Supplementary Table S1. Diagnostic value of KRA^{G12/13} mutation detection by ddPCR, Sanger sequencing, and PNA-clamping assay, and NGS panel sequencing (IonS5, Thermo Fisher Scientific).

	Detection of KRAG12/13 Mutation			
	ddPCR	Cancarcaguancina	DNIA clamping access	NGS
	uurck	Sanger sequencing	PNA-clamping assay	panel sequencing
Sensitivity	96.43%	100%	89.29%	39.29%
Specificity	98.11%	92.45%	84.91%	88.68%
PPV	96.43%	87.50%	75.76%	64.71%
NPV	98.11%	100%	93.75%	73.44%

PPV, positive predictive value; NPV, negative predictive value. * For comparative analysis, cases in which $KRAS^{G12/G13}$ mutation was detected by three or more methods were defined as positive references.