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Use of multiple acoustic reflections to enhance SAW UV photo-detector sensitivity

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Abstract

A new approach to enhance the sensitivity of existing SAW UV-visible photo-detectors based on the monitoring of the multiple reflections of the acoustic waves in the Fourier transform of the frequency response (S_{21}) is demonstrated. By using this concept, it is possible to monitor the UV-visible light in a wide intensity range from very high to ultralow. We present a strategy to obtain an ultrafast SAW UV photo-detector with millisecond response by tuning the deposition conditions of the ZnO film and using the multiple-reflections concept.

Keywords: SAW, surface acoustic wave, UV detector, ZnO

(Some figures may appear in colour only in the online journal)

1. Introduction

Ultraviolet surface acoustic wave (UV SAW) photo-detectors allow us to use the benefits of resistive type photo-detectors to build more complicated SAW-based devices. Typically, a semiconductor layer is deposited onto a channel where SAW propagates, or a layer of the piezoelectric semiconductor itself is a SAW propagation channel [1–5]. An incident light with photon energy higher than a semiconductor band gap is absorbed with generating electron–hole pairs. An electric field of a SAW propagating on the surface of the acoustic channel penetrates the semiconductor and interacts with carriers generated by light. The acousto-electronic interaction results in (i) the increase of insertion loss for the propagating wave and (ii) a reduction in surface-wave velocity, which is observed as a phase shift and a SAW arrival delay [2, 5].

Attenuation, velocity/phase or frequency shift in SAW-based devices are nonlinear functions of an acoustic wavelength, incident light wavelength and intensity. The acoustic wavelength is defined by the geometry of the interdigital transducers (IDTs) and properties of the acoustic channel. Whereas the wavelength selectivity and sensitivity of the ZnO-based SAW photo-detector are defined by the properties of the semiconductor.

There are several approaches to enhance the sensitivity of a SAW photo-detector, namely by designing a device working at higher frequencies [5, 6], by the use of semiconductor nanowire arrays [7] and nanoparticles [8] and/or plasmon nanoparticles [9] to increase the light harvesting. Indeed, the performance of a SAW photo-detector is normally considerably enhanced at higher frequencies. In particular, it was shown that for the ZnO/ZnMgO/ZnO/a-Al₂O₃ UV photo-detector the insertion loss change in the Sezawa wave mode at 711.3 MHz is higher than the one for the Rayleigh wave mode at 545.9 MHz [5]. The high sensitivity observed as an extreme SAW oscillator frequency shift of 1017 kHz was obtained under illumination at 385 nm, 551 $\mu\text{W cm}^{-2}$ [6]. However, the number of oxygen vacancies and structural peculiarities of a light-sensitive ZnO layer has a major effect on the sensitivity and response/relaxation time of a photo-detector [8, 10]. For instance, the SAW oscillator presented by Kumar *et al* [11] was capable of detecting ultralow UV-light intensity of 450 nW cm⁻². However, the detector suffered from relatively slow photo-response (tens of seconds), caused by oxygen deficiency of the ZnO film. Moreover, the majority of the previously reported SAW photo-detectors suffered from inadmissibly slow photo responses and long recovery times of 2.4–30 s [7, 8, 10, 12–14] and only in rare studies were faster responses achieved [9].

