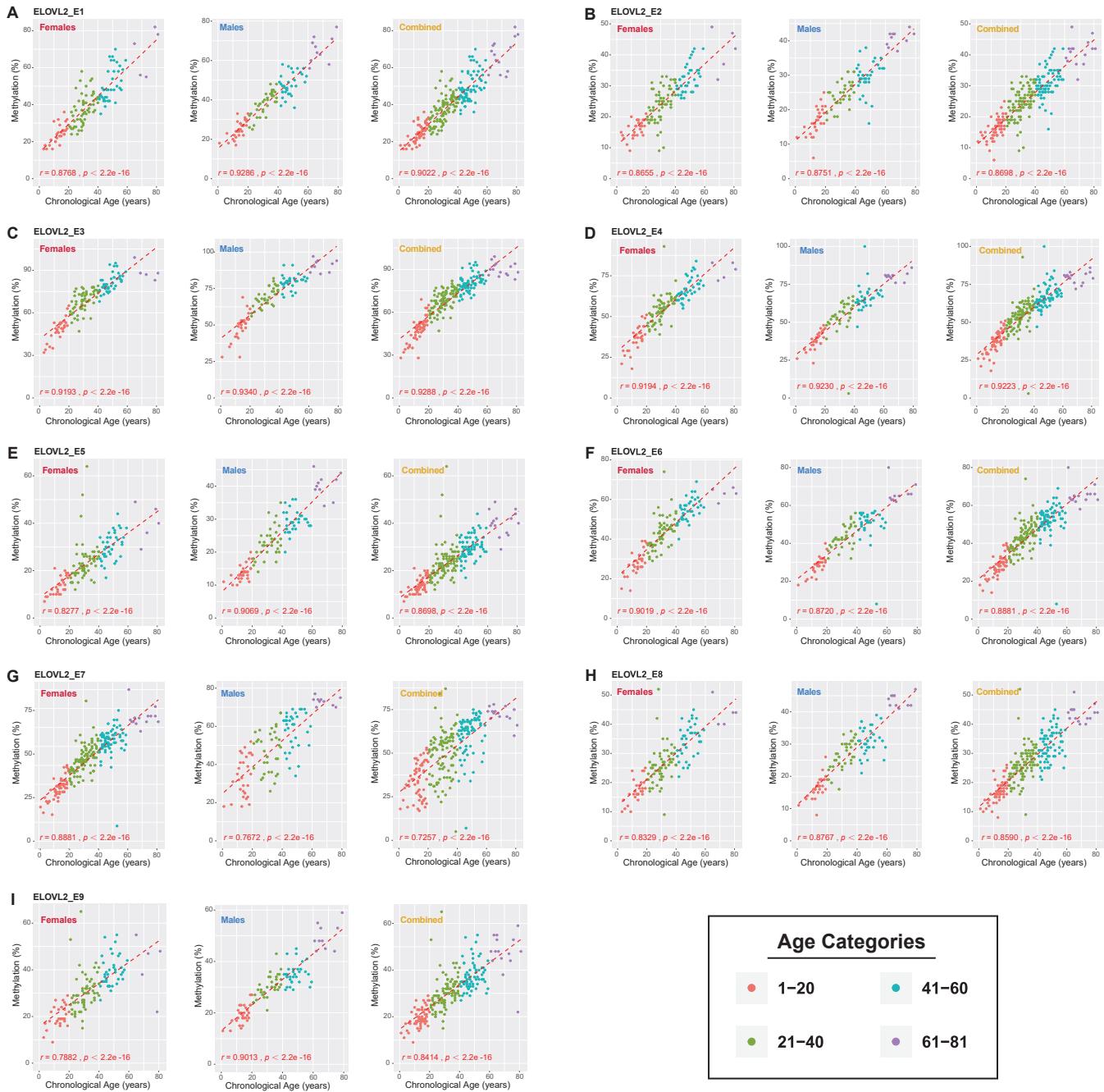
## Supplementary Figure S1



**Supplementary Figure S1** Detailed correlation coefficient ( $r$ ) meta-analysis forest plots of nine candidate DNA methylation biomarkers in meta cohort of 7084 individuals.

