

Genotypic differentiation of *Monilinia* spp. populations in Serbia using a High-Resolution Melting (HRM) analysis

ANTONIOS ZAMBOUNIS^{1*}, ELENI STEFANIDOU², PANAGIOTIS MADEIS^{2,3},
JOVANA HRUSTIĆ⁴, MILICA MIHAJLOVIĆ⁴, BRANKICA TANOVIĆ⁴

¹Institute of Plant Breeding and Genetic Resources, Department of Deciduous Fruit Trees,
ELGO-DEMETER, Naoussa, Greece

²Institute of Applied Biosciences, CERTH, Themi, Thessaloniki, Greece

³Laboratory of Molecular Biology of Plants, School of Agricultural Sciences, University of Thessaly,
Thessaly, Greece

⁴Laboratory of Phytopathology, Institute of Pesticides and Environmental Protection, Belgrade, Serbia

*Corresponding authors: antonios.zamb@gmail.com

ELECTRONIC SUPPLEMENTARY MATERIAL (ESM)

The authors are fully responsible for both the content and the formal aspects of the electronic supplementary material. No editorial adjustments were made.

Table S1. Sequences and amplicon lengths of the *Monilinia* species-specific primers used in this study

<i>Monilinia</i> spp.	Primer name	Primer sequence	Amplicon length
<i>Monilinia laxa</i>	ILaxaS	TGAGCACGAGTGAATGTATAG	226 bp
	ILaxaAS	TGAGCACGAGGGCATATC	
<i>Monilinia fructicola</i>	IColaS	GAGACGCACACAGAGTCAG	336 bp
	IColaAS	GAGACGCACATAGCATTGG	
<i>Monilinia fructigena</i>	IGenaS	TGCTCTGCCCCGTACCCAG	734 bp
	IGenaAS	GGATTATTGTGATGTAGTTTCG	

Table S2. Analysis of molecular variance for the three *Monilinia* spp. populations based on the HRM analysis

<i>Monilinia</i> spp.	Source	SS	MSS	Variance component	variation (%)
<i>Monilinia laxa</i>	among populations	2.604	1.302	0.06	7
<i>Monilinia fructicola</i>		6.000	3.000	0.423	44
<i>Monilinia fructigena</i>		4.418	2.209	0.425	44
<i>Monilinia laxa</i>	within populations	18.473	0.803	0.803	93
<i>Monilinia fructicola</i>		8.000	0.533	0.533	56
<i>Monilinia fructigena</i>		5.429	0.543	0.543	56
<i>Monilinia laxa</i>	total	21.077		0.863	
<i>Monilinia fructicola</i>		14.000		0.956	100
<i>Monilinia fructigena</i>		9.846		0.968	

