An MCM-Based Microsystem for Colorimetric Detection of Biomolecules in Biological Fluids

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Abstract—This paper presents a multichip module microsystem for biological fluid analysis. It is composed of three parts, namely 1) CMOS silicon optical detection microsystem, which includes a photodetector and a light-to-frequency converter for readout; 2) dielectric thin-film based optical filter on top of the photodetector fabricated using IC-compatible postprocessing; and 3) glass die on top containing the microchannels fabricated using SU-8 techniques. The application is in low-cost quantitative measurement of the concentration of biomolecules in biological fluids. Its operation is based on optical absorption in the part of the visible spectrum that is defined by the specific biomolecule. Signals proportional to the intensity of the light transmitted through the biological fluid are available at the output in the form of bit streams, which allow simple computer interfacing. The quantitative measurement of the total protein in urine is successfully demonstrated. The photodiode responsivity is 224 mA/W with a full-width at half-maximum of 10 nm at $\lambda = 600$ nm. The optical system sensitivity is 1 kHz/W m$^{-2}$ at $\lambda = 670$ nm.

Index Terms—Biomolecule analysis, multichip module (MCM), optical absorption, SU-8 techniques.

I. INTRODUCTION

For diagnostic reasons, patients in a hospital are often subjected to biochemical analysis of their body fluids. Usually, the analyses are carried out in clinical laboratories, and the results become available after several hours, sometimes days. Therefore, a reliable diagnosis cannot be performed within the consultation time. Mistakes in the logistics, such as lost samples and mislabeling, may further delay diagnosis [1]. The automated equipment used in a state-of-the-art laboratory reduces errors, but use of high sample and reagent volumes makes the analysis expensive and does not contribute to patient comfort [2]. Outside the laboratory environment, reagent strips are commercially available [3]. Such strips are intended for a limited set of biomolecules to be analyzed and qualitative color readout. The need for rapid and on-line measurements with low sample volumes has led to the development of microsystems with the fluidic and detection and readout systems integrated in a single module.

This paper describes a multichip module (MCM)-based microsystem for application in clinical analysis of biological fluids using spectrophotometry. It allows the measurement of the concentration of biomolecules in those fluids. The measurement is based on colorimetric detection by optical absorption in a part of the visible spectrum defined by the reaction of the specific molecule with a specific reagent. The microsystem combines in an MCM the detection and readout system, the optical filtering system, and the microfluidic system.

II. MICROSYSTEM DESIGN

The optical detection microsystem avoids the need for expensive readout optics and may enable low-cost disposable devices, which would improve the use of spectrophotometric analysis in clinical diagnostics. The optical channel is composed of the fluidic channel, a thin-film optical filter, and a silicon photodetector (see Fig. 1). The structure is optimized for optical response at the absorption peak of a particular biomolecule. The optical absorption intensity is proportional to the biomolecule concentration.

A. Detection and Readout System

The design involves selection of the photodetector type using the active layers available in a CMOS process and selection...
of the dielectric layers for suitable optical response. The photodetector selected is a p-n junction photodiode based on the n+ layer/p-epilayer junction. This structure is chosen because it provides the best possible quantum efficiency in the desired spectral range of photodiodes available in a CMOS process [4]. In a standard CMOS process, the junction depth of the photodiode is fully defined and cannot be altered. The thickness of the n+ layer is 350 nm. The thickness of the epilayer is 12 μm, and the doping concentration is $10^{16}$ cm$^{-3}$. However, the quantum efficiency can be improved by a suitable arrangement of dielectric layers on top of the photodiode surface. These act as a thin-film interference filter and influence the optical transmittance for each wavelength independent of the selective measurement of the intensity at the desired wavelength. SiO$_2$ thin films. The multi-layer is composed of a stack of TiO$_2$ and SiO$_2$ thin films. The thickness of the films determines the tuned wavelength. SiO$_2$ has been selected because the wavelength dependence of its refractive index for the spectral band between 500 and 700 nm is almost constant (1.465 and 1.457, respectively). TiO$_2$ has been selected due to fabrication constraints (the deposition process is well characterized).

**B. Optical Filtering System**

An optical filter on top of the photodiode is required with a narrow passband around the wavelength for which the biomolecule being analyzed has its absorption maximum. This enables the selective measurement of the intensity at the desired wavelength transmitted through the mixture. The optical interference filter is based on a nine-layer dielectric thin-film design, which offers high reflectivity with low absorption losses. The multilayer is composed of a stack of TiO$_2$ and SiO$_2$ thin films. The comparator detects $V_{\text{comp}} < V_{\text{ref}}$, which causes $V_{\text{out}}$ to change to the high logic level again and the cycle to repeat (see Fig. 4). The output bit stream frequency is a function of the charge change in the capacitor $\Delta Q$, which is proportional to the input photocurrent [6] and hence to the biomolecule concentration, with $f_{\text{bitstream}} = I_{\text{photodiode}}/\Delta Q$.

