



Chromosome-level genome and methylome of vine tea suggest roles for tandem duplication and CHH hypomethylation in high dihydromyricetin accumulation

Yingmei Wu¹, Hanrui Hu¹, Qing Ye¹, Xueqing Chen², Miyuan Tian³,
Jing Li⁴, Li Lu⁵, Yifei Liu¹, Yi Shang⁶, Yanling Li^{7*}, Xingyao Xiong^{7, 8*}, and Junbo Gou^{1*}

¹ Hubei Shizhen Laboratory, Hubei Key Laboratory of Resources and Chemistry of Chinese Medicine, College of Pharmacy, Hubei University of Chinese Medicine

² Shenzhen Huayi Xinchuang Biotechnology Co., Ltd

³ Laifeng County Rural Investment and Construction Co., Ltd

⁴ State Key Laboratory of Plant Diversity and Specialty Crops, Wuhan Botanical Garden, Chinese Academy of Sciences

⁵ State Key Laboratory of Hybrid Rice, School of Pharmaceutical Sciences, Wuhan University

⁶ Engineering Research Center for Valorization of Unique Bio-Resources in Yunnan, Ministry of Education, Yunnan Normal University

⁷ College of Horticulture, Hunan Agricultural University

⁸ Shenzhen Branch, Guangdong Laboratory for Lingnan Modern Agriculture, Shenzhen Key Laboratory

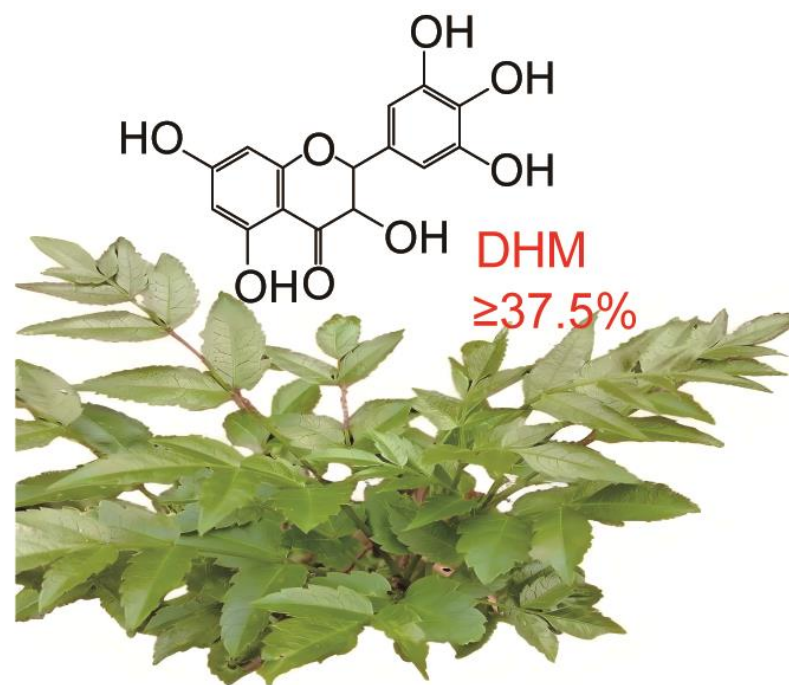


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Introduction



Nekemias grossedentata (Vine tea)

- ❑ **High medicinal value:** Young leaves contain up to 37.5% DHM and exhibit diverse bioactivities.
- ❑ **Strong research foundation:** Genome sequencing and biosynthetic gene variation analyses have been completed.
- ❑ **Key question:** The methylation and gene-duplication mechanisms underlying high DHM accumulation in young leaves remain unclear.