- A. ELOVL2** ( $n = 5847$ ,  $r = 0.89$ ); **B. C1orf132** ( $n = 2419$ ,  $r = 0.85$ ); **C. FHL2** ( $n = 3768$ ,  $r = 0.81$ );  
**D. TRIM59** ( $n = 2206$ ,  $r = 0.81$ ); **E. ASPA** ( $n = 1421$ ,  $r = 0.74$ ); **F. EDARADD** ( $n = 846$ ,  $r = 0.71$ );  
**G. ITGA2B** ( $n = 1215$ ,  $r = 0.71$ ); **H. KLF14** ( $n = 2812$ ,  $r = 0.70$ ); **I. ZNF423** ( $n = 1405$ ,  $r = 0.59$ ).

## Supplementary Figure S2



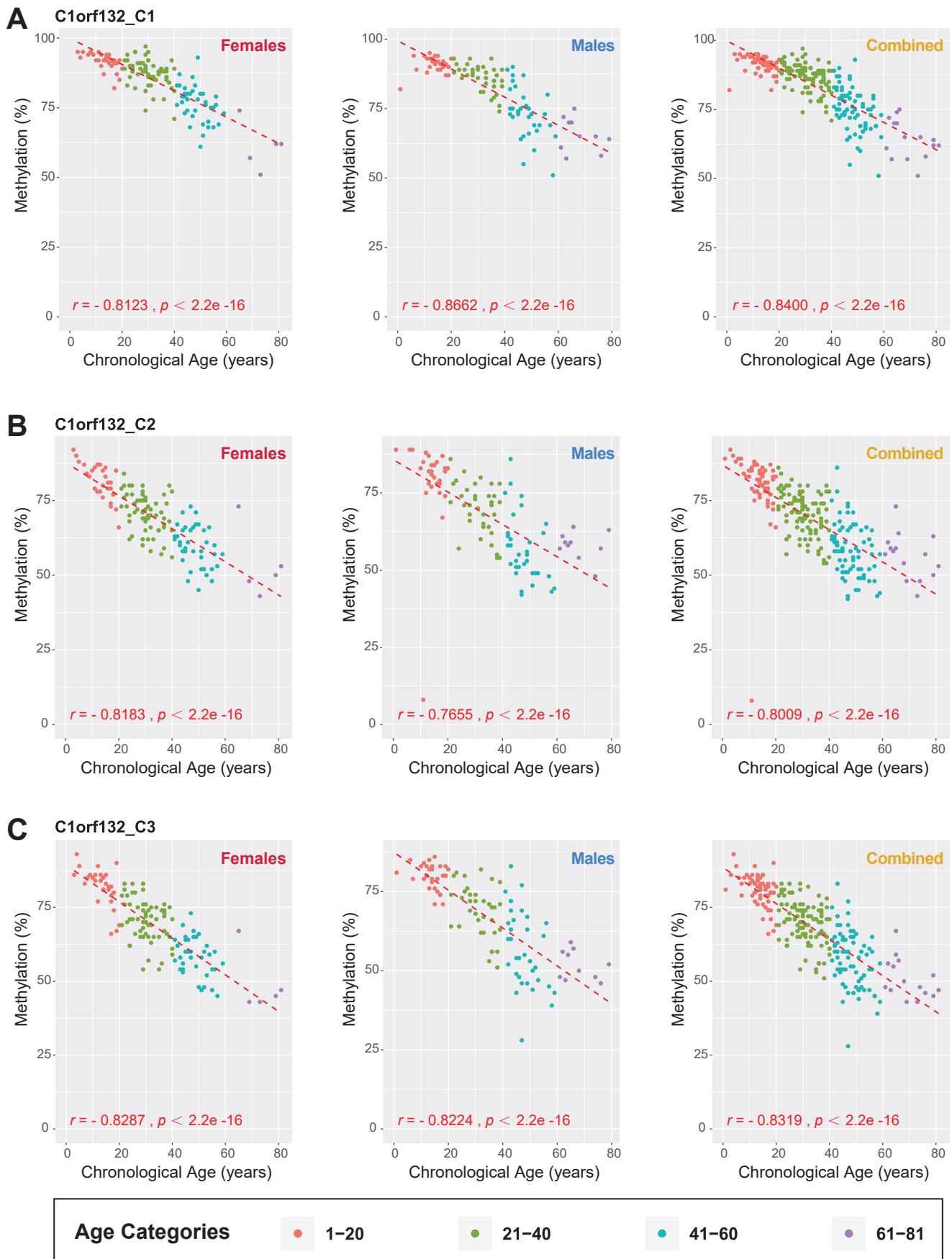
**Supplementary Figure S2** Spearman correlation analyses between different CpG methylation levels of ELOVL2 and the chronological ages of different gender datasets in CHS cohort ( $n = 240$ , blood samples).

