High Quantum Efficiency Ultraviolet Avalanche Photodiode

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This thesis is dedicated

to my parents

Abstract

Ultraviolet (UV) radiation covers the wavelength range from 10 nm to 400 nm. The detection of UV radiation presents a wide range of civil and military applications, such as chemical and biological analysis, flame detection, inter-satellite communications and astronomical studies. For many decades, the detection of UV light has been accomplished primarily with photomultiplier tubes (PMT), which have the beneficial characteristics of high sensitivity with low noise. But they are also fragile, need large power sources, which make systems bulky and heavy, and are relatively expensive. For some application, avalanche photodiodes, which are semiconductor detectors, are potential replacements for PMTs. Among all the material candidates, 4H-SiC has the most promising material properties and APDs based on SiC have achieved excellent performance with high sensitivity and good robustness. However their relatively narrow spectral response has limited their use for deep-UV and near-UV detection. This dissertation describes my work on the deep-UV enhancement of SiC-based APDs and a study of GaAs/AlGaAs-based APDs to achieve comparable dark current, multiplication gain and excess noise performance to SiC but with enhanced responsivity for near-UV detection.

From my study of the spectral response of SiC PIN-structure APDs, the weak deep-UV response was found to be related to the high surface recombination velocity at the top surface and the short diffusion length in the top highly doped layer. Two different approaches were investigated to enhance the deep-UV response of these SiC APDs. First,

a SiC metal-n⁻-p structure was examined. By replacing the highly doped top layer with a semitransparent metal contact in the active area, deep-UV photons are absorbed within the high electric field depletion region. Photodiodes with external quantum efficiency (EQE) greater than ~40% at wavelength between 200 nm and 235 nm were demonstrated with this structure. Second, an NIP structure with a graded doped top layer was studied. The high-low junction formed due to the graded doping in the top layer is expected to provide an electric field that pushes carriers away from the surface and, thus, suppresses surface recombination. However no significant improvement in the deep-UV response was observed. The possible reason is that the electric field generated by the high-low junction is not large enough to compensate the electric field in the opposite direction induced by the surface band bending.

As an alternative method to achieve high deep-UV response, while keeping SiC as multiplication layer material, an AlGaN/SiC heterogeneous structure photodiode was explored. As early effort, a separate absorption and multiplication (SAM) structure was utilized so that the spectral response could be tuned by changing the Al mode fraction in the AlGaN absorption layer. However, it was found that the electric field is confined in the SiC multiplication region by the polarization-induced charge at the AlGaN/SiC interface. Since there is a lattice mismatch between AlGaN and SiC, high defect densities were found in the AlGaN layers. Consequently, an electric field is needed in this layer to achieve good collection efficiency. An AlN inter-layer was simulated to be effective in introducing an electric field in the AlGaN layer, but experimental results showed that the AlN acted as a barrier that prevented carriers in the AlGaN absorption layer from entering the SiC multiplication layer. To overcome this problem, an AlGaN top layer with higher Al fraction

was used. With a bandgap of over 5.38 eV, this layer is transparent to the deep-UV photons. The photodiodes with this structure achieved high peak external quantum efficiency of $\sim 76\%$ at 242 nm.

The second part of my work focused on near-UV detection using the GaAs/AlGaAs material system. Al_{0.8}Ga_{0.2}As, with a band gap of 2.23 eV, is a promising candidate to replace Si for near-UV detection. An Al_{0.8}Ga_{0.2}As PIN-structure APD was designed, fabricated, and characterized. These APDs have exhibited very low dark current of 179 nA/cm² at gain as high as 200. Also the measured excess noise factor was very low. The k factor, which is the ratio of the electron ionization coefficient, α , to that of the holes, β , is about 0.15, which is comparable to Si. The peak external quantum efficiency was ~26% at 470 nm. The response in the near-UV was lower than the expectation. Secondary ion mass spectrometry (SIMS) measurements showed that the doping in the p- layer is significantly higher than the specification that was sent to the foundry. The relatively low near-UV response is due to the fact that a large fraction of the carriers were generated in the undepleted p- layer.

My final research project was to utilize recessed window and surface texturing techniques to enhance the near-UV response of GaAs/AlGaAs photodiodes. The surface texturing using nanosphere natural lithography provided a photodiode surface with very low reflectivity (<5%) across a wide spectrum. Combined with a recessed window structure, the GaAs/AlGaAs photodiode exhibited an external quantum efficiency in the range 45% to 55% from 300 nm to 850 nm. These photodiodes were fabricated into arrays and sent to NASA for a project to reduce the number of cameras in satellites.

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Chapter 1. Introduction

1.1 Ultraviolet radiation detection

The ultraviolet (UV) spectrum, corresponding to photon energies from 3.1 eV to 124 eV is divided into the following subdivisions [1.1]:

Near UV	NUV	400 nm – 300 nm
Middle UV	MUV	300 nm – 200 nm
Far UV	FUV	200 nm – 122 nm
Extreme UV	EUV	122 nm – 10 nm

Ultraviolet light has high photon energy, which can activate many chemical processes. The most important natural UV source is the sun. Approximately 9% of the energy emitted by the sun is in the UV range. But the stratospheric ozone layer prevents wavelengths shorter than 280 nm from reaching the Earth's surface. Also UV radiation with wavelength shorter than 200 nm will be absorbed by oxygen in the atmosphere.

The detection of UV radiation is important for a wide range of civil and military applications, such as chemical and biological analysis (ozone, pollutants, and most organic compounds have absorption lines in the UV range) [1.2], flame detection (fire alarm and missile plume), and short-range secure communication [1.3].

The primary requirements for UV detectors are high quantum efficiency, low dark current, high gain, and low noise. In additions, for some applications, solar-blind operation is very important to eliminate the background noise. Solar-blind operation requires a cutoff in detection for wavelengths longer than 280 nm, since the radiation shorter than 280 is strongly absorbed by the stratospheric ozone layer. Traditionally, UV detection has been achieved using photomultiplier tubes (PMTs). They exhibit high sensitivity in the UV and large detection area. However, they are bulky, fragile, subject to magnetic fields, expensive, and they have limited dynamic range. As an alternative, semiconductor photodetectors have the advantages of smaller size and good reliability. For some applications avalanche photodiodes (APDs), which utilize impact ionization to achieve the internal gain, could be a potential replacement for PMTs.

1.2 Material candidates for UV detection

Due to its well-established technology, Si is the most common material choice for APDs for UV detection, although commercial GaAs-based UV photodiodes are also available. However, owing to its narrow bandgap, the peak quantum efficiency of a Si APD is in the visible range. For UV-detection, visible light from the environment will result in a high background noise and expensive filters are required. Also, for high-sensitivity applications, the detector must be cooled to reduce the dark current [1.4].

UV photodetectors based on wide-bandgap semiconductors such as SiC and III-nitrides have the potential to exceed the performance of Si APDs in some areas [1.5-1.6]. The wide bandgap itself is an important advantage for photodetectors, since it enables roomtemperature operation and provides intrinsic visible-blindness. In addition, the thermal conductivity of wide bandgap materials is, in general, significantly higher than silicon, which makes them suitable for high temperature and high power applications.

1.3 Review of SiC PIN-structure APDs

4H-SiC, with an indirect bandgap of 3.2 eV, is particularly suitable for UV detection. Previous students in our group have done extensive studies on 4H-SiC APDs and developed various technologies to improve the performance [1.7-1.9]. State-of-art SiC APDs exhibit extremely low dark current of 0.18 μ A/cm² at a gain of 1000. By incorporating a recessed, window a peak external quantum efficiency of 65% has been achieved [1.10]. This structure is shown in Fig. 1-1. However, the photo response decreases rapidly below 260 nm and has a long tail out to 380 nm, which limits their use for solar-blind applications. This relatively narrow spectral response limits the applications in which SiC APDs can be deployed. The measured external quantum efficiency of the structure in Figure 1-1 is shown in Figure 1-2.

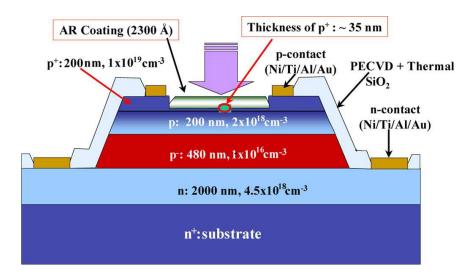


Figure 1-1. Structure of SiC PIN APD with recess window

Beginning from the top, the structure consists of a 200 nm p+ layer (1.0×10^{19} cm⁻³), a 200 nm p layer (2.0×10^{18} cm⁻³), a 480 nm p⁻ layer (1.0×10^{16} cm⁻³) and a 2000 nm

n buffer layer (4.5×10^{18} cm⁻³). The p+ contact layer was etched from an initial thickness of 200 nm to about 35 nm in the central active region to form a recessed window. Also a 230 nm SiO₂ was deposited as an anti-reflection coating (optimized for 260 nm).

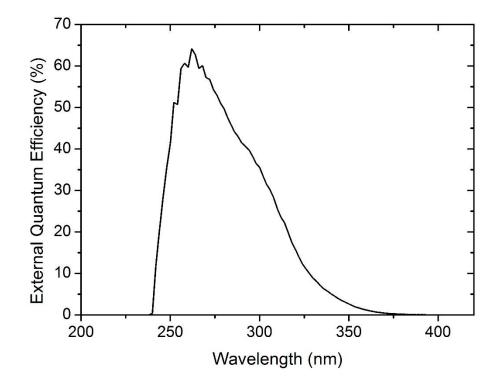


Figure 1-2. Measured external quantum efficiency of SiC PIN APD

My work focused on developing new photodetectors with high efficiency in the spectral regions where current SiC photodiodes have poor response. The discussion will be divided into two parts, the deep-UV (200 nm - 250 nm) and the near-UV (300 nm - 400 nm). The work in the deep-UV will be SiC-based and the work in the near-UV spectrum will utilize GaAs and AlGaAs.

1.4 Organization of dissertation

This dissertation is divided into 7 chapters. In chapter 1, I have briefly introduced the background of UV detectors and the performance of SiC PIN APDs. In chapter 2, I describe the general process of fabricating AlGaN/SiC- and GaAs/AlGaAs-based APDs and present the experimental characterization techniques for photodiodes. Chapter 3 focuses on the different approaches that I have investigated to improve the deep-UV response of 4H-SiC detectors. In chapter 4, AlGaN/SiC heterogenous structure photodiodes are demonstrated as an alternative way to achieve good deep-UV detection. Both a separate absorption and multiplication (SAM) structure and a transparent window structure were studied. In chapter 5, Al_{0.8}Ga_{0.2}As, as a material candidate to replace SiC and Si for near-UV detection, is described. In Chapter 6 I discuss my work using a recessed window structure combined with a surface texturing technique to achieve high quantum efficiency from near-UV to near-IR in GaAs detectors. Chapter 7 concludes my work and discusses possible future research directions closely related to this work.

Chapter 2. Fabrication and characterization of photodetectors

The first part of this chapter will discuss fabrication of III-nitride/SiC photodiodes and GaAs/AlGaAs photodiodes. Since these two material systems are very different in their fabrication processes, they will be discussed separately. In the later part of this chapter I describe several of the measurement setups used to characterize the devices developed in this work.

2.1 Fabrication of AlGaN/SiC Photodiodes

2.1.1 Sample preparation

SiC wafers purchased from Cree Inc. are often four-inch double-side-polished wafers. Usually there will be markers on the back side of the wafer. But sometimes there are no markers on either side of the wafer. In this case, a standard sheet resistance measurement using a four-point probe setup is used to determine on which side the epitaxial layers were grown. The resistivity of the epitaxial layer is usually several orders of magnitude higher than the highly doped substrate. For the AlGaN/SiC sample, the epitaxial layer side can be determined from the color.

After this step, the wafer is scored with a diamond scribe in order to cleave it into small pieces. The relative locations of the small pieces are recorded. Then one of the pieces is cleaned using the following procedure: First the sample is submerged in an ultrasonic cleaner for 5 minutes, and then rinsed with methanol and de-ionized (DI) water. After cleaning, the sample is blown dry using a nitrogen gun and baked at 90 °C on a hot plate

for 1 minute to make sure the water is fully evaporated. This step is necessary since water will affect adhesion of photoresist in the following steps.

2.1.2 Mesa patterning

Both SiC and III-Nitride are very hard materials, which makes it difficult to pattern them. No standard wet etch technique with acceptable etching rate at room temperature is available. So dry etching using reactive ion etch (RIE) and inductively coupled plasma (ICP) etch were used.

Depending on the required etch depth, different types of photoresist can be used to pattern mesas. Usually, AZ4620, a thick photoresist, is used to achieve etch depths deeper than 1µm. Prior to coating with photoresist, Hexamethyldisiloxane (HMDS), an adhesion promoter for photoresist, is dropped on the wafer and spun at 4000 revolutions per minute (rpm) for 30 s. Next AZ 4260 photoresist is applied to the wafer and spun at 4000 rpm for 40 s. The wafer is then baked for 3¹/_h minutes at 90 °C. Then the sample is exposed with a Suss MicroTech MJB-4 aligner using the "mesa" mask. The exposure time is different for different photomask materials, e.g., 160 seconds is used for a quartz mask and 250 seconds is used for soda lime glass mask. The exposed photoresist is then developed in a solution of AZ 400K diluted to a ratio1:3 AZ 400K: de-ionized water. The total time for this step can range from 40 s to 1 min. Following confirmation that the photoresist has been completely developed by observing under a microscope, the wafer is baked again at 130 °C for five minutes. This will make the photoresist reflow and harden, so that a small bevel angle at the edge of the photoresist is formed. This step is critical for the AlGaN/SiC mesa

etch, because this bevel angle will be transferred to the device during the dry etch, which lowers the electric field at the sidewall. The profile of the patterned photoresist is measured with a profile meter. Typically the thickness of the photoresist is $\sim 7 \,\mu$ m and the bevel angle is less than 20 degrees.

