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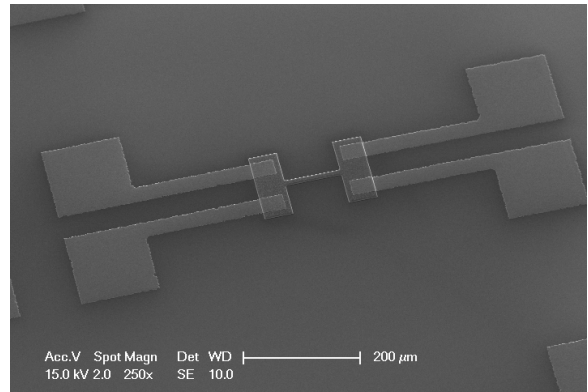
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# Surface-micromachined Silicon Carbide Pirani Gauges for Harsh Environments

J. Mo, L.M. Middelburg, B. Morana, H.W. van Zeijl, S. Vollebregt and G.Q. Zhang

**Abstract**—The application of pressure sensors in harsh environments is typically hindered by the stability of the material over long periods of time. This work focuses on the design and fabrication of surface micromachined Pirani gauges which are designed to be compatible with state-of-the-art Silicon Carbide CMOS technology. Such an integrated platform would boost harsh environment compatibility while reducing the required packaging complexity. An analytical model was derived describing the design variables of the Pirani gauges followed by Finite Element Analysis. The Pirani gauges were fabricated in a CMOS compatible cleanroom with a process employing only three masks, thus suitable for mass production. The SiC-based Pirani gauge is far more competitive than the traditional Si-based Pirani gauge in terms of endurance in high-temperature environments. From 25°C to 650°C, the gauge shows a reproducible response to pressure changes and has a maximum sensitivity of 17.63  $\Omega$ /Pa at room temperature, and of 1.23  $\Omega$ /Pa at 650°C. Additionally, some of the gauges were demonstrated to operate at temperatures up to 750°C.



**Index Terms**—Pressure sensor, MEMS, harsh environments, vacuum gauge, silicon carbide, surface micromachining.

## I. INTRODUCTION

PRESSURE monitoring is an important topic for several industries, such as automotive, aerospace, and semiconductor. In terms of vacuum level monitoring, the Pirani gauge is, without doubt, one of the most popular choices for measuring rough and/or medium vacuum. Its success relies on its simple structure and the absence of moving parts. This results in excellent mechanical robustness and high reliability.

Not until recently, MEMS (micro-electro-mechanical systems) technology has been employed for the realization of silicon-based Pirani gauges. By leveraging on the mature silicon manufacturing technology, it is possible to transform conventional macroscopic pressure sensors into micro-fabricated devices. Miniaturized devices have advantages in terms of size, power consumption, and costs compared with conventional devices. Additionally, Si-based Pirani gauges can be integrated with a readout circuit to further reduce the size and enhance the reliability of the overall system. In 1991, Mastrangelo et al. fabricated a MEMS Pirani gauge based on poly-Si with an integrated on-chip circuitry able to process the sensing signal [1].

In the past decade, the improvement of the Pirani gauge

is still ongoing. Santagata et al. optimized the traditional structure of a Pirani gauge by creating a tube-shaped device that resulted in improved stiffness and higher sensitivity [2]. Also, advanced materials have been introduced to build Pirani sensors. In 2013, Yu et al. employed dielectrophoretic (DEP) assembly technique to prepare a single-walled carbon nanotube Pirani gauge, which demonstrated an ultra-low power consumption (15 nW) and high sensitivity in a large dynamic range (0.8 to 80000 Pa) [3]. In 2018, the world first graphene-based Pirani gauge was designed by J. Romijn et al. The graphene-based Pirani gauge they fabricated has a 100 times smaller footprint than the existing Pirani gauge while maintaining a comparable sensitivity to the state-of-the-art micro-fabricated Pirani gauge [4].

In recent years, more scenarios require measurement systems to work in extremely high temperatures, demanding harsh environment compatible sensors. Application examples are, for instance, low pressure chemical vapor deposition furnaces, oil drilling, combustion monitoring, and the electronics for space exploration [5]. A variety of sensors are designed to fit the high-temperature application. Table 1 summarizes such progress in the last two decades.