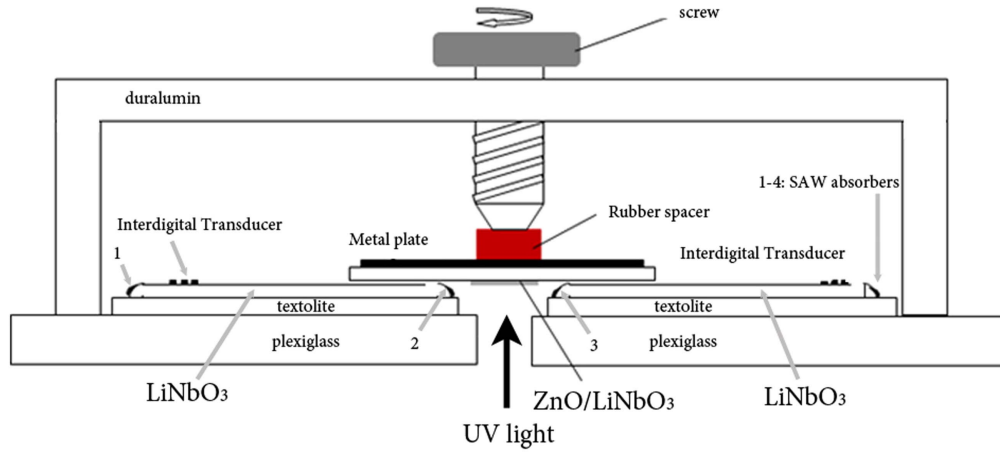


Figure 1. Experimental setup to study the properties of the SAW UV photo-detector.

In this paper, we report on a new approach to enhance the sensitivity of existing ZnO/LiNbO₃ SAW UV-visible photo-detectors. In contrast to previous approaches applied to SAW-based light detection, we use second, third and higher-order reflections of the acoustic wave to monitor low UV-light intensity. We also demonstrate that a ZnO layer prepared under different conditions may drastically modify the major characteristics of a photo-detector. We report on the strategy to build a SAW UV detector, in which both ultrafast photo-response and high sensitivity may be achieved.

2. Experimental details

ZnO films were deposited onto the central part of a (1 × 2) cm 128° YX LiNbO₃ substrate by pulsed laser deposition at an oxygen pressure of 2–5 · 10⁻² mbar and a temperature of 500 °C (Sample 1) and 550 °C (Sample 2), similar to the procedure reported elsewhere [15]. Laser ablation was done by focusing a KrF (248 nm) laser beam on a rotating ZnO ceramic target to give a fluence of 2 J cm⁻². The target to substrate distance was set to be 5 cm and a ~250 nm thick film was deposited for 4000 laser pulses.

To avoid the SAW propagation losses inserted by the edges of the ZnO film, we do not use a mask during the deposition. A homogeneous deposition area of ~1 × 1 cm is defined by laser-plume geometry. Our approach to excite and detect a SAW differs from those applied in previous studies, where IDTs are deposited onto the UV-sensitive film. We use two 128° YX LiNbO₃ substrates with one-directional IDTs on each piece to guide the SAW to the sample and detect the corresponding signal. The examined ZnO/128° YX LiNbO₃ sample is mounted on the top of these two substrates near their edges, as shown in figure 1.

The acoustic channels of the sample transmitter and receiver substrates are positioned on the same line to allow the SAW to propagate from one substrate to another. The SAW generated by the IDT on the first substrate passes to the ZnO/LiNbO₃ sample and then passes to the third substrate with the detecting IDT. The polished LiNbO₃ surfaces are tightly pressed to each other using a metal plate and rubber

spacer, which guarantee a homogeneous press. Thus, a substrate break is avoided. IDTs have a central frequency of 120.7 MHz. The LiNbO₃ substrate edges are rounded and are covered by absorbers. SAW absorbers should not touch the examined sample so that it does not block the propagating wave. We deliberately separate the ZnO/ LiNbO₃ sample from the IDTs. This approach ensures that we do not deposit the IDTs onto the examined film, thus providing more flexibility for technological procedures, such as heat treatment and further growth of nanostructures onto the semiconductor film, without the risk of the metallic constructions being damaged. Note that a high-quality ZnO crystalline film needed as a high-quality detector is normally obtained at an elevated temperature of 450 °C–650 °C. The possible contamination of the ZnO film surface with chemicals or film degradation during the lithography step are also excluded in this way. The use of this approach is a forced arrangement and a good alternative to the lift-off procedure often used to overcome the technological difficulties [16].