**A. ELOVL2\_E1; B. ELOVL2\_E2; C. ELOVL2\_E3; D. ELOVL2\_E4; E. ELOVL2\_E5;**

**F. ELOVL2\_E6; G. ELOVL2\_E7; H. ELOVL2\_E8; I. ELOVL2\_E9;**

Detailed CpG information in Supplementary Table S1.

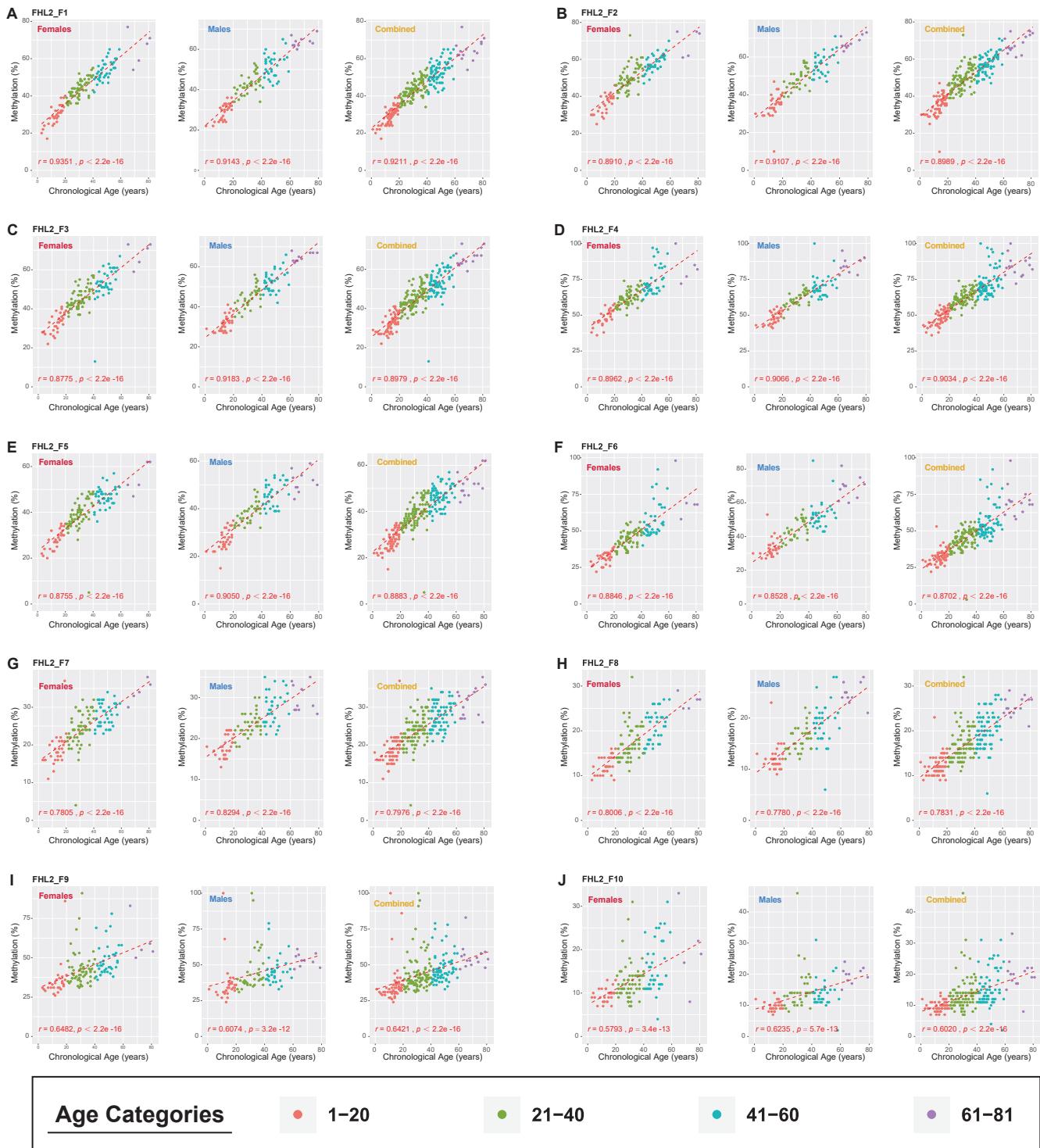
## Supplementary Figure S3



**Supplementary Figure S3** Spearman correlation analyses between different CpG methylation levels of C1orf132 and the chronological ages of different gender datasets in CHS cohort ( $n = 240$ , blood samples).

