This is an Accepted Manuscript, which has been through the Royal Society of Chemistry peer review process and has been accepted for publication.

Accepted Manuscripts are published online shortly after acceptance, before technical editing, formatting and proof reading. Using this free service, authors can make their results available to the community, in citable form, before we publish the edited article. We will replace this Accepted Manuscript with the edited and formatted Advance Article as soon as it is available.

You can find more information about Accepted Manuscripts in the Information for Authors.

Please note that technical editing may introduce minor changes to the text and/or graphics, which may alter content. The journal’s standard Terms & Conditions and the Ethical guidelines still apply. In no event shall the Royal Society of Chemistry be held responsible for any errors or omissions in this Accepted Manuscript or any consequences arising from the use of any information it contains.
Implementing a strategy for detection of cell-free DNA fragments using magnetoresistive sensors: A translational application in cancer diagnostics using ALU elements

T.M. Dias a,b, F.A. Cardoso a, S.A.M. Martins a, V.C. Martins a, S. Cardoso a, J.F. Gaspar c, G. Monteiro b, P.P. Freitas a

a INESC-MN – Microsystems and Nanotechnologies, Rua Alves Redol, 9 1000-029 Lisbon, Portugal.
b IBB–Institute for Bioengineering and Biosciences, Av. Rovisco Pais, 1049-001 Lisboa, Portugal.
c Toxomics, NOVA Medical School / Faculdade de Ciências Médicas Universidade Nova de Lisboa, Edifício CEDOC II, Rua Câmara Pestana, 6 1150-082, Lisbon, Portugal.

Corresponding author: T.M. Dias (tdias@inesc-mn.pt)
Phone: +351213100237 | Fax: +351213145843

Abstract

Cell-free DNA (cfDNA) is foreseen as a promising source for liquid biopsies in cancer diagnostics. Despite its promise, methods available for its evaluation lack in robustness or, in the case of Next-Generation sequencing (NGS), are extremely sensitive but overly complex for routine operation. Contrary to NGS, integrated Lab-on-Chip devices, offer advantages particularly in terms of automation, cost and speed. These devices, however, have rarely demonstrated the detection of biologically relevant DNA fragments originated from blood. For this end, we present a strategy for the magnetic labeling and detection of cfDNA fragments, using an array of 30 magnetoresistive (MR) sensors integrated in a portable biochip platform. As proof-of-concept, we selected the fragments ALU115 and ALU247, recently identified as promising cancer targets in cfDNA integrity assessment. This work reveals a rational optimization of the DNA probes design and density at the surface which allowed achieving specific target detection and increased inhibition of unspecific interactions, without the need of blocking agents. The developed strategy is adaptable for the detection of mutations, homologous or truncated sequences such as the case of ALU115 and ALU247, sequences that share great similarity. Upon optimization, the MR sensors detected a concentration of the ALU elements within the picomolar range, showing potential for cfDNA analysis in cancer diagnostics.

Keywords: Biochip; Cancer diagnostics; Cell-free DNA; Lab-on-Chip; Magnetoresistive sensors, Liquid biopsies
Introduction

The assessment of circulating cell-free DNA (cfDNA) is being targeted as a promising non-invasive methodology in cancer diagnostics [1]. The cfDNA accumulates in blood mostly from cells dying either by apoptosis or necrosis, a process extensively described elsewhere [2, 3]. With cfDNA acting as a liquid biopsy, somatic mutations that were originated at an individual’s tumor site can be potentially genotyped without having to perform invasive biopsies [4]. The detection of cfDNA derived from tumors, however, carries substantial challenges, as circulating tumor DNA (ctDNA), often only represents a very small fraction (< 1.0%) of the total cfDNA fraction [5]. This presents a limitation for standard sequencing approaches such as Sanger sequencing and pyrosequencing which only detect mutant alleles in a sample down to 15% and 5 %, respectively, of the total cfDNA fraction. In both cases, not enough to detect tumor-derived mutant fragments unless the patients are already affected with tumors in advanced stages [5, 6]. In contrast, techniques that use next-generation sequencing (NGS) [7] or novel PCR approaches such as Scorpion Amplification Refractory Mutation Systems (ARMS-Scorpion) [8], digital PCR [9], Pyrophosphorolysis-activated polymerization (PAP) assays [10], beads, emulsion, amplification, and magnectics (BEAMing) [11] or Tagged-Amplicon deep Sequencing (TAM-Seq) [12] provide a sensitivity for mutants at 1% down to 0.01 % or lower, though with a substantial extra financial cost allied to a complex operation [5] [13]. A midpoint should be met at the frontier between these high throughput NGS technologies and “ultra-low cost” diagnostic systems such as the conventional paper-based assays, so that patients can benefit from affordable, user-friendly approaches that are simultaneously robust and sensitive. Lab-on-chip technologies meet these criteria as they can be easily integrated with electronic components and highly sensitive detectors, requiring only diminutive amount of test samples for analysis. As important as the development of new technologies, the discovery of relevant biomarkers is essential. Besides the identification of somatic mutations in ctDNA, the analysis of the total cfDNA levels in blood (either of normal cfDNA or tumor DNA) alongside with its integrity (size of the cfDNA fragments) have shown to be promising in the assessment of cancer, tumor staging and metastatic potential [14].