The mesa is etched by inductively coupled plasma (ICP) with an Oxford ICP 100 system using the following recipe. RF power 160 W, ICP power 1000 W, and pressure of 6 mTorr using 10 sccm Ar and 40 sccm Cl_2 at -25 °C. A small drop of fomblin oil is placed on a silicon wafer carrier with a thick SiO₂ layer; the wafer is placed on top of the oil. This prevents the wafer from overheating during the etch process, which can result in uneven etching or difficulty removing the photoresist. The etch rate is approximately 5 nm/s for SiC, and 10 to 20 nm/s for AlGaN. The etching selectivity of SiC relative to photoresist is approximately 1:3. After etching, the photoresist is removed by ultrasonic cleaning in acetone. An optional oxygen plasma with power 200 to 300 W can be used if the acetone clean does not work well. This is followed by a spin clean with acetone, methanol, and deionized water.

2.1.3 Passivation

In order to remove ICP etch residues, the wafer is first immersed in 25% ammonium hydroxide in an ultra-sonic bath for 10 minutes. Then a modified RCA cleaning process is performed. The sample is first dipped in 1:1:4 mixture of NH₄OH, H₂O₂, and DI water at 75 °C for 5 minutes to clean the organic residues. Then it is dipped in buffered oxide etchant (BOE) for 1 minute to remove native oxides and some of the ionic contaminants. Finally

the sample is dipped in 1:1:4 mixture of HCl, H_2O_2 , and DI water at 75°C for 5 minutes to remove the remaining traces of ionic contaminants. If the wafer has an AlGaN layer, the sample is loaded onto the heated shelf of the March oxygen plasma cleaner immediately after the RCA clean to perform thermal oxidation. An oxidation of 15 minutes followed by immersion in BOE for 1 minute is repeated 3 times to remove the surface damage. The thin layer of oxide from the last oxidation is kept as a passivation layer. If the wafer is pure SiC, an oxidation in the furnace at higher temperature is preferred. Instead of loading the wafer into an oxygen plasma cleaner, the wafer is placed in a quartz boat and loaded into the center in the oxidation furnace. A thermal oxidation at 1050 °C for 3.5 hours is used to form a thin silicon oxide passivation layer. This is followed by SiO_2 deposition by plasma enhanced chemical vapor deposition (PECVD) to achieve the final anti-reflection coating thickness. An SiO₂ layer is then deposited by PECVD using the following recipe: temperature = 285 °C, RF power = 33, reflected RF power = 0, pressure = 800 mTorr and gases; 105 sccm SiH₄, 400 sccm N₂O and 500 sccm N. The rate of deposition is approximately 10 nm/min. The same recipe is used for SiO_2 deposition for all fabrication described in this dissertation. Based on the reflectance calculations, a specific thickness is used to create an anti-reflection coating optimized at a desired wavelength.

2.1.4 Contact formation

AZ nLOF 2020 negative photoresist is used to define the pattern for contact formation. The photoresist is applied to the sample and spun at 5.5 krpm for 30 seconds and then baked at 110 °C on a hotplate for 1 minite. The pattern is defined by exposure for 1.5 seconds using the "contact" mask. This is followed by 1 minute post-exposure bake at 110 $^{\circ}$ C and developing in AZ 300 MIF for about 30 seconds. Oxygen plasma etch using a March plasma cleaner at power 150 W for 1 min is used to remove the undeveloped photoresist and ensure that the contact window is open. A chemical etch in BOE removes the SiO₂ in the contact window for access to the semiconductor. The etch rate is approximately 110 nm/min; over etching by 15 seconds ensures all SiO₂ is removed. The wafer is rinsed thoroughly in de-ionized water and blown dry with the nitrogen gun. The wafer is then placed in the March plasma cleaner again for oxygen plasma etch at the same power and time. This etch is to remove any residue left behind following the SiO₂ removal and it also ensures a clean surface for metal deposition.

The wafer is then loaded into the electron beam evaporator. Four layers of metal are deposited for both p and n contacts on SiC in the following sequence: Ni 25 nm / Ti 35 nm / Al 100 nm / Ni 80 nm. For the n contact on AlGaN, V 15 nm / Al 80 nm / V 20 nm / Au 100 nm is used. The photoresist and unnecessary metal is removed by metal lift-off. Initially the wafer is soaked in acetone for 30 mins and then light ultra-sonic agitation is used to remove any remaining metal. The wafer is spin cleaned with acetone, methanol, DI water and dried. A contact anneal by rapid thermal anneal (RTA) for 3 minutes at 850 °C in vacuum is then performed to achieve ohmic contacts and lower the contact resistance.

If no contact pad is required the device is ready for testing. If a contact pad is required nLOF 2020 is again used to pattern the contact pad. A two layer metal stack is used for the contact pad: Ti 20 nm / Au 150 nm. Metal lift-off completes the fabrication and the devices are ready for testing.

2.2 Fabrication of GaAs/AlGaAs Photodiodes

There are some similarities between the fabrication steps for III-nitride/SiC and GaAs/AlGaAs photodiodes. Therefore I will only discuss the steps that are unique to the GaAs/AlGaAs photodiode fabrication process. The photolithography process for mesa definition is similar, but instead of using AZ4620, a thinner photoresist AZ5214 is used. A 1:1:10 mixture of H₃PO₄, H₂O₂ and DI water is used to etch the mesas. The etch rate for GaAs is about 300 nm/s. For AlGaAs, the etch rate will increase with Al mole fraction. The typical etch rate for Al_{0.8}Ga_{0.2}As is 1000 nm/s.

The mesa etch is followed by sulfur sidewall passivation. For this procedure, the wafer is soaked in 5% ammonium sulfide solution for 5 minutes and then rinsed with DI water and blown dry using a nitrogen gun. After that, a 300 nm-thick layer of SiO₂ is deposited by PECVD to isolate the passivated sidewall from the air. The contact window is formed in the same manner as above but the metal contact stack is Ti 30 nm / Pt 50 nm / Au 100 nm. The wafer is then annealed using RTA at 450 °C for 30 seconds in nitrogen. All the other steps are the same as the fabrication of AlGaN/SiC photodiodes.

2.3 Characterization

The following characterization techniques are used to determine photodiode and avalanche photodiode performance.

2.3.1 Current-voltage and capacitance-voltage measurements

DC current-voltage (I-V) measurements are made using a low-noise probe station capable of low current measurement (as low as 1fA with sufficient integration time) and an HP 4156B semiconductor parameter analyzer. A 30 W deuterium lamp is used as a broad spectrum UV light source for the photocurrent measurement. Photocurrent and dark current are measured versus bias voltage. Avalanche gain at a particular voltage is calculated using the flowing formula:

$$M = \frac{I_{photo} - I_{dark}}{I_{photo_unity} - I_{dark_unity}}$$

where I_{photo} and I_{dark} are the photocurrent and dark current, respectively. I_{photo_unity} and I_{dark_unity} are the photocurrent and dark current at a predetermined bias where the avalanche gain is insignificant. A bias that is ~50% of the breakdown voltage is usually used as the unity gain point for a PIN structure.

The capacitance-voltage (C-V) measurement utilizes the same probe station. The semiconductor parameter analyzer is replaced by an HP 4275A multi-frequency LCR meter. Open-circuit and short-circuit calibrations are performed prior to measurements. The capacitance–voltage measurement is useful in determining the reach-through voltage of a separate absorption and multiplication (SAM) structure APD. It can also be used to determine the depletion width of photodiodes.

2.3.2 External quantum efficiency measurements

The external quantum efficiency is the ratio of the number of carriers collected to the number of photons incident on the photodiode at unity gain. In some publications, responsivity is used instead of quantum efficiency. Responsivity, which has a unit of amperes per watt, is the unity-gain photocurrent per incident power at a given wavelength from the photodiode. The responsivity can be calculated from quantum efficiency by the following formula:

$$R_{\lambda} = \eta_{\lambda} \lambda \cdot \frac{q}{hc} \approx \eta_{\lambda} \lambda \cdot \frac{1}{1240 \, W \cdot nm/A}$$

where λ is the wavelength, η_{λ} is is the quantum efficiency at given wavelength, q is the elementary charge, c is the speed of light in vacuum, h is Planck's constant, and R is the responsivity.

A laser driven light source, which has significantly higher power in the UV spectrum than other wide spectrum light sources such as a Xenon lamp is used in our quantum efficiency setup. The light from the light source is coupled into a monochromator, which functions as a tunable, narrow spectral width source from the UV to the visible. The original blazed grating in the spectrometer, which has a blaze wavelength of 1 μ m was replaced by a new blazed grating with a blaze wavelength of 250 nm to further enhance the UV light power from the spectrometer. The output from the monochromator is coupled into a 300 μ m-core UV enhanced fiber that has low attenuation in the UV. From the fiber output two uncoated microscope objectives placed back-to-back collimate the beam and focus the light to a small spot on the active area of the device. It is important that all the light is focused on the device in order to get an accurate measurement. A chopper placed in front of the fiber input provides frequency modulation to the signal (usually 200 – 300 Hz). A Stanford Research SR-850 lock-in amplifier measures the photocurrent of the device. The photocurrent response of a calibrated UV-enhanced silicon detector is also measured. The quantum efficiency and responsivity of the device under test (DUT) can then be calculated from the following formula:

$$\eta_{DUT} = \eta_{ref} \cdot \frac{I_{DUT}}{I_{ref}}$$

where η_{ref} is the quantum efficiency of the calibrated silicon detector. I_{DUT} and I_{ref} are the test device and reference photocurrents.

2.3.3 Excess noise measurements

The excess noise measurement characterizes the noise that arises from fluctuations in the gain of an avalanche photodiode. Avalanche gain results from impact ionization of the carriers. The carriers gain energy from the high electric field and initiate impact ionization. However, impact ionization is a stochastic process. Not all the carriers experience the same gain. Excess noise can be characterized by the excess noise factor F(M).

The excess noise factor is measured by a HP8970B noise figure analyzer. An Ar-ion laser that operates at 351/363 nm is used as the light source. The device is biased by a Keithley 2400 source meter through a bias tee and the AC component is fed into the noise figure analyzer. The center frequency of the HP8970 is chosen so that ambient noise is minimized.

The excess noise figure, F(M), of an APD can be calculated from the spectral noise density, S, where S is given by Eqn. 2-1 below. In the equation I₀ is the photocurrent at unit gain, $R(\omega)$ is the frequency dependent impedance of the system and M is the gain.

$$S=2qI_0 R(\omega)M^2 F(M)$$
 Eqn. 2-1

The gain can be calculated based on the DC current read from the Keithley source meter. $2qR(\omega)$ is determined by measuring the noise versus different photocurrent, I₀, at unity gain. Also, the effective impact ionization coefficient ratio k is can be calculated using McIntyre's local mode equation [2.1]:

$$F(M) = kM + (1-k)(2-1/M)$$
. Eqn. 2-2

2.3.4 Spatial uniformity measurements

A spatial uniformity scan provides a picture of the spatial response of the detector, which is useful to determine if there are "hot spots" such as edge breakdown or microplasmas. It can also reveal low response in the active area due to high lateral resistance. For this measurement, the chopped UV light from the Ar-ion laser is focused onto the APD by a UV lens with a spot size less than 5 μ m. The lens is mounted on a motorized micropositioner that can be controlled by a computer. A labview program controls the movement of the micropositioner and collects photocurrent from the lock-in amplifier.

2.3.5 Surface Reflection

Owing to the significant refractive index difference between the semiconductor and air, part of the incident optical signal is reflected, a factor that limits the quantum efficiency. Therefore it is important to minimize the light reflection at the operating wavelength. The reflectance is characterized with a Perkin Elmer Lambda 950 spectrometer. The measurement steps are similar to the quantum efficiency measurement. First the optical power reflected by the semiconductor wafer is measured. Then a calibrated mirror with known reflectance is measured under the same conditions. The reflectance of the sample can be calculated by multiplying the reflectance reading from the spectrometer by the reflectance of the calibrated mirror.

Chapter 3. Deep-UV Enhanced 4H-SiC Avalanche Photodiode

As discussed in the first chapter, 4H-SiC is very suitable for UV detection. The wide bandgap results in very low dark current. It also has a favorable impact ionization coefficient ratio. High performance SiC APDs with high peak responsivity, low dark current, low excess noise, and high gain with diameters up to 250 µm have previously been demonstrated by our group. However, the narrow spectral response limited its applications in deep-UV detection. In this chapter, I discuss the source of the poor deep-UV response of SiC PIN APD and different approaches to enhance the deep-UV performance.

3.1 Spectral response study of SiC PIN APD

3.1.1 Spectral response modeling and analysis

In order to better understand the spectral response curve that is shown in Figure 1-2, the quantum efficiency of the 4H-SiC APD with PIN structure shown in Figure 1-1 is modeled using a simple method that is described in Ref [3.1]. The idea of the modeling is that for the PIN structure, the total photocurrent measured externally is due to i) electrons collected from the p layer at the p-i depletion edge, ii) electron-hole pairs collected from within the depletion region, and iii) holes collected from the n layer at the i-n depletion edge. It is assumed that all carriers generated within the depletion region are collected, while only a fraction of the carriers generated in the undepleted p and n regions are

collected. The collection efficiency depends on the layer thickness, diffusion length, and surface recombination velocity.

The modeled spectral response was fitted to the measured data by adjusting the fitting parameters. The fitted external quantum efficiency curve and the measured curve are shown Figure 3-1.