As for Pirani gauges, despite the fact that there are many excellent Pirani gauge designs, their performance at high temperatures and corrosive environments is a huge concern. As stated in the literature, Si-based electronics cannot operate at temperatures higher than 125°C [13]–[15]. Beyond this temperature, the minority carrier generation rate dramatically

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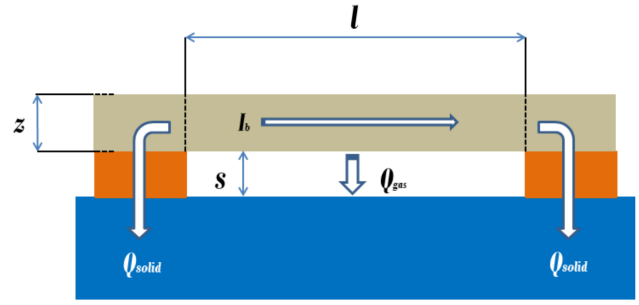
**TABLE I:** Sensors for the high-temperature environment in the recent 20 years.

Researcher [ref.]	Sensor type/Mechanism	Functional material	Max. operating temperature
D.J. Young et al. [6]	Capacitive pressure sensor	SiC	400°C
M. Werner et al. [7]	Piezoresistive pressure sensor	Diamond	170°C
S. Li et al. [8]	Piezoresistive pressure sensor	SOI	350°C
Okojie et al. [9]	Piezoresistive pressure sensor	SiC	800°C
S. Koide et al. [10]	Hall effect sensor	AlGaIn/GaN	400°C
Y. Zhu et al. [11]	Fiber-optic temperature sensor	Sapphire	1600°C
W.J. Pulliam et al. [12]	Fiber-optic pressure sensor	SiC	1100°C
This work	Pirani gauge	SiC	750°C

increases so the leakage current both in the device and in the circuitry becomes significant, which limits the operating temperature of the Si-based smart sensor. As the temperature goes beyond 125°C, the intrinsic carrier concentration will keep increasing and eventually reach the doping concentration, therefore the PN junction will not work. Additionally, when the temperature exceeds 500°C, silicon starts to show plastic deformation under mechanical stress. Beyond 800°C, Si will react with water and/or oxygen, resulting in the formation of an oxide layer on its surface. These properties make traditional Si-based devices show limited feasibility of applying to high-temperature environments. Although some of the micro-fabricated Pirani gauges reported in the literature are made of platinum [16], which is more compatible with high temperatures, the fabrication process shows less CMOS compatibility compared to common semiconductor-based devices and still would be limited by the maximum operating temperature of the Si electronics.

Silicon Carbide (SiC) is one of the most promising wide bandgap semiconductors for the development of integrated circuits and sensors intended for harsh environments. Compared to Si, SiC possesses a much wider bandgap, and it has a much lower intrinsic carrier concentration. As a result, SiC-based devices suffer less from leakage current and generation of minority carriers at elevated temperatures. For this reason, considerable effort has already been invested to develop SiC-based integrated circuits (ICs) aiming for high-temperature stability [17]–[19]. For instance, D.J. Spry et al. demonstrated a SiC IC with more than 100 transistors per chip which can work at 500°C for over 5000 hours [19].

SiC also has excellent mechanical properties. SiC is not only one of the hardest materials on earth, but it also has Young's modulus that ranges from 390 to 690 GPa while Si has Young's modulus of 160 GPa, therefore SiC is advantageous for the realization of suspended structures. Thanks to the strong covalent bond between silicon and carbon atoms (4.6 eV), SiC is highly chemically inert making it an excellent candidate for corrosive environment application. Pressure sensors for



**Fig. 1:** Schematic illustration of a MEMS Pirani gauge. It consists of a suspended heating filament and two anchors to support the filament. This configuration allows to translate an amount of heat loss by the filament through the path  $Q_{gas}$  into a pressure.

over-pressure monitoring based on poly-SiC with maximum operating temperature up to 800°C are reported in the literature [20], illustrating the prospect of SiC as a harsh environment compatible sensing material. It can be concluded that there is potential to incorporate high-temperature sensors with high-temperature readout electronics, eventually achieving harsh-environment compatible smart sensors.

This paper presents the first poly-SiC based Pirani gauges, which is meant to achieve pressure monitoring at high temperatures. Surface micromachining is used as a starting point to enable future monolithic integration with SiC CMOS technologies. A modular design is one of the objectives, enabling tunable device performance, such as measurement range and sensitivity. Moreover, the device design is aimed to be independent from the substrate used, thus favoring its integration possibilities.

