



# Reflectance pulse oximetry: Practical issues and limitations

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## Abstract

The demand for reflective-mode pulse oximetry to monitor oxygen saturation has been continuously increasing because it can be used at diverse measurement sites such as the feet, forehead, chest, and wrists. For the wrists, in particular, pulse oximeters are easily available in the form of a band or watch. In this study, we developed a reflectance pulse oximeter and used it to measure oxygen saturation levels at the fingertips and the wrist. We analyzed the performance of this oximeter to address the challenges and limitations associated with using reflective-mode oximeters at the wrist for clinical purposes.

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**Keywords:** Pulse oximetry; Reflective mode; Wrist; Wearable sensor

## 1. Introduction

Pulse oximetry is a noninvasive method for accurately estimating oxygen saturation ( $\text{SaO}_2$ ) by reading the peripheral oxygen saturation ( $\text{SpO}_2$ ). As  $\text{SaO}_2$  and  $\text{SpO}_2$  are sufficiently correlated and pulse oximetry has the advantages of being safe, convenient, inexpensive, and noninvasive, this approach is clinically accepted for monitoring oxygen saturation [1,2].

Pulse oximetry is simple to carry out; it only uses two different light sources and a photodiode [3–5]. Depending on the measurement site, either the transmissive or the reflective mode can be used. In the transmissive mode, the light sources and photodiode are opposite to each other with the measurement site between them. Light then passes through the site. In the reflective mode, the light sources and photodiode are on the same side, and light is reflected to the photodiode across the measurement site.

Currently, the transmissive mode is the most commonly used method because of its high accuracy and stability. Nevertheless, the demand for reflective-mode oximetry is continuously increasing because it does not require a thin measurement site. It can be used at diverse measurement sites such as the feet, forehead, chest, and wrists. In particular, if the wrist is the available

measurement site, pulse oximeters can be conveniently used in the form of a band or watch. To the best of our knowledge, reflectance pulse oximetry, specifically for monitoring oxygen saturation at the wrist, is currently not practiced clinically. In recent years, many reflectance pulse oximeters have become commercially available, but they are only for personal monitoring of oxygen saturation and for entertainment purposes. Obviously, the focus is on developing oximeters for medical purposes, and research is being carried out in this regard. Furthermore, most research papers discuss only the basic principle of pulse oximetry and its utilization in smart devices; only a few papers discuss the challenges associated with reflective pulse oximetry.

In this study, we developed a reflectance pulse oximeter and addressed the practical issues and limitations associated with its use. Using this oximeter, we investigated the results of AC amplitudes and DC offsets of the red and infrared signals, including modulation ratio, which are critical factors to estimate  $\text{SpO}_2$ . We first tested our device by measuring oxygen saturation at a fingertip and wrist and analyzed its performance. Finally, we summarized all issues and discussed the feasibility of using the device for clinical purposes.

## 2. Methods

### 2.1. Principle of $\text{SpO}_2$ estimation

Pulse oximetry measures arterial oxygen saturation based on the light absorption properties of blood. When it combines with

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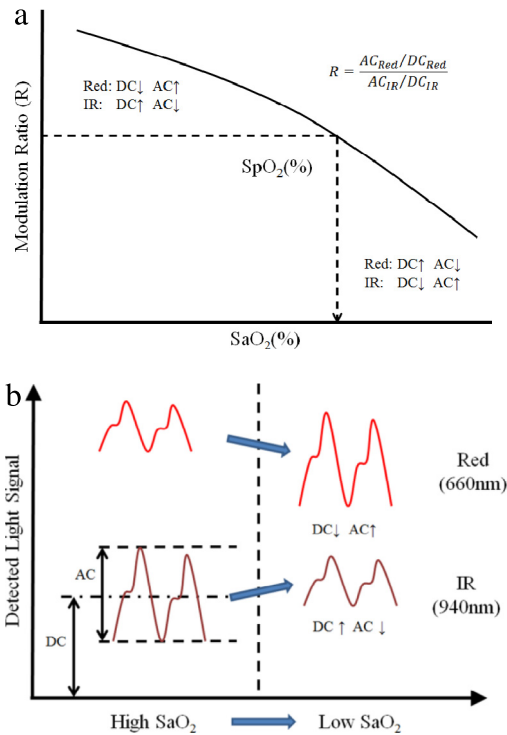


