Diabetes screening by telecentric digital holographic microscopy

A. DOBLAS*, E. ROCHE†, F. J. AMPUDIA-BLASCO‡, M. MARTÍNEZ-CORRAL*, G. SAAVEDRA* & J. GARCIA-SUCERQUIA§

*3D Imaging and Display Laboratory, Department of Optics, University of Valencia, E-46100 Burjassot, Spain
†Department of Applied Biology-Nutrition, Institute of Bioengineering, University of Miguel Hernandez, E-03203 Elche, Spain
‡Diabetes Reference Unit, Department of Endocrinology and Nutrition, Valencia Clinic University Hospital, E-46010 Valencia, Spain
§School of Physics, Universidad Nacional de Colombia Sede Medellin, A.A: 3840-Medellin-050034- Colombia

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Summary

Diabetes is currently the world’s fastest growing chronic disease and it is caused by deficient production of insulin by the endocrine pancreas or by abnormal insulin action in peripheral tissues. This results in persistent hyperglycaemia that over time may produce chronic diabetic complications. Determination of glycated haemoglobin level is currently the gold standard method to evaluate and control sustained hyperglycaemia in diabetic people. This measurement is currently made by high-performance liquid chromatography, which is a complex chemical process that requires the extraction of blood from the antecubital vein. To reduce the complexity of that measurement, we propose a fully-optical technique that is based in the fact that there are changes in the optical properties of erythrocytes due to the presence of glucose-derived adducts in the haemoglobin molecule. To evaluate these changes, we propose to perform quantitative phase maps of erythrocytes by using telecentric digital holographic microscopy. Our experiments show that telecentric digital holographic microscopy allows detecting, almost in real time and from a single drop of blood, significant differences between erythrocytes of diabetic patients and healthy patients. Besides, our phase measurements are well correlated with the values of glycated haemoglobin and the blood glucose values.

Introduction

Diabetes mellitus is a disease that includes a series of metabolic alterations that affect different organs and tissues (Maahs et al., 2010; Chen et al., 2011). In 2013, 382 million people around the world were affected with diabetes and according to the International Diabetes Federation (IDF) and the World Health Organization (WHO) this metabolic disorder is expected to affect 592 million people in 2035 (Maahs et al., 2010; Chen et al., 2011). Diabetes is mainly due to a deficient production of insulin by the endocrine pancreas or by abnormal insulin action in peripheral tissues. Diabetes is generally classified into four broad categories (Maraschin, 2012): type 1, type 2, gestational diabetes and others. Type 1 and type 2 are the most prevalent and therefore responsible for the majority of the health care cost attributed to this disease. Type 1 diabetes mellitus (T1DM) is characterized by the almost total loss of insulin due to an autoimmune destruction of pancreatic beta cells (Boitard, 2012). Therefore, type 1 diabetic patients require lifelong insulin injections, since there are no alternative hormones in the organism that can mimic the effect of insulin.

In any case, insulin injections cannot replace the exquisite control exerted by the pancreatic beta-cells that secrete the hormone in response to glucose variations in blood in a dose-dependent manner (Malik et al., 2014). For this reason, patients with T1DM present prominent fluctuations in blood glucose, with longer periods of hyperglycaemia compared to nondiabetic individuals (Korytkowski, 2013; Lee, 2013; Todi, 2014). This may result over time in the development of secondary complications that affect mainly the retina, kidneys, nerves and cardiovascular system (Forbes et al., 2013). Therefore, hyperglycaemia control is one of the main objectives to minimize the risk of developing retinopathy, nephropathy, neuropathy and cardiovascular complications (Korytkowski, 2013; Todi, 2014). Plasma glucose measurements based on dry chemistry and performed by a glucometer are quite accurate and minimally invasive, requiring only a drop of blood. However, these determinations, performed 4–6 times per day by the own patient, provide only a prompt index of the circulating glucose.