## II. PHYSICS & ANALYTICAL MODELLING

The key component of a Pirani gauge is a suspended filament, of which the gaseous conduction of heat to its environment is affected by the ambient pressure ( $P$ ). As shown in Fig. 1, a constant current is injected into the suspended thin filament. This causes its temperature to rise as a result of Joule heating. A temperature distribution will be developed over the filament after the system reaches the thermal equilibrium. Typically, the total heat losses ( $Q_{total}$ ) consist of four parts: thermal convection ( $Q_{convection}$ ), thermal radiation ( $Q_{radiation}$ ), thermal conduction through the anchor parts ( $Q_{solid}$ ) and through the gas ( $Q_{gas}$ ). Among these terms,  $Q_{radiation}$  and  $Q_{solid}$  are not related to pressure directly, so they are both treated as constants [21].  $Q_{convection}$  is negligible because no bulk movement of gas molecules exists at very low pressures [21].  $Q_{gas}$  is the thermal conduction through the gas beneath the suspended filament and is the only heat loss source that is pressure-dependent.

The pressure dependency can be described by Knudsen number ( $K_n$ ):

$$K_n = \frac{\lambda}{d} \quad (1)$$

where  $\lambda$  is the mean free path of gas molecules which increases as pressure drops, and  $d$  is the characteristic length

of a system, which in our case corresponds to the gap size. When  $K_n < 0.01$ , gas molecules will collide with each other intensively. In this condition, referred as continuum regime in fluid mechanics,  $Q_{gas}$  gets saturated and shows no sensitivity to pressure change.

As the pressure becomes lower,  $\lambda$  starts to increase and becomes comparable to the gap size. When  $K_n > 1$ , the sensor will enter the molecular regime, where  $Q_{gas}$  is linear to the number of molecules that participate in the gaseous conduction. If the pressure decreases, the number of molecules that participate in the gaseous conduction will decrease, causing a decrease in  $Q_{gas}$ . Therefore, the temperature of the suspended heater will increase until a new thermal equilibrium is reached. The pressure drop can be transduced into a resistance change of the filament employing the following equation:

$$R_{filament}(P) = R_0[1 + \alpha\bar{T}(P)] \quad (2)$$

where  $R_{filament}(P)$  is the pressure-dependent resistance of the filament, and  $R_0$  is the value of the resistance of the filament measured at standard atmospheric pressure at room temperature (usually 300 K).  $\alpha$  is the temperature coefficient of resistance (TCR) of the filament material.  $\bar{T}(P)$  is the average value of the temperature that exceeds the room temperature. From equation (2), it can be seen that the pressure-dependent average temperature change is needed to determine the filament resistance.  $\bar{T}(P)$  is derived by Mastrangelo *et al.*, therefore  $R_{filament}(P)$  can be further expressed as follows [1]:

$$R_{filament}(P) = R_0[1 + \frac{\delta\alpha}{\epsilon}(1 - \frac{\tanh(\sqrt{\epsilon}\frac{l}{2})}{(\sqrt{\epsilon}\frac{l}{2})})] \quad (3)$$

in which,

$$\delta = \frac{I_b^2 R_0}{\lambda_b w l z}; \quad \epsilon = \frac{\eta \lambda_g(P)}{\lambda_b s z} - \frac{I_b^2 R_0}{\lambda_b w l z} \quad (4)$$

From the model, it can be seen that the dimensions ( $l$ ,  $w$ ,  $z$ , and  $s$ ) of the Pirani gauge determine the sensor performance. The lower detection limit ( $P_L$ ) is limited by the sum of  $Q_{convection}$ ,  $Q_{radiation}$  and  $Q_{solid}$ . At very low pressures,  $Q_{convection}$ ,  $Q_{radiation}$  and  $Q_{solid}$  will exceed  $Q_{gas}$ , and the device will lose sensitivity to the pressure below that point, despite of a large  $K_n$ . On the other hand, the higher detection limit ( $P_H$ ) is limited by the saturation of  $Q_{gas}$  due to the decrease in the mean free path of gas molecules. In order to lower the  $P_L$ , the area of gaseous conduction should be increased by making the filament as long as possible. Besides, the anchor area should be kept as small as possible to minimize  $Q_{solid}$ . In such way, a large  $Q_{gas} / (Q_{convection} + Q_{radiation} + Q_{solid})$  can be obtained thus achieving a lower  $P_L$ . In terms of  $P_H$ , the gap size should be minimized to make the saturation occur at a higher pressure. Generally, the most common ways to design a Pirani gauge with a large dynamic range are: (1) maximize the length of the filament; (2) minimize the gap distance between the filament and the substrate; (3) minimize the solid contact area. However, these design rules usually lead to a suspended filament with low stiffness.