SS –sum of squares; MSS – mean sum of squares

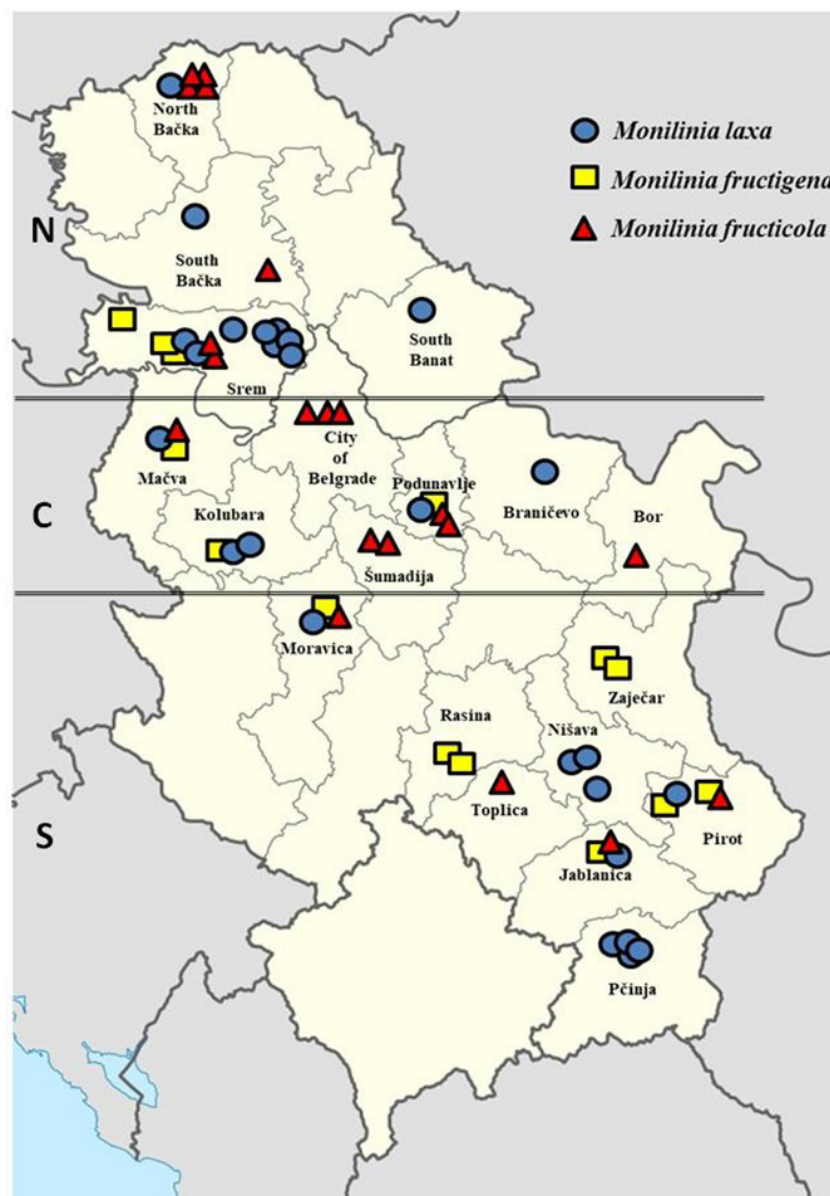


Figure S1. Geographical map indicating the sampling locations across the three geographical regions of Serbia