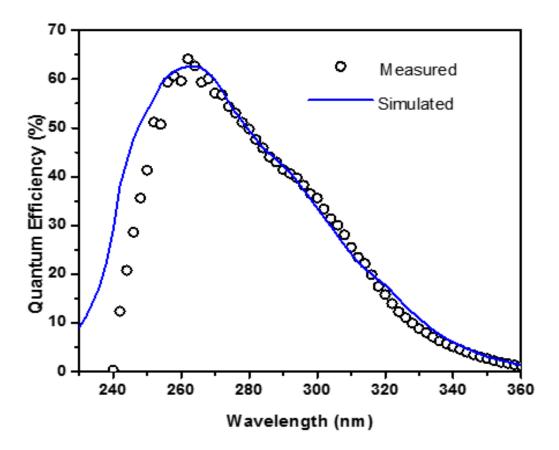
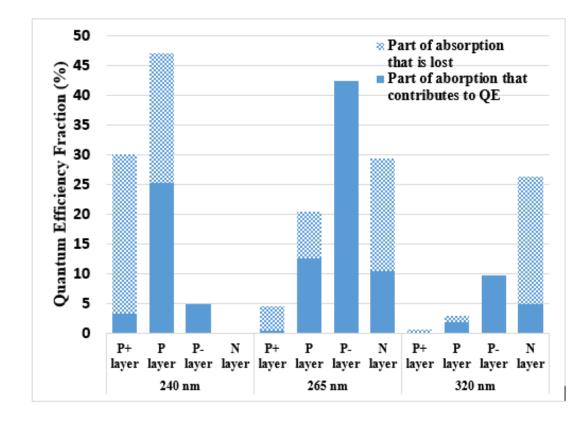
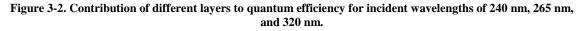


Figure 3-1. Measured and Simulated Quantum Efficiency of SiC APD

To better illustrate this, the contribution of different layers to the external quantum efficiency is compared in Figure 3-2 at three typical wavelengths (240 nm, 265 nm, and 320 nm). Since SiC has an indirect bandgap, the absorption near the band edge is very weak. Therefore at 320 nm, most of the photons are absorbed in the n buffer layer and the

substrate. Due to the short diffusion length in the highly doped material, many of the carriers are absorbed in the n buffer layer and the substrate. Owing to the short diffusion length in the highly doped material, many of the carriers recombine before reaching depletion region. The absorption profile was optimized for operation at 265 nm, so that most of these carriers are absorbed in the depletion region and, thus, high quantum efficiency was achieved.





At 240 nm, most of the photons are absorbed in the top two layers. Similar to the carriers generated in the n layer, the photons generated in the highly doped p region also have short diffusion lengths. But there is another effect that also limits the deep UV

response, surface recombination. Surface recombination is characterized by the surface recombination velocity.

3.1.2 Surface band-bending

Surface recombination is caused by band bending at the surface [3.2]. This is illustrated in Figure 3-3. Surface defects such as dangling bonds, pin the Fermi level near the middle of the band gap at the surface. This creates a region where photo generated carriers are pushed toward the surface where they quickly recombine through trap states instead of diffusing to the depletion region where they contribute to the photocurrent and are collected. Surface recombination is characterized by the surface recombination velocity. The reported surface recombination velocity of SiC ~1 × 10⁸ cm/s [3.3], which is about 6 orders of magnitude higher than that of Si [3.4].

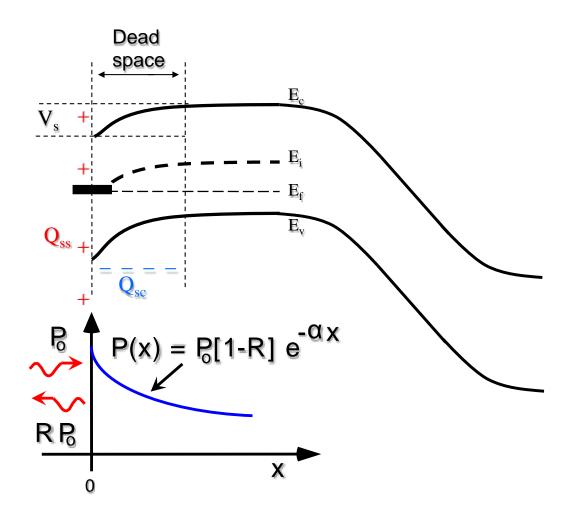


Figure 3-1. Band diagram showing surface band-bending (top) and the spatial distribution of absorbed photons (bottom)

In summary, SiC has a narrow spectral response. The poor response at wavelengths shorter than 260 nm is due to the shallow absorption, the high surface recombination velocity, and short diffusion length. The slowly decreasing response at wavelength longer than 300 nm is due to fact that the absorption is weak at the band edge for the indirect bandgap material.

3.1.3 Approaches to enhance the deep-UV response of SiC photodiode

From the study of the spectral response of the SiC PIN APDs, we know that the deep-UV response is greatly affected by three parameters: top doped layer thickness, diffusion length, and surface recombination velocity.

Firstly, a thinner top highly-doped layer will reduce the percentage of photons that are absorbed in this layer, which will minimize the loss of the photon-generated carriers due to the short diffusion length and high surface recombination velocity. Usually this is accomplished using a recessed window structure, which has already been studied by previous students in our group and proved to be effective if the recessed depth is optimized. Theoretically greater deep-UV enhancement will be achieved with deeper recessed windows. But as the top highly-doped layer becomes thinner, the lateral resistance will increase to the extent of causing non-uniform response across the active area. In order to overcome this problem, a thin layer of semitransparent metal can be deposited on the active area of the photodiode. A metal-n⁻-p structure utilizing this approach is discussed below in 3.2.

Secondly, it will also be beneficial to increase the diffusion length. However, since the diffusion length strongly depends on material quality and the doping level, modifying it is not straightforward. Instead of increasing the diffusion length, a high–low junction can be incorporated into the highly-doped layer to provide an effective built-in electric field. This electric field can assist drift into the depletion region and increase efficiency. Also the electric field created by the high-low junction near the top surface will block carriers from reaching the top surface, thus suppressing surface recombination. This approach is

described in section 3.3

Since high surface recombination velocity is the most detrimental factor for the deep-UV performance of SiC, I have also analyzed surface band bending and found that it is due primarily to the dangling bonds that arise from termination of the periodic crystal structure. There are two ways to solve this problem: a) Use a transparent window layer on top of the absorption material. This is widely used in InP/InGaAs photodiodes in which InP acts as transparent window layer at telecom wavelengths [3.5]. A similar structure with AlGaN acting as window layer on SiC is discussed in chapter 4. b) Choose suitable material to passivate the top surface in order to lower the surface state density. For example, sulfur is a good passivation material on GaAs [3.6] and hydrogen is found to be effective on Si [3.7]. Similar surface passivation treatment on SiC discussed in chapter 7.

3.2 SiC metal-n⁻-p photodiodes

From the analysis of the photo-response of SiC in the last section, we know that if most of the photons are absorbed in the depletion region, high quantum efficiency can be achieved. Therefore different Schottky [3.8] and metal-semiconductor-metal (MSM) structures [3.9] have been studied by other groups. As an alternate approach, we examined SiC a metal-n⁻-p structure, where the n- absorption region is fully depleted at low bias.



Figure 3-4. Device structure of the SiC metal-n⁻-p photodiode

The SiC metal-n⁻p diode structure is shown in Figure 3-4. From bottom to top, it consists of a 2 μ m-thick p+ SiC layer and a 350 nm n- SiC layer. A thin semi-transparent metal contact consisting of Ni 5nm/Au 5nm was deposited in the central active region to ensure that the electric field is uniformly distributed across the active area and to make it semitransparent in the 200 nm to 400 nm spectral range. A semitransparent metal with the same thickness was also deposited on a sapphire wafer to measure the transmission spectrum. The measurement was done with the Perkin Elmer Lambda 950 spectrometer. The measured transmission of the semitransparent contact is shown in Figure 3-5. For comparison, a conventional p-n-n+ SiC photodetector was fabricated consisting of a 2- μ m p+ SiC layer, a 480 nm n- SiC layer, and a 100 nm n+ SiC layer.

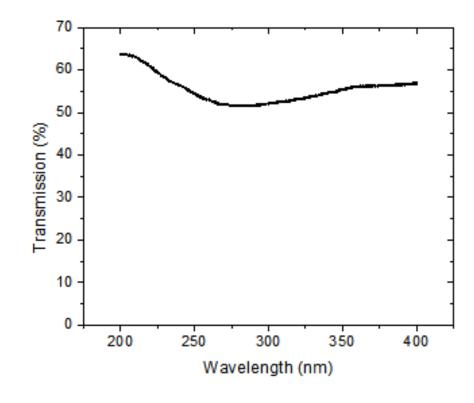


Figure 3-5. Transmission of the semitransparent contact

I

The current-voltage characteristic of a typical 50 µm-diameter metal-n⁻-p photodiode is shown in Figure 3-6. Dark current less than 0.02 nA is observed for reverse bias voltage less than 60 V. Gain of 50 is estimated at reverse bias of 90 V.

The measured external quantum efficiency at unity gain of typical metal-n⁻-p and p-nn+ diodes are shown in Figure 3-7. The response of the p-n-n+ photodiode peaks at 270 nm and sharply decreases at shorter wavelength. In contrast, the response of the metal-n⁻p photodiodes is nearly flat from 270 nm to 225 nm with a slight decrease below 225 nm and a peak external quantum efficiency of ~45%. This response is greater than that observed for the p-n-n+ diode for wavelength shorter than 240 nm despite losses due to the semi-transparent metal contact. After accounting for these losses, the internal quantum efficiency at 255 nm is 81%, which is significantly higher than that for p-n-n+ devices.

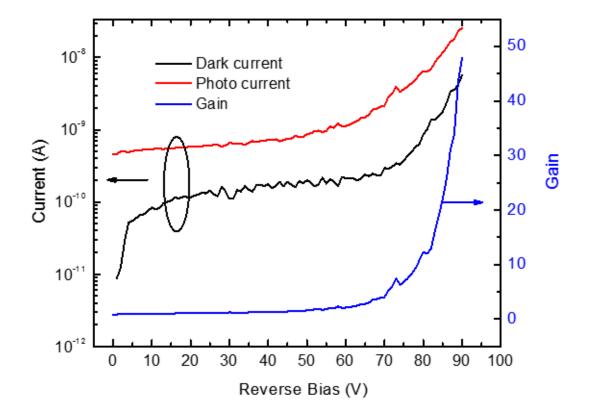


Figure 3-2. I-V of SiC metal-n⁻-p photodiode.

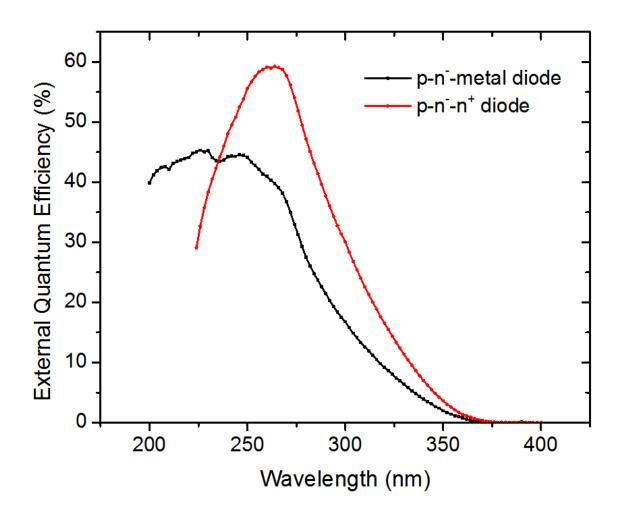


Figure 3-7. QE of metal-n⁻-p photodiode compared with n+-n⁻-p photodiode.

The enhancement of photo-response at short wavelength is attributed to improved collection of photo-generated carriers through drift within the high electric field region near the surface of the lightly-doped SiC absorption region despite the presence of surface states, while the observed decrease in photo-response at wavelengths shorter than 255 nm indicates that surface recombination still plays an important role.

3.3 SiC NIP photodiode with graded doping n layer

As noted above, modeling of the spectral response of the SiC PIN APD shows that the high surface recombination velocity is responsible for the poor deep-UV response. Most of the carriers generated near the surface will move toward the surface and recombine through the states in the forbidden band. In order to stop the carriers from reaching the surface, a high-low junction utilizing a graded doping profile is incorporated into the highly-doped layer. Figure 3-8 shows the device structure design of the SiC NIP photodiode with graded doping in the top n-type region.

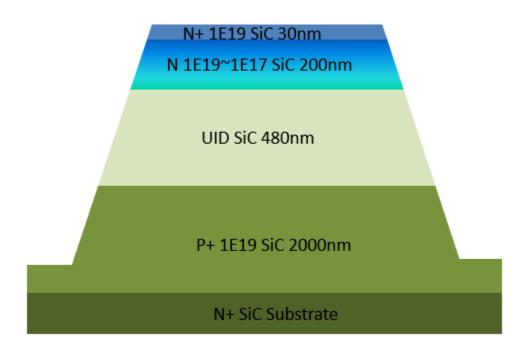


Figure 3-8. Cross section of the SiC NIP photodiode design

The wafer was grown by Etc-Epi Inc. From bottom to top, the structure I designed consists of a SiC n+ substrate with 2 µm p+ layer, a 480 nm unintentionally doped (UID)

layer, a 200 nm n layer with graded doping density changing from 1×10^{17} cm⁻³ to 1×10^{19} cm⁻³, and a 30 nm n+ top layer. The 30 nm top layer was added because approximately 30 nm of SiC will be consumed during the thermal oxidation step in the photodiode processing. We also sent a small piece of this wafer for Secondary Ion Mass Spectrometry (SIMS) analysis to verify the wafer doping profile and layer thickness. The SIMS result is shown in Figure 3-9. Al is the p type dopant for SiC and N is the n type dopant. The SIMS analysis shows that most of the layer thicknesses and the doping profile, including the graded doping, were well within the target range. However, the top layer thickness was measured to be 150 nm instead of the targeted 30 nm.