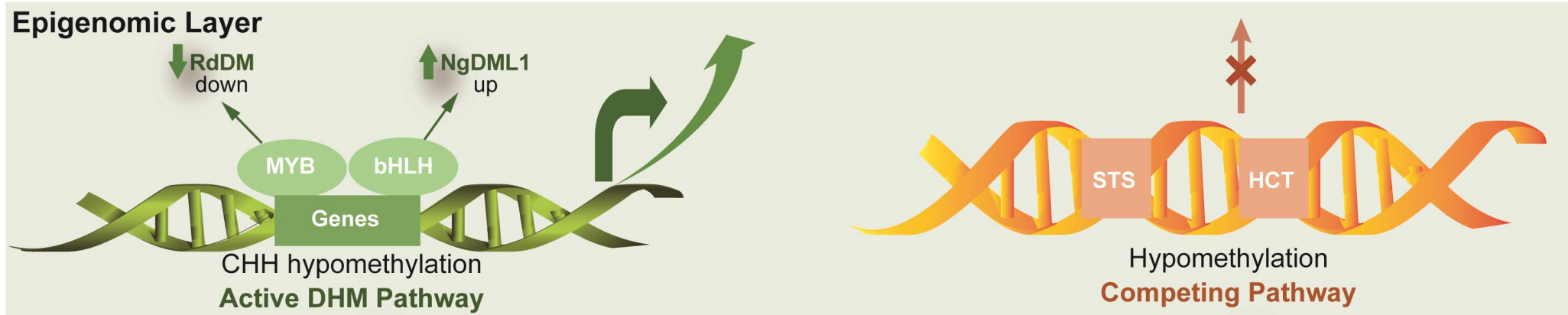


Highlights

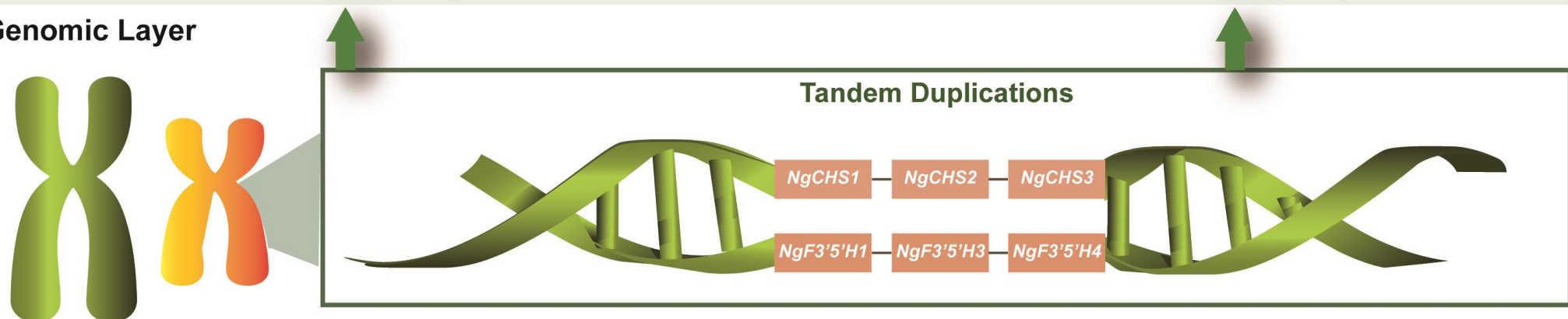
Metabolic Layer



Epigenomic Layer

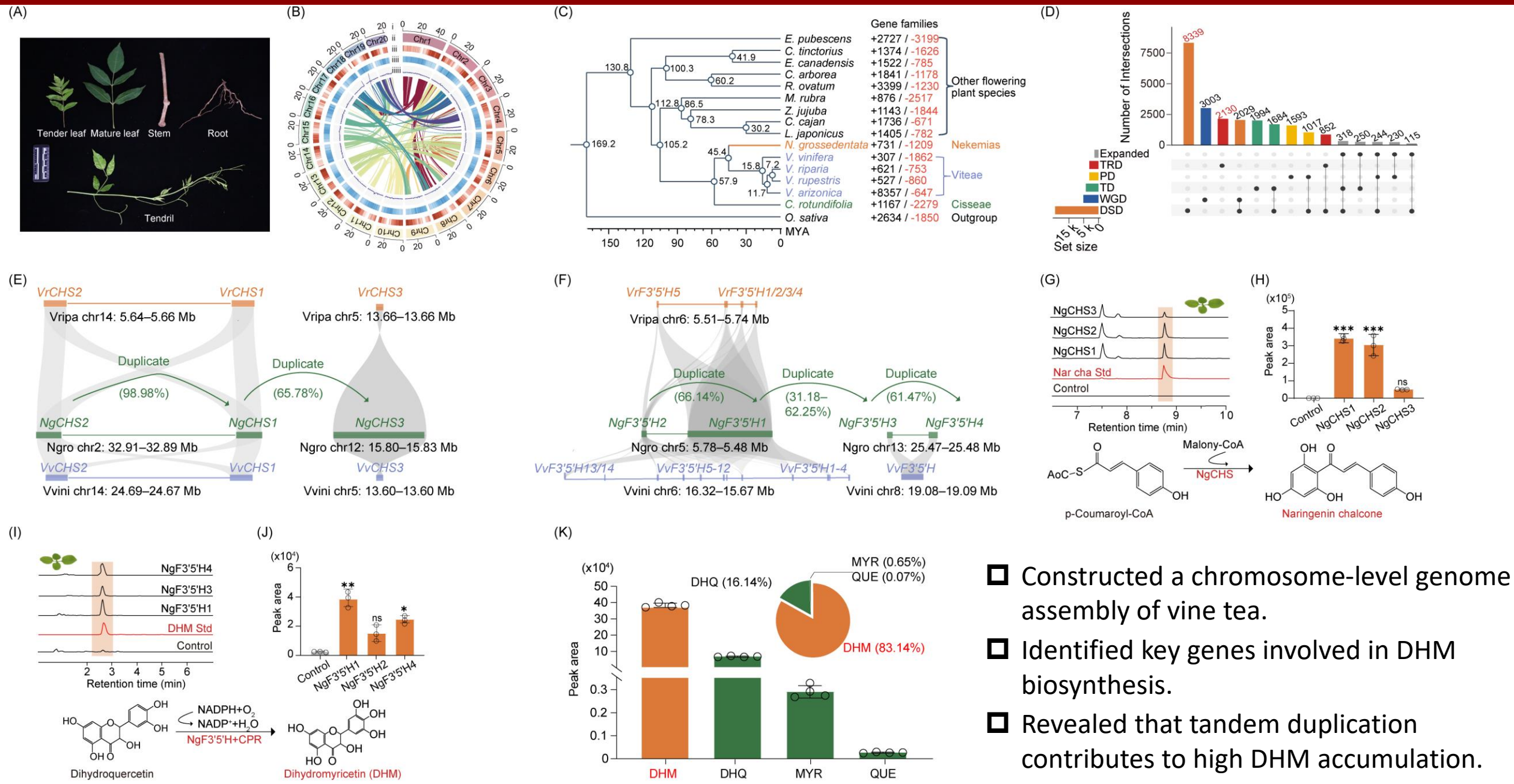