The analysis of the concentration and integrity of the cfDNA is promising as the whole DNA content is assessed surpassing the limitations in sensitivity required (< 1 %) for the identification of somatic alterations. Regarding to cfDNA integrity, in healthy individuals, apoptosis is highly predominant over necrosis leading to the release of small and uniform DNA fragments of around 180-200 bp into the
bloodstream [15, 16]. In patients with cancer, the increased cellular turnover and number of cells dying by necrosis, lead to the release of elevated amounts of undigested DNA into the circulation [17]. With these concepts, Umetani et al. designed a method to measure the ratio of longer to shorter DNA fragments (DNA integrity index) based on the assessment of two ALU repeats by quantitative PCR (qPCR) [18]. The fragments assessed with this method were a sequence of 115 bp (ALU115) and a sequence of 247 bp (ALU247), of which, the ALU115 sequence was truncated within the ALU247. Various studies have reported this method, for example, in the diagnosis of gliomas [19], lobular breast cancer [20] and in the detection and monitoring of colorectal cancer patients in both early and late stages by qPCR [21].

This is the first time, however, that these ALU elements, or in fact, up to our knowledge, blood-based cfDNA sequences are targeted for detection using a portable biochip system. The designed biochip comprises an array of 30 magnetoresistive (MR) sensors and offer higher sensitivity and increased portability when compared, for instance, to standard fluorescence techniques [22]. Elucidation on the mechanism by which MR sensors detect biological entities is described elsewhere [23-25]. MR sensors have been used in the multiplexed detection of proteins, magnetically labeled cells and short sequences of single-stranded (synthetic) DNA as proof-of-concept for demonstration of GMR and TMR-based magnetoresistive platforms [26-31]. Still to demonstrate remains the detection of biologically relevant DNA fragments originated from blood as presented here. The methodology followed in this work is divided into three main sections. The first one addresses the generation of magnetically-labeled DNA targets, from the stages of blood collection and cfDNA extraction to the amplification of ALU115 and ALU247 by PCR, digestion of the generated amplicons into single-stranded products and labeling with magnetic nanoparticles. The second section addresses the optimization strategy used to promote the specific binding between SH-capture115 (DNA probe designed to capture ALU115) with ALU115 and of SH-capture247 (DNA probe designed to capture ALU247) with ALU247 at the surface of the sensors. The third section corresponds to the fabrication and characterization of the MR sensors used in this work and the detection and discrimination of ALU115 and ALU247.
Material and Methods

Plasma preparation and cell-free DNA purification

An average of 2.5 mL blood samples were collected from healthy donors into ethylenediaminetetraacetic acid (EDTA) Vacutainer tubes (BD®), and plasma was isolated via a double centrifugation process of 2500 rpm for 10 min, followed by additional 2500 rpm for 10 min, after a tube transfer following the first spin. Samples were processed within 30 min after collection to minimize the release of potential contaminants into the plasma, such as leukocyte genomic DNA. From 1 mL plasma samples, collected after centrifugation, cfDNA was isolated using the QIAamp Circulating Nucleic Acid Kit, and resuspended in 50 µL of Tris-EDTA (TE) buffer, pH 8.0. To validate the DNA extraction and to generate a sufficient amount of DNA amplicons for use in posterior biochip assays, Real-Time PCR was performed.