Fig. 1. (a) Changes in  $AC_R$ ,  $AC_{IR}$ ,  $DC_R$ , and  $DC_{IR}$  with decreased  $SaO_2$ . (b) Empirical relationship between modulation ratio  $R$  and  $SaO_2$ .

oxygen, deoxyhemoglobin (Hb) changes its light absorption characteristics. Pulse oximetry exploits the light absorption difference between Hb and oxygenated hemoglobin ( $HbO_2$ ).  $HbO_2$  absorbs more infrared light (660 nm wavelength) and lesser red light (940 nm wavelength) than Hb. In the transmissive mode, light from a pair of red and infrared light-emitting diodes (LEDs) is transmitted through a fingertip. Then, the transmitted light is received by a photodiode on the opposite side. The transmitted light signals consist of a direct current (DC) component and pulsatile alternating current (AC) component. Pulse oximetry calculates the modulation ratio  $R$  by using the DC and AC components of the red and infrared signals as follows:

$$R = \frac{AC_R/DC_R}{AC_{IR}/DC_{IR}}, \quad (1)$$

where  $AC_R$  and  $AC_{IR}$  are the AC amplitudes of the red and infrared signals, respectively [6,7].  $DC_R$  and  $DC_{IR}$  are the DC offsets of the red and infrared signals, respectively. Then, empirically derived calibration curves are used to estimate  $SaO_2$  based on the modulation ratio  $R$ , as seen in Fig. 1(a).  $AC_R$  and  $DC_{IR}$  increase with decreasing  $SaO_2$ , as seen in Fig. 1(b). On the other hand,  $AC_{IR}$  and  $DC_R$  decrease.

## 2.2. Reflectance pulse oximeter

We developed a reflective pulse oximeter where the LED transmission unit and photodiode receiving units are on the

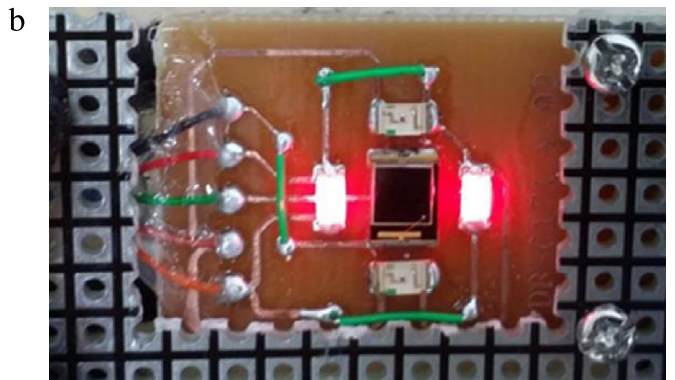
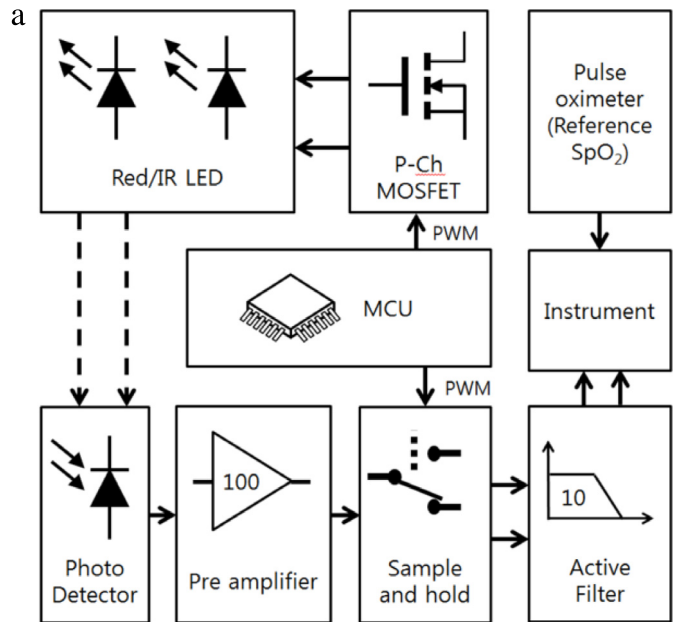