Genomic Layer

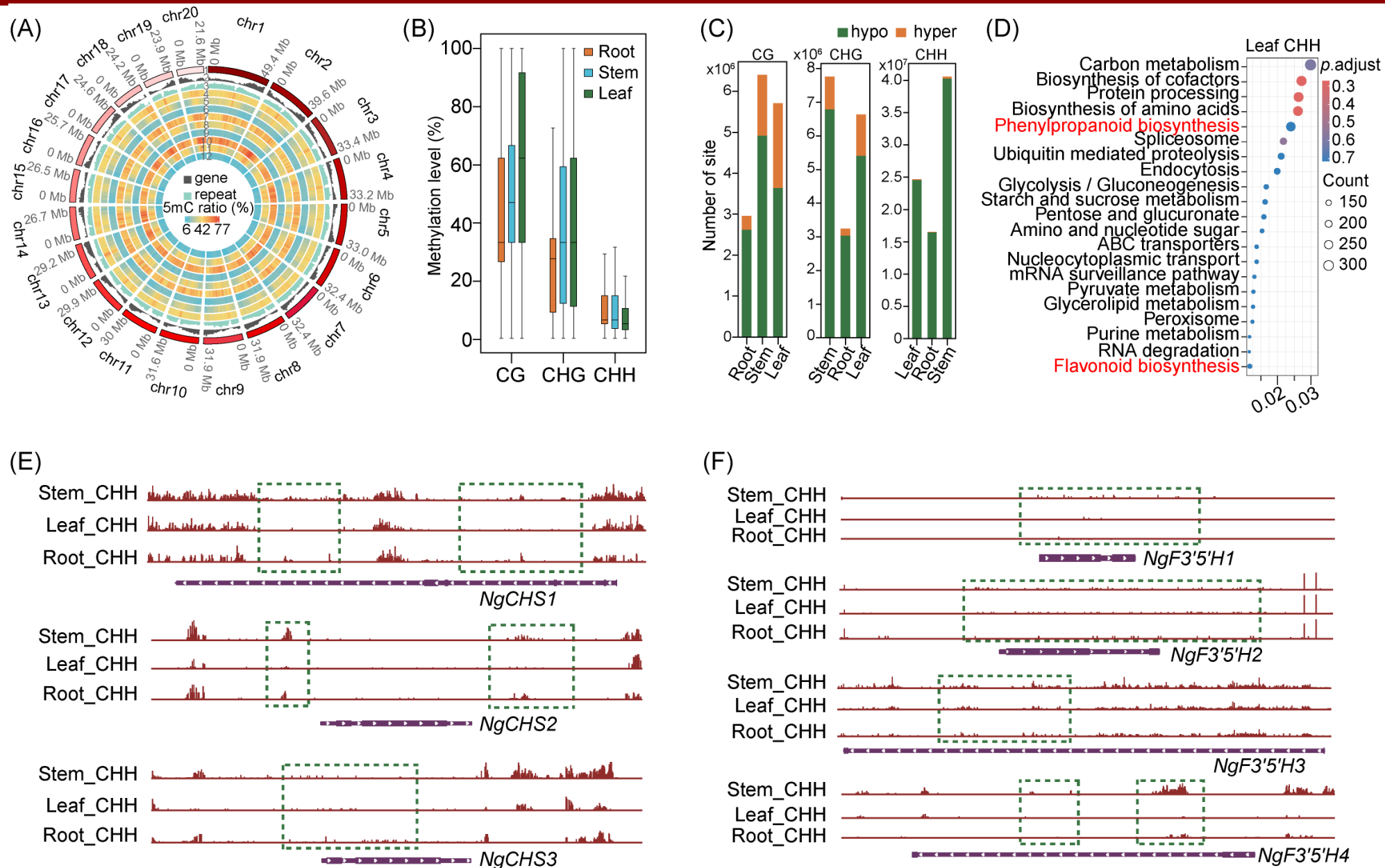




Chromosome-level genome assembly reveals tandem duplication of DHM biosynthetic genes