Amplification of target fragments by PCR

Two sets of primers were designed for amplification of ALU115 and ALU247. The sequences of the primers were adapted from Umetani’s work [18] with modifications that include the addition of a 5’-phosphorylated terminal group in both ALU115 and ALU247 forward primers and a biotin moiety at the 5’-end of both reverse primers. Additionally, the forward primer used for amplification of ALU115 was modified with a non-homologous overhang of 8 base pairs (bp) (GTAGGTAC) upstream to the homologous region of the primer. Generated ALU115 amplicons incorporate, therefore, the 8-bp overhang within its own sequence and assimilate, with this strategy, a distinctive signature from the generated ALU247 amplicons. The primers were purchased from StabVida, Portugal and the sequences, given the above considerations, were as follows: ALU115 forward: Phosphate-5’-GTAGGTACCCTGAGGTCAGGAGTTC-3’; ALU115 reverse: Biotin-5’-CCCGAGTAGCTGGGATTACA-3’; ALU247 forward: Phosphate-5’-GTGGCTCACGCCTGTAATC-3’; ALU247 reverse: Biotin-5’-CAGGCTGGAGTGCA GTGG-3’. The full ALU sequence (Fig. S1), reaction mixture and PCR program are presented in the Supplementary data. To validate the cfDNA extraction, a negative control (without template DNA) was run along with a positive control containing 20 ng of genomic DNA in the reaction mixture, extracted from Human Embryonic Kidney (HEK) 293T
cells (ATCC-LGC Nr: CRL-11268) (additional information on the supplementary methods). PCR
products were electrophoresed on a 2% agarose gel for size confirmation.

**Generation of single-stranded targets and labelling with magnetic nanoparticles**

After size confirmation, the generated amplicons were purified from contaminants using the DNA
Clean & Concentrator™-25 from Zymo research and diluted in TE (10 mM; pH 7.4) buffer for
further use in labelling assays. The amount of generated fragments was quantified by UV absorption
at 280 nm using a NanoDrop Spectrophotometer (NanoValue Plus from Bioxtra). A quantity of 22
ng/µL for ALU115 and of 53 ng/µL for ALU247 was obtained, which corresponds to a concentration of
approximately 0.6 µM and 0.7 µM respectively.

Lambda exonuclease was used in this work for the digestion of the generated double-stranded
amplicons into single-stranded products that can be used in the hybridization assays with
complementary DNA probes. The preferential digestion of one of the strands within the DNA double
helix occurs when one of the strands is modified with a phosphate terminal group [32]. The reaction
used for the digestion consisted of 250 ng of DNA template ALU115 and 500 ng of ALU247 in
separated solutions with 0.5 µL (2.5 Units) of Lambda exonuclease (supplied by New England
Biolabs). The reaction mixture and digestion program used is provided in the supplementary
information (1.5 Lambda exonuclease digestion). The digestion was assessed on a 2% gel
electrophoresis. After digestion, dilutions were performed reaching to a target concentration of 3 nM
and 300 pM for ALU115 and 2 nM and 200 pM for ALU247 based on the initial concentration of
dsDNA quantified by UV absorption. The PCR products were finally bound to streptavidin-coated
superparamagnetic 250 nm magnetic beads (prod. code 09-19-252, Micromod) through streptavidin-
biotin interaction. At this step, 1 µL of target DNA at different concentrations were reacted with a 10 µL
suspension of the beads, 10 x diluted from the stock suspension, for 45 min at room temperature and
under agitation. With the elution in the suspension of beads, the concentration of targets was diluted
10 x from its initial solution. A detailed explanation is provided in the supplementary data (1.6 Labelling
of DNA targets with 250 nm magnetic beads).

**Preparation of gold substrates functionalized with DNA capture probes**

Single-stranded DNA probes (SH-capture115 and SH-capture247), complementary to ALU115 and
ALU247, were synthesized by StabVida Portugal. The base sequences of SH-capture115 and SH-
capture247 were 5’SH-TTTTTTTTTTTTTTGTAGGTACCCTGAGGTCAGG-3’ and 5’SH-TTTTTTTTTTTTTTGATCTGCAGTGGCTCACGCCTGT-3’, respectively. The probes were designed with a 15-bp thymine spacer to provide better mobility and availability for interaction with target fragments and were terminally thiolated to ensure its immobilization on gold fabricated structures. Various concentrations of the DNA probes were prepared (1 µM; 3 µM; 5 µM; 10 µM) and brought into contact with gold substrates by manual spotting. The gold layer on the substrates consisted of Ti 5 nm/Au 40 nm, deposited by sputtering (Alcatel, SCM-450) over a ¼ of 6-inch silicon wafer and cut into small pieces of 7 x 5 mm². Before the spotting of the DNA probes, the gold substrates underwent through a meticulous cleaning process, as described elsewhere [30]. A volume of 1 µL was dispensed for the spotting of each of the concentrations tested. Immobilization for each case proceeded for 2 h at room temperature. Unbounded DNA probes were washed with TE buffer after immobilization. All the steps were performed inside a Petri-dish in a humid atmosphere to prevent evaporation. The composition of TE and PB buffers is provided in the supplementary information.