Fig. 2. (a) Control and data flow. (b) Developed reflectance pulse oximeter.

same side. To switch between red and infrared LED lights, the duty cycle was set to 50% with a switching frequency of 500 Hz, which generated a pulse width modulation signal through the microcontroller unit (MCU). Then, each reflected light beam was converted to an electrical signal, which was subsequently amplified with a gain of 100. A sample-and-hold circuit consisting of an analog switch and operational amplifier then separated each amplified signal with a sampling rate of 500 Hz from red and infrared LEDs. Each separated signal was then filtered using a low-pass filter with a cutoff frequency of 10 Hz and an additional gain of 10. For this study, we used an MCP6004 operational amplifier, a TM4C123GHPM MCU, an NJL5310R photodiode, an SML-LX0805SRC-TR red LED, and a KP-2012F3C infrared LED. Fig. 2 shows our design and its implementation in reflectance pulse oximetry. The maximum LED current was 20 mA, and the light intensity was set to the maximum level of the LED capability.

To determine  $AC_R$ ,  $AC_{IR}$ ,  $DC_R$ , and  $DC_{IR}$ , we first detected the pulse peak by incorporating a filter bank with variable cutoff frequencies, spectral estimates of the heart rate, a rank-order

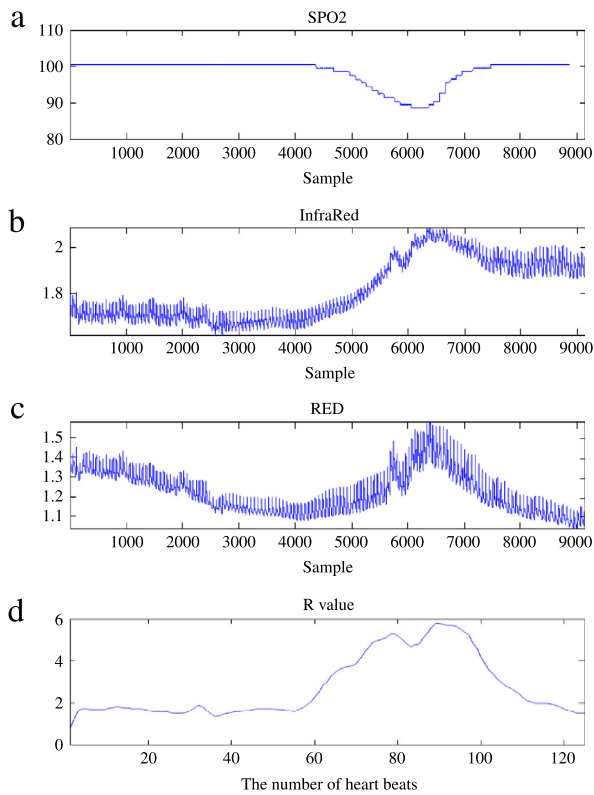


Fig. 3. Results for fingertip: (a) Reference SpO<sub>2</sub> (%), (b) reflected infrared raw signal, (c) reflected red raw signal, and (d) the modulation ratio *R*.

nonlinear filter, and decision logic [8]. Next, we calculated the average values of a 1-s segment signal with each detected peak as its center. Finally, *DC* was calculated from each average value, and *AC* was calculated by subtracting *DC* from each peak value. This method was applied to both red and infrared signals to determine *AC<sub>R</sub>*, *AC<sub>IR</sub>*, *DC<sub>R</sub>*, and *DC<sub>IR</sub>*.