3. Results and discussion

Figure 2 shows that an x-ray diffraction (XRD) curve of the deposited ZnO film revealed peaks at 34.5°, 32.9°, 68.8°, corresponding to ZnO (002), LiNbO₃ (104) and (208) crystal orientations with ZnO (002)∥LiNbO₃ (104) figure 2. The full width at half maximum (FWHM) for the ZnO (002) peak is 0.3°, showing the high crystalline quality of the film.

When the ZnO/LiNbO₃ structure is exposed to UV light, extra carriers are generated in the ZnO layer. The SAW velocity change (Δv) and wave intensity attenuation (Γ) induced by acousto-electric interaction are given by ¹⁶

$$\frac{\Delta v}{v_0} = \frac{k_{ef}^2}{2} \frac{1}{1 + \sigma^2/\sigma_m^2} \quad \Gamma = k_{ef}^2 \left(\frac{\pi}{\lambda} \right) \frac{\sigma/\sigma_m}{1 + \sigma^2/\sigma_m^2},$$

where λ , σ , $k_{ef}^2 = \frac{e_0^2}{\epsilon c} \ll 1$ and $\sigma_m = \omega_m \epsilon \epsilon_0$ denote a SAW wavelength, ZnO film sheet conductivity, effective electro-mechanical coupling coefficient and the material constant, respectively. Thus, when the sheet conductivity of the ZnO

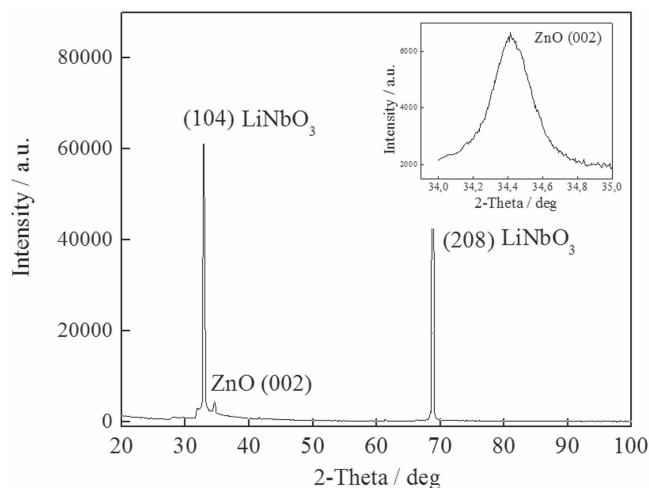


Figure 2. XRD spectrum of a typical ZnO/LiNbO₃ film. The ZnO (002) peak is shown in the inset.

film is limited to σ_m value, the attenuation Γ of a SAW reaches a maximum. The initial sheet resistance of the ZnO layer for Sample A grown at 500 °C was as high as $\sim 6.75 \times 10^3 \text{ Ohm} \cdot \text{cm}$ and is several times higher than the one for Sample B ($1.375 \times 10^3 \text{ Ohm} \cdot \text{cm}$) for which the temperature was increased to 550 °C. Under illumination by UV light of a HeCd laser at 325 nm with intensity of 3.2 mW cm^{-2} the sheet resistance decreased to $\sim 6.52 \times 10^3$ and $\sim 1.1 \times 10^3 \text{ Ohm} \cdot \text{cm}$ for Sample A and B, respectively. The frequency response (S_{21}) of the ZnO/ LiNbO₃ SAW photo-detector (Sample A) is shown in figure 3(a). Once a SAW starts propagating, the response reveals an interference picture. The distance between neighboring maxima is equal to $\sim 70 \text{ kHz}$, which corresponds to a time delay of $\sim 14.3 \mu\text{s}$, which is the time needed for a SAW with a velocity of 3980 m s^{-1} to propagate the distance of $\sim 5.7 \text{ cm}$ between IDTs. The SAW detector-time response, which is a Fourier transform of the S_{21} characteristic is shown in figure 3(b). When a SAW is reflected from the IDTs, second, third and higher-order reflections are observed at 28.4, 42.6 μs , etc. The absence of the additional reflections due to intersections of pieces was guaranteed by the absence of extra peaks in the Fourier transform of the frequency response. The attenuation of a SAW propagating in Sample A with a low conductivity of the ZnO layer is very low and, thus, up to seven SAW reflections are detected, as shown in figure 3(b). More specifically, a ‘native’ SAW attenuation induced by the ZnO layer in the dark can be roughly estimated from the first to second peak intensity ratio. Note that a SAW attenuation caused by reflection from the IDT also contributes to this value. In particular, in Sample A the SAW is attenuated for 9.4 dB, because of the acousto-electronic effect caused by carriers present in the ZnO film and a reflection from the IDT. Even at the highest applied incident UV-light intensity, the photo-detector does not show complete attenuation of the S_{21} characteristics, as shown in figure 3(a). Two opposite cases of peak attenuation, corresponding to the high/low incident UV-light intensity of 280 and 2.8 mW cm^{-2} are illustrated in