**Hybridization of magnetically labelled DNA targets with immobilized DNA probes**

10 µL of the DNA targets conjugated to the magnetic beads, and at a concentration of 0.5 µM, were dispensed over the gold substrates previously functionalized with the DNA probes and left to settle and react for 45 minutes at room temperature inside a Petri-dish in a humid atmosphere to prevent evaporation. At this step, each of the generated conjugates was reacted with the correspondent complementary probes (i.e. ALU115 with SH-capture115 and ALU247 with SH-capture247) and cross-reacted (i.e. ALU115 with SH-capture247 and ALU247 with SH-capture115) for the various concentrations of DNA probes tested. A schematic representation of the procedure is presented in Fig. 1. These experiments were performed in order to select the condition that offered the highest specific binding between complementary pairs of probes-targets along with reduced cross-reactivity between the non-complementary pairs. After 45 min of incubation, unbounded or weekly bounded DNA-beads conjugates were removed by washing with PB buffer. Finally, the gold substrates were analysed by optical microscopy with a total magnification of 40x. The agglomerated magnetic nanoparticles over the surface exhibited a brownish colour that could be visualized depending on the density of particles agglomerated in each spot. To further and more accurately analyze the density of magnetic nanoparticles, ImageJ (image software analyzer) was used. The density of beads on spotted areas was calculated and subtracted to background reference values (areas outside the spots) to obtain the
relative surface coverage. The values obtained from analysis with ImageJ must not be seen as absolute but used solely for comparison between different substrates subjected to the same conditions.

**Sensors and Biochip microfabrication**

The biochip configuration comprises an array of 30 U-shaped spin-valve (SV) sensors with the following stack deposited by Ion Beam deposition in a Nordiko 3000 tool: Ta 1.5/ NiFe 3.6/ CoFe 2.5/ Cu 2.1/ CoFe 3.3/ MnIr 11/ Ta 10 (thicknesses in nm) [33]. Here, NiFe, CoFe, and MnIr stand for Ni\textsubscript{80}Fe\textsubscript{20}, Co\textsubscript{90}Fe\textsubscript{10} and Mn\textsubscript{76}Ir\textsubscript{24} film compositions in %. The sensors were defined by direct write laser photolithography and ion milling, resulting in U-shaped sensors with a final active area of 2.5×80 µm\textsuperscript{2} (Fig. 2III inset). Contact leads (AlSiCu 300 nm /15 nm TiW(N)) were made by magnetron sputtering (Nordiko 7000 tool) and were defined by lift-off. The chip was further passivated with 300 nm thick SiN layer deposited by CVD (at 350 °C). The passivation layer was opened by reactive ion etching at the contact pads. Finally, a thin gold pad (Ti 5 nm/Au 40 nm, 43×13 µm\textsuperscript{2}) was sputtered and patterned by lift-off on top of the sensors. The sensors were arranged in six sensing regions each one including four active sensors (with gold) plus a reference sensor (without gold) (Fig. 2 I and 2 II). With the thiol-gold biochemistry, the gold pads determine where the biological probes will chemically bind. After fabrication, the biochips were wire-bonded on a printed circuit board (PCB) and the wires were protected from the external environment with a layer of silicone gel (Elastosil E41). Encapsulated sensors had an average magnetoresistance, minimum resistance and sensitivity (S) of (7.65±0.14) %, (750±30) Ω and (-1.27±0.11) mV/Oe, respectively (Fig. 2 III). For the biochip readout, the PCB was connected to the portable measurement platform and the signal variation during each experiment was acquired in real-time for each of the sensors used.