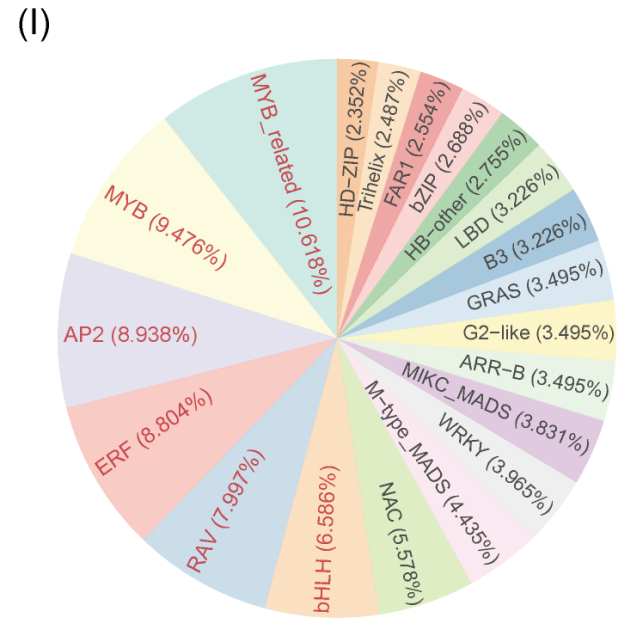
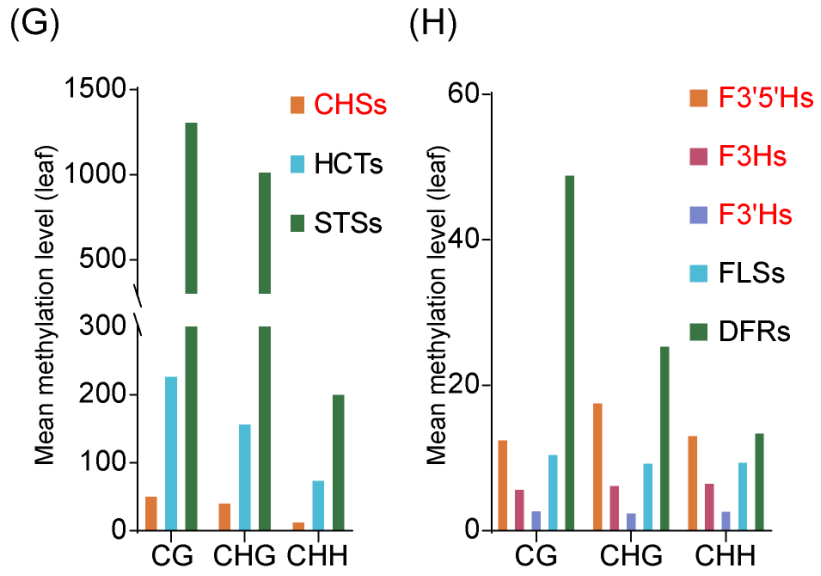
Tissue-specific CHH hypomethylation is associated with the activation of DHM pathway genes