Four photodiodes are used in each measurement. The first photodiode is for the reagent (without biomolecules), the second is for the biological fluid being analyzed, the third is for measuring the photodiode dark current, and the fourth is for the calibrator, a standard with a well-known concentration of the biomolecule that is being analyzed. The dark current is the current that flows in a photodiode when there is no optical radiation incident on the photodiode. It is usually measured and subtracted from the flux. As the dark current is temperature dependent, one measurement at the beginning of the experiment is usually not sufficient. Thus, in the reported circuit, a dark current compensation channel is implemented using photodiode 3. This photodiode is completely covered with metal.

The logic circuit drives the analog switches $S_{2a}$ through $S_{2d}$ for selecting the channel to be measured. These switches are identical complementary p-n/MOS switches. The comparator is a clocked high-speed regenerative comparator with a rail-to-rail input circuit. For reliable operation, a two-phase nonoverlapping clock is used for the analog switch $S_1$ and for the comparator. It is possible to use a digital counter for counting the output pulses of the comparator during a fixed time period, producing the digital value correspondent to the photocurrent intensity. Alternatively, a microcontroller can substitute the counter, generating all the control logic and performing additional calculations as well.

![Fig. 2. Cross section of the basic structure of the photodiode for fabrication in a standard n-well CMOS process.](http://ieeexplore.ieee.org)
bands by adjusting the thickness of one or more layers without affecting the microsystem layout.

C. Microfluidic System

The microfluidic system contains the microchannels and the detection chambers (see Fig. 6). There are three detection chambers, each 2 mm wide, 3 mm long, and 500 \( \mu \text{m} \) deep. The high depth is crucial for the optical absorption measurements. The detection chamber A contains only the reagent, and it is needed to obtain the baseline reference and to calibrate the light source. The detection chamber B allows the mixed solution analysis; it contains the reagent plus the sample with biomolecules. The detection chamber C is needed to calibrate the biomolecule concentration (with a well-known concentration standard). The main channel is 500 \( \mu \text{m} \) wide, 70 mm long, and 500 \( \mu \text{m} \) deep, with a liquid volume quantity of 20 \( \mu \text{L} \). It has two inlets, one for the reagent, R, and the other for the sample, S. The chosen structure allows a complete and homogeneous mixing of the reagent with the sample. The mixing process was described in [7].

III. MICROSYSTEM FABRICATION

A. Detection and Readout System

The CMOS-compatible photodetectors and readout circuits have been fabricated through a double-metal, single-polysilicon, 1.6-\( \mu \text{m} \) n-well CMOS process. The area of each optical channel is \( 500 \times 500 \ \mu \text{m}^2 \). Fig. 7 shows a photograph of the fabricated optical detection microsystem.

B. Optical Filtering System

The optical filter is postprocessed, by evaporation, on top of the photodetector using the same mask that was used for the selection of the dielectric layers available in the standard CMOS process. The filter fabrication starts with the deposition of an 80-nm \( \text{TiO}_2 \) layer (layer 1 in Table I) after completion of the standard CMOS process, including the metalization and the etching of the two oxide layers on top of the photodiode. Then, the eight subsequent layers of \( \text{SiO}_2 \) and \( \text{TiO}_2 \) are deposited with the thicknesses described in Table I. A scanning electron microscopy (SEM) photograph of the cross section of the optical channel is shown in Fig. 8. A commercially available passband optical filter on top of the MCM-based microsystem is used to block the nonvisible part of the spectrum.

C. Microfluidic System

Fig. 9 shows a photograph of the microchannels fabricated using SU-8 techniques. The SU-8-based fabrication is a low-cost process, biocompatible, UV lithography semiconductor compatible, and does not require expensive masks. Moreover, SU-8-based processing enables the fabrication of deep microchannels with very low sidewall roughness and is suitable for optical absorption measurement [8].
A negative mask of the microchannels’ die layout is fabricated from a regular transparency foil (like the one used in printed circuit boards). The SU-8 photosresist chosen is the SU-8 100. The microchannel fabrication starts with spinning of the SU-8 photosresist on a glass substrate at 700 r/min for 100 s. Soft bake prior to exposure is performed at 90 °C for 3000 s. The second spun is carried out at 700 r/min for 100 s, and a soft bake for 300 s at 50 °C is again performed. The temperature is ramped up to 90 °C for 5400 s and subsequently ramped down again. The exposure takes 900 s on an Electronic Vision EV-420 mask and bonding aligner. Postexposure bake is performed for...
Fig. 10. Oscilloscope capacitor voltage (1) and inverted comparator output voltage (2) for a photodiode current of 137.0 nA.