Long-term glycaemic control is evaluated in terms of glycated haemoglobin (HbA1C) concentration (International Expert Committee, 2009). High glucose concentration in blood results in nonenzymatic glycation of side-chain amino groups
of the amino acid residues, such as lysine, of circulating proteins. Glycation rate increases with higher concentrations of the sugar. The concentration of HbA1C indicates the level of exposure to persistent hyperglycaemia over the last 2–3 months, coinciding with the half-life of haemoglobin (approximately 120 days). Therefore, HbA1C is the gold standard parameter to evaluate periodically glycaemic control in patients with diabetes (International Expert Committee, 2009). HbA1C is measured currently by high-performance liquid chromatography, and expressed both in percentage and in International Federation of Clinical Chemistry (IFCC) units (mmol/mol; Consensus Committed, 2007). Recently, the American Diabetes Association (ADA) as well as other institutional bodies have adopted HbA1C levels as diagnostic of diabetes if above 6.5% (American Diabetes Association, 2010).

In this work, we propose a novel, fully-optical technique for the easy evaluation of HbA1C concentrations, and therefore for obtaining a fast screening of T1DM levels. This study is motivated by two previous published works (Mazarevica et al., 2002, Okamoto et al., 2000) which show that the refractive properties vary in presence of diabetes mellitus. In fact, Mazarevica et al. demonstrated a relationship between the refractive index, which is measured using a Nomarski polarizing-interference microscope, and pH level for diabetic and healthy donors. On the other hand, Okamoto et al. verified that variations in plasma glucose causes refractive index changes in eyes of diabetic subjects and, as a result, transient myopia or hyperopia is derived. Based on their findings, it seems logical to hypothesize that the phase of red blood cells (RBCs) could also change. Our method is based in the use of a telecentric digital holographic microscopy (DHM) for the acquisition of quantitative phase maps of erythrocytes samples obtained from patients affected by diabetes, and also from nondiabetic (named as controls) individuals. After analysing the alterations in the phase distributions we have found that there is a strong correlation between the absolute phase value of RBCs and the corresponding values of HbA1C. This correlation permits us to suggest the use of telecentric DHM to quantify persistent hyperglycaemia states. Besides, since only a small drop of blood is required and the phase measurement can be performed in any time, we can adopt this technique as a first-screening method for the diagnosis of abnormal hyperglycaemia.

**Material and methods**

**Sample preparation**

Samples were obtained from both nondiabetic individuals (controls) and also from patients with T1DM who have been under treatment with insulin injections over a period of 23 ± 10 years (mean ± standard deviation). Capillary blood specimens from patients with T1DM were collected during routine ambulatory visits at the Diabetes Reference Center from the Clinic Hospital of Valencia (Spain). Control samples were obtained in a certified clinical analysis laboratory, Elche (Spain). All individuals participated voluntarily in this study, being adequately informed about the conditions and the purpose of the assessment. Participants signed an informed consent approved by the Ethics Committee of the University Miguel Hernández (Elche, Spain), fulfilling the requirements of the Declaration of Helsinki regarding human research.

Clinical measurements of HbA1C were performed using high-performance liquid chromatography of blood extracted from the antecubital vein. On the other hand, telecentric DHM was performed, at the 3D Imaging and Display Laboratory (University of Valencia), on a capillary blood drop smearing on a glass slide. The smeared blood was dried out at room temperature in a dust-free environment. After the sample preparation, each slide was placed properly in the optical object path of our off-axis transmission telecentric DHM and a hologram was captured for each sample. The time required for the DHM operation and data processing did not exceed 1 min per sample.

**Telecentric DHM**

The telecentric DHM was mounted, in open configuration, on an optical table. The scheme is shown in Figure 1. A He-Ne laser (wavelength = 633 nm, 50 mW output power, random polarization, by Edmund Optics) was used as the illumination source. The light proceeding from the laser was collimated and split to produce the reference (R) and object waves (O). A Nikon Plan Epi 50/0.55 NA microscope objective and a tube lens of focal length $f_{TL} = 200$ mm were placed in the object arm of the microscope. A CCD camera with 1024 1024 square pixels of 6.9 μm in side, recorded the holograms formed by the interference between the wavefield diffracted by the sample and a reference plane wave R. In order to operate in single-shot mode, the DHM was arranged in off-axis geometry by slightly tilting the reference wave R. The beam-splitter BS2 and the mirror M2 controlled the tilt angle. Note that the DHM operated in a telecentric regime in order to provide accurate quantitative phase measurements (Doblas et al., 2013; Doblas et al., 2014). A detailed description of the image formation procedure and the reconstruction process have been carefully given in references Doblas et al. (2014) and Sánchez-Ortiga et al. (2014).