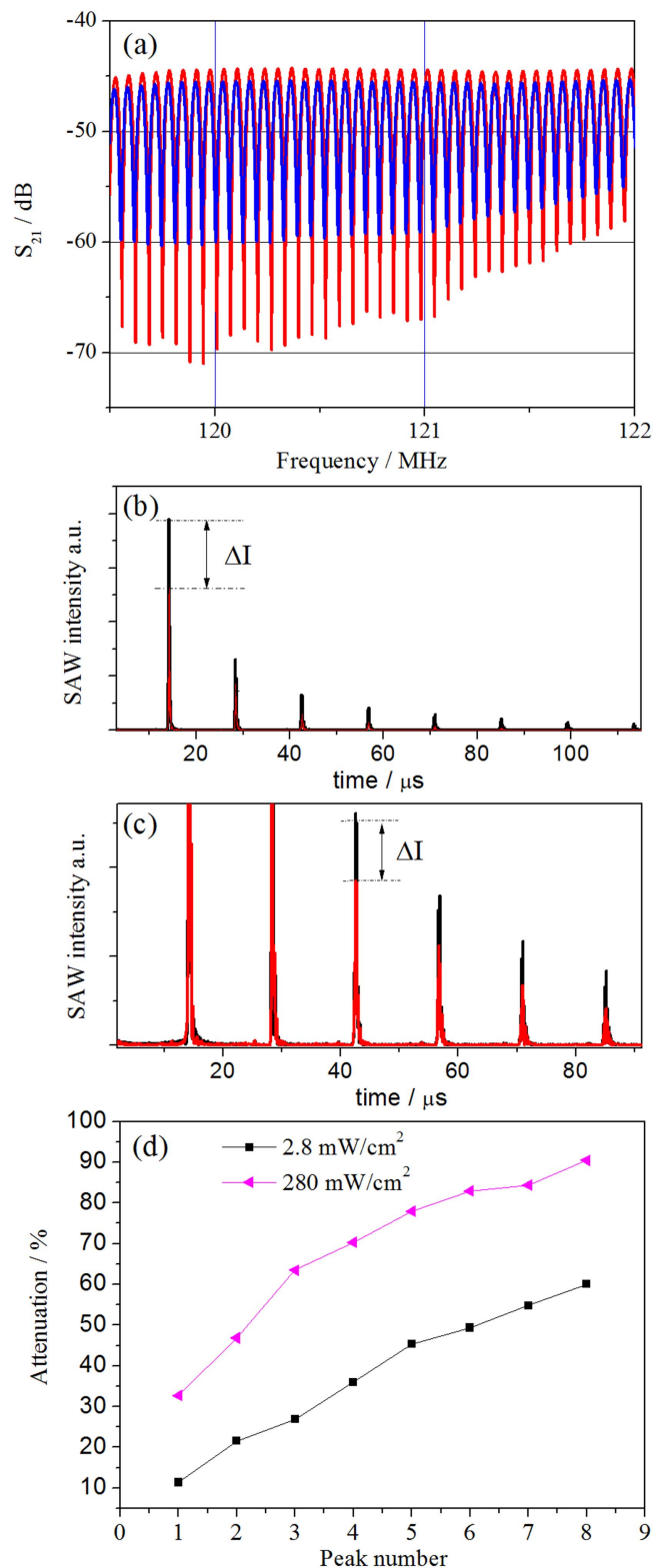


Figure 3. Frequency response (S_{21}) of the ZnO/LiNbO₃ SAW detector (Sample A) in dark (red) and under illumination by UV light at 325 nm and intensity of 280 mW cm^{-2} (blue) (a); Fourier transform of S_{21} characteristics with (red)/without (black) UV light at intensity of 280 mW cm^{-2} (b) and 2.8 mW cm^{-2} (c); attenuation of peaks in time response characteristics of the SAW detector (d).

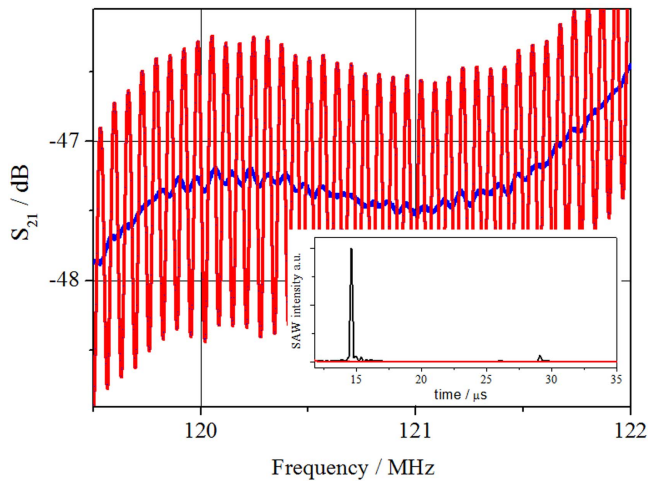


Figure 4. Frequency response (S_{21}) of Sample B in the dark (red) and under illumination by UV light at 325 nm and 280 mW cm^{-2} (blue) and corresponding Fourier transform of S_{21} characteristics with (red)/without (black) UV light.

figures 3(b) and (c). Attenuation of the peaks, corresponding to the SAW propagating through the Sample A several times, is presented in figure 3(d). When the light intensity is as high as 280 mW cm^{-2} the first peak of the S_{21} Fourier transform characteristics is attenuated by $\sim 30\%$ and, thus, may be used to measure the UV-light power, as shown in figure 3(b). For low incident intensity of 2.8 mW cm^{-2} the first peak at $14.3 \mu\text{s}$ does not change much and it cannot be used for light detection any more. However, the attenuation of higher-order SAW reflections is feasible to monitor the light intensity. In particular, the attenuation of $\sim 