Olabode, Olaitan; Kosunen, Marko; Halonen, Kari

A current controlled oscillator based readout front-end for neurochemical sensing in 65nm CMOS technology

Published in:
ISCAS 2016 - IEEE International Symposium on Circuits and Systems

DOI:
10.1109/ISCAS.2016.7527290

Published: 11/08/2016

Document Version
Peer reviewed version

Please cite the original version:
A Current Controlled Oscillator Based Readout Front-end for Neurochemical Sensing in 65nm CMOS Technology

Olaitan Olabode, Marko Kosunen and Kari Halonen
Department of Micro and Nanosciences
Aalto University School of Electrical Engineering, Espoo, Finland
Email: olaitan.olamilehin@aalto.fi

Abstract—This paper presents the design of an integrated current-controlled oscillator (CCO) based readout front-end for neurochemical sensing applications. The readout front-end chip is implemented in 65 nm CMOS technology and occupies an area of 0.059 mm². The proposed design supports an input current range of 1.2 µA (±600 nA) and can also be configured to support wider current range. The CCO-based structure utilized in this design results in noise averaging of the detected neurochemical input signal due to its inherent ∆Σ first-order noise shaping and anti-alias filtering characteristics. Thus, the prototype chip achieves a current resolution of 100 pA and can detect dopamine concentrations as small as 10 µMol based on measured data from novel diamond-like carbon electrodes. In addition, the digital codes obtained from the readout front-end attain a signal-to-noise (SNR) of 82 dB and linearity limited effective-number-of-bits (ENOB) of 8 at full current range input, without employing any calibration or linearization techniques. The proposed read-out front-end consumes 33.7 µW of power in continuous operation.

I. INTRODUCTION

Sensing and real-time monitoring of neural activities within the central nervous system (CNS) has become a fast-growing area of research due to the need to understand more about how neurons communicate as well as the emerging needs related to personalized healthcare and brain machine interfaces. In addition, further knowledge on how neurons transmit information within the CNS is of significant value to researchers in the field of neuroscience for improving treatment of neurological disorders and neurodegenerative diseases.

Neurons in the CNS are connected by synapses and communicate through electrical and chemical impulses or signals. Transmission of neurochemical signals occurs over short distances in the order of (20 – 30) nm, across chemical synapses; as a result of discharge and absorption of biochemical molecules also known as neurotransmitters [1]. The region or gap between chemical synapses forms a chemical synaptic junction also known as synaptic cleft. The synaptic cleft is filled with an extracellular fluid which aids the chemical reactions that occur during neurochemical transmission. Neurochemical signals are responsible for controlling cognitive, learning and memory functions in the brain. Thus, several neurological disorders such as Parkinson’s disease, Schizophrenia, Alzheimers and Epilepsy have been reported to be associated with abnormal concentration levels of neurotransmitters such as glutamate and dopamine [2]. Hence, the readout and analysis of neurotransmitter concentration levels from the brain provides insight into neurochemical signalling and plays a vital role in the development of more effective treatments for patients suffering from neurological disorders.

Neurochemicals such as dopamine, histamine, norepinephrine, and serotonin; are primarily monitored with the help of potentiostats which operate based on electrochemical transduction principle [3]. Electrochemical transduction principle is the process of applying an electrical potential (Vcell) across an electrochemical cell and measuring the induced reduction-oxidation (redox) current within the cell (Icell) as illustrated in Fig. 1. Thus, this paper proposes a readout front-end for the measurement of neurochemical induced currents based on redox reactions of neurotransmitters within the synaptic cleft. The detected current profiles from the readout chip (DORSI) represents change in concentration levels of neurotransmitters at the neurochemical sensor interface as depicted in Fig. 1. As a result, the detected oxidation and reduction peak potentials help to regulate the voltage applied by neurostimulation electrodes when used in deep-brain stimulation of patients suffering from dopamine-deficient disorders such as Parkinson’s disease [4].