**A.** C1orf132\_C1; **B.** C1orf132\_C2; **C.** C1orf132\_C3; Detailed CpG information in Supplementary Table S1.

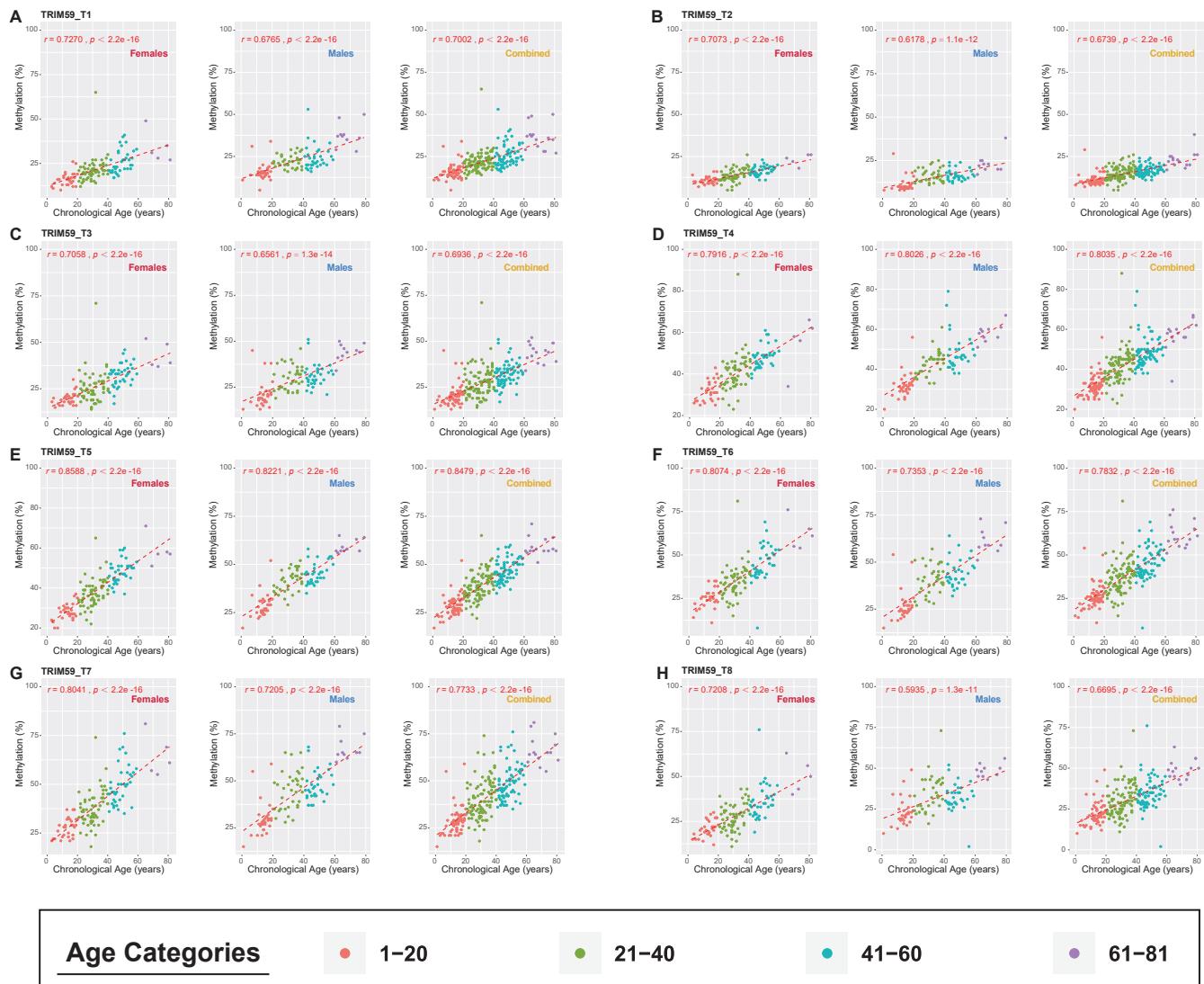
## Supplementary Figure S4



**Supplementary Figure S4** Spearman correlation analyses between different CpG methylation levels of FHL2 and the chronological ages of different gender datasets in CHS cohort (n = 240, blood samples).