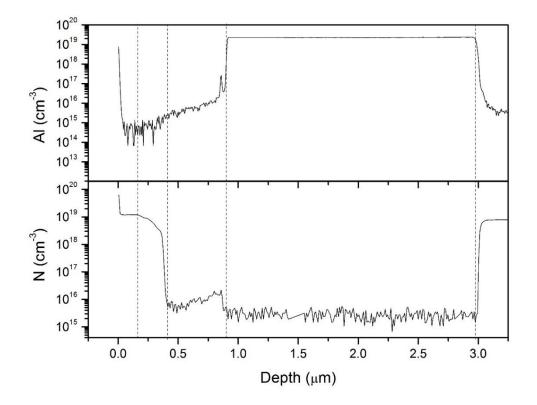


Figure 3-9. SIMS analysis of the SiC NIP wafer

The wafer was fabricated into photodiodes using the process flow described in chapter 2. However, due to the thick top layer, a recessed window was added after mesa patterning; 120 nm of the top layer in the active area was removed by ICP etching. The etch depth was confirmed by the profile meter. Also the thickness of the SiO₂ anti-reflection (AR) coating was optimized for 240 nm.

The I-V characteristic of a 50 μ m-diameter device is shown in Figure 3-10. The photodiode exhibited very low primary current near breakdown and high gain over 1000 was achieved. Dark current of ~1 pA was observed at gain of 100.

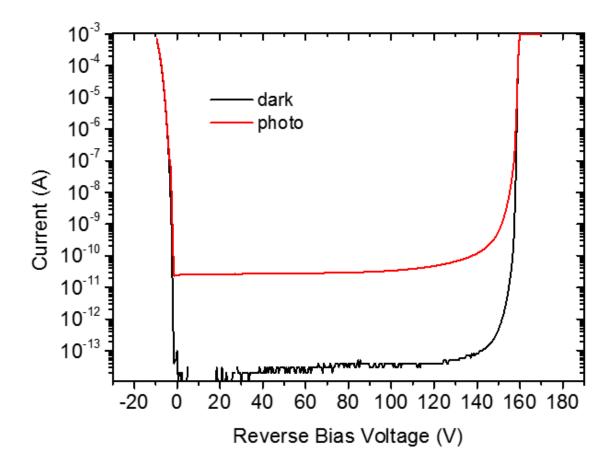


Figure 3-10. I-V characteristics of the SiC NIP photodiode

External quantum efficiency measured at unity gain compared with a SiC PIN photodiode that was described earlier in this chapter with same i layer thickness is plotted in Figure 3-11. The measurement yielded peak quantum efficiency of ~55% at 270 nm, which is about 7% lower than the PIN structure. This might be due to the fact that the AR coating thickness was optimized at 240 nm instead of the peak quantum efficiency wavelength. It was also observed that for wavelength shorter than 240 nm the efficiency is enhanced, which is consistent with the motivation of this design. The quantum efficiency at 240 nm was improved from ~1% to ~30%, which is believed to be a result of the built-in electric field in the n layer created by the graded doping profile.

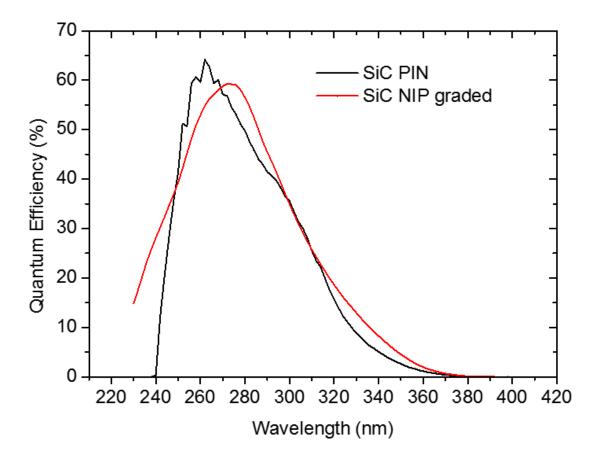


Figure 3-11. External quantum efficiency of the SiC NIP APD with graded n layer

3.4 Conclusion

In this chapter, the spectral response of the SiC APDs with PIN structure has been studied by external quantum efficiency modeling. The contributions from carriers generated in different layers are analyzed. The results show over 75% of the deep-UV photons are absorbed in the two highly-doped top layers, but less than half of these carriers diffuse into the high electric field region and get collected. This is due to the fact that the diffusion length in the highly-doped SiC is relatively short and to the high surface recombination velocity. The surface band bending, which is the physical explanation of the high surface recombination, is also discussed.

Different approaches to enhance the deep-UV response of the SiC APD are discussed and two of them are investigated in this chapter. The SiC metal-n⁻-p photodiode demonstrated flat external quantum efficiency of ~45% from 270 nm to 225 nm with a slight decrease below 225 nm. This response is greater than that observed for the p-n-n+ diode for wavelength shorter than 240 nm despite losses due to the semi-transparent metal contact. The deep-UV response enhancement results from the improved collection of photo-generated carriers through drift within the high electric field region near the surface of the lightly doped SiC absorption region despite the presence of surface states. However, these devices exhibited high dark current due to the high electric field at metal/SiC interface.

The SiC NIP photodiode with graded doping profile in the n layer utilizes a high-low junction to provide an effective electric field that can restricts carriers from reaching the top surface. An enhanced deep-UV response was observed with the external quantum efficiency of ~30% at 240 nm. Also the dark current at high bias voltage is as low as the SiC p-i-n APD.

Chapter 4. AlGaN/SiC Photodiode for Deep-UV detection

In the previous chapter, I have investigated various approaches to enhance the deep-UV response of homojunction SiC APDs. In this chapter, I will focus on the AlGaN/SiC heterojunction structure with AlGaN heteroepitaxially grown on SiC epitaxial layers. The goal of this project is to extend the spectral response of SiC APD into shorter wavelength utilizing the optical properties of AlGaN. This work was also done in collaboration with researchers at the Army Research Laboratory in Adelphi, MD.

4.1 AlGaN/SiC SAM avalanche photodiode

As discussed in previous chapters, SiC APDs exhibit high avalanche gain, low dark current, and low excess noise, but suffer from poor deep-UV response due to a shallow absorption profile and surface recombination. Conversely, aluminum gallium nitride (AlGaN) alloys have a direct bandgap that can be engineered for efficient absorption from 3.4 eV to 6.2 eV, but they exhibit large leakage current [4.1]. Furthermore, AlGaN has a k-factor, a figure of merit for excess noise in APDs, close to 1, a signature of high excess noise [4.2]. To combine the advantages of SiC and AlGaN, a separate absorption and multiplication (SAM) APD has been designed using AlGaN as the absorption layer and SiC as the multiplication layer. The SAM structure allows the peak response to be tuned below 260 nm by adjusting the AlN mode fraction in the AlGaN layer. AlGaN/SiC SAM APDs differ from traditional APDs in several important respects, including: (1) The AlGaN

absorption region has a wider bandgap than the SiC multiplication region and therefore the photo-response will include contributions from absorption in both layers. (2) The AlGaN is grown on lattice mismatched SiC, which introduces a high density of defects in the AlGaN [4.3]. As a result, the carrier diffusion length is short and an electric field in the AlGaN absorption region is required for carrier collection. (3) Both SiC and AlGaN are polar materials and the smaller spontaneous polarization of SiC as compared to AlGaN results in the formation of significant positive polarization induced charge at the interface between SiC and AlGaN [4.4]. This interfacial charge will confine the electric field in the SiC layer.

One way to extend the electric field into the AlGaN absorption layer is to insert an AlN layer, which has larger spontaneous polarization than both the AlGaN absorption and SiC multiplication regions [4.5]. This layer creates a two-dimensional electron gas (2DEG) in the SiC multiplication region and two-dimensional hole gas in the AlGaN absorption region, with the effect of reducing the interface charge density. However, the AlN layer also forms a barrier for hole injection from AlGaN to SiC and electron injection from SiC to AlGaN. Under reverse bias, the height of the barrier is significantly reduced. It is expected that the carriers will tunnel though the reduced barrier if the AlN layer is thin enough. Figure 4-1 shows a simulation of the band diagrams of the SiC/AlGaN heterojunction with and without an AlN layer.

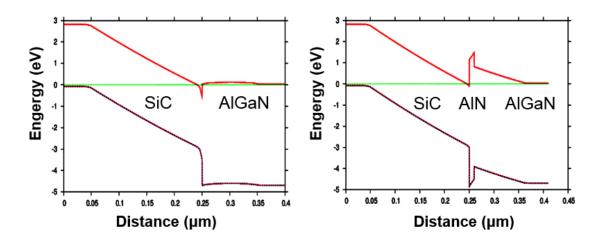


Figure 4-1. Comparison of a) thick i-layer PIN and b) SACM structure.

The influence of the AlN layer on the interface charge and device behavior is studied by comparing the photo response of an Al_{0.62}Ga_{0.38}N/SiC SAM APD with and without AlN layers. The structure of the Al_{0.62}Ga_{0.38}N/SiC APD is shown in Figure 4-2. From top to bottom, it consists of a 50 nm n-doped Al_{0.62}Ga_{0.38}N contact layer, a 300 nm lightly ndoped Al_{0.62}Ga_{0.38}N absorption layer, a 470 nm lightly n-doped SiC multiplication layer, and a 2000 nm p-doped SiC buffer layer. The SiC substrate with two epitaxial layers was purchased from Cree, Inc. The Al_{0.62}Ga_{0.38}N epitaxial layer was grown by Molecular Beam Epitaxy (MBE) by our collaborators at the Army Research Laboratory. The AlGaN/SiC photodiodes were fabricated using the process flow described in chapter 2.

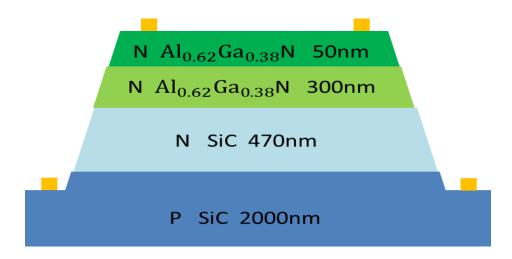


Figure 4-2. Cross section of the AlGaN/SiC SAM APD

Figure 4-3 shows the external quantum efficiency of the devices with and without the AlN layer. The photo response of these devices is expected to consist of two components, one arising from absorption in the $Al_{0.62}Ga_{0.38}N$ region and the other arising from the SiC multiplication region. The bandgap of $Al_{0.62}Ga_{0.38}N$ is 4.84 eV, which results in a cutoff wavelength at 255 nm. Most of the light with wavelength shorter than 255 nm is absorbed in the $Al_{0.62}Ga_{0.38}N$ layer.

From Figure 4-3a, the response of the device without AlN layer looks similar to homojunction SiC APDs. The shape of the curve did not change with bias voltage. The sharp cut-off near 250 nm indicates that the photo-generated carriers in the Al_{0.62}Ga_{0.38}N absorption layer are not collected due to the short diffusion length. The Al_{0.62}Ga_{0.38}N layer principally acts as long pass optical filter.

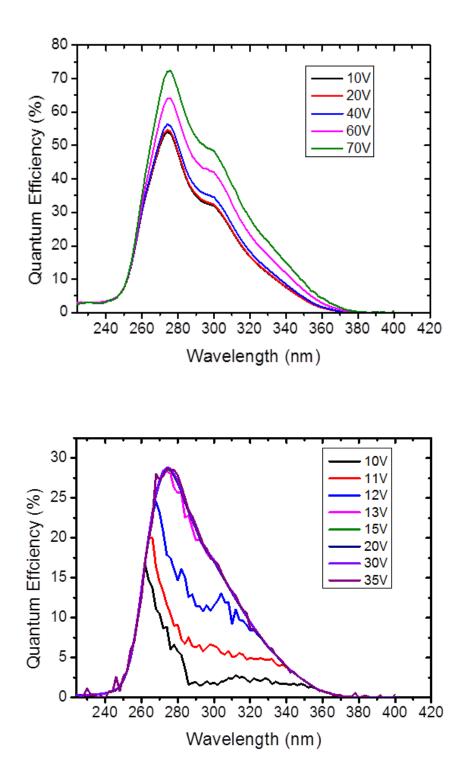


Figure 4-3. External quantum efficiency of AlGaN/SiC SAM APD without (a) and with AlN layer (b) [You need to label the figures (a) and (b).]

In contrast, the response of the device with 10 nm AlN layer shown in Figure 4-3b has a bias-dependent behavior. This is an indication that here is a barrier at the interface. The increase in long-wavelength response results from reduction in the barrier height that is associated with the AlN layer at the interface. As the applied reverse bias is increased, the barrier is lowered and electrons generated from lower energy photons in the SiC are able to surmount it and contribute to the photo-response. We note that the curve also drops sharply at 250 nm, which means that the carriers generated in the Al_{0.62}Ga_{0.38}N layer are still lost. This is due to the fact that the photo-generated holes in the Al_{0.62}Ga_{0.38}N do not have enough energy to tunnel through the 10 nm AlN layer to SiC. The lower peak quantum efficiency can also be attributed to recombination through defects at SiC/AlN interface.

Overall, the AlN layer is effective in reducing the polarization-induced interface charge, but optimization of the thickness is required for higher injection efficiency of carriers generated in the Al_{0.62}Ga_{0.38}N absorption region.