27\%$ and $\sim 60\%$ for the 2nd and 7th SAW reflections were detected, respectively, as illustrated in figure 3(d). The UV-light-induced SAW delays did not exceed $\sim 0.02\text{--}0.04 \mu\text{s}$ even at the highest used light power and, thus, they cannot be seen in figures 3, 4 and may hardly be used for reliable light detection.

The ZnO film of Sample B with higher conductivity shows only $\sim 2\text{--}3$ SAW reflections from the IDTs, which evidences that a propagating SAW is considerably attenuated even in the dark. From the first to second peak intensity ratio, one can find that the SAW attenuation in the ZnO film induced by acousto-electric effect in the dark is already as high as 28,1 dB. The sheet conductivity of the ZnO layer for Sample B is much higher than the one for Sample A and, according to equation (1), is much closer to σ_m value even in the dark. When the detector is exposed to UV light, the film conductivity is further increased and is limited to σ_m value, which is accompanied by additional SAW attenuation. In contrast to Sample A, the frequency response (S_{21}) of Sample B shows complete saturation at UV-light intensity of 280 mW cm^{-2} , as shown in figure 4. As can be seen from the Fourier transform of the S_{21} characteristic in the inset of figure 2, the SAW propagating in Sample B is nearly completely attenuated already after the first pass.