### 2.3. Performance evaluation

We performed two experiments: one at a fingertip and the other at a wrist. In each experiment, the subjects breathed regularly for 40 s and then held their breath for as long as they could. Subsequently, an additional 1 min was given for their SpO<sub>2</sub> levels to return to the normal state. PowerLab 8/35 (ADINSTRUMENTS, Sydney, Australia) was used with an oximeter pod (ADINSTRUMENTS, Sydney, Australia) to measure the reference SpO<sub>2</sub>.

## 3. Results and discussion

### 3.1. Results for the fingertip

Fig. 3 shows the results for the fingertip: the reference SpO<sub>2</sub>, reflected infrared raw signal, reflected red raw signal, and modulation ratio *R*. As expected, the ratio *R* increased with decreasing SaO<sub>2</sub>. Fig. 4 shows the resultant *AC<sub>IR</sub>*, *AC<sub>R</sub>*, *DC<sub>IR</sub>*, and *DC<sub>R</sub>*. When SpO<sub>2</sub> decreased, *AC<sub>IR</sub>* decreased and

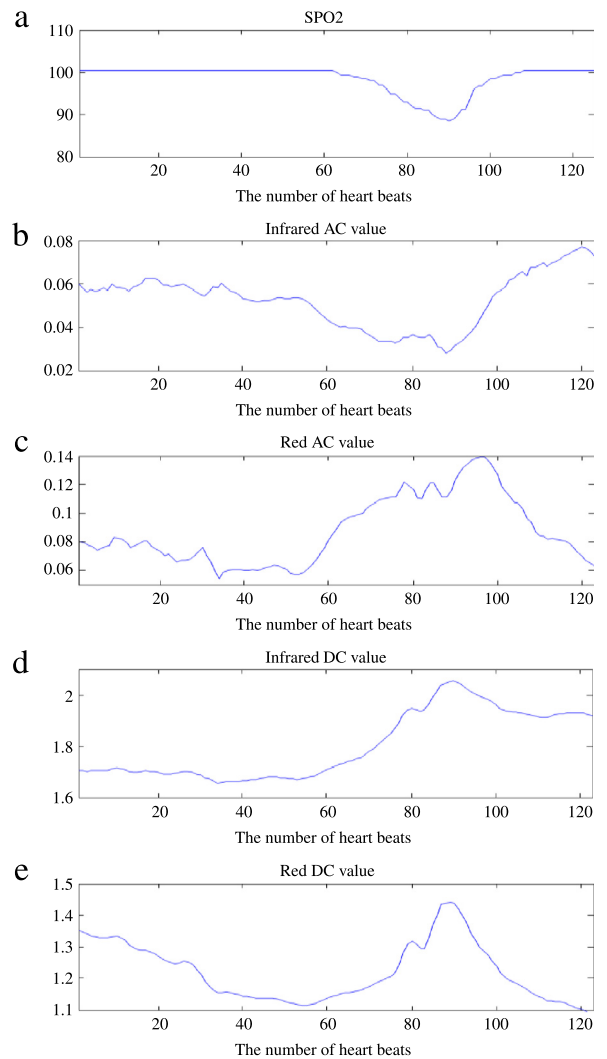


Fig. 4. Results from fingertip: (a) Reference SpO<sub>2</sub> (%) measured at fingertip, (b) resultant *AC<sub>IR</sub>*, (c) resultant *AC<sub>R</sub>*, (d) resultant *DC<sub>IR</sub>*, and (e) resultant *DC<sub>R</sub>*.

*AC<sub>R</sub>* increased, which sufficiently match the light absorption characteristics described in Fig. 1(a) and (b). On the other hand, the reflected DC components partially reflected these characteristics. As SaO<sub>2</sub> decreases, *DC<sub>IR</sub>* should increase and *DC<sub>R</sub>* should decrease. However, in the results, only *DC<sub>IR</sub>* followed this pattern. In fact, *DC<sub>R</sub>* showed an increase. This behavior can be attributed to the fact that the reflective mode is more sensitive to pressure and ambient light sources, which leads to DC instability.