4.2 SiC avalanche photodiode with AlGaN window layer

An alternative approach to increase the absorption of the deep-UV photons in the highelectric-field, lightly-doped SiC absorption region while reducing absorption losses in the deep UV is to employ an $Al_{0.8}Ga_{0.2}N$ n-type doped contact window layer that is transparent to light with wavelength longer than 230 nm. Figure 4-4 shows a cross section of the $Al_{0.8}Ga_{0.2}N/AlN/SiC$ device.

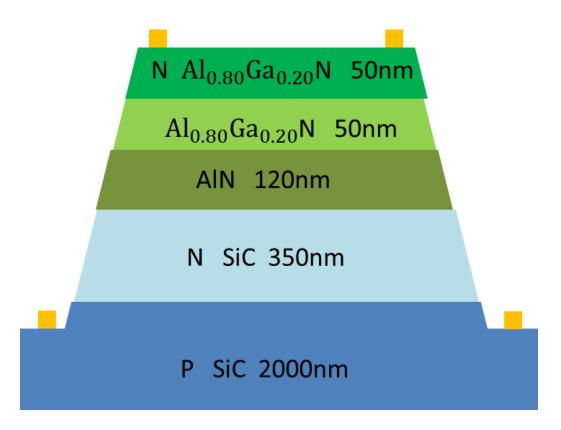


Figure 4-4. Cross section of Al_{0.8}Ga_{0.2}N /AIN/SiC photodiode

The external quantum efficiency versus bias voltage is shown in Figure 4-5. The spectral response of this detector shows a strong cut-off at short wavelength and a peak that shifts from 230 nm to 242 nm when the bias voltage is increase from 10 V to 40 V. A peak external quantum efficiency of ~80% is observed at 242 nm, which is significantly higher than the conventional homojunction PIN SiC detector. This can be explained by the shift of the absorption in Al_{0.8}Ga_{0.2}N. The band gap of Al_{0.8}Ga_{0.2}N is 5.38 eV [4.6], which corresponds to an absorption edge at approximately 230 nm. So this enhancement results from increased absorption of the deep-UV photons in the depleted SiC layer. Carriers generated in this spectral range are then collected more efficiently by drift in this region.

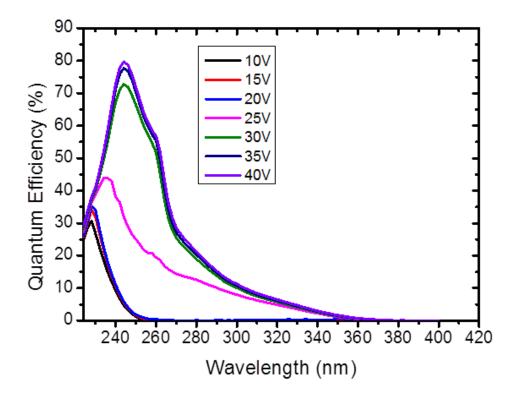


Figure 4-5. External quantum efficiency of Al_{0.8}Ga_{0.2}N /AlN/SiC photodiode

The increase and red-shift of the photo-response with increasing reverse bias and the sharp cutoff at 242 nm is explained by the presence of a barrier associated with polarization-induced charge at the hetero-interfaces as shown if Figure 4-6. The Al-polar AlN grown on SiC give rise to positive polarization-induced charge at the AlN/SiC interface. The Al_{0.8}Ga_{0.2}N grown on AlN results in genitive polarization-induced charge at the Al_{0.8}Ga_{0.2}N/AlN interface. Considering the thickness of the AlN barrier layer, this barrier is significantly larger than the band offsets between these materials and prevents holes generated within the Al_{0.8}Ga_{0.2}N window layer from being collected regardless of reverse bias. As a result, this layers acts as an optical filter that cuts off the short-

wavelength response of the photodiode above the band gap energy of the $Al_{0.8}Ga_{0.2}N$ window layer.

The relative lack of bias dependence on the response at wavelength shorter than 224 nm indicates that this barrier has minimal impact on electrons that are generated in higher of 4H-SiC [4.7], as these carriers can have sufficient energy and the relaxation time allows them to traverse the barrier. Increasing the reverse bias reduces the barrier, resulting in a red-shift in photo-response associated with more electrons that are excited to lower bands having sufficient energy to cross it. The strong suppression of the photo response at wavelength longer than 260 nm for all reverse bias indicates that the barrier is still sufficient at the highest bias investigated to reduce the collection of the associated photo-excited carriers in the bottom of M and lower L valleys.

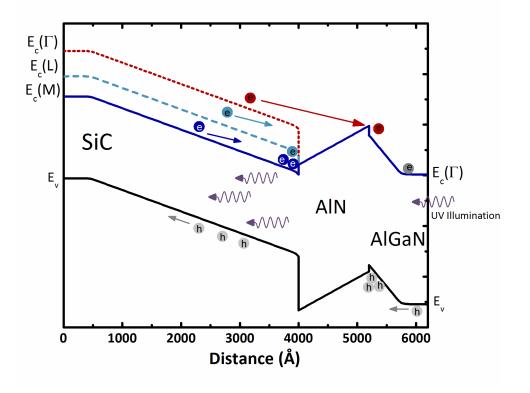


Figure 4-6. Band diagram of SiC/AIN/Al_{0.8}Ga_{0.2}N for different band in SiC

Given the fact that the Al_{0.8}Ga_{0.2}N window layer has proved to be effective in enhancing the deep-UV response of the SiC APD, a modified structure was designed to further improve the performance. A cross section of that structure is shown in Figure 4-7. The material of the window layer was changed from Al_{0.8}Ga_{0.2}N to Al_{0.9}Ga_{0.1}N. The motivation is to increase the bandgap energy of the window layer by increasing the Al mole fraction.. As a result, more photons with higher energy are able to pass through the window layer and get absorbed in the SiC n- layer, with a projected blue shift of the spectral response. Also the AlN layer between the AlGaN and SiC layer was removed so that there will be no barriers for the electrons that in the lower L and M valleys. The suppression of the photo response at longer wavelength discussed in the structure above is expected to be greatly reduced.

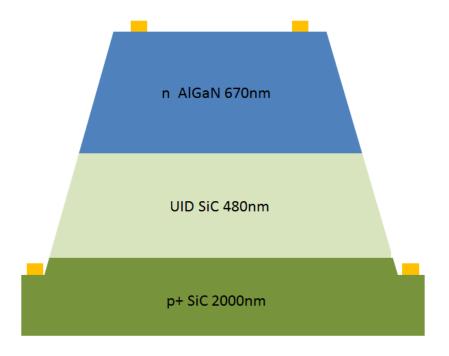


Figure 4-7. Cross section of Al_{0.9}Ga_{0.1}N /SiC photodiode

The photoresponse of the Al_{0.9}Ga_{0.1}N/SiC photodiode and homojunction SiC photodiodes are shown in Figure 4-8. The recessed window SiC photodiode has a peak quantum efficiency of ~60% at 260 nm and a sharp drop off in response at shorter wavelength. In contrast, the Al_{0.9}Ga_{0.1}N/SiC photodiode has a flatter response in the deep-UV that exceeds 60% over a spectral range from 215-256 nm and a sharp cut-off at shorter wavelengths. This enhanced deep-UV response is attributed to the generation of carriers within the depletion region of the heterojunction, where these carriers are more efficiently collected through drift as shown in Figure 4-9. Importantly, the presence of fixed, positive, polarization-induced charge at the hereto-interface arising from the larger spontaneous polarization of AlGaN over that of SiC should act to repel holes from the interface so as to further suppress interface recombination. However, additional studies are required to understand the relative contributions of each of these effects. The sharp short-wavelength cutoff observed at 210 nm is attributed to the absorption of photons in the $Al_{0.9}Ga_{0.1}N$ window layer. The short carrier diffusion lengths within this region associated with the presence of dislocations arising from lattice mismatch inhibits the collection of photogenerated carriers through diffusion.

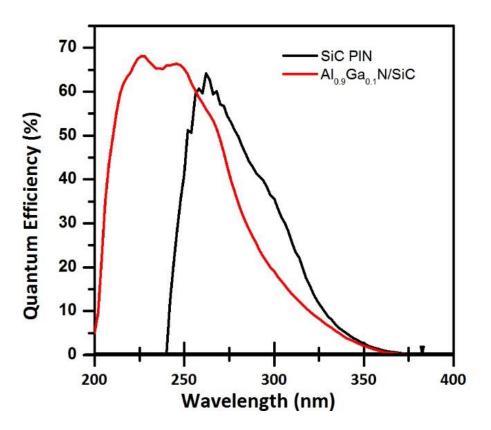


Figure 4-8. Quantum efficiency of Al_{0.9}Ga_{0.1}N /SiC and SiC PIN photodiode

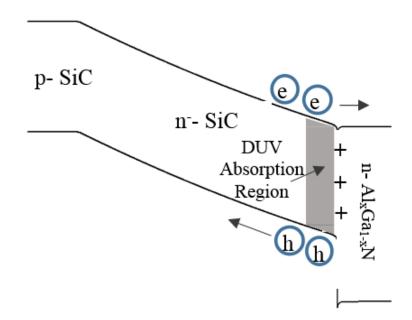


Figure 4-9. Band diagram of Al_{0.9}Ga_{0.1}N /SiC photodiode

4.3 Conclusion

AlGaN/SiC heterogeneous photodiodes were investigated for deep-UV detection in this chapter. Both reach-through and non-reach-through structure were studied.

The AlGaN/SiC SAM structure is investigated as the early effort to extend the spectral response of SiC into deep-UV. However, the large polarization induced charge formed at the hetero-interface is detrimental to device performance, because it confines the electric field to the SiC multiplication layer. An AlGaN/AlN/SiC SAM structure with Al_{0.62}Ga_{0.38}N as absorption layer was fabricated and characterized. The AlN layer proved to be effective in reducing the polarization-induced charge and extending the electric field into the AlGaN. However the bias dependent behavior of the AlGaN/AlN/SiC indicates that the AlN acts as a barrier layer, thus a further optimization of the AlN layer thickness is needed.

The SiC with AlGaN transparent window layer provides another way to achieve high deep-UV response by absorbing the deep-UV photons in the depleted SiC layer. The Al_{0.8}Ga_{0.2}N/AlN/SiC photodiodes exhibited a peak quantum efficiency of ~80% at 242 nm. However, the AlN layer formed a barrier between the AlGaN and SiC, especially for carriers with lower energy in the SiC L and M valleys. In order to remove the barrier effect and further enhance the deep-UV response, photodiodes an Al_{0.9}Ga_{0.1}N/SiC structure were studied. The Al_{0.9}Ga_{0.1}N/SiC photodiode showed a flat response in the deep-UV that exceeded 60% over a spectral range from 215-256 nm, a significant enhancement in efficiency compared to a SiC PIN photodiode.

Chapter 5. Al_{0.8}Ga_{0.2}As Avalanche Photodiodes for Near-UV detection

In the previous chapters I discussed the performance of SiC-based APDs. They have exhibited decent external quantum efficiency from 200 nm to 350 nm. However, due to the fact that SiC has an indirect bandgap the absorption coefficient is very small near the band edge. Several micrometers of absorption layer are needed to achieve strong absorption in the near UV, which will result in very high breakdown voltage and very long transient time. Among the other material candidates are Si, GaP and AlGaAs. Si APDS have, for most applications, been the preferred detectors owing to their high efficiency, low dark current, and low excess noise [5.1]. GaP APDs have demonstrated low dark current and high quantum efficiency. However, their excess noise is high as evidenced by the nearunity effective impact ionization coefficient ratio k (0.6~0.7) [5.2]. Another promising candidate is AlGaAs, which is used widely in semiconductor photonic devices, due to its tunable bandgap and refractive index.

5.1 Material property

There have been numerous published works on AlGaAs-based photodiode with Al fraction less than 60% [5.3-5.5]. However, due to their relatively narrow bandgap, the dark current is higher than that of the wide-bandgap materials discussed in previous chapters [5.6]. Also the low avalanche gain and poor impact ionization coefficient ratio limit their performance as avalanche photodiodes [5.7]. However, AlGaAs with a high Al mode fraction is a promising material candidate for high-performance APDs. Low impact ionization coefficient ratio between 0.1 and 0.2 has been reported for Al_{0.8}Ga_{0.2}As [5.8].

With 80% Al mole fraction, the bandgap is 2.23 eV, which is expected to yield low dark current and a better absorption profile in the UV.

5.2 Photodiode design and fabrication

The first structure I investigated is shown in Figure 5-1. The doping of the p- layer that we requested from the foundry is as low as possible to ensure a fully depleted layer and uniform electrical field in the multiplication layer. The i layer thickness was designed to be 1 μ m in order to keep the breakdown electric field below that at which tunneling becomes significant. The top layer is GaAs since it is easier to form low resistance ohmic contact layer on GaAs than Al_{0.8}Ga_{0.2}As. In order to ensure minimal absorption in the contact layer, its thickness is only 10 nm.

