

Table S2: Herein described new primers and PCR conditions

Gene locus	Primer name: sequence ^a	Amplicon size (bp)	PCR conditions
<i>PRP8</i>	fw: CTTCAGTCGTTTCCCCTG	560	Initial denaturation step (3min) at 95°C
	rev: TGCTGCTCTGTAGAAACACT		DNA denaturation step (15s) at 94°C
	fw FFPE: GAAGCCGATGAAATCCAGGG	111	Annealing step (30s) at 65°C (-0.7°C/cycle) for the next 12 cycles, thereafter at 56°C for 20 cycles
	rev FFPE: TATCAGGAGTGCCAACAGGT		Extension step (60s) at 72°C for 35 cycles (45 cycles for FFPE and fresh biopsy)
<i>CYP51pA</i>	fw: AGACTACCGTGTTTCTTGGA	646	Final extension step (5min) at 72°C
	rev: GCAATCTCGATGTCTGGGA		Initial denaturation step (3min) at 95°C
	rev FFPE: TCCATCAGCTTCGAGTTTGG	110	DNA denaturation step (15s) at 95°C
			Annealing step (30s) at 55°C
<i>CYP51pB</i>	fw: GCACGGCGACATCTTCAC	596	Extension step (60s) at 72°C for 35 cycles (45 cycles for FFPE and fresh biopsy)
	rev: CTGGCCGTTCTTGTAAGTGC		Final extension step (5min) at 72°C
	rev FFPE: TACTTGATGAACTTCTTCTGCT	170	Initial denaturation step (3min) at 95°C
			DNA denaturation step (15s) at 95°C

^a All primer sequences are in 5' to 3' configuration. fw: forward; rev: reverse; *PRP8*: *PRP8* intein; *CYP51pA*: cytochrome P450 enzyme lanosterol 14 α -demethylase A; *CYP51pB*: Cytochrome P450 enzyme lanosterol 14 α -demethylase B; FFPE: formalin-fixed paraffin-embedded samples)