**Immobilization of the DNA probes on the gold pads and readout of the sensors**

1 µL of the DNA probes, SH-capture115 and SH-capture247, were manually spotted over the sensors at a concentration of 5 µM in TE buffer. SH-capture115 was always spotted on the left column of the chip covering all the sensors in that area while the SH-capture247 was always spotted on the sensors at the right column of the chip (Illustration in Fig.2 IV). Immobilization proceeded for 2 hours at room temperature inside a Petri-dish in a humid atmosphere to prevent evaporation of the spotted drops. After immobilization, unbounded DNA probes were removed by washing with TE buffer. The
PCB carrier was then inserted in the measurement platform and integrated with a microfluidic module to allow the controlled injection of the magnetically labelled DNA targets onto the sensing areas of the chip. The process for fabrication of the microfluidic device is described in the supplementary data.

**Results**

In this work, the optimization strategy for the detection of ALU115 and ALU247 using an array of magnetoresistive sensors integrated in a portable electronic platform is presented and discussed.

**Efficiency of cfDNA isolation and generation of single-stranded biotinylated DNA targets**

The purification of cfDNA from plasma was validated by qPCR, using two calibrators per set of primers for ALU115 and ALU247, one positive control containing 20 ng of genomic DNA and a negative control without DNA. The amplification curves obtained for ALU115 and ALU247 are presented in the supplementary data (Figs. S3 and S4). The crossing points (Cp) for the reactions was derived and, as expected, the positive controls presented a lower Cp in comparison to the negative controls while the samples containing cfDNA purified from plasma led to a Cp value in between, but closer to the Cp of the positive controls. This indicates that a comparable amount of DNA was present in the initial sample (Fig. 3 III). The specificity in the product formation was verified by the melting curves obtained after PCR where only one peak was observed per run (Fig. 3 II). The generated products were within the expected molecular sizes as observed in Fig. 3 I (ALU115 in lane 2 and ALU247 in lane 4).

Additionally, after digestion with lambda exonuclease, the ALU products migrated further in the gel in comparison with the non-digested fragments.

**Surface Biochemistry assays**

SH-capture115 and SH-capture247, at different concentrations (1 μM, 3 μM, 5 μM and 10 μM), were individually spotted over gold fabricated substrates and hybridized with the generated amplicons ALU115 and ALU247, digested *a priori* with lambda exonuclease and labelled with the 250 nm magnetic beads. Before proceeding with analysis in the biochip platform, the hybridization between the capture probes and target DNA, in these assays, was assessed through optical microscopy and data treatment with ImageJ. When magnetically labelled ALU115 and ALU247 were spotted over gold substrates functionalized with TE (without DNA probes), a relative surface coverage of respectively 15
.7 ± 0.7 % and of 27 ± 6% were obtained (Fig. 4 I), suggesting that unspecific adsorption of the targets occurred, with a higher extent for ALU247, possibly due to its higher mass and increased number of nitrogen atoms in the DNA chain available for the unspecific adsorption on gold [34]. The surface coverage for the interaction between complementary probe-target pairs (ALU115 with SH-capture115 and ALU247 with SH-capture247) at 1 µM concentration of DNA probes was identical to the surface coverage obtained when the pairs were cross-reacted (ALU115 with SH-capture247 and ALU247 with SH-capture115). These results are comparable to when no functionalization with DNA probes was performed. For a concentration three times higher of the DNA probes (3 µM), an increase in the surface coverage for ALU115 when paired with SH-capture115 was verified from 17.5 ± 2.5% to 21 ± 3.6 %. For the cross-reaction of ALU115 with SH-capture247, the surface coverage decreased to 10 ± 0.9 %. In the case of ALU247, the hybridization with SH-capture247 resulted in a coverage density of 27.0 ± 7.8 % and a substantial decrease was verified when interacted with SH-capture115, from 32.7 ± 10.5 to 20.7 ± 5 %. By increasing the concentration of the DNA probes to 5 µM, a surface coverage of 18.5 ± 2.6 % and 6.0 ± 0.6 % was obtained for ALU115 with SH-capture115 and with SH-capture247, respectively. For ALU247, the surface coverage was 26.6 ± 6.0 % with SH-capture247 and 6.4 ± 0.5 % with SH-capture115. To conclude, at a concentration of 5 µM, a proportionally higher surface coverage for complementary pairs of DNA probes and target DNA was obtained when compared to the non-specific pairs when cross-reacted. The concentration of probes was tested further at 10 µM and at this concentration, the unspecific interactions were shown to be practically inexistent, however, the specific interactions started to be compromised. The hybridization of ALU115 with SH-capture115 resulted in a surface coverage of 5 ± 0.2%, nearly three times less than for 5 µM. For the case of ALU247, the hybridization with SH-capture247 at this concentration, resulted in a surface coverage of 18.0 ± 4.0%, also reduced in comparison to a concentration of 5 µM. The values reported here are specific to this system and assay conditions used in this study, however the general trends presented in Fig. 4 should be expected for a wide range of parameters, including but not limited to the initial DNA target concentrations, probe-target kinetics, buffer’s ionic strength or incubation times and temperature. An illustration summarizing the prevailing events for the different concentration of probes is provided in Fig. 4 II. Optical images from which the surface coverage percentages were calculated using ImageJ are shown in Fig. S5 as an example of the formed spots for ALU115 when hybridized with SH-capture115 and SH-capture247.
On-chip detection of ALU115 and ALU247