**A. FHL2\_F1; B. FHL2\_F2; C. FHL2\_F3; D. FHL2\_F4; E. FHL2\_F5; F. FHL2\_F6; G. FHL2\_F7; H. FHL2\_F8; I. FHL2\_F9; J. FHL2\_F10;**  
Detailed CpG information in Supplementary Table S1.

## Supplementary Figure S5

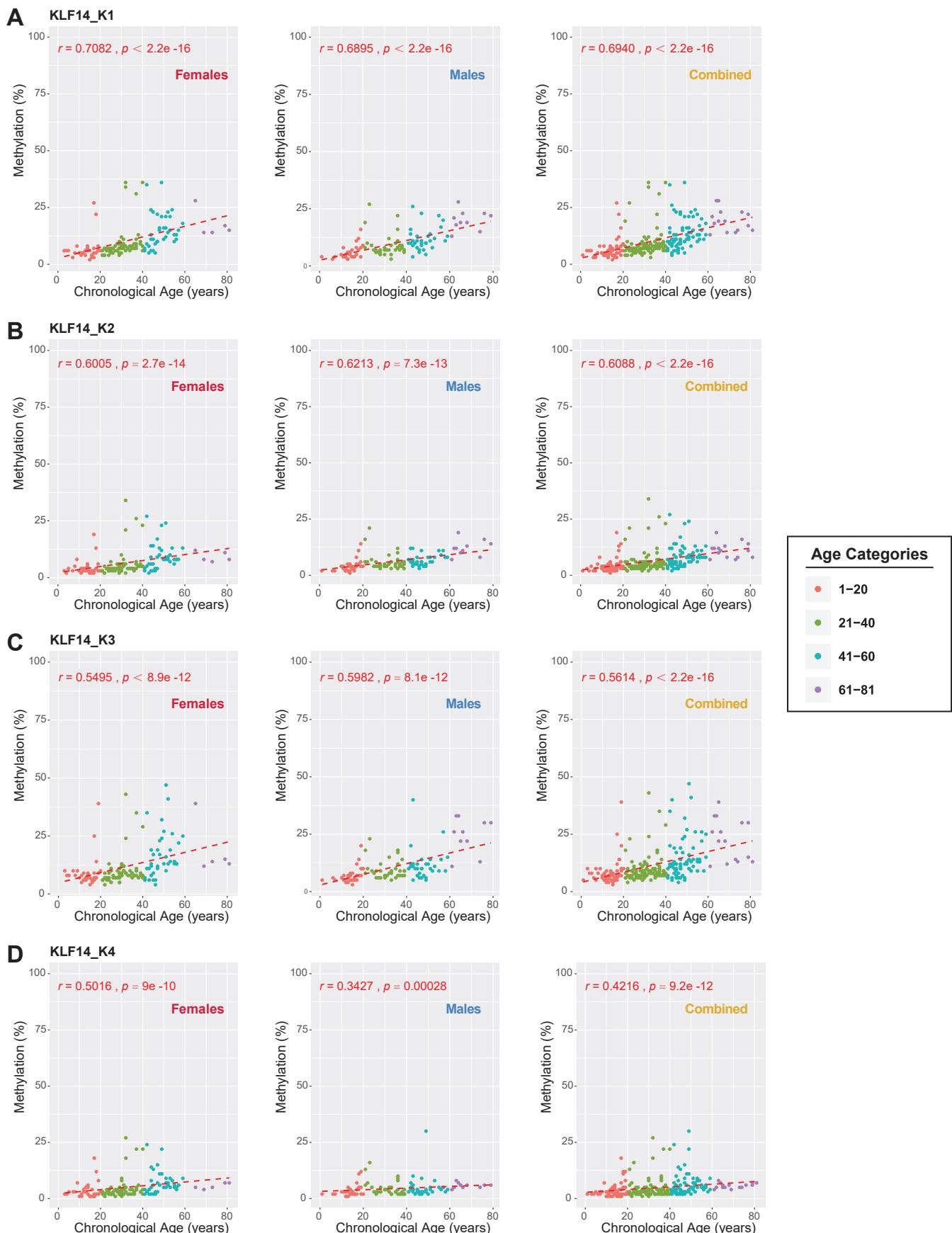


**Supplementary Figure S5** Spearman correlation analyses between different CpG methylation levels of TRIM59 and the chronological ages of different gender datasets in CHS cohort ( $n = 240$ , blood samples).

**A.** TRIM59\_T1; **B.** TRIM59\_T2; **C.** TRIM59\_T3; **D.** TRIM59\_T4;  
**E.** TRIM59\_T5; **F.** TRIM59\_T6; **G.** TRIM59\_T7; **H.** TRIM59\_T8;

Detailed CpG information in Supplementary Table S1.