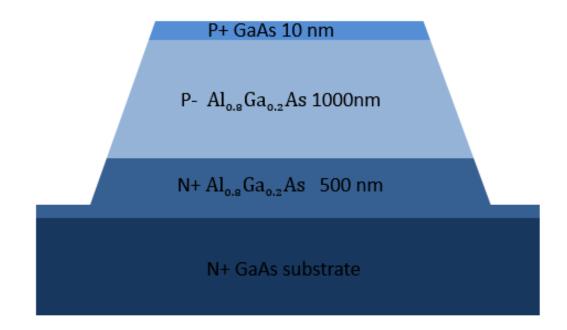


Figure 5-1. Cross section of the Al_{0.8}Ga_{0.2}As APD

The Al_{0.8}Ga_{0.2}As APDs were fabricated using the process flow described in chapter 2. The sidewall passivation is critical for Al_{0.8}Ga_{0.2}As detectors. For mesa structure APDs, atoms at the side wall that are not fully bonded will result in the formation of electrically active defects that degrade device performance through increased dark current or premature breakdown. In order to reduce the impact of these sidewall surfaces on device performance, mesa structure photodiodes are generally coated with a passivation material to reduce the density of active surface states. Sulfur passivation has been widely used to improve the electronic properties of GaAs devices by reducing the high density of states in the bandgap due to the native oxide layer, which is generally considered to be an unsuitable dielectric, having high defect densities [5.9]. The passivation procedure can cause unpinning of the Fermi level and inhibit the surface from further interaction with atmospheric contaminants. The sulfur passivation is typically achieved by soaking in 5% ammonium sulfide $((NH4)_2S_x)$ solution before the deposition of SiO₂ as sidewall passivation. This will leave a monolayer of sulfur that is bonded with the gallium atoms on the sidewall, without introducing any disorder or defects near the surface.

5.3 Device performance

The I-V characteristics of a 150 μ m-diameter device are shown in Figure 5-2. The dark current is 31.7 pA (179 nA/cm²) when biased at 99% of the breakdown voltage, which yields a gain of 200. For comparison, a low-dark-current commercial Si APD from Laser Components, Inc. exhibits dark current density of ~500 nA/cm² at gain = 250. In order to determine the primary effects that contribute to the dark current, the I-V characteristics of

Al_{0.8}Ga_{0.2}As with different diameters were measured. The dark current at three different bias voltages is plotted versus device area in Figure 5-3. The linear relationship between the dark current of the device area at all the bias voltages implies that the dark current of these APDs is bulk-dominated both at low electric field and high electric field, which indicates that the sulfur passivation effectively suppresses surface leakage. For comparison, 150 μ m-diameter devices were fabricated with the same process but without sulfur passivation. Figure 5-2 shows that the dark current with sulfur passivation is one order of magnitude lower than the one without sulfur passivation near breakdown.

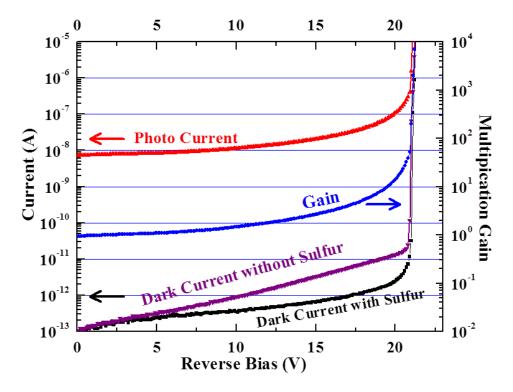


Figure 5-2. I-V characteristics of the Al_{0.8}Ga_{0.2}As APD

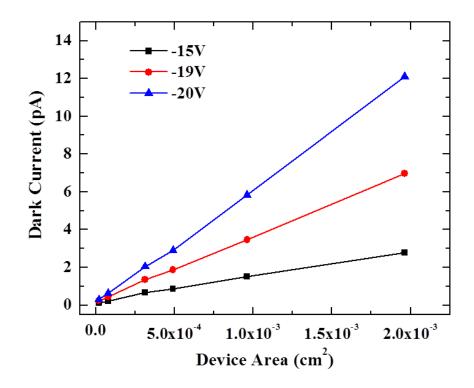


Figure 5-3. Dark current vs device area of Al_{0.8}Ga_{0.2}As APD

The external quantum efficiency at unity gain is plotted versus wavelength in Figure 5-4. The peak quantum efficiency is 27% at 465 nm. The cut-off wavelength is approximately 550 nm, which corresponds to the bandgap of Al_{0.8}Ga_{0.2}As. However, an abrupt decrease in the efficiency dropping from 10% to 5% from 400 nm to 350 nm is observed, The relatively low external quantum efficiency can be explained by the capacitance measurement shown in Figure 5-6. The depletion width of the photodiode is calculated from the capacitance. The result shows that only 1/3 of the p- layer is depleted. A further SIMS analysis shown in Figure 5-6 revealed that the doping of the p- layer is 2×10^{17} cm⁻ ³, which is much higher than specified. The poor UV response is due the fact that a large fraction of the carriers were generated in the undepleted p- layer. Some carriers are generated as far as 600 nm away from the depletion region. Combined with short diffusion length, this results in poor efficiency. A simulation of the external quantum efficiency shown in Figure 5-4 yields a diffusion length of 120 nm for electrons in the p-doped $Al_{0.8}Ga_{0.2}As$ layer, which is much shorter than the distance most electrons need to diffuse before reaching the edge of the depletion region.

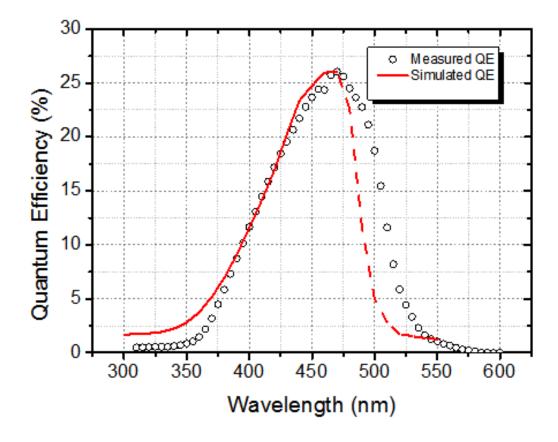


Figure 5-4. Measured and simulated external quantum efficiency of Al_{0.8}Ga_{0.2}As APD

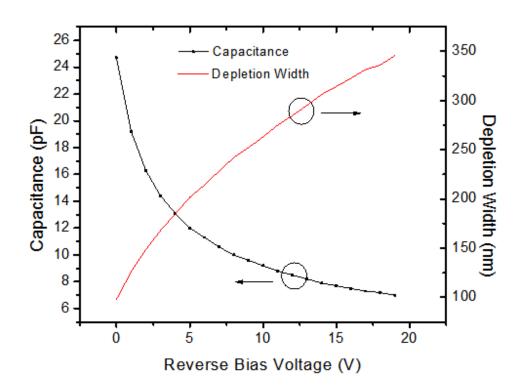


Figure 5-5. Capacitance and depletion width vs voltage of the Al_{0.8}Ga_{0.2}As APD

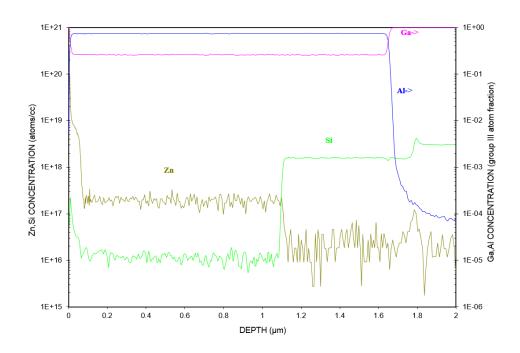


Figure 5-4. SIMS analysis of Al_{0.8}Ga_{0.2}As wafer

The excess noise factor was measured following the procedure described in chapter 2. The device was illuminated with an argon laser at 351/363 nm. Figure 3-16 shows a plot of the excess noise factor versus gain for the Al_{0.8}Ga_{0.2}As APD. The curves for k values ranging from 0.1 to 0.2 are included for reference. A k value of ~0.15 provided the best fit to the data. This measurement agrees well with previous reports on Al_{0.8}Ga_{0.2}As APD.

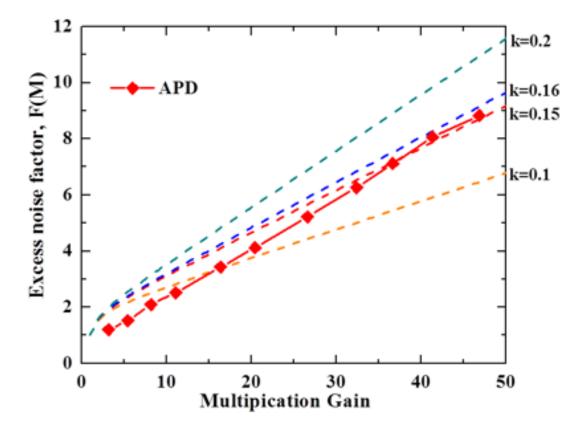


Figure 5-7. Excess noise of Al_{0.8}Ga_{0.2}As APD

For the spatial uniformity measurements, an Al_{0.8}Ga_{0.2}As APD with diameter of 200 μ m was biased near breakdown at a gain of 10. Light from an argon laser at 351/363 nm was scanned across the detector while the photocurrent was measured with a lock-in amplifier. The spot size was ~5 μ m, which is smaller than the contact ring of the device allowing us to resolve the features of the device. In Figure 5-5 the center region represents

the active or operational region of the device. The valley directly outside the center region is the top contact ring. This feature is due to the fact that the metal of the contact ring blocks the incident signal. The outer ring is the edge of the mesa. Shadowing of the probe tip causes the dip in the upper region of the ring. It is observed from the raster scan that the $Al_{0.8}Ga_{0.2}As$ APD has a very uniform spatial response.

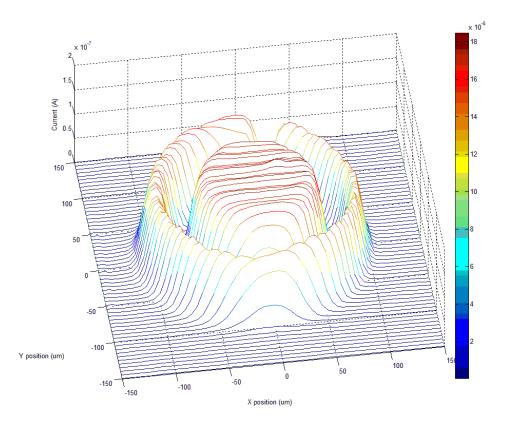


Figure 5-8. Raster scan Al_{0.8}Ga_{0.2}As APD at gain = 10

5.4 Conclusion

In this chapter, Al_{0.8}Ga_{0.2}As APD was studied as a candidate to replace Si APDs for the weak light detection in the near-UV spectrum. Al_{0.8}Ga_{0.2}As APDs with PIN structure were

designed, fabricated and characterized. Sulfur passivation using 5% ammonium sulfide was used to passivate the sidewall of the photodiode. The dark current of the $Al_{0.8}Ga_{0.2}As$ APD was 31.7 pA (179 nA/cm²) at gain of 200, which is lower than the state of the art Si APD. The dark current of the photodiode was found to be proportional to the device area, which indicates that the dark current is bulk dominated and the sulfur passivation is effective. The excess noise of the $Al_{0.8}Ga_{0.2}As$ APD is low with a k value of ~0.15. The raster scan at a gain of 10 showed a uniform spatial response.

The Al_{0.8}Ga_{0.2}As APD yielded a peak quantum efficiency of 27% at 465 nm. However, an abrupt decrease of the efficiency at shorter wavelength is observed, decreasing from 10% to 5% for wavelengths 400 nm to 350 nm. This is due to the fact that the doping in the p- layer is higher than the design. A large fraction of the carriers were generated in the undepleted p- layer. A lower doping in the lightly-doped layer is needed to achieve higher quantum efficiency in the near-UV spectrum.

Chapter 6. GaAs/AlGaAs Photodiode for Wide Spectrum Detection

GaAs photodetectors are widely used in for visible and near infrared detection [6.1]. However, there have been few reports on the performance in the UV spectrum. In this chapter, I will discuss different techniques to enhance the UV response of the GaAs photodiode. In this project I developed a photodiode that can achieve high quantum efficiency from UV to near infrared. This work was funded by NASA and motivated by the goal of reducing the number of cameras in some satellites.

6.1 Wafer Design

A schematic cross section of the first structure studied is shown in Figure 6-1. The wafer was purchased from Landmark Optoelectronics, Inc. The i-layer thickness was chosen to ensure strong absorption for wavelengths as long as 850 nm. The Al concentration in the AlGaAs layer is graded from 30% at the top to 10% at the AlGaAs/GaAs interface. This graded bandgap structure creates an electric field that assists carrier transport into the depletion region instead of relying exclusively on diffusion.



Figure 6-1. Wafer structure for GaAs/AlGaAs photodiode

GaAs/AlGaAs photodiodes were fabricated using the process flow described in chapter 2. The current-voltage (I-V) characteristics of a 200-µm diameter device are shown in Figure 6-2. The photodiode has a breakdown voltage of 50 V. The dark current is less than 3 pA for bias voltage lower than 30 V. The dark current near breakdown at a gain of 10 is 500 pA.

The initial measurement of the external quantum efficiency yielded low efficiencies in the UV spectrum. This is due to two reasons. First, the absorption depth in this spectrum region is very shallow. Most of the photons are absorbed within 100 nm of the surface. Owing to surface recombination and the short diffusion length, a significant fraction of the carriers recombine prior to being collected. Second, the reflectance of GaAs increases rapidly with the photon energy. For photons with wavelength shorter than 400 nm, the reflectance is higher than 40%.

In order to improve the quantum efficiency in the UV spectrum, the recessed window structure and surface texturing techniques were investigated.

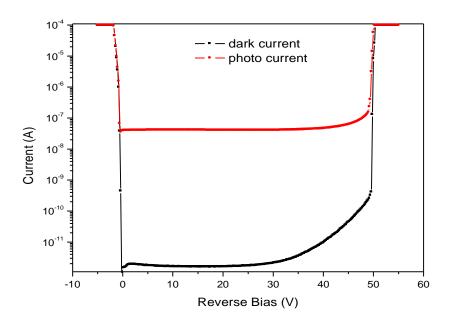


Figure 6-2. Current - voltage characteristics of a 200 µm-diameter GaAs/AlGaAs photodiode

6.2 Recessed window structure

For the recessed window approach, the top GaAs p+ cap layer was removed by chemical etching in the active portion of the device. The etch process was optimized such that the bottom of the window extended into the 100 nm p AlGaAs layer. SEM top-view and side-view photographs after etching the window are shown in Figure 6-3. The recessed window enhances detection at short wavelengths by decreasing the distance to the collection region, which is beneficial for UV radiation.

