

Supplementary Information

Fluorescence detection of single-nucleotide differences using aptamer-forming binary DNA probes.

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Fig. S1 Structure of deoxythymidine labeled with fluorescein

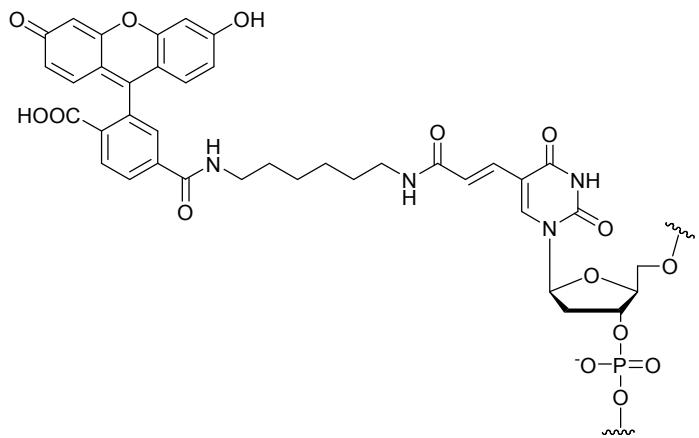
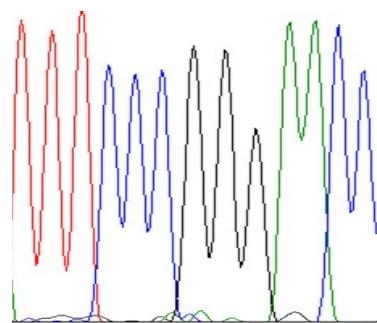


Fig. S2 Sequencing data for the PCR products amplified from human genomic DNA, which include CYP-681 (a) or CYP-636 (b). Data around the SNP sites are shown. Nucleotides with SNP are shown in capital letters.

(a) CYP-681: t t a **t t t c c c G g g a a c c c a t a**



(b) CYP-636: g c a c c c c **t g G a t c c a g a t a**

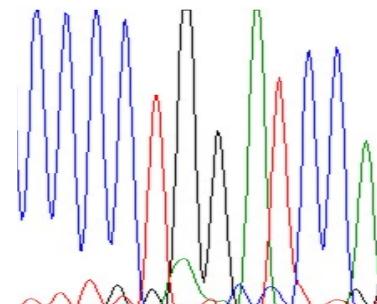


Table S1 DNA sequences used in this study. “T” in bold indicates deoxythymidines labeled with fluorescein.

DNAs	Sequence
CYP-681 (X = G, A, T, or C)	5' -TTATTTCCC X GGAAACCCATA-3'
BP 1 (Y = T or C)	5' -CCAGTTGG T GYGGGAAATAA-3'
BP 2	5' -TATGGGTTCCCACCAACTGG-3'
CYP-636 (X = G, A, T, or C)	5' -GCACCCCC X TCCAGATA-3'
BP 3	5' -CCAGTTGG T GCAGGGGTGC-3'
BP 4 (Y = T or C)	5' -TATCTGGATYCACCAACTGG-3'
PCR primer for 96mer target containing CYP-681 (forward)	5' -TATGCAATAATTTC CC ACTATC-3'
PCR primer for 96mer target containing CYP-636 (reverse)	5' -TCTCCAAAATATCACTTTCCATAA-3'
PCR primer for 81mer target containing CYP-636 (forward)	5' -GAATGAAA AC ATCAGGATTGTAA-3'
PCR primer for 81mer target containing CYP-636 (reverse)	5' -ACTGTAAGTGGTTCTCAGGAAG-3'

Table S2 Sequences of ssDNAs amplified from *CYP2C19* gene by an asymmetric PCR method.

These sequences were determined by a standard dideoxy method. The sequences of CYP-681 and CYP-636 are shown in bold letters. Nucleotides with SNP are shown in red letters. The PCR primer regions are underlined.

ssDNA target	Sequence
96mer target containing CYP-681	5' – <u>TATGCAATAATTTC</u> CCACTAT <u>CATTGA</u> TTATTTCCC GGAAACCC CATAA CAAATTACTAAAAACCTTGCTT <u>TATGGAAAGTGATATTTGGAGA</u> –3'
81mer target containing CYP-636	5' – <u>GAATGAAAACATCAGGATTGTAA</u> GCACCCCC TG GA TCCAGGT A AGGCCAAGTTTGCTTCCTGAGAACCA <u>CTTACAGT</u> –3'