Figure S1. Detection of *EGFR* mutations using ddPCR in lung cancer cell lines harboring *EGFR* mutations. (A and B) ddPCR using *EGFR* L858R primer sets. *EGFR* L858R was amplified in the serially diluted DNA of H3255 cells with *EGFR* L858R. (C and D) ddPCR using *EGFR* exon 19 deletion primer sets. *EGFR* exon 19 deletion was amplified in the serially diluted DNA of RPC-9 cells harboring *EGFR* exon 19 deletion and T790M mutations. (E and F) ddPCR using *EGFR* T790M primer sets. *EGFR* T790M was amplified in the serially diluted DNA of RPC-9 cells harboring *EGFR* exon 19 deletion and T790M mutations. (E and F) ddPCR using *EGFR* T790M primer sets. *EGFR* T790M was amplified in the serially diluted DNA of RPC-9 cells harboring *EGFR* exon 19 deletion and T790M mutations. (EGFR exon 19 deletion and T790M mutations. EGFR exon 19 deletion and T790M mutations.



Figure S2. ddPCR in negative controls. (A) ddPCR in elution buffer or EBC samples from healthy volunteers. (B) ddPCR using primer set for *EGFR* L858R. No positive droplets were observed in elution buffer or EBC samples from healthy volunteers. (C) ddPCR using *EGFR* exon 19 deletion primer sets. Positive droplets were observed in elution buffer and EBC samples from healthy volunteers. (D) ddPCR using *EGFR* T790M primer sets. No positive droplets were observed in elution buffer or EBC samples from healthy volunteers. (D) ddPCR using *EGFR* T790M primer sets. No positive droplets were observed in elution buffer or EBC samples from healthy volunteers. (D) ddPCR using *EGFR* T790M primer sets. No positive droplets were observed in elution buffer or EBC samples from healthy volunteers. ddPCR, droplet digital PCR; EBC, exhaled breath condensate; Ex19del, *EGFR* exon 19 deletion; HD802, EGFR gene-specific multiplex gDNA reference standard.



Figure S3. ROC analysis to determine the threshold for epidermal growth factor receptor exon 19 deletion. ROC, receiver operating curve.

