

## CORRIGENDUM

## A novel method of identifying genetic mutations using an electrochemical DNA array

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The authors would like to apologize for the omission of Tomohiro Urata, from TUM Gene, Inc., 3-1 Kazusa-Koito Kimitsu, Chiba 292-1149, Japan, from the author list of this paper.

The authors would also like to apologize for two incorrect probes listed in Table 1. The 10th and 14th probes should be A261T-M and W382X-M instead of A261T-W and W382X-W, respectively. The complete corrected Table 1 is given below.

**Table 1.** Primers (a and b) and probe (c) sequences

(a) First PCR primer	
Ex3-FP	CTGTGCCAATGGGTTTCCA
Ex3-RP	CACTGTTTTGGACACATAAGTCTC
Ex5-FP	GAAATTTACAAATCTGTGTTCTGCT
Ex5-RP	CATTGGGTCAATAAGGGTTAAGGA
Ex6-FP	AGACATGCCAAATGAAACACTCT
Ex6-RP	ACTCCTTGGTTTCCTTATTTACAACA
Ex7-FP	TTCATAAAGATTGATCAACATGTTCTGA
Ex7-RP	ACTGGTGCCATGATGACCG
Ex8-FP	GAGAGCTGATCTCTATAACTAACCA
Ex8-RP	CTCTGATCTTCTGAATGGCGA
(b) Second PCR primer	
Y61X-CP	TAGGTGGGTATTTAAGAAAGCTTGT
Y61X-SP	<u>GAGAGTTGGGTGCCTCTCTTGTACAGGGCGGCC-ACAAG</u>
V200A-CP	GTAGACGTCTTACACACATTCACCAGAGGG
V200A-SP	<u>TGGGCATGTTGACATCTGGCTGAAAAGTACCTCC-ATTCGGG</u>
A221-del-CP	CAGTTGGGCATGTTGACATTTACCCG
A221-del-SP	<u>CTATCCGCGTGATTCTTCTAAATAATATTTACCTCC-AAGTCTCTCTCTGC</u>
R243C-CP	GCACCTGTAGGCCTTACTTGGATTTTCT
R243C-SP	<u>CTCGTGGGAGCACACCCAGATGTGGACCAGCTAG-TGAA</u>
A261T-CP	CCCACGAGCGCTCCATTCATCTCT
A261T-SP	<u>CCTACAGGTGCAGTAGAGCCCTTTCTCAAAGGCTT-CCTGG</u>
A334T-CP	CCTCCCAACAGTCTTCCATTACCAAGTAAAG
A334T-SP	<u>CCTTTGAGATTTCTCTGGGATGTTCTCACTCTCGGC-CACGGTGCCAT</u>
W382X-CP	AGACCTACTCCTCCTAATTTACACAGAGGTAG
W382X-SP	<u>AAGAGTGATTCATACTTTCTGCTCCACCAGTCTGAC-CAGC</u>

**Table 1.** Continued

(c) Probe	
Y61X-W probe	AAAGCTGTGTGCATCATCTTCAGGTAACAGGA-ATGTAT
Y61X-M probe	AAAGCTGTGTGCATCATCTTCAGGTAACAGGA-ATGTAA
V200A-W probe	GGGTCCCCTGGTCGAAGCATTGGAATCCAGAA-ACCAGT
V200A-M probe	GGGTCCCCTGGTCGAAGCATTGGAATCCAGAA-ACCAGC
A221-del-W probe	TGGAGGTACTTTTCAGCCAGGATGTAACATTGG-AGAAG
A221-del-M probe	ATGGAGGTACTTTTCAGCCAGGATGTAACATTGG-GAGAA
R243C-W probe	TCATTCAACAGAGAGTCGATGAAGAGATGAATG-GAGCG
R243C-M probe	TCATTCAACAGAGAGTCGATGAAGAGATGAATG-GAGCA
A261T-W probe	CATCGACTCTCTGTTGAATGAAGAAAATCCAAG-TAAGG
A261T-M probe	CATCGACTCTCTGTTGAATGAAGAAAATCCAAG-TAAGA
A334T-W probe	TTTTTCTGGGACTGAGAGTGAAACCCATACCAA-TCAGG
A334T-M probe	TTTTTCTGGGACTGAGAGTGAAACCCATACCAA-TCAGA
W382X-W probe	TAGATATTGGAGAACTACTCATGTTGAAGCTCA-AATGG
W382X-M probe	TAGATATTGGAGAACTACTCATGTTGAAGCTCA-AATGA

In (a) FP denotes 'forward primer' and RP denotes 'reverse primer'. In (b) CP and SP indicate 'counter primer' and 'special primer', respectively. The underlined nucleotides in the SP are the tag sequences essential to make a stem structure in a self-loop formation (cf. Figure 3D). In (c) W probe and M probe indicate 'wild type probe' and 'mutant type probe', respectively.