Droplets of 1 µL of SH-capture115 and SH-capture247 at a concentration of 5 µM (corresponding to 3x10^12 DNA probes/µL) were brought into contact with the biochip over the two columns of sensors, separately, before assembling with the microfluidic module. The measurement platform can detect in a single experiment the output signal of 24 bioactive sensors and of 6 reference sensors present in the biochip. For each sensor, a real-time acquisition of the signal was performed. In Fig. 5 it is shown the signal acquisition obtained for six sensors, for detection of magnetically-labeled ALU115 at a concentration of 300 pM. Three of these sensors were functionalized with SH-capture115 and other three with SH-capture247. For each of the sensors, the data presented in Fig. 5 corresponds to the signal measured in real-time (V^{measured}) subtracted to the sensor's baseline signal (V^{baseline}), acquired before the loading of the magnetic beads, and divided by the sensor's baseline (V^{baseline}) (equation 1).

\[ \Delta V_{output} = \frac{V^{baseline} - V^{measured}}{V^{baseline}} (V_{rms}/V_{rms}) \]

With these adjustments it was possible to normalize and compare in the same plot the signal output variation for various sensors (V^{output}), as they all possess a different baseline signal but all vary proportionally to the presence of magnetic beads. In Fig. 5 it is emphasized the various phases of the measured signals, starting with the I) baseline acquisition (V^{baseline}) of the sensor, acquired before the loading of the magnetic beads, followed by the II) injection of the magnetically labelled DNA targets and III) incubation to allow the targets to react with the DNA probes at the surface of the sensors. After incubation, IV) a washing step is performed to remove unbounded or weakly bounded beads from the surface and, finally, V) the binding signal (V^{binding}), originated by specifically bounded beads, is acquired and subtracted to the baseline signal of the sensor (V^{baseline}) to obtain the output signal variation (\Delta V^{output}). The \Delta V^{output} for each sensor is further subtracted to the output signal generated by the reference sensors (biologically inert) (\Delta V^{output reference}) to account for signal variations that may have been originated by thermal drifts at the surface of the sensors (\Delta V^{corrected}) (equation 2).

\[ \Delta V^{corrected} = \Delta V^{output} - \Delta V^{output reference} (V_{rms}/V_{rms}) \]

The average of the normalized and corrected output signals (\Delta V^{corrected}) for each of the interactions performed on-chip between the probes and targets are presented in Fig. 6. For each interaction a
mean of at least 6 sensors was considered. The standard error of the mean for each case was calculated. When ALU115 (300 pM) was interacted with SH-capture115, a signal of $0.048 \pm 0.0010$ (Vrms/Vrms) was achieved, significantly higher than when interacted with SH-capture247, with a signal of $0.003 \pm 0.0002$ (Vrms/Vrms). The hybridization of ALU247 (200 pM) with SH-capture247 resulted in a signal of $0.029 \pm 0.0012$ (Vrms/Vrms) while the same concentration of ALU247 with SH-capture115 resulted in $0.008 \pm 0.0003$ (Vrms/Vrms). Decreasing the concentration of ALU115 to 30 pM resulted in a signal of $0.018 \pm 0.0050$ (Vrms/Vrms) upon interaction with SH-capture115 while the interaction with SH-capture247 led to a signal of $0.001 \pm 0.0001$ (Vrms/Vrms). For a concentration of 20 pM of ALU247, a signal of $0.019 \pm 0.0060$ (Vrms/Vrms) was obtained for the pairing with SH-capture247. Finally, for the pairing with SH-capture115 the signal obtained was $0.004 \pm 0.0010$ (Vrms/Vrms). In Fig.S7 it is presented a group of five sensors exhibiting the presence of magnetic particles. The reference sensor is bioinert while the other four correspond to specific biointeraction between probe and target DNA after on-chip hybridization.