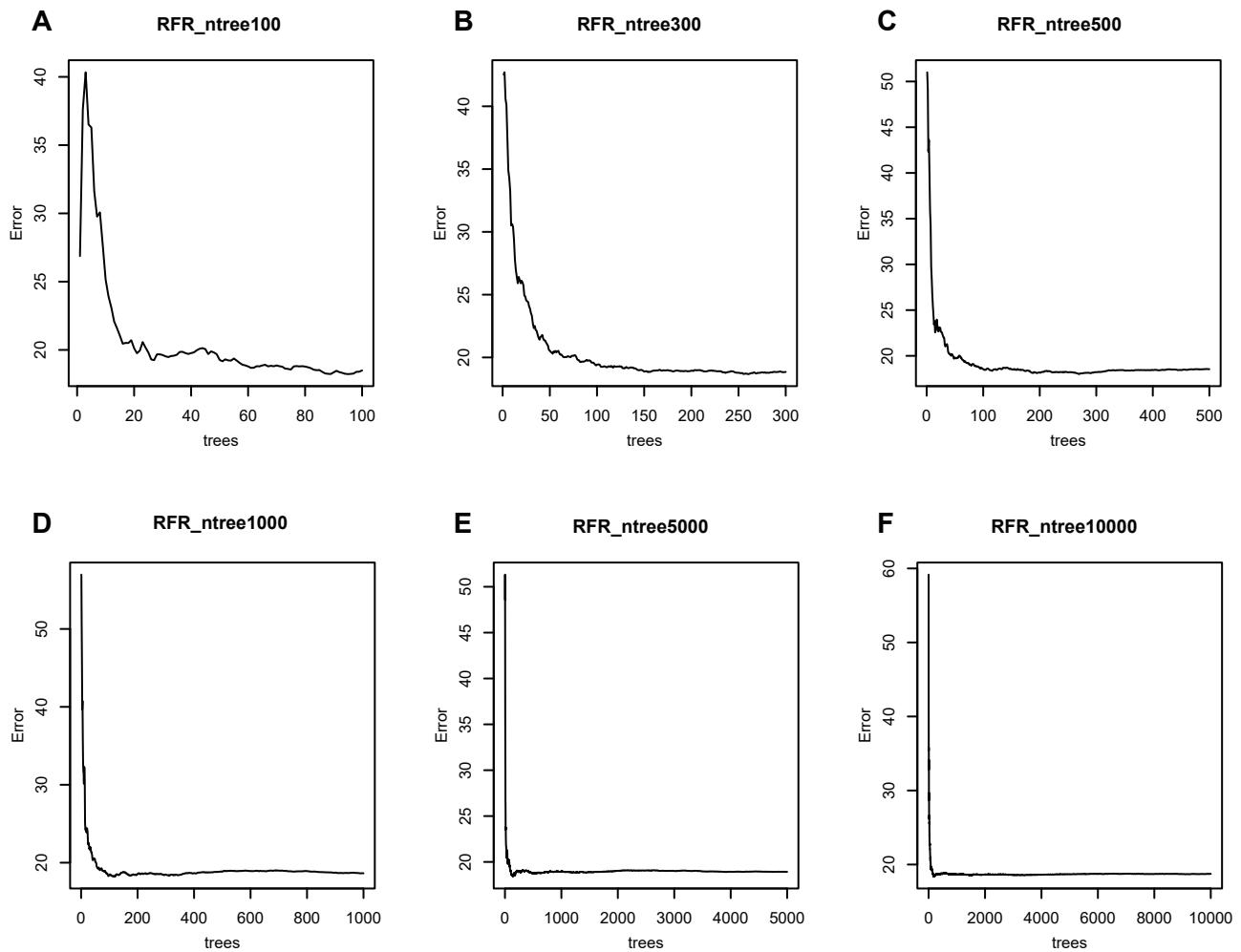
## Supplementary Figure S6



**Supplementary Figure S6** Spearman correlation analyses between different CpG methylation levels of KLF14 and the chronological ages of different gender datasets in CHS cohort ( $n = 240$ , blood samples).

**A. KLF14\_K1; B. KLF14\_K2; C. KLF14\_K3; D. KLF14\_K4;** Detailed CpG information in Supplementary Table S1.

## Supplementary Figure S7

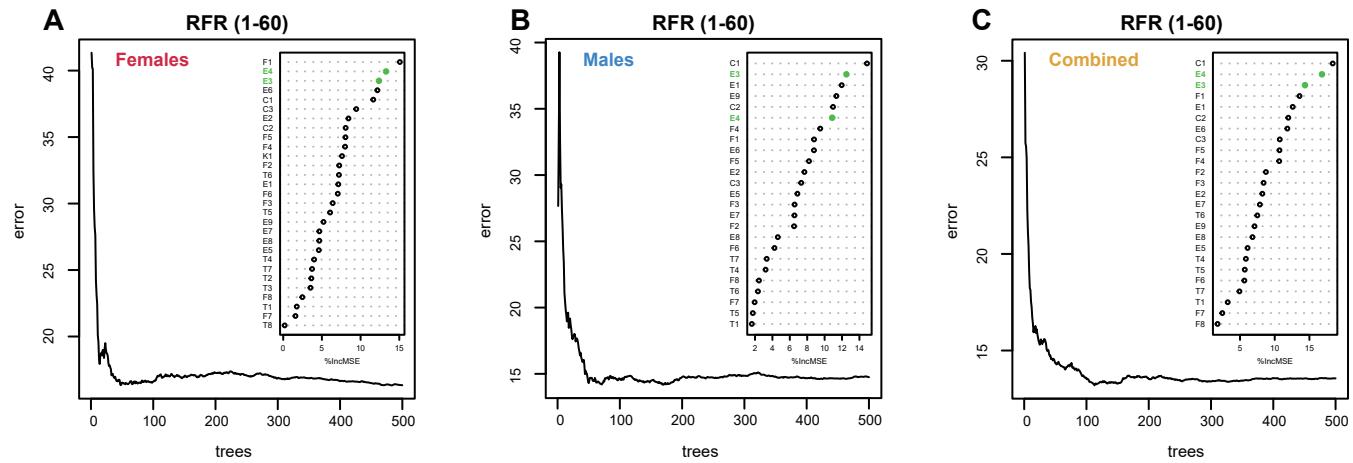


**Supplementary Figure S7** Error rates of random forest regression (RFR) model at six different *ntree* features (100-10000) in CHS cohort ( $n = 240$ , blood samples).

A.  $ntree = 100$ ; B.  $ntree = 300$ ; C.  $ntree = 500$ ; D.  $ntree = 1000$ ; E.  $ntree = 5000$ ; F.  $ntree = 10000$ .

(**ntree**, number of trees to grow, which should not be set to too small a number, to ensure that every input row gets predicted at least a few times.)

