

A modified Phenol-chloroform extraction method for isolating circulating cell free DNA of tumor patients

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Abstract

Searching for new cancer biomarkers, circulating cell-free DNA (cfDNA) has become an appealing target of interest as an elevated level of cfDNA has been detected in the circulation of cancer patients in comparison with healthy controls. Since cfDNA can be isolated from the circulation and other body fluids of patients without harming their physical condition, cfDNA is becoming a promising candidate as a novel non-invasive biomarker for cancer. The challenge in the diagnostic analysis of cfDNA is its very low presence in human plasma/serum and its partially strong fragmentation. Here we evaluated a modified phenol/chloroform extraction method for the isolation of cfDNA and compared it with published standard methods for cfDNA isolation.

Protocol

Searching for new cancer biomarkers, circulating cell-free DNA (cfDNA) has become an appealing target of interest as an elevated level of cfDNA has been detected in the circulation of cancer patients in comparison with healthy controls. Since cfDNA can be isolated from the circulation and other body fluids of patients without harming their physical condition, cfDNA is becoming a promising candidate as a novel non-invasive biomarker for cancer.¹⁻³

The challenge in the diagnostic analysis of cfDNA is its very low presence in human plasma and its partially strong fragmentation.²

Here we evaluated a modified phenol/chloroform extraction method for the isolation of cfDNA in human plasma and compared it with published standard methods for cfDNA isolation.⁴ Although cfDNA was about 5-fold higher in serum samples compared to plasma (data not shown).⁵ We only show cfDNA levels determined from plasma samples which allows a better comparison with previously published protocols.⁶⁻⁸ Therefore, plasma samples from healthy individuals (n=10) and patients with colon carcinoma (n=15) and breast cancer (n=15) were analyzed in parallel using our modified phenol/chloroform method in comparison to the Maxwell[®] 16 LEV

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Figure 1. Control gene PCR for the assessment of amplifiability and integrity of DNA samples. Five control genes exons and the five primer sets for obtaining PCR products of 100, 200, 300, 400, and 600 bp⁹. 100bp ladder marker, samples from cancer patients 1-4; healthy controls 5, 6 and no template control, primer used as described in van Dongen *et al.* 2003.

Table 1. Results obtained by real-time PCR quantification of cfDNA, using hTERT obtained from 1 mL of plasma, respectively (mean \pm std. error).

Methods	Healthy individuals pg/ μ L	Patients with carcinoma pg/ μ L	P value
Maxwell [®] 16 LEV DNA Purification Kit	0.89 \pm 0.33	nd	0.0001
QIAamp [®] DNA Mini and Blood Mini kit	1.22 \pm 0.80	nd (13-309*)	0.0001
NucleoSpin [®] Plasma XS8	1.87 \pm 0.99	nd (10-423*)	0.0001
Modified Phenol-chloroform extraction	86.91 \pm 13.04	755.4 \pm 199.6	
Phenol-chloroform extraction by Schmidt <i>et al.</i> ⁶	23.78 \pm 3.91	190.3 \pm 52.14	0.008 (healthy) 0.035 (carcinoma)

P-values indicate significance of results in comparison to the modified phenol-chloroform extraction. (nd: not determined). *Data according to publications; Silva *et al.* 2002, Stemmer *et al.* 2003, Xie *et al.* 2004, Lui *et al.* 2001, Lecomte *et al.* 2002, Gal *et al.* 2004, Gautschi *et al.* 2004, Di *et al.* 2003, Ito *et al.* 2003.