Fig. 11. Oscilloscope capacitor voltage (1) and inverted comparator output voltage (2) for a photodiode current of 138.1 nA.

300 s at 50 °C. The temperature is again ramped up to 90 °C for 1200 s and ramped down. The fabrication ends with the development during 2700 s in the SU-8 developer.

IV. EXPERIMENTAL RESULTS

The electrical characteristics and spectral responsivity were measured using a Keithley 487 picoamperemeter (full-scale range from 10 fA to 2 mA and a resolution of 5 1/2 digit), and a 250-W quartz tungsten halogen lamp with the Oriel Cornerstone 130 monochromator was used as light source. The measurements were calibrated with a calibrated commercially available photodiode as reference (Hamamatsu S1336-5BQ).

The oscilloscope traces in Figs. 10 and 11 show the electrical characteristics of the readout circuit. These signals were obtained for a photodiode current of 137.0 and 138.1 nA, which represents 15 and 1 mg/dl of the total protein concentration in urine, respectively. The output frequency of the light-to-frequency converter for different values of the photodiode current is shown in Fig. 12. The photodiode dark current is 0.38 pA (1.52 × 10⁻¹⁸ A/µm²) at 0 V, its responsivity is 224 mA/W at λ = 600 nm (the wavelength at which the total protein in urine has its absorption maximum), and the sensitivity achieved is 1 kHz/W m⁻² at λ = 670 nm (using the TLS230 from Texas Instruments as a reference).

The total protein concentration in a urine sample was measured as a test of this MCM-based microsystem. The reagent used in the measurements was the Microprotein-PR reagent from Sigma-Aldrich. It reacts with a sample of urine containing total protein in a 50:1 ratio and produces an absorption maximum at a specific wavelength (λ = 600 nm).

Fig. 13 shows the measured transmittance through the entire optical channel as a function of protein concentration. Measurements were performed in a range between 1 and 50 mg/dl, which includes normal and abnormal values in a human being (normal values are typically less than 20 mg/dl). The transmittance is defined as \( T = I/I_0 \), where \( I \) is the measured photodiode current for each mixture, and \( I_0 \) is the measured photodiode current of the reagent. The difference in the transmittance value for different concentrations is small once the total protein appears in small quantities in human urine despite its composition, which includes hundred of proteins [2]. The reproducibility variation coefficients for ten different measurements and ten different assays are less than 10%. The
transmittance at $\lambda = 600$ nm as a function of the different total protein concentrations is shown in Fig. 14.

Fig. 15 presents the spectral responsivity for the optical channel (optical effect of the photodiode and the optical filter) measured without the total protein concentration (only with the reagent). The presence of the optical filter limits the light transmitted to the underlying photodiode. However, once it is composed of high-performance dielectric layers, the light transmitted through the optical filter is 92% of the incident light.

The output frequency of the light-to-frequency converter when the MCM-based microsystem is tested for measuring the different total protein concentrations in urine is presented in Fig. 16. This frequency is proportional to the input photocurrent (see Fig. 12) and, hence, to the total protein concentration.

The measurements confirm the direct proportional dependence between the intensity of the color produced by the mixture and the total protein concentration. The calibration curve of Fig. 14 shows a linear behavior at the peak of the absorption spectrum at 600 nm. These results agree with macroscopic measurements performed with well-known total protein standards and using state-of-the-art laboratory equipment [9]. It can also be concluded that the fabricated optical channel is sensitive to a narrow spectral band centered at the wavelength for which the biomolecule being analyzed has its absorption maximum and with a FWHM = 10 nm.

V. CONCLUSION

The MCM microsystem presented here offers a new approach for clinical analysis, especially in biological fluid analysis, due to its portability, ensuring that the analysis can be performed at any location with instantaneous results. Moreover, its optical detection and filtering system avoids the need for expensive readout optics and opens the door to low-cost disposable devices. The performance was successfully demonstrated in the quantitative measurement of the total protein in urine. The photodiode responsivity is 224 mA/W with a FWHM of 10 nm at the wavelength at which the total protein concentration has its absorption maximum, i.e., at $\lambda = 600$ nm. The photodiode sensitivity was 1 kHz/W m$^{-2}$ at $\lambda = 670$ nm, using the TLS230 from Texas Instruments as reference.

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REFERENCES

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