A typical photo-electric response of Sample A and B at moderate intensity of 3.2 mW cm^{-2} is shown in figure 5. Sample A shows fast response and conductivity recovery

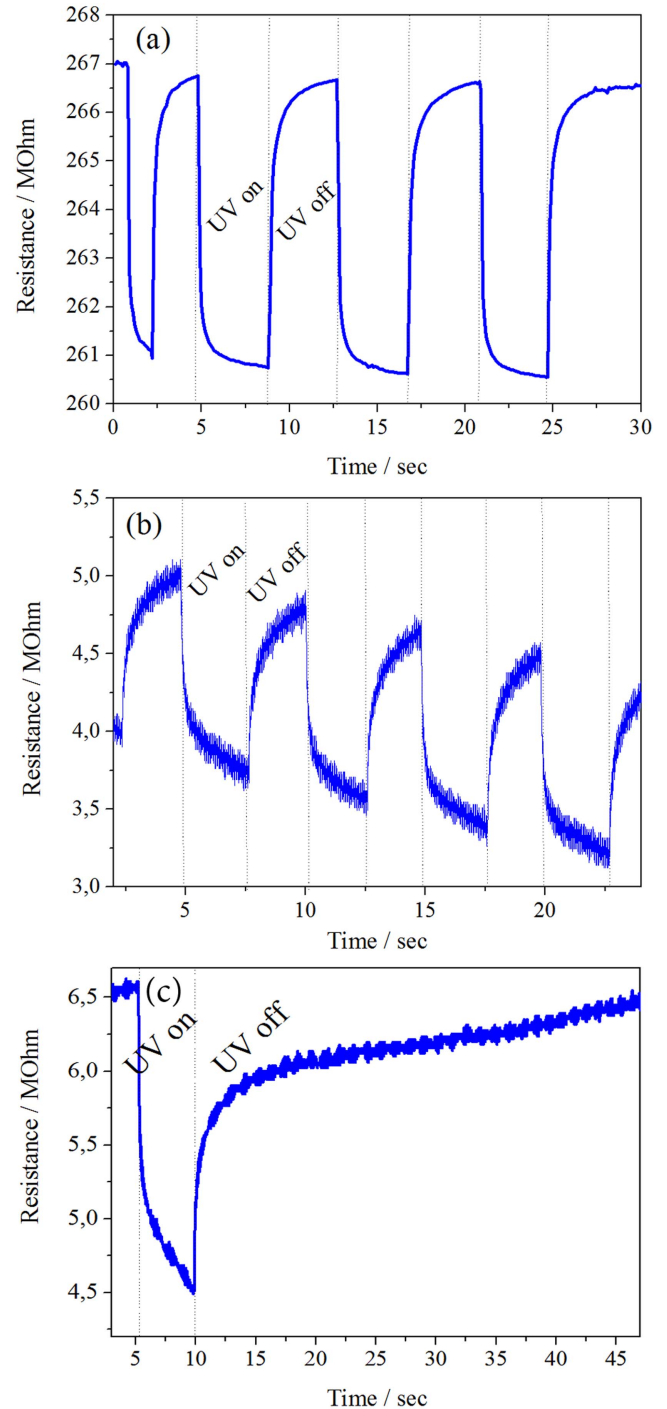


Figure 5. ZnO resistance change in Sample A (a) and B (b), (c), measured with/without illumination by UV light 325 nm (HeCd laser) and intensity of 3.2 mW cm^{-2} .

time, as presented in figure 5(a), whereas the resistance of Sample B drops and recovers very slowly ($\sim 40 \text{ s}$), as shown in figures 5(b), (c). Note that the photo-response kinetics shown in figure 3 was measured using a mechanical chopper and presented here as a qualitative illustration of the difference in response of the two detectors. The time-resolved photo-response of the ‘fast’ photo-detector A is characterized more accurately below. The long recovery time of the ZnO film for Sample B is similar to those reported previously