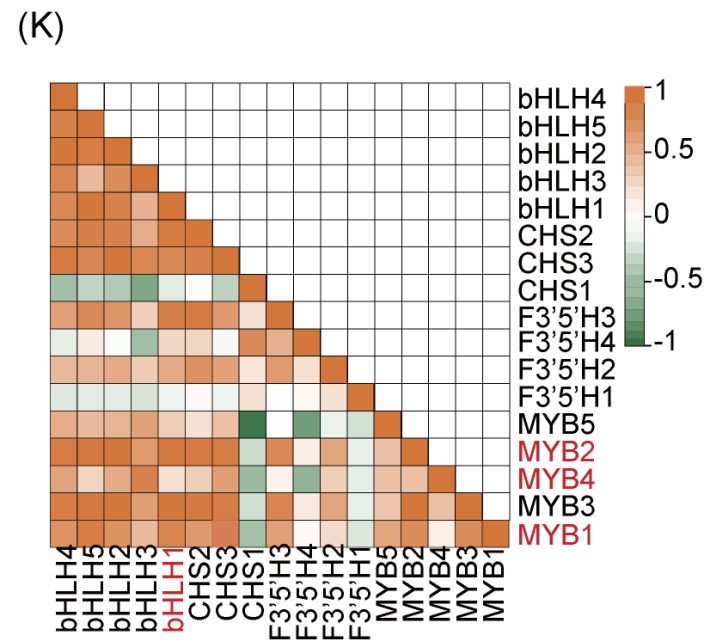
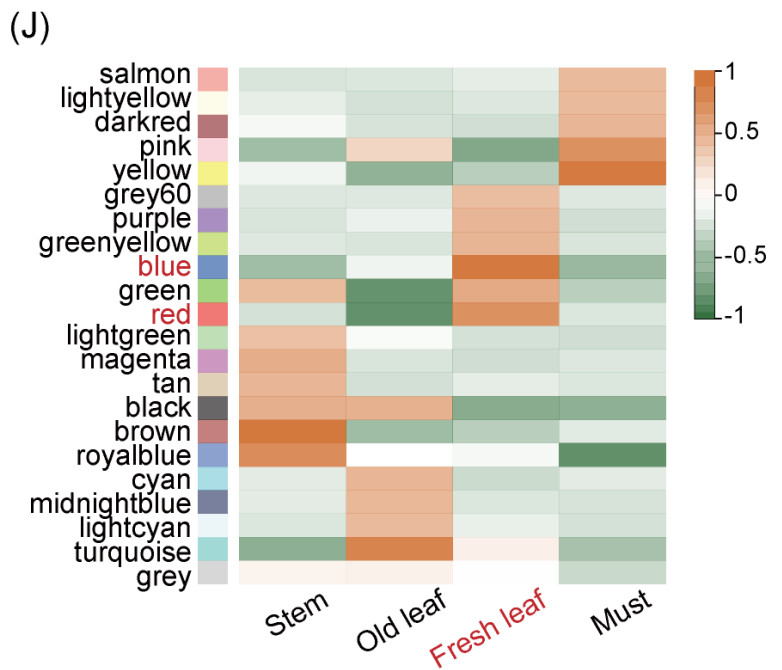
□ Young leaves show reduced CHH methylation.

□ CHH hypomethylation on activates DHM biosynthetic genes.

Tissue-specific CHH hypomethylation is associated with the activation of DHM pathway genes



- CHH hypomethylation activates DHM biosynthetic genes.
- Gene duplication and MYB/bHLH promote DHM accumulation.





Summary

- ❑ Genetic basis: Tandem duplication of CHS and F3'5'H supports high DHM biosynthesis.
- ❑ Key tissue: Young leaves show the strongest DHM biosynthetic activity.
- ❑ Epigenetic regulation: CHH hypomethylation may activate DHM pathway genes.
- ❑ Duplication-associated activation: Gene duplication, CHH hypomethylation, and MYB/bHLH factors jointly promote DHM accumulation.

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