

Figure S1. Agarose gel (1.0% w/v) electrophoresis for total RNA of representative samples (N=6). The presence of distinct ribosomal bands (28S, 18S, 5S from top to bottom) and the absence of degradation products shows that total RNA was intact, suitable for downstream analyses. .

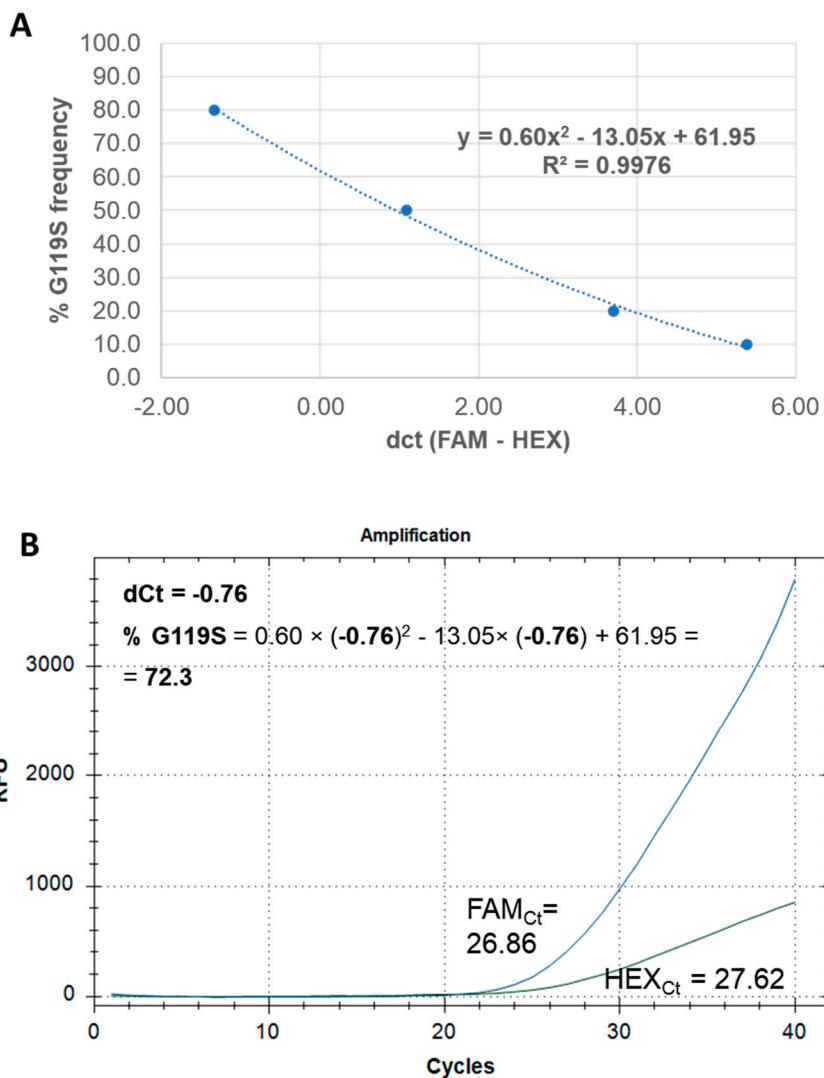


Figure S2. Standard curve for G119S using plasmid sequences with known % frequencies (A) and calculation of % G119S for an unknown population of the study as an example (B).

Table S1 List of primers and probes used for the multiplex resistance gene expression analyses.

Oligo	Assay	Sequence (5'-3')	Final Concentration (nM)
RPS7_F	Dtx (A)-(D)	CCACCATCGAACACAAAAGTTGA	100
RPS7_R	Dtx (A)-(D)	TGCTGCAAACCTCGGCTATT	200
RPS7_P	Dtx (A)-(D)	FAM-CCGTGACGTTACGTTCGAATTCCCA-BHQ1	250
CYP6P3_F	Dtx (A)	ACAATGTGATTGACGAAACCCT	400
CYP6P3_R	Dtx (A)	GGATCACATGCTTGTGCCG	500
CYP6P3_P	Dtx (A)	HEX-ACCCCGTACCGTCTGTGGACT-BHQ1	350
CYP6M2_F	Dtx (A)	CTGGCGTTGAATCCAGAGGT	600
CYP6M2_R	Dtx (A)	GATACTTGCGCAGTGATTCAATTAG	400
CYP6M2_P	Dtx (A)	ATTO647N-AGAGAAATCCTGCAAAAGCACAACGGAGA-BHQ3	250
CYP9K1_F	Dtx (B)	CCGACACGTGGTATGGATAC	200
CYP9K1_R	Dtx (B)	CGTCGTCGGTCCAGTCAAC	400
CYP9K1_P	Dtx (B)	HEX-CAATCTCTGATGCAGGCCGCAA-BHQ1	300
CYP6P4_F	Dtx (B)	CTGGACAACGTTATCAATGAAACC	400
CYP6P4_R	Dtx (B)	GCACGGTGTAAATCACCGCATC	500
CYP6P4_P	Dtx (B)	ATTO647N-CCGATCGAGTCACTTCGCGCG-BHQ3	300
CYP6Z1_F	Dtx (C)	CCCGCAACTGTATCGGTCTG	100
CYP6Z1_R	Dtx (C)	TTCGGTGCCAGTGTGATTGA	600
CYP6Z1_P	Dtx (C)	HEX-TGATGCTGTCCCGATTAACTTTCGGC-BHQ1	250
GSTE2_F	Dtx (C)	CCGGAATTGTGAAGCTAAACC	100
GSTE2_R	Dtx (C)	GCTTGACGGGTCTTCGG	400
GSTE2_P	Dtx (C)	ATTO647N-CGGTACGATCATACCGAGAGCCAC-BHQ3	300
CYP6P1_F	Dtx (D)	ACAGGTGGTGAACGAAACCC	100
CYP6P1_R	Dtx (D)	GGTGTAACTCTGTCCCGCAA	500
CYP6P1_P	Dtx (D)	HEX-CCGCTCGAAACGACGCTGCG-BHQ1	300
CYP4G16_F	Dtx (D)	GTCCAAGAAGTTGCGTCGGAC	200
CYP4G16_R	Dtx (D)	TCTTCGATTGCGTTGACGTG	200
CYP4G16_P	Dtx (D)	ATTO647N-CTGCAGGCCGACATCATTGAGC-BHQ3	300