Discussion

This paper reports the optimization process for detection of ALU115 and ALU247 with an integrated system based on magnetoresistive sensors as a promising lab-on-chip technology for cfDNA analysis. Its key features are portability, versatility for manufacturing and capability for integration with various electronic components and microfluidics, providing the system a superior autonomy when compared, for instance with conventional laboratory procedures [35].

The purification and amplification of the target fragments is still not integrated with the detection system but the strategy here employed serves as the basis for a total integrative device having in mind the required steps to generate magnetically labelled single-stranded.

In the particular case of detection of the ALU elements, since the fragment ALU115 is truncated within ALU247, the detection and discrimination of both DNA targets required a particular optimization. ALU115 was generated by PCR but using a forward primer modified with a non-homologous 8-bp (GTAGGTAC) overhang, as a strategy to assimilate in ALU115 a distinctive signature from the generated ALU247 amplicons. SH-probe115 was designed with the complement of the 8-bp signature in order to target that region of ALU115, which in turn, was absent in the ALU247 products. With this
strategy, the complementarity between SH-capture115 and ALU115 was maintained while the complementarity between SH-capture115 and ALU247 was decreased by 40%. The 8-bp signature was balanced in terms of GC and AT content to ensure stable binding between the probe and the complementary target. The reaction efficiency for ALU115 amplification was not affected by the 8-bp overhang as demonstrated by the amplification curves (Fig. S3 and S4), product formation (verified on an agarose gel) and melting curve analysis (Fig. 5).

Regarding to the hybridization tests, it was important to optimize the biofunctionalization of the sensors so that specific complementary targets could be recognized at the surface of the sensors, simultaneously avoiding unspecific adsorption or cross-reactions. Especially knowing that DNA adsorbs non-specifically to gold via interaction with nitrogen atoms from the DNA chains, this optimization was essential [36]. Usually, blocking agents, such as bovine serum albumin (BSA), are used to diminish the amount of free sites at the surface and, therefore, used to prevent the unspecific interaction of target DNA with the gold atoms of the surface [37]. As an alternative, in this work, it was investigated a simple strategy based on the optimization of the DNA probes density at the surface so that no free gold spaces were available for the non-specific adsorption of DNA. It is well known that the density of DNA probes at the surface highly affect the amount of target DNA that can be captured [38]. A highly dense monolayer of probes may inhibit the capture of complementary target DNA due to steric hindrance effects while a low dense monolayer may not be enough to specifically capture a good proportion of target DNA [34, 39]. These concepts were applied in this work so that the DNA probes itself could act as blocking agents to non-complementary DNA but still used at a concentration that allowed the recognition of complementary target DNA. At 1 µM, an equivalent surface coverage was verified for complementary probe-target pairs (ALU115 with SH-capture115 and ALU247 with SH-capture247) and for cross-reacted pairs (ALU115 with SH-capture247 and ALU247 with SH-capture115), indicating that the spots of beads may have been originated mainly through adsorption of the DNA targets on the gold substrates. An increased concentration of probes (3 µM) led to a decreased adsorption and increased specific binding as the surface coverage for the non-complementary pairs diminished in comparison to the complementary pairs. At 5 µM, the disparity between both complementary and cross-reacted pairs increased, meaning that the specific binding was favoured in comparison to the adsorbed fraction. The packaging of DNA probes, at this condition, seemed to have blocked the unspecific adsorption of DNA, at a greater extent, and limited the
interaction of ALU247 with SH-capture115. At 10 µM, the amount of probes at the surface started competing with the specific hybridization amongst complementary probe-target pairs.

After optimization of the concentration of probes, the magnetic detection of both targets was validated in the picomolar range, using an array of magnetoresistive sensors as demonstrated in Figs 5 and 6. Although in the past, the detection of DNA in the femtomolar range was performed, using magnetic fields to direct the magnetic particles towards the sensors, in this work, the limit of detection required for cfDNA analysis was circumvented by using pre-amplification of target DNA [25]. This was performed to analyse the signal variation of the system for known concentrations of the targets, for the conditions tested in these assays. For evaluation in a clinical setting, the PCR would have to be stopped at the exponential phase of the reaction so that a correlation with initial DNA concentrations can be performed upon detection.