This paper is organized as follows: Section II describes the system level design of the proposed readout front-end. Section III presents post-layout simulation results based on measured data from the neurochemical sensor electrodes. Finally, performance of the proposed design is summarized in Section IV.

II. PROPOSED DESIGN

The main challenge in the design of readout front-ends for neurochemical sensing is the required support of a wide range of input currents while achieving current resolution in pA or less range. Hence, the proposed design of the readout front-end is based on a mixed-signal architecture for minimizing
the effect of noise and achieving high resolution of detected current signals. In addition, sensitivity and selectivity of the sensor electrodes to neurochemicals play an important role in achieving good current resolution from the readout front-end. Thus, the novel diamond-like carbon electrodes used in this work ensures more stable detection of neurochemicals and provides lower background current \( (I_{bg}) \). The input signal is processed within the readout front-end along three main stages, across analog and digital domains as shown in Fig. 2. First, the neurochemical induced current is acquired in the IA stage. Then, the acquired current from the IA stage is converted to frequency in the I-F stage. Lastly, the pulses generated from the I-F stage are quantized in the ID stage such that the digital output from the readout front-end can be further processed by neurostimulation circuitry or externally outside the brain.

A. System Architecture

This section describes the internal structure of the proposed design and noise averaging technique of the current-controlled oscillator (CCO) based architecture. The induced cell current \( (I_{cell}) \) from the neurochemical sensor is processed in the IA stage in order to generate the control current \( (I_{ctrl}) \) for the I-F stage. Then, a simple RC-filter is applied to \( I_{ctrl} \) in order to limit the noise bandwidth. The low-pass filtered \( I_{ctrl} \) signal is conveyed to the CCO which converts the current signal to frequency domain with I-F conversion gain \( (K_{cco}) \) as illustrated in Fig. 3. Thus, the frequency of the oscillator \( (F_{cco}) \) is modulated by changes in the detected input current. The frequency-domain information at the output of the I-F gain stage is integrated to generate changes in phase domain \( \phi_{cco} (t) \). Hence, the continuous change in the phase of signal \( x(t) \) is quantized to the amplitude domain in the ID stage with an integrate-and-dump algorithm.

The quantizer is implemented as an up-counter which is triggered at the rising edge of the generated pulses from the CCO. In addition, the accumulated counter codes \( (\Sigma(n)) \) at the output of the quantizer are sampled to generate discrete representation of the accumulated phase change \( \phi[n] \). As a result, the ID stage is controlled by two clocks namely: \( F_{cco} \) and \( F_s \), which divides the ID block into increment and readout clock domains respectively. The increment clock domain is controlled by an asynchronous clock from the CCO since \( F_{cco} \) varies with changes in \( I_{cell} \). On the other hand, the readout clock domain is controlled by a fixed sampling clock \( F_s \) which synchronizes subsequent processing by the discrete-time derivator and determines the effective data output rate of the system. Hence, the counter utilizes gray-coding in order to mitigate the effect of possible timing violations that may occur at the clock domain crossing (CDC) between the two clock domains as depicted in Fig. 3. In addition, the use of gray-code counters ensures that the minimum error due to possible metastability in the digital circuitry is limited to 1 LSB.

Furthermore, the discrete-time phase sampler defines the integration time \( (T_s) \) of the changes in phase and the output of the sampler \( \phi[n] \) is differentiated in order to obtain the digital output codes \( (D_{OUT}) \). Hence, the CCO-based analog-to-digital (A/D) conversion with integrate-and-dump digital interface results in a cascaded-integrator-comb (CIC) filter which has a continuous-time sinc frequency response. As a result, the integrate-and-dump structure effectively averages the noise over the sampling interval \( T_s \) and prevents aliasing of wide-band noise into the signal band [5]. Thus, the frequency response of this system is given as follows with images on integer multiples of \( F_s \) [6].

\[
[H(f)] = \frac{K_{cco} \ast \sin \left( \frac{\pi \ast f}{F_s} \right)}{\pi \ast f} \quad (1)
\]

\[
[H(0)] = \frac{K_{cco}}{F_s} = K_{cco} \ast T_s \quad (2)
\]

B. Circuit Level Design

This section describes the system level design of the readout front-end and circuit implementations of each signal processing stage of the proposed design. Fig. 4 presents the top-level schematic of the readout front-end based on the system architecture described in the previous Section II-A. The following sub-sections elaborate on the design of each processing block in the system.