N – north; C – central; S – south

<https://doi.org/10.17221/35/2020-PPS>

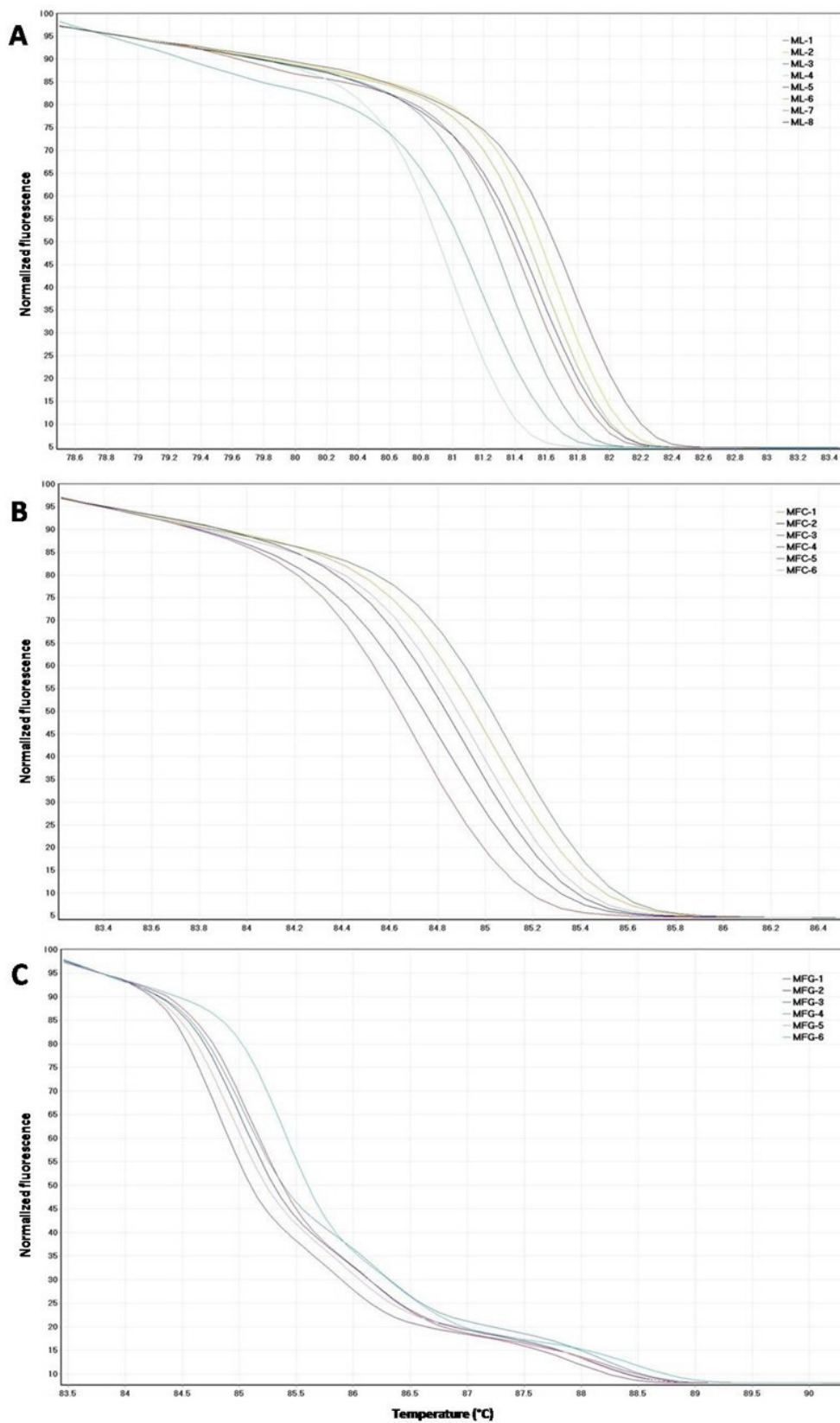


Figure S2. Discrimination of *Monilinia* spp. genotypes upon their normalized melting curves profiles generated by the HRM analysis of (A) *M. laxa*, (B) *M. fructicola* and (C) *M. fructigena*