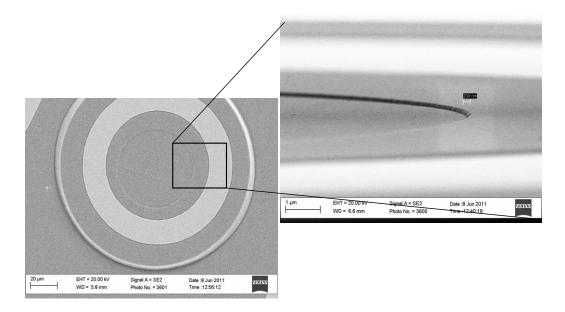


Figure 6-3. SEM image of the recessed window on GaAs/AlGaAs photodiode

6.3 Nanosphere surface texturing

At present, dielectric anti-reflection (AR) coatings are widely used to reduce the surface reflectivity and increase the responsivity of photodiode near the wavelength of interest. However, the reflectance of an AR coating is very sensitive to the wavelength and thus not able to achieve low reflectance across a wide spectrum. Surface texturing is another technique to reduce the surface reflectance loss. It has been demonstrated to successfully reduce reflectance in solar cells, light emitting diodes, and photodiodes [6.2-6.5]. A previous student in our group has investigated laser surface texturing on Si photodiodes. High efficiency over a broad spectrum was achieved [6.6]. However, the devices exhibited higher dark current after the texturing, possibly due to the laser induced damage. Alternatively, a texturing technique referred to as nanosphere natural lithography texturing can create a random pattern with feature size ~100 nm, while limiting the damage to the top contact layer [6.7].

For the texturing, 5% solution of 100 nm-diameter SiO₂ spheres in water was purchased from Corpuscular Inc. A solution of surfactant Triton X-100 and Methanol was added to the suspension. The volume ratio of the SiO₂ suspension to Triton X-100 solution was 10:1. The SiO₂ suspension was spun onto the wafer with spin rate of 100 rpm. This created a uniform monolayer of SiO₂ spheres that covered most of the surface area after the solvent evaporated. Then an Ar/BCl₃ inductive coupled plasma etch was performed to transfer the nanosphere monolayer random pattern onto the wafer surface. The etch depth was ~100 nm in the areas between the SiO₂ spheres. This depth kept the plasma from etching into the lightly-doped layer. A buffered oxide etch was used to remove any SiO₂ residue. A welltextured surface will look very dark. A scanning electron microscope (SEM) image of the textured surface is shown in Figure 6-4. The reflectance of the textured surface was measured using the Perkin Elmer Lambda 950 spectrometer. The detailed measurement steps are discussed in Chapter 2. Figure 6-5 compares the spectral reflectance of the textured GaAs/AlGaAs wafer and the SiO₂ AR coating deposited on the same wafer using PECVD. The AR coating was optimized for ~ 400 nm. The minimum reflectance for the AR coating is the same as that for the textured surface. However, the textured surface provides low reflectivity across a very wide spectrum from 300 nm to 600 nm.

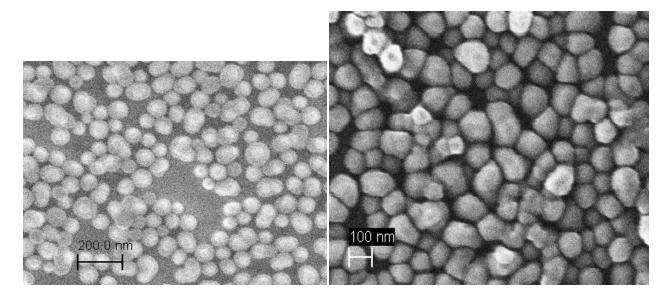


Figure 6-4. SEM image of the textured surface on GaAs/AlGaAs photodiode

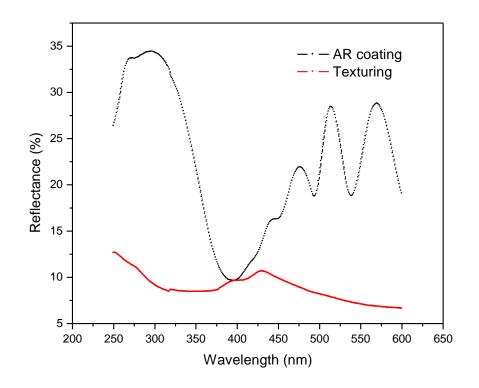


Figure 6-5. Measured reflectance of the textured surface compared with SiO₂ AR coating

6.4 Device performance

The GaAs/AlGaAs photodiodes were fabricated with both the recessed window and surface texturing technique using the processing steps described in chapter 2. The recessed window and surface texturing were finished before the mesa patterning so that the ICP plasma would not damage the sidewall. The same I-V characteristics were achieved after incorporating the surface texturing, showing that the surface texturing did not increase the dark current of the photodiode. The external quantum efficiency of the GaAs/AlGaAs photodiode with recessed window and surface texturing is plotted in Figure 6-6 and compared with a photodiode that was fabricated without these techniques. A much flatter curve with efficiency over 45% across a wide spectrum from 350 nm to 850 nm was achieved. The external quantum efficiency at 400 nm increased from 5% to 45%.

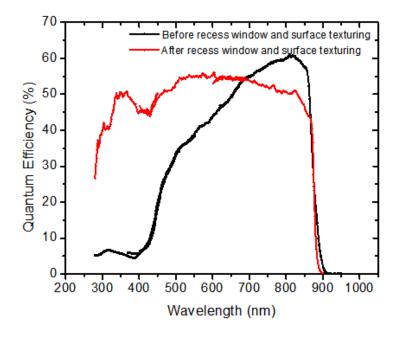
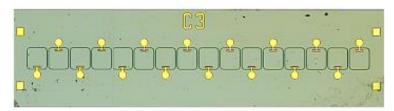


Figure 6-6. External quantum efficiency of GaAs/AlGaAs photodiode

6.5 High-efficiency UV to near IR arrays

The photodiodes described in the previous section represent new state-of-the-art sensitivity across a broad spectral range. Given this success, these devices were fabricated into 1x16 and 1x32 photodiode arrays. For the array fabrication, new photomask was designed using a round-corner square mesa in order to achieve higher fill factor. The current voltage characteristics for each element in the array were measured. Figure 6-7(a) is a photograph of a completed 1x16 linear array and Fig. 6-7(b) is an enlarged picture of three 200 μ m x 180 μ m diodes. Figure 6-8 shows the dark current of each of the 16 photodiodes at -1 V bias. From the dark current distribution we can find the uniformity among the elements in the linear array is very good. The dark current of the single element ranges from 0.7 pA to 3.4 pA. Figures 6-11 and 6-12 show similar results for the 1x32 linear arrays. Because of the smaller size of the element in the 1 x 32 array, the dark current for the element is lower and better uniformity is achieved.



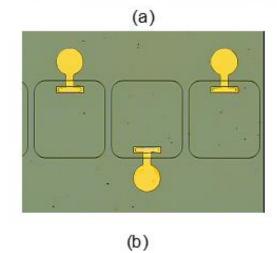


Figure 6-7. (a) Photograph of a 1x16 UV to near IR linear array and (b) enlarged picture of three 200 μm x 180 μm diodes.

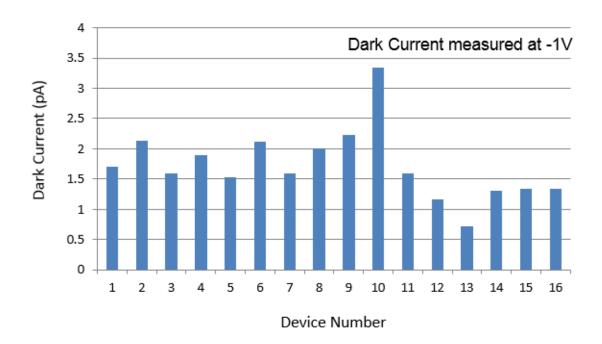
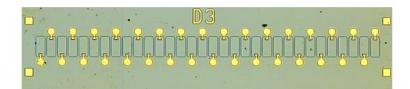


Figure 6-8. Dark current of each element in the 1x16 UV to near IR linear array at -1 V bias.



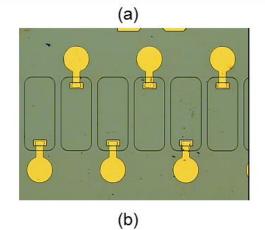


Figure 6-9. (a) Photograph of a 1x32 UV to near IR linear array and (b) enlarged picture of three 200 μm x 85 μm diodes.

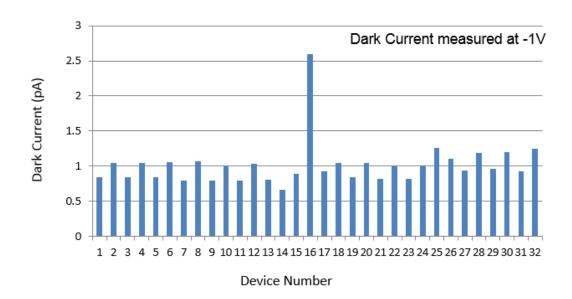


Figure 6-10. Dark current of each element in the 1x32 UV to near IR linear array at -1 V bias.

6.6 Conclusion

GaAs/AlGaAs photodiodes were studied for wide spectrum detection from the near-UV to the near infrared. The photodiodes exhibited low dark current of less than 10 pA at bias voltage lower than 30 V and 500 pA near breakdown at gain of 10.

In ordered to enhance the near-UV response and achieve a flat quantum efficiency curve over the wide spectrum, a recessed window and surface texturing were employed. In the recessed window structure, part of the highly doped layer in the active region is removed by chemical etching. The recessed depth was optimized such that the bottom of the window extended into the 100 nm p AlGaAs layer. Following the recessed window, the surface was textured using the nanosphere natural lithography etching, which created a random pattern on the top surface with feature size of ~ 100 nm. The etch depth was controlled at around 100 nm to keep prevent etching the lightly-doped layer.

With recessed window and surface texturing, the GaAs/AlGaAs photodiode yielded external quantum efficiency over 45% from 350 nm to 850 nm. At 400nm, the responsivity showed relative enhancement of 300%. These photodiodes were fabricated into 1x16 and 1x32 UV-to-near infrared linear arrays and delivered to NASA.

Chapter 7. Conclusion and future work

7.1 Summary

My work has focused on developing high performance avalanche photodiodes (APD) in the deep-UV (200 nm -250 nm) and near-UV (300 nm – 400 nm) spectrum for weak light detection.

Previously in our group, 4H-SiC APDs with low dark current, low noise, high gain and high peak quantum efficiency have been developed. However, the narrow spectral response has limited their use for applications in the deep-UV and near-UV detection. Initially I studied the spectral response of the SiC APD with PIN structure by modeling the external quantum efficiency. The results showed that over 50% of the carriers that were generated in the highly-doped top layer were did not diffuse into the depletion region, which resulted in poor deep-UV response. This is due to the fact that the diffusion length in highly-doped SiC is relatively short and the surface recombination velocity is high. Different approaches utilizing both the homojunction and heterojunction structures to enhance the deep-UV response of the SiC APD have been reported here.

A SiC metal-n⁻-p photodiode was first studied in order to minimize the absorption in the undeleted region. The fabricated device demonstrated flat external quantum efficiency of ~ 45% from 270 nm to 225 nm with a slight decrease below 225 nm. This response is greater than that observed for the p-n-n+ diodes for wavelengths shorter than 240 nm despite losses due to the semi-transparent metal contact. The deep-UV response enhancement resulted from the improved collection of photo-generated carriers through drift within the high electric field region near the surface of the lightly-doped SiC absorption region despite the presence of surface states.

A SiC NIP photodiode with graded doping profile in the top n layer was also investigated. The goal of this structure was to utilize the high-low junction to provide an effective electric field that can stop the carriers from reaching the top. An enhanced deep-UV response was observed with the external quantum efficiency of ~30% at 240 nm.

The AlGaN/SiC SAM structure illustrates another promising way to achieve high, tunable UV response with an AlGaN absorption layer. An AlGaN/AlN/SiC SAM structure photodiode with Al_{0.62}Ga_{0.38}N as the absorption layer was fabricated and characterized. An AlN layer proved to be effective in reducing the polarization-induced charge and extending the electric field into the AlGaN. However the bias dependent behavior of the AlGaN/AlN/SiC indicated that the AlN acts as a barrier layer, thus a further optimization of the AlN layer thickness was needed.

A SiC detector with AlGaN transparent window layer achieved high deep-UV response by absorbing the deep-UV photons in the depleted SiC layer. The Al_{0.8}Ga_{0.2}N/AlN/SiC photodiodes exhibited a peak quantum efficiency of ~80% at 242 nm. A modified structure Al_{0.9}Ga_{0.1}N/SiC structure without the AlN layer was shown to further enhance the deep-UV response, The Al_{0.9}Ga_{0.1}N/SiC photodiode showed a flat response in the deep-UV that exceeded 60% over a spectral range from 215-256 nm, which is a significant enhancement compared with a SiC PIN photodiode

For near-UV detection, GaAs/AlGaAs was investigated as an material candidate to replace Si. Al_{0.8}Ga_{0.2}As APDs with PIN structure were designed, fabricated and characterized. Sulfur passivation using the 5% ammonium sulfide was used to passivate

the sidewall of the photodiode. The dark current of the $Al_{0.8}Ga_{0.2}As$ APD was 31.7 pA (179 nA/cm²) at gain of 200, which is lower than state-of-the-art Si APDs. The dark current of the photodiode was found to be proportional to the device area, which indicates that the dark current is bulk dominated and the sulfur passivation is effective. The excess noise of the $Al_{0.8}Ga_{0.2}As$ APD is low with a k value of ~0.15. The raster scan at a gain of 10 showed a uniform spatial response. The $Al_{0.8}Ga_{0.2}As$ APD yielded a peak quantum efficiency of 27% at 465 nm. However, an abrupt decrease of the efficiency from 10% to 5% for 400 nm to 350 nm was observed. This is due to the fact that the doping in the p- layer is higher than the design. A large fraction of the carriers were generated in the undepleted p- layer. A lower doping in the lightly-doped layer is needed to achieve higher quantum efficiency in the near-UV spectrum.