DNA Purification Kit (Promega), QIAamp® DNA Mini and Blood Mini kit (Qiagen), NucleoSpin® Plasma XS (Macherey-Nagel)⁸ and a published Phenol-chloroform extraction method.⁶

The samples were processed to obtain plasma within an hour after the withdrawal of blood from donors. The plasma was obtained by two centrifugations of whole blood EDTA-tubes

at 1500 g at 4 °C for 10 min and at 3000g at 4°C for 10min, and was then preserved at -80°C. To 1 mL of plasma, 100 µL of a solution containing 250 mmol/L EDTA and 750 mmol/L NaCl, 100 µL of 100 g/L sodium dodecyl sulfate and 20 µL of proteinase K (final concentration 20 mg/mL) was added. The samples were incubated for 2 hours at 56°C, and the proteins were precipitated with 200 µL of saturated 6M NaCl solu-

tion (final concentration, 0.86 mol/L). The cfDNA was extracted with a 1:1 phenol-chloroform mixture at room-temperature. After incubation time of 5 min at room temperature the solution was centrifuged for 15 min at 14,000 g. The cleared supernatant was transferred into a new tube and the DNA was precipitated by adding the same volume of absolute ethanol and incubating overnight at -20°C. The DNA was first centrifuged for 15 min at 14,000 g, then washed with 70% ethanol and dissolved in 50 µL water. We used 5 µL of template DNA for quantification, and each sample was analyzed in duplicate. The DNA was quantified by a real-time PCR analysis using the sequence of hTERT, the human telomerase reverse transcriptase.⁷ The amplicon size of the hTERT single copy gene was 98 bp. As shown in Table 1, our modified cfDNA extraction method was superior to all tested kit-based standard extraction methods and led also to a DNA yield which was about four times higher compared to the recently published method by Schmidt *et al.*,⁶ which is similar to our presented protocol. However, Schmidt *et al* describe the isolation of cfDNA within 2 working days, whereupon we decided to set up a protocol aiming at completing cfDNA purification within only one working day. To achieve this, we performed a single DNA precipitation, whereas Schmidt *et al* performed two consecutive rounds of precipitations. As DNA gets lost with every round of precipitation, omitting the second precipitation step may at least in part explain the higher amount of cfDNA gained by our protocol. In addition, the slightly different salt concentration used for cfDNA precipitation may also account for the higher efficiency of our method.

Regarding the fragment-length of the isolated cell free DNA we were able to detect fragments from 100 pb up to 600 bp via PCR,⁹ isolated with all methods but noticed a higher fragmentation of cfDNA in patients compared to healthy controls (Figure 1 and Table 2).¹⁰

However, the presence of sequences shorter than 98 bp as early marker for the attendance of tumors was not examined in this study (Moulière *et al.* personal communication).

Regarding the clinical aspects we were able to detect tumor-specific mutations (KRAS codon 12-13 and BRAF V600E point mutations)¹¹⁻¹⁴ and we found significantly increased levels of cell-free DNA in patients compared to controls, which is in line with former published data (Figures 2 and 3).^{3,15-20} In summary, we were able to show a simple and robust method for extraction, isolation and analysis of cfDNA, suitable for a routine clinical-oncology laboratory. In addition, this method was suitable for further PCR-based characterization sequencing applications.

Table 2. Control gene PCR for the assessment of amplifiability and integrity of DNA samples. Five control genes exons and the five primer sets for obtaining PCR products of 100, 200, 300, 400, and 600 bp⁹. PCR positive number of patients in comparison to total number of samples for controls and patients set in parentheses.

	PCR product control samples	PCR positive PCR positive patient samples
100 bp	10 (10)	27 (30)
200 bp	10 (10)	26 (30)
300 bp	10 (10)	24 (30)
400 bp	10 (10)	23 (30)
600 bp	10 (10)	21 (30)