### 3.2. Results for the wrist

Fig. 5 shows the results for the wrist: the reference SpO<sub>2</sub>, the modulation ratio *R*, *AC<sub>IR</sub>*, *AC<sub>R</sub>*, *DC<sub>IR</sub>*, and *DC<sub>R</sub>*. As SpO<sub>2</sub> decreased, the modulation ratio *R* increased as expected. However, the modulation ratio *R* was less stable than that for the fingertip. More specifically, as SpO<sub>2</sub> decreased, *AC<sub>IR</sub>* did not reflect the light absorption characteristics. Furthermore, the

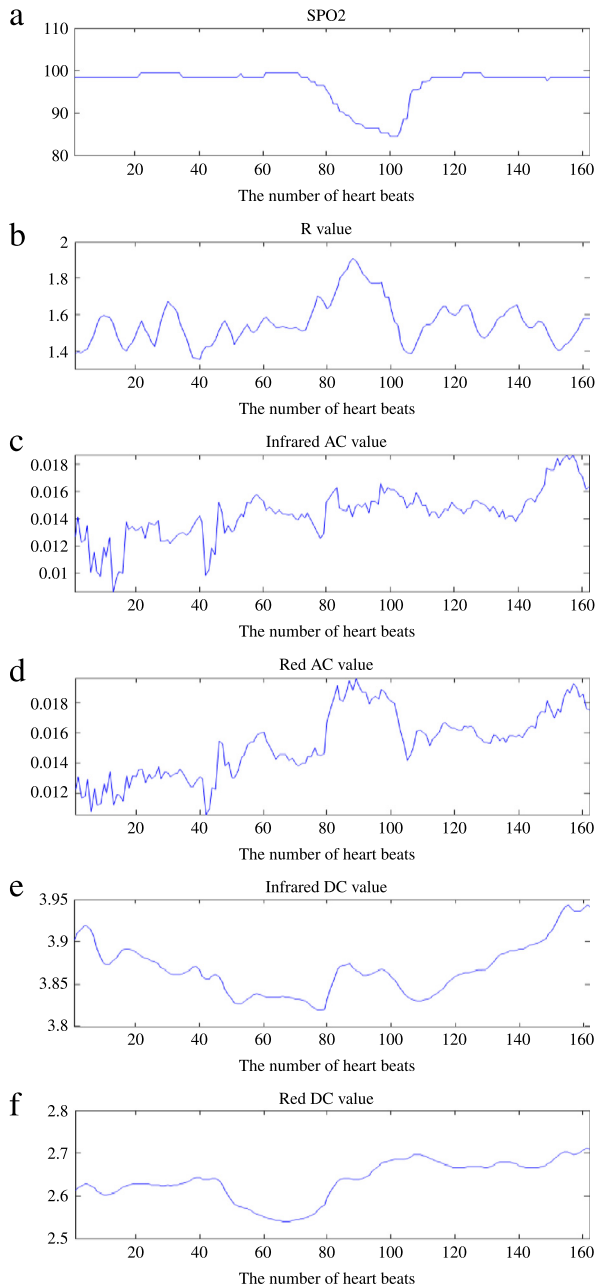


Fig. 5. Results for wrist: (a) Reference  $\text{SpO}_2$ , (b) modulation ratio  $R$ , (c) resultant  $\text{AC}_{IR}$ , (d) resultant  $\text{AC}_R$ , (e) resultant  $\text{DC}_{IR}$ , and (f) resultant  $\text{DC}_R$ .

change in  $\text{AC}_R$  was less pronounced than that observed for the fingertip. Similarly, the reflected DC components did not

reflect the characteristics well. This variation may be because the reflective mode has a low signal-to-noise ratio (SNR) and is sensitive to pressure and ambient light sources.

### 3.3. Discussion

Based on the performance of the oximeter, monitoring  $\text{SpO}_2$  at the wrist using the reflective mode presents challenges with regard to clinical use. Another limitation is that the reflected red and infrared pulses can only be used for specific areas, such as a radial artery; thus, most areas of the wrist are not available for monitoring. In addition, a slight position change at the measurement site significantly affects the performance of the oximeter. Thus, the focus of research studies involving oximetry should be on choosing appropriate measurement sites, and optimizing pressure, ambient light, and SNR.

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