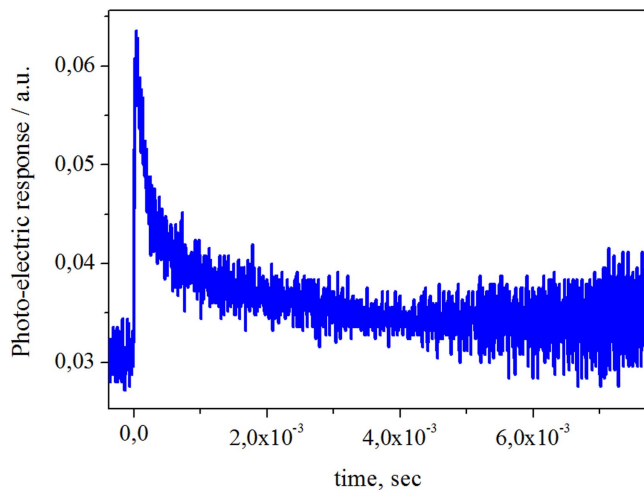


Figure 6. Time-resolved photo-electric response of ZnO film in Sample A, measured under illumination by 15 ns pulse of the third harmonic of the YAG:Nd³⁺ laser at 355 nm with fluence of 23 mJ cm⁻².

[7, 8, 10, 12–14] and is likely caused by oxygen-related long-living trap states inside of the semiconductor band gap.

To accurately characterize the photo-electric response kinetics of the ZnO film for Sample B, we measured a resistance change kinetics under excitation by 15 ns laser pulse of the third harmonics of the YAG:Nd laser.

The measured photo-response shown in figure 6 was fitted with a double exponential function as:

$$I = A_1 \exp\left(-\frac{t}{\tau_{\text{slow}}}\right) + A_2 \exp\left(-\frac{t}{\tau_{\text{fast}}}\right)$$

The corresponding fast and slow components of the kinetics were found to be $\tau_{\text{fast}} = 0.14 \pm 0.08$ ms and $\tau_{\text{slow}} = 1.3 \pm 0.01$ ms. The recovery time is several orders of magnitude shorter than those for other UV SAW photo-detectors previously presented, where recovery times as long as 2.4–30 s were reported [7, 8, 10, 12–14]. The time, which the SAW with velocity of 3980 m s⁻¹ spends to propagate 1 cm under the ZnO film is ~ 2.5 μ s. As the photo-response relaxation time is more than ~ 160 times longer than the time during which the SAW propagates in the ZnO layer, one may consider that the process is stationary, i.e. the ZnO sheet conductivity does not change during the acousto-electronic interaction of carriers with the wave. The relaxation time of the SAW photo-detector with the ZnO film, similar to that of Sample A, is limited by the slowest process, i.e. by the relaxation time of the photo-generated carriers. As shown above, the ZnO film in Sample A (with higher sheet resistance) shows fast photo-electric response, but lower UV-light induced change in resistance. Therefore, a relative change in the intensity in the fundamental peak of the S₂₁ Fourier transform is lower than the one for Sample B. If using a fundamental peak change for photo-detection, one could consider that the ZnO film similar to the one in Sample B is more advantageous. However, the high sensitivity of the ZnO film in Sample B towards UV light is accompanied by very slow response. To overcome this ‘slow response’ limitation,

one may design a photo-detector with a ZnO layer of the Sample A type and use one of the SAW reflections for detection as a better alternative. In this way, a SAW propagates several times back and forth and the resulting effect of the SAW interaction with light is larger. Thus, the drawbacks related to low sensitivity of ZnO film can be easily compensated by using our approach to monitor multiple SAW reflections of a photo-detector device.

4. Conclusions

In summary, we presented a new approach to enhance the sensitivity of the existing SAW UV-visible photo-detectors by using multiple reflections of the acoustic waves. We show that the ultrafast SAW UV photo-detector with millisecond response may be obtained by tuning the deposition conditions of the ZnO film. Applying the concept of monitoring the intensity of the multiple SAW reflections in Fourier transform of the frequency response (S₂₁) an ultrahigh sensitivity of a SAW photo-detector may be achieved. To detect high incident light intensity, one may monitor only the fundamental peak of a time response.

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