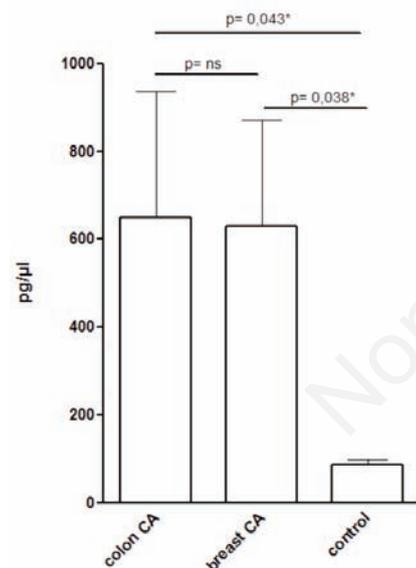
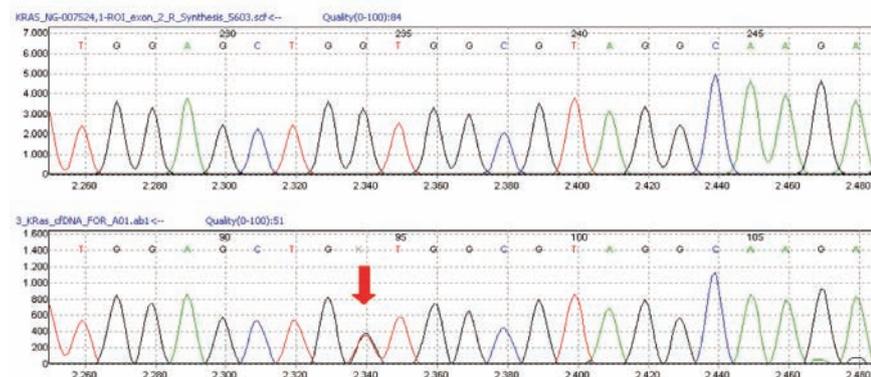


Figure 2. A) Comparison of cfDNA levels obtained from plasma from colon-CA patients (n=15), breast-CA patients (n=15) and healthy controls (n=10), P-values indicate significance of results. B) Results obtained by sequencing the K-ras gene in cfDNA. The same mutation (red arrow) as in the primary tumour was found in the cfDNA. (K-Ras codon 12) of patient with colorectal cancer; PCR and sequencing performed as described in Karapetis *et al.* 2008.¹⁹



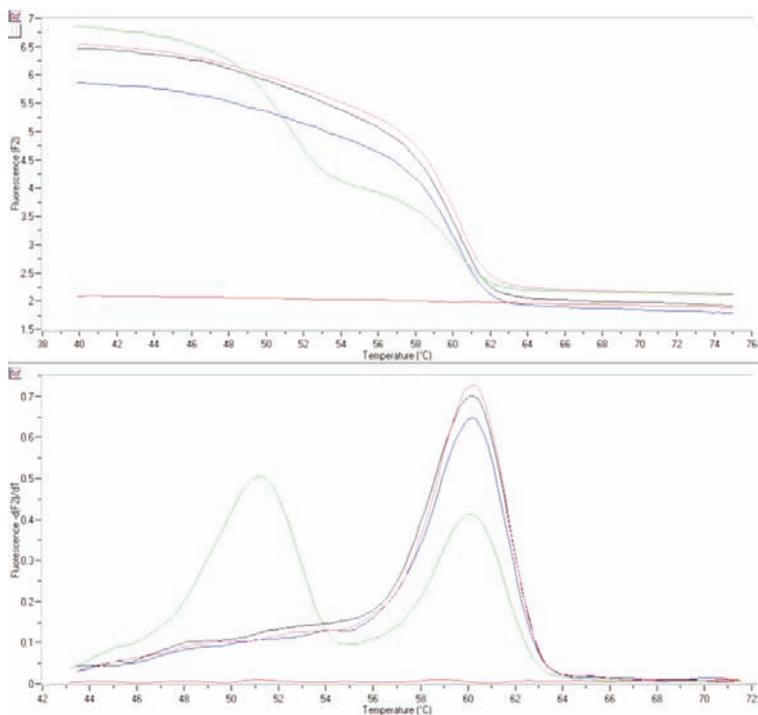


Figure 3. Results obtained by real-time PCR of B-Raf V600E point mutation in cfDNA of patient with breast cancer (same mutation in primary tumour). Melting curve analysis of a B-Raf V600E-mutated patient (green graph) with hybridization probe, PCR and detection method performed as described in Nikiforova et al 2003.²⁰

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