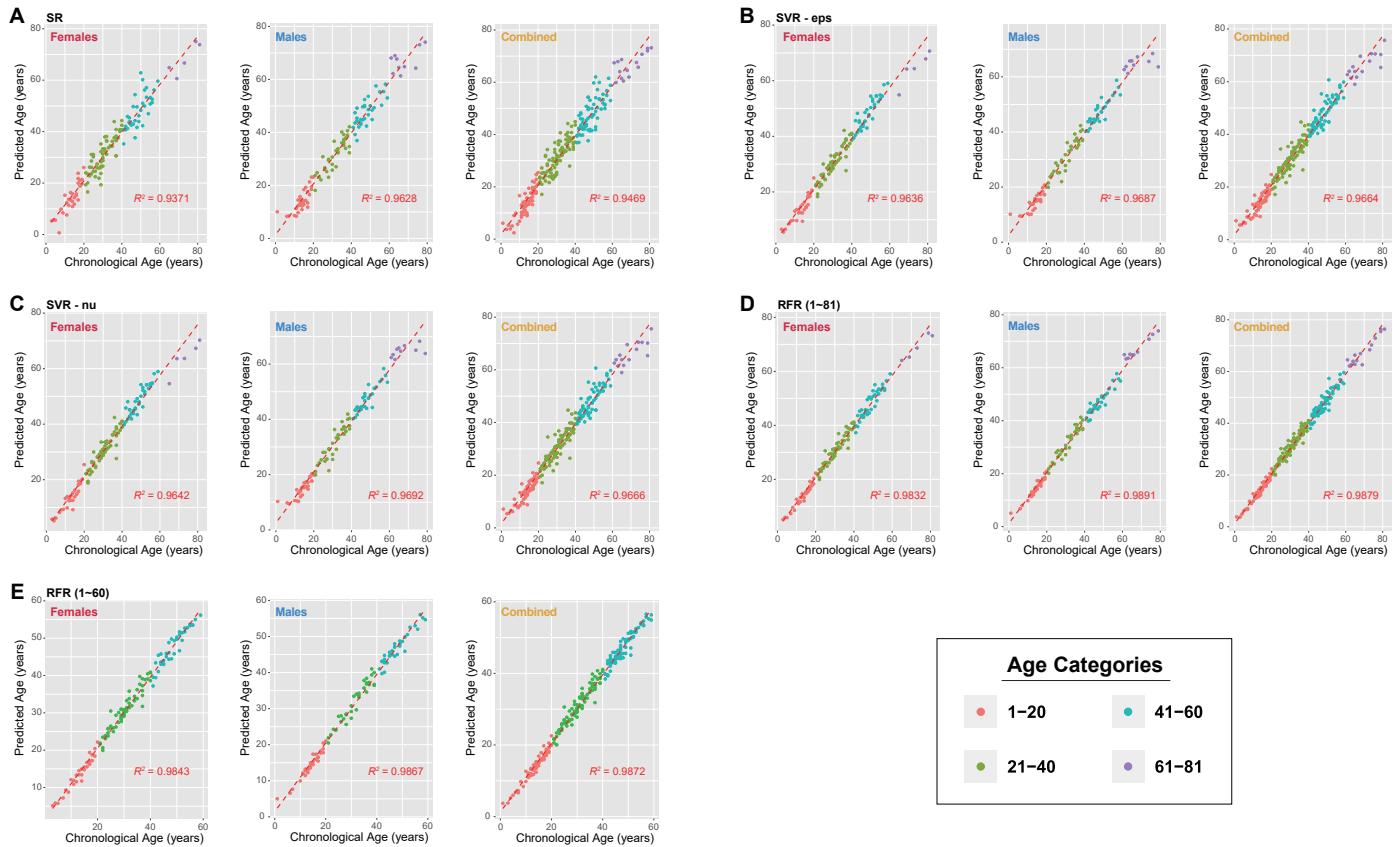
## Supplementary Figure S8



**Supplementary Figure S8** Error rates ( $n_{tree} = 500$ ) and AR-CpG importance ranking of random forest regression (RFR) model at three different gender datasets of 1-60 age categories (n = 225, blood samples).

**A.** Female dataset of 1-60 age categories (n = 127); **B.** Male dataset of 1-60 age categories (n = 98); **C.** Combined dataset of 1-60 age categories (n = 225). ( $n_{tree}$ , number of trees to grow; %IncMSE, increase in mean squared error.)

## Supplementary Figure S9



**Supplementary Figure S9** Linear relationships between predicted ages and chronological ages in different machine learning models. ( $R^2$ , coefficient of determination/goodness of fit)

- A. Stepwise Regression (SR) model;
- B. Support Vector Regression eps-regression (SVR-eps);
- C. Support Vector Regression nu-regression (SVR-nu);
- D. Random Forest Regression (RFR) at 1-81 age categories of CHS cohort;
- E. Random Forest Regression (RFR) at 1-60 age categories of CHS cohort.