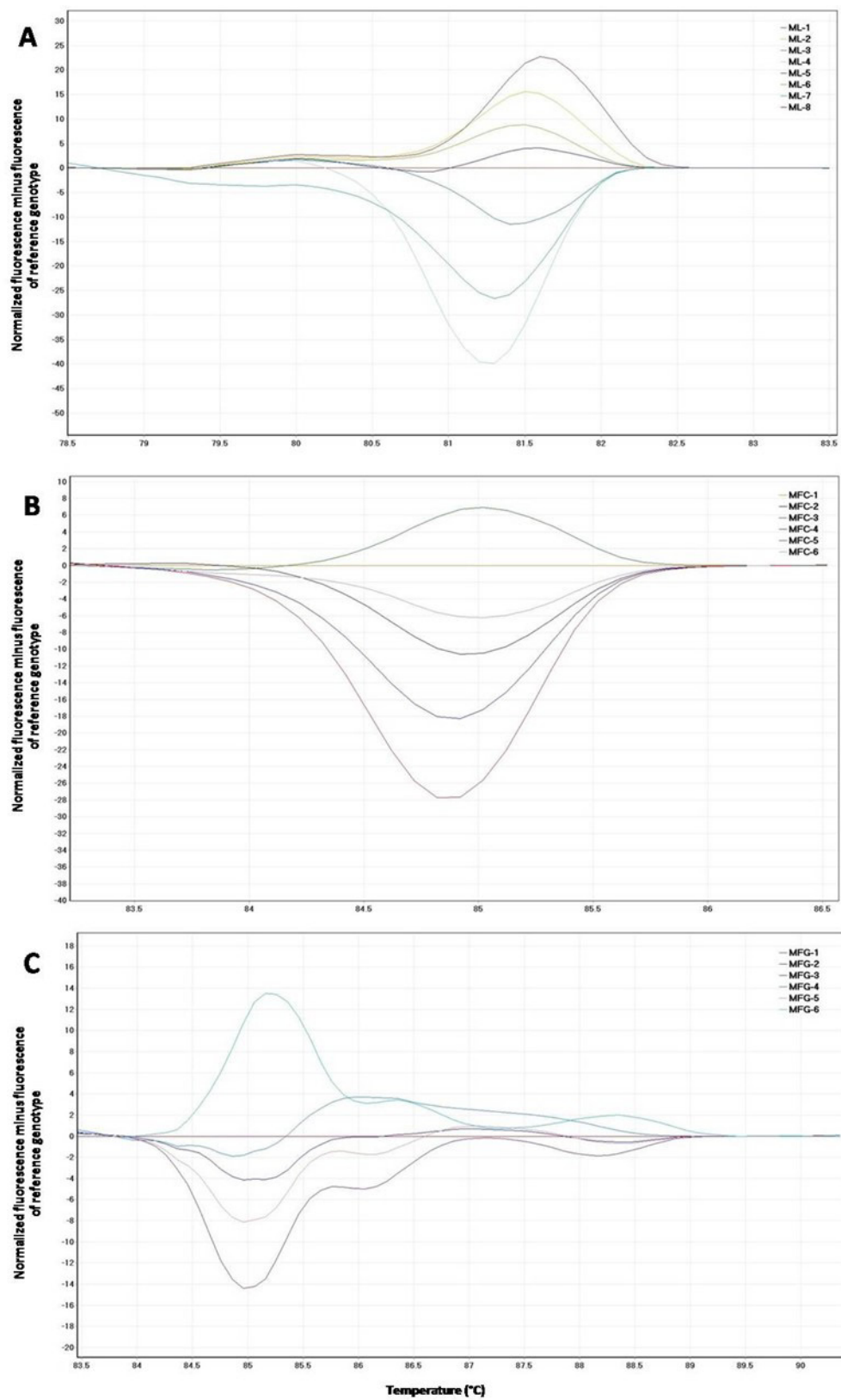


Figure S3. Difference graphs of the *Monilinia* spp. genotypes using reference genotypes converted to a horizontal lines upon the HRM analysis of (A) *M. laxa*, (B) *M. fructicola* and (C) *M. fructigena*

https://doi.org/10.17221/35/2020-PPS



Figure S4. Nucleotide alignments of the HRM genotypes for each *Monilinia* spp. *M. laxa* (ML⁻¹ up to ML⁻⁸)
The SNPs positions among the HRM genotypes are indicating with colored annotations

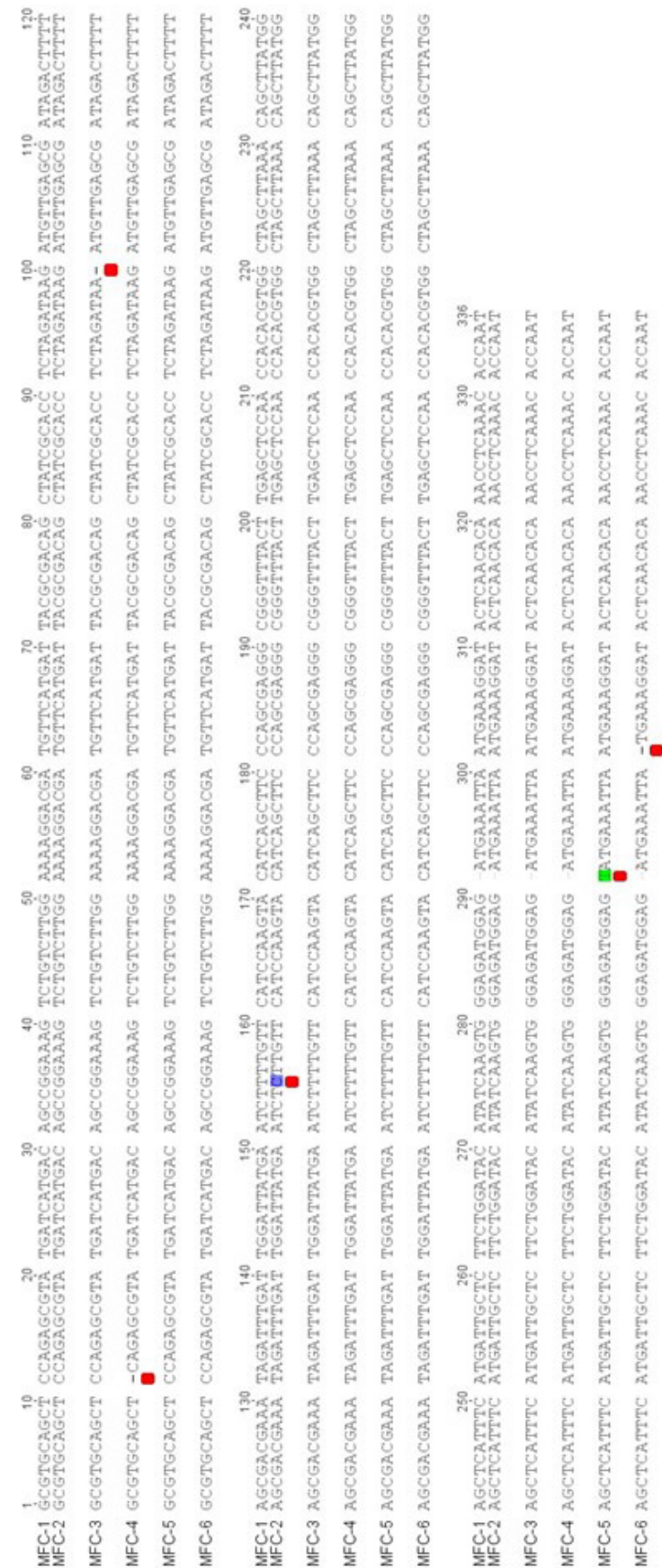


Figure S4: Nucleotide alignments of the HRM genotypes for each *Monilinia* spp. *M. fructicola* (MFC-1 up to MFC-6). The SNPs positions among the HRM genotypes are indicating with colored annotations