In conclusion, the strategy here utilized for the manipulation of the DNA target strands and, consequently, of the DNA capture probes, along with the optimization of their density at the surface, was effective in preventing the majority of the unspecific interactions without affecting specific target-probe recognition events. This strategy can be adapted to the detection of other truncated sequences or applied for the detection of polymorphisms or mutations as a method to decrease the specificity between cross-reactant species. Plus, the blocking of unspecific interactions solely by controlling the density of probes at the surface, in general, decrease the time and cost of any assay as no blocking solutions are required. Above all, these results demonstrate that MR sensors can effectively be used at the Point-of-Care for detection of blood-based DNA biomarkers upon proper optimization of each particular case-study. For future work, a module for PCR amplification is expected to be implemented into the system so that the integration between amplification and detection can be achieved. Additionally required is a microfluidic module for purification of target DNA from the contaminants in the plasma which affect the amplification of targets by PCR and ultimately the detection. Finally, we aim to detect other biomarkers, characteristic to specific types of cancer, in combination to ALU115 and ALU247, for a more inclusive analysis in cancer diagnostics, hopefully surpassing NGS devices in terms of speed, costs and portability.

Acknowledgments
The author Tomás Dias, wants to thanks FCT for his PhD grant SFRH / BD / 81537 / 2011. INESC-MN acknowledges FCT funding through the IN associated laboratory (Pest-OE/CTM/LA0024/ 2011), and also the projects FCT -EXCL/CTM-NAN/0441/2012, PTDC/EEA-ELC/108555/2008, PTDC/CTM-NAN/110793/2009 and EXPL/EEI-ELC/1029/2012.


Fig. 1: Scheme with steps for the spotting of DNA probes and biomolecular recognition of magnetically-labeled targets. I) Bare gold substrates; II) Manual spotting of 1 µL of DNA probes; III) 10 µL spotting of magnetically labeled targets over previously immobilized DNA probes (complementary or non-complementary probes); IV) The specific recognition of targets by the complementary probes spotted at the surface creates a dense spot while the hybridization with non-complementary DNA probes results in the removal of the particles, after washing.
Fig. 2: I) Biochip comprising six sensing regions. II) Each one including four active sensors (gold pad on top of the sensors) plus a reference sensor (without gold). III) Transfer curve of a U-shaped spin-valve sensor (2.5×80 µm²). Inset: Optical microscope image (400 x magnification) of a sensing unit comprising the spin-valve surrounded by an aluminium current line. IV) Spotting of DNA probes and integration with microfluidics. 1) SH-capture115 was always spotted on the left column of the chip covering all the sensors in that area while the SH-capture247 was always spotted on the sensors at the right column of chip. 2) The biochip was integrated with a microfluidic channel to allow the 3) loading of the magnetically-labeled targets into the system.
Fig. 3: DNA product analysis for ALU115 and ALU247. I) Agarose gel with ALU115 and ALU247 before and after digestion with lambda exonuclease. 1- HyperLadder IV (ranging from 10kb to 200 bp). 2- ALU115 before digestion. 3- ALU115 after digestion. 4- ALU247 before digestion. 5- ALU247 after digestion. II) Melting curve analysis of the PCR products. III) Cp values obtained from the amplification by PCR of ALU115 and ALU247, including the positive and negative controls. The Cp values were derived using the Fit Points Method.
Fig. 4 I) Surface coverage (%) for various concentrations of probes SH-probe115 and SH-probe247 in interaction with magnetically labelled ALU115 and ALU247 (n ≥ 3). II) Illustrations representing each situation and summarizing the prevailing event for each of the concentration of DNA probes.
Fig. 5: Signal response for detection of ALU115 (300 pM) by sensors functionalized with complementary SH-probe115 (red curves) and sensors functionalized with non-complementary SH-probe247 (black curves).
Fig. 6: On-chip detection of ALU115 and ALU247. Data obtained from the average of different sensors for each of the measurements (min = 8 sensors; max = 12 sensors).
Table of contents

Optimization of a strategy for detection of cell-free DNA using an array of magnetoresistive (MR) sensors integrated in a portable readout platform.