1) Current Acquisition (IA): This block represents the analog front-end of the system and controls the cell voltage \( (V_{cell}) \) that is applied across the working electrode (WE) and
As a result, the reference current \( I_{\text{ref}} \) as expressed in equation (4). The reference current \( I_{\text{ref}} \) and reduction \( I_{\text{red}} \) current peaks of neurochemicals during the forward and reverse sweep of the input voltage as depicted in Fig. 5. As a result, the current-to-digital code conversion gain of this block is limited by the signal bandwidth. In addition, the current-to-digital conversion gain of this block is defined by \( F_{\text{cco}} \) and \( F_s \) based on the number of pulses or phase transitions \( (N_p) \) within each sampling interval \( T_s \), where \( N_p \) changes as \( F_{\text{cco}} \) is modulated by \( I_{\text{cell}} \). Therefore, tunability of the \( F_{\text{cco}} \) range offers flexible control of the dynamic range \( (DR) \) and digital code resolution \( (n) \) of the system as shown in the following equations based on equation (2). However, the maximum \( F_{\text{cco}} \) tuning range is limited by non-linearity of the CCO which significantly degrades the effective-number-of-bits (ENOB) of the ADC [5].

\[
\text{Dynamic range } (DR) \approx \frac{F_{\text{cco}}(max) - F_{\text{cco}}(min)}{F_s} \\
\Rightarrow \text{Resolution } (n) \approx \log_2(DR) 
\]

Finally, it should be noted that the derivation stage is evaluated off-chip during post-processing in Matlab. The ID block occupies an area of 0.035 mm\(^2\) and draws an average current of 11.2 \( \mu \)A from 1 V supply.

### III. POST-LAYOUT SIMULATION RESULTS

The readout front-end is implemented in 65 nm CMOS technology and Fig. 6 shows the layout of the fabricated chip. Fig. 7 shows the performance of the I-F and ID blocks based on acquired \( I_{\text{cell}} \) from the IA block. The I-F block and the ID block achieve current sensitivity of 13 kHz/\( \mu \)A and 100 pA/LSB respectively. The I-F block is optimized to provide 16 MHz frequency range from the differential CCO as shown in Fig. 7a, but it should be noted that the current sensitivity of the I-F block can be increased by tuning the CCO to provide wider frequency range. Fig 7b shows that 13.3-bits digital code resolution was obtained from the ID block which could be further increased by reducing the sampling frequency. Fig. 8
The readout of concentration levels of neurochemicals from the brain contributes to the realization of fully-implantable closed-loop interfaces for stimulation of degenerative neurons and control of neural activities. This paper described the design of a CCO-based readout front-end for neurochemical sensing applications. The proposed design was implemented in a 65 nm CMOS process. Post-layout simulation results shows that the readout front-end provides current resolution of 100 pA and could detect minimum dopamine concentration of 10 µMol based on measured data from novel diamond-like carbon electrodes. Higher and lower dopamine concentration than 10 µMol can also be detected from the readout front-end due to its support for a wide current range of 1.2 µA(±600 nA). The digital code representation of the detected dopamine has a resolution of 13.3-bits with RMS conversion error of 0.18 LSB which results in an SNR of 82 dB at full current range input. The achieved resolution of the readout front-end provides good sensitivity of released neurochemicals in the brain which is useful for further understanding of neurotransmitters and fostering research into improved treatments of related neurodegenerative diseases.

IV. CONCLUSION

The authors would like to thank Academy of Finland and Aalto University for funding this research work.

REFERENCES