Since DHM is a holographic method, the computational reconstruction of the object wave follows well-established and exact methods. Specifically, to retrieve the object information, it is necessary to filter out the Fourier transform of the object information from the hologram spectrum, and then to perform the inverse Fourier transform. It is worth to mention that in our scheme the CCD is placed at the image plane of the imaging system, see Figure 1, and therefore not back-propagation algorithms are required (Kreis, 2004; Kim, 2011; Popescu, 2011; Picart & Li, 2012). The computational processing of the digital holograms to provide a quantitative phase map of the sample was performed using Matlab (Mathworks, Natick, MA, USA).
Fig. 1. Experimental configuration of the off-axis DHM. The DHM operates in a telecentric regime and the CCD is placed at the image plane. CL is a collimating lens; BS1 and BS2 are beam-splitters; M1 and M2 are mirrors; MO is a microscope objective; and, TL is a tube lens.

Stability analysis
As it is well known, phase measurements are strongly dependent on ambient fluctuations. To provide a controlled environment during the holograms acquisition, the complete microscope has been enclosed in a polymethymethacrylate cage with its temperature controlled. To exclude any incidence of ambient perturbations on the results of the experiment, we initially tested the stability of the set-up by recording 25 holograms of a test sample over a period of 5 min. From the 25 holograms, we calculated the corresponding 25 phase maps. Using these maps, the mean value of the phase over an area free of RBCs information was measured. It is worth to mention that the size of the area used to compute the mean phase was 41 x 41 μm². For this region, the obtained value was 1.591 ± 0.013 rad. From the small value of the standard deviation (0.013 rad), we concluded that our DHM system was fairly isolated from ambient fluctuations to guarantee trustable measurements for at least 5 min and, thus, any divergence between the measurements would be due to differences on the samples.

Statistical analysis
Nonparametric tests were used since the sample was small and, consequently, the variables were not normally distributed. Mann–Whitney (Hollander et al., 1999; Randolph et al., 2013) test was used to assess differences between control and diabetes groups with each one of the diagnostic methods (DHM and high-performance liquid chromatography). Correlations between methods were later computed by means of the Spearman rho test [23, 24]. Note that the statistical tests can be interpreted using p-values (Hollander et al., 1999; Randolph et al., 2013). The p-value is the probability of obtaining a test statistic result at least as extreme as the one that was actually observed, assuming that the null hypothesis is true. If the p-value is less than the required significance level, then one says the null hypothesis is rejected at the given level of significance. The lower significance level the stronger presumption against null hypothesis. All data were statistically analysed using the SPSS software (version 22.0).

Results and discussion
Table 1 summarizes the main characteristics of the population participating in the study. The total number of participants was 43, separated into 14 controls (nine men and five women) and 29 T1DM patients (13 men and 16 women). The results shown in Table 1 indicate that blood glucose and HbA1C values of the control group are much more homogeneous, in contrast to the values obtained from the T1DM group.

Using our telecentric DHM we measured the phase maps of RBC samples from both the healthy control group and the T1DM patients. As illustrating example of the results we obtained, in Figure 2 we show the phase maps corresponding to three controls and to three patients. For better visualization, the images have been pseudocoloured using the same colour scale. From these samples, it is apparent that phase values of healthy RBCs are significantly smaller than those obtained for the T1DM group. The phase maps were used to calculate a mean phase value for each participant. For each RBC the phase was obtained as the average value over a square area of approximately 1.12 x 1.12 μm² surrounding the centroid of the RBC. The phase values of 20 erythrocytes were averaged to obtain the phase value for each individual of the test. Owing to the sample preparation, the measured phase includes contributions not only from the RBCs but also from the plasma components. To remove the latter, the average value of the phase over the areas free of RBCs is subtracted from the total phase. This procedure was indeed applied to the phase maps show in Figure 2, hence the phase value over the RBCs is essentially provided by the optical path length along the blood cells. The stripe-like pattern seen over the images of the RBC in Figure 2, are derived from the limited accuracy of the digital holographic microscope that introduces these artefacts. The limited size of the recording camera, the numerical processing of the recorded holograms via fast Fourier transforms, are some of the facts that could produce these
Table 1. Basic characteristics (mean, standard deviation and range) of the sample under research