<https://doi.org/10.17221/35/2020-PPS>

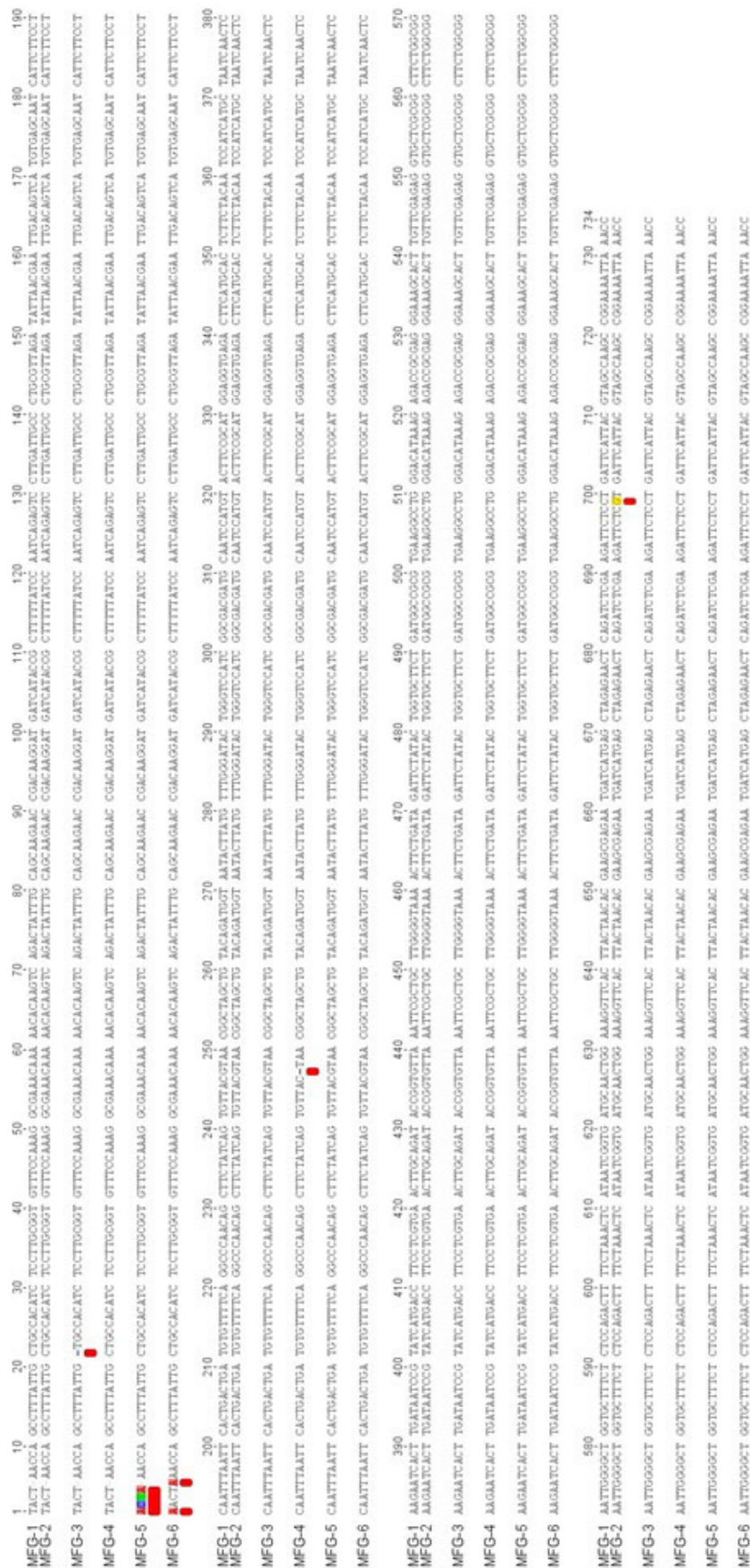


Figure S4. Nucleotide alignments of the HRM genotypes for each *Monilinia* spp. *M. fructigena* (MFG-1 up to MFG-6)

The SNPs positions among the HRM genotypes are indicating with colored annotations