A GaAs/AlGaAs photodiode was studied for wide spectrum detection from the near-UV to the near infrared. The photodiodes exhibited low dark current of less than 10 pA at bias voltage lower than 30 V and about 500 pA near breakdown at gain of 10. In ordered to enhance the near-UV response and achieve a flat quantum efficiency over a broad spectrum, recessed window and surface texturing were employed. In the recessed window structure, part of the highly doped layer in the active region was removed by chemical etching. The recessed depth was optimized such that the bottom of the window extended into the 100 nm p AlGaAs layer. Following the recessed window, the surface was textured using the nanosphere natural lithography etching, which created a random pattern on the top surface with feature size of about 100 nm. The etching depth was controlled to approximately 100 nm in order to protect the lightly-doped layer. With the recessed window and surface texturing, the GaAs/AlGaAs photodiode yielded external quantum efficiency over 45% from 350 nm to 850 nm. At 400nm, the responsivity improvement was 300%. These photodiodes were fabricated into 1x16 and 1x32 arrays that exhibited excellent device-to-device uniformity.

7.2 Future work

7.2.1 Nitrogen passivation on SiC avalanche photodiode

As discussed in chapter 2, surface band bending is detrimental to the deep-UV response of SiC. Different approaches have been investigated in chapter 2 and chapter 3 to overcome this problem. I have shown with the SiC metal-n⁻-p structure and AlGaN/SiC heterojunction structure that the deep-UV response can be greatly enhanced. These APDs still exhibit high dark current at high bias voltage, which makes their performance in the high gain region inferior to SiC PIN APDs. The source of the dark current is believed to be related to high defect densities at the metal/SiC or AlGaN/SiC interfaces. A direct chemical passivation on the SiC surface without introducing another materials in the high electric field region would be the best solution.

A very significant advantage of SiC, over other wide bandgap semiconductors, is the high quality of the SiO₂ layers grown by thermal oxidation. Usually, this thermally-grown SiO₂ is used as the passivation material in the SiC APD fabrication. But the high surface recombination velocity indicates that there is still a high density of defects present at the SiO₂/SiC interface, which result in a Fermi level pinning. Since SiC is widely used for high power metal-oxide-semiconductor field effect transistors, the researcher have done studies on the defects in the SiO₂/SiC interface, which result in low channel mobility [7.1-7.3].

Studies have shown that there is large and broad density of states fixed at approximately 2.9 eV above the valence band edge [7.4]. This high density of states in the forbidden band is attributed to a residue carbon complex that is released during thermal oxidation and remains at or near the SiO₂/SiC interface. Different chemical treatments have been investigated to reduce the interface states density [7.5-7.6]. Among them, post-oxidation annealing in nitric oxide (NO) ambient is the most widely adopted method. It has been shown that nitric oxide annealing is effective in increasing the channel mobility and device reliability for SiC MOSFETs [7.7]. It is believed that it is the nitrogen atoms that passivate the interface by removing the carbon complex and creating strong Si-N bonds [7.8].

It appears that both the low channel mobility in SiC MOSFETs and the high surface recombination velocity in SiC APDs are related to the high density of interface states between SiO₂ and SiC. It follows that a similar nitrogen passivation technique could be introduced in the SiC APD fabrication process. The tentative passivation process modification that I have designed is listed here:

- 1) RCA clean of the sample after mesa patterning
- Load the sample into the oxidation furnace at a temperature of 1050 C for an hour with a constant oxygen as flow
- Change the gas from oxygen to nitric oxide and maintain the temperature for another hour
- 4) Cap the thermally grown oxide with ~300 nm PECVD grown oxide

7.2.2 InGaP/AlGaAs UV-enhanced avalanche photodiode

In chapter 5 I have demonstrated Al_{0.8}Ga_{0.2}As APDs with low dark current, high gain and low noise. The good linear-mode performance of these APDs confirmed that Al_{0.8}Ga_{0.2}As is a promising multiplication region material candidate to replace Si for the near-UV detection. However, the high surface recombination velocity of GaAs and AlGaAs prevented the photodiode from achieving high quantum efficiency [7.9]. Though sulfur passivation has proved to be effective in reducing the sidewall leakage, there is no evidence that this improves the quantum efficiency.

Inspired from the AlGaN/SiC photodiode that is discussed in chapter 4, a heterojunction structure could be used to achieve better surface recombination velocity would utilize $In_{0.49}Ga_{0.51}P$ which is lattice matched to GaAs/AlGaAs as the top surface layer. First, $In_{0.49}Ga_{0.51}P$ has a reported surface recombination velocity of ~10⁴ cm/s [7.10], which is two orders of magnitude lower than GaAs. Second, the lattice match between $In_{0.49}Ga_{0.51}P$ and GaAs/AlGaAs will eliminate dislocation-related defects. An InGaP/GaAs photodiode utilizing a similar structure has been reported with quantum efficiency as high as 70% in the 360 -870 nm range [7.11]. Despite the favorable electrical properties of $In_{0.49}Ga_{0.51}P$, it also has one disadvantage for our purpose, which is a relatively high absorption coefficient (~ 5.5×10^5 cm⁻¹ at 400 nm) [7.12]. In order to minimize the absorption in the $In_{0.49}Ga_{0.51}P$ layer, a very thin layer is needed. The initial design of the structure is shown in Figure 7-1. A 50 nm GaAs contact layer is used for better ohmic contact formation, but the GaAs in the active area can be easily removed by the selective chemical etchant.

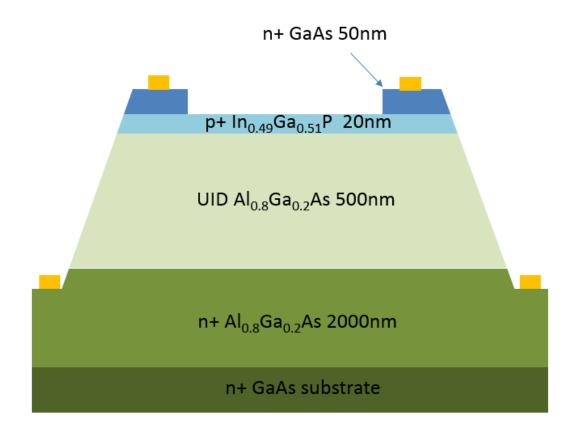


Figure 7-1. Cross section of the InGaP/AlGaAs APD

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Appendix. Quantum Efficiency Modeling Code

Total_QE.m: Set all the parameters and device structure, calculate the total QE

Top_Layer_QE.m: Calculate the QE due to absorption in the undepleted top layer

Depletion_Layer_QE.m: Calculate the QE due to absorption in the depleted layer

Bottom_Layer_QE.m: Calculate the QE due to absorption in the undepleted bottom layer

Get_AbsorptionCoefficient.m: Get absorption coefficient for each wavelength Get_Reflectance.m: Get reflectance for each wavelength.

Total_QE.m

%% Set each layers' thickness

X1 = 260e-7;X2 = 170e-7+X1;X3 = 500e-7+X2;

%% Set the diffusion length and diffusion constant for electrons and holes, here are the

numbers for Al_{0.8}Ga_{0.2}As

Diff_Length_H = 200e-7; Diff_Length_E = 80e-7; Diff_Const_H =1.84; Diff_Const_E = 5.28;

%% Set surface recombination velocity Surf_Recom_V = 9e8; %% Set start and stop wavelength Wavelength_start=300; Wavelength_end=550; N = (Wavelength_end-Wavelength_start)/10+1;

%% Create space for QE QE1=[1:N]; QE2=[1:N]; QE3=[1:N];

```
%% Create space for wavelength, reflectance and absorption coefficient
Wavelength=Wavelength_start:10:Wavelength_end;
Reflect = ones(1,(Wavelength_end-Wavelength_start)/10+1);
Absorp_Coeff=ones(1,(Wavelength_end-Wavelength_start)/10+1);
for m_W=1:(Wavelength_end-Wavelength_start)/10+1
Absorp_Coeff(m_W) = get_absorptioncoefficient(Wavelength(m_W));
Reflect(m_W)= get_reflectance(Wavelength(m_W));
```

```
end
```

%% Calculate QE due to absorption from different layers for n = 1:(Wavelength_end-Wavelength_start)/10+1

%% Top_Layer_QE(Absorp_Coeff, X1, Diff_Length,Diff_Const, Surf_Recom_V) QE1(n) = Top_Layer_QE(Absorp_Coeff(n), X1, Diff_Length_E, Diff_Const_E,

```
Surf_Recom_V);
```

%% Depletion_Layer_QE(Absorp_Coeff, X1, X2) Absorption Layer QE2(n) = Depletion_Layer_QE(Absorp_Coeff(n), X1, X2); %% Bottom_Layer_QE(Absorp_Coeff, X2, X3, Diff_Length, Diff_Const, Surf_Recom_V) QE3(n) = Bottom_Layer_QE(Absorp_Coeff(n), X2, X3, Diff_Length_H, Diff_Const_H, 0);

end

%% Plot the graph simulatedQE=(QE1+QE2+QE3).* (1-Reflect)*100; plot(Wavelength,simulatedQE, 'black');

Top_Layer_QE.m

%% Calculate QE from top layer thickness, absorption coefficient, diffusion length, %% diffusion constant and surface recombination velocity function QE = Top_Layer_QE(Absorp_Coeff, X, Diff_Length, Diff_Const, Surf_Recom_V)

aa = Absorp_Coeff*Diff_Length/(Absorp_Coeff^2*Diff_Length^2-1);

bb = (Absorp_Coeff*Diff_Const + Surf_Recom_V) - exp(-Absorp_Coeff*X)*(

Surf_Recom_V* cosh(X/Diff_Length) + Diff_Const/Diff_Length*sinh(X/Diff_Length)

);

cc = (Surf_Recom_V* sinh(X/Diff_Length) +

Diff_Const/Diff_Length*cosh(X/Diff_Length));

dd = Absorp_Coeff*Diff_Length*exp(-Absorp_Coeff*X);

QE = aa * (bb/cc - dd);

Depletion_Layer_QE.m

%% Calculate the QE from the depleted layer, assuming all the photo generated carriers %% in this layer contribute to the QE

function QE = Depletion_Layer_QE(Absorp_Coeff, X1, X2)

 $QE = exp(-Absorp_Coeff * X1) * (1 - exp(-Absorp_Coeff * (X2 - X1)));$

Bottom_Layer_QE.m

%% Calculate the QE due to absorption in the bottom layer using the all three layers'%% thickness, absorption coefficient, diffusion length and surface recombination%% velocity

function QE = Bottom_Layer_QE(Absorp_Coeff, X2, X3, Diff_Length, Diff_Const, Surf_Recom_V)

 $Kn_aa = Surf_Recom_V * (exp(-Absorp_Coeff*(X3-X2)) - cosh((X3-X2))) - co$

X2)/Diff_Length));

Kn_bb = Absorp_Coeff * Diff_Const * exp(-Absorp_Coeff * (X3-X2));

Kn_cc = Diff_Const / Diff_Length * sinh((X3-X2)/Diff_Length);

Kn_dd = Diff_Const / Diff_Length * cosh((X3-X2)/Diff_Length);

Kn_ee = Surf_Recom_V * sinh((X3-X2)/Diff_Length);

 $Kn = (Kn_aa - Kn_bb - Kn_cc) / (Kn_dd+Kn_ee);$

QE_aa = Absorp_Coeff * Diff_Length;

 $QE = QE_aa * exp(-Absorp_Coeff*X2) / (QE_aa^2-1) * (QE_aa + Kn);$

Get_AbsorptionCoefficient.m

%% Get the absorption coefficient at given wavelength, data is from published literature function absorption=get_absorptioncoefficient(Wavelength)

Lamda=[300	310	320	330	340	350	360	370	380	390	400
410	420	430	440	450	460	470	480	490	500	510
520	530	540	550];							

Alfa=[8.95e5 8.37e5 8.00e5 7.16e5 5.99e5 4.69e5 3.51e5 2.63e5 2.06e5 1.65e5 1.35e5 1.13e5 9.44e4 7.55e4 5.91e4 5.17e4 4.20e4 3.24e4 1.94e4 6.67e3 2.5e3 1.37e3 8.22e2 7.19e2 6.29e2 5.75e2];

absorption=exp(interp1(Lamda,log(Alfa),Wavelength,'linear',1));

Get_Reflectance.m

%% Get reflectance spectrum, data is from reflectance measurement in the lab

function R=get_reflectance(Wavelength)

Lamda=[300	310	320	330	340	350	360	370	380	390	400
410	420	430	440	450	460	470	480	490	500	510
520	530	540	550];							

Reflectance=[45.5		45.6	46.0	46.3	45.9	45.0	43.3	41.6	40.1	38.9
37.9	37.0	36.3	35.7	35.0	34.4	33.9	33.6	33.3	32.8	32.0
31.4	30.9	30.5	30.2	29.9];						

R=interp1(Lamda,Reflectance,Wavelength,'linear',1)/100;