<table>
<thead>
<tr>
<th></th>
<th>Control (N = 14)</th>
<th>T1DM Patient (N = 29)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>Standard deviation</td>
</tr>
<tr>
<td>Age (year)</td>
<td>33</td>
<td>14</td>
</tr>
<tr>
<td>Height (cm)</td>
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<td>10</td>
</tr>
<tr>
<td>Weight (kg)</td>
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<td>15</td>
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<tr>
<td>Glucose (mg/dL)</td>
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<td>12</td>
</tr>
<tr>
<td>HbA1C (%)</td>
<td>5.2</td>
<td>0.3</td>
</tr>
</tbody>
</table>

Fig. 2. Pseudocoloured image of the phase maps of healthy RBCs (first row) and T1DM-RBCs (second row), measured by a telecentric-DHM. In all cases, the image area is $13.8 \times 13.8 \mu m^2$.

patterns, which however do not perturb the measurements of the RBCs.

Figure 3 shows the histograms of the RBC phases from healthy individuals and for diabetic patients. The mean values were $3.15 \pm 0.13$ rad and $3.72 \pm 0.15$ rad, respectively. Clearly, the phase values in the T1DM group were significantly higher than in the healthy control group. In addition, there is a big gap between the phase values of the two groups, see in Figure 3 that the phase values ranged 2.94–3.30 rad for healthy RBCs and 3.51–4.01 rad for T1DM RBCs. It is then reasonable to claim that the phase measurement made with telecentric DHM permits to divide the population into groups of healthy or T1DM patients and, moreover, phase values above 3.40 rad are an indication of hyperglycaemia. Another way to ensure this conclusion, and perhaps even the most formal way, is by applying the Mann–Whitney test, which tells us if the phase values from both groups are independent of each other. It is important to realize that this test shows
greater efficiency than the regular t-test (Hollander et al., 1999; Randolph et al., 2013) on non-normal distributions, see in Figure 3 that our phase distributions are not Gaussian. Using the Mann–Whitney test, we found that the phase distributions of both groups are statistically independent with a very high significance level ($p < 0.001$).

As a proof of the validity of the diabetes screening with telecentric DHM, we have contrasted the DHM results with the HbA1C values, which is the gold standard to assess diabetic patients. In Figure 4 we have plotted the phase versus the HbA1C values for all the population under study. We measured the rho correlation coefficient to obtain rho = 0.739, with $p < 0.01$. The high correlation coefficient and the clear separation between the two populations confirm that both parameters may potentially be used to diagnose diabetes states as well as to evaluate long-term glycaemic control in patients with diabetes. This result can be explained by accepting the findings by Barer et al. (Barer, 1952) which demonstrated that haemoglobin concentration is directly related to phase values. Although, from Figure 5, it seems that both DHM and high-performance liquid chromatography should be individually sufficient to separate out the two populations, it is worth to highlight that using both approaches would allow for even better screening of diabetes mellitus. Besides, DHM technique could be used as the first method since only a capillary blood drop is needed, it can be performed in any time and the phase measurement is obtained in a fairly instantaneous way.

**Conclusions**

In this paper is demonstrated, for the first time to our knowledge, the use of telecentric DHM for diabetes screening. The utility of the DHM results from the possibility of providing quantitative phase maps as indicators of long-term blood sugar control similar to HbA1C. Although a strictly medical research would require the analysis of much more individuals, our results should be viewed as a proof-of-concept study. In addition, the proposed method shows great potential as optional tool to screen for diabetes because: (1) it is a minimally invasive technique since only a small drop of capillary blood is needed; (2) it can be performed at any time and the results can be obtained almost in real time; and (3) it is a wide-field technique which can be easily implemented in a conventional microscope (Sánchez-Ortega et al., 2015) and can be used to analyse illnesses in which the refractive index or/and the morphology are distorted (Park et al., 2008; Kemper et al., 2010; Byun et al., 2012; Pavillon et al., 2012; Sridharan et al., 2015).

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**References**