As stated by Mitchell *et al.* [22] and Santagata *et al.* [23], long and thin suspended heaters are more vulnerable

TABLE II: The definitions of the variables in all the equations

Symbol	Physical meaning
$K_n$	Knudsen number
$\lambda$	Mean free path of gas molecules
$d$	Characteristic length of a system
$P$	Pressure
$R_{filament}(P)$	Pressure-dependent resistance of the filament
$R_0$	Initial resistance of the filament
$\alpha$	Temperature coefficient of resistance of filament material
$\bar{T}(P)$	Average value of the temperature that exceeds the environment temperature
$l$	Length
$w$	Width
$z$	Thickness
$s$	Gap size
$\lambda_b$	Thermal conductivity of SiC
$\eta$	Correction factor for the fringing heat flux
$\lambda_g(P)$	Thermal conductivity of air
$I_b$	Injected current
$\rho$	Material density
$C_p$	Heat capacity of the material
$Q$	Heat generation per unit volume of the medium
$P_0$	Transition pressure
$\lambda_{g0}$	Thermal conductivity of the air at a standard atmospheric pressure
$T_a$	Ambient pressure
$v$	Average velocity of gas molecules
$R_{fractional}(P)$	Fractional resistance change

to deflection brought by external interferences. As a result, more heat is going to drain towards the substrate directly, and therefore the sensor will have a smaller dynamic range. In addition, putting the Pirani gauge in a high-temperature operating environment will amplify this effect because of the possible additional thermal buckling which can occur due to thermal expansion of the beam [24]. In order to overcome the buckling issue, researchers have tried different structures to improve the stiffness of long filament, for example, the ladder structure proposed by Mitchell *et al.* [22] and the tube-shaped Pirani gauge proposed by Santagata *et al.* [23].

In order to design a Pirani gauge that is capable to work in at elevated temperatures while showing high sensitivity over a large dynamic range, a MEMS Pirani gauge based on the micro-bridge structure in SiC is implemented. By the usage of poly-SiC, the stiffness of the structure can be greatly enhanced. This allows a longer and thinner Pirani gauge to be built even without dedicated supporting structures. Another advantage of SiC is that the tensile residual stress of the used poly-SiC layers results in flat (non-buckled) suspended beams and cancel out thermal expansion to some extent.

### III. SIMULATION AND DESIGN

A finite element analysis (FEA) was implemented with COMSOL to verify the analytical model in section II. To simulate the operation of the Pirani gauge, the Joule Heating multiphysics interface which couples the "AC/DC" module and the "Heat Transfer" module was used. With the AC/DC module, an electrical current is injected to one end of the heater, and the other end is grounded. The resistivity at the room temperature and the TCR of SiC are also required as inputs of this module to determine the temperature-dependent



(thus the pressure-dependent) filament resistance. To do this, the resistance of an electrical linewidth measurement (ELM) structure was measured from room temperature to 200°C. The resistivity of the poly-SiC layer is  $52.39 \times 10^{-3} \Omega\cdot\text{cm}$ . As indicated by the linear fit in Fig. 2, the poly-SiC layer obtained has a negative TCR of around -1770 ppm/°C in this temperature range.

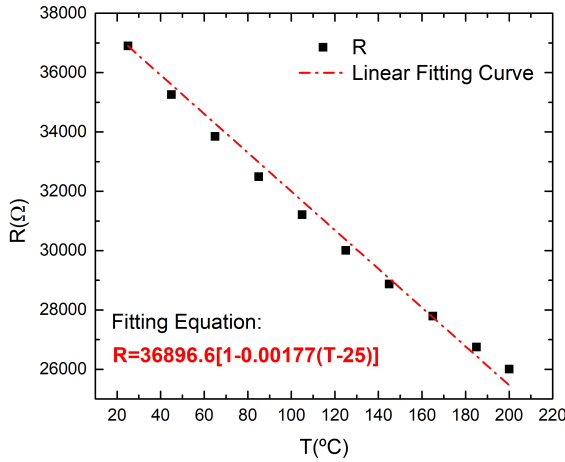


Fig. 2: The black dots show the TCR measurement result using the ELM structure, and the red line indicates the linear fitting result.

In the Heat Transfer module, the temperature distribution on the filament ( $T$ ) is defined by the following equation:

$$\rho C_p \cdot \nabla T = \frac{\partial}{\partial x}(\lambda \frac{\partial T}{\partial x}) + \frac{\partial}{\partial y}(\lambda \frac{\partial T}{\partial y}) + \frac{\partial}{\partial z}(\lambda \frac{\partial T}{\partial z}) + Q \quad (5)$$

where  $Q$  is the heat source.  $\rho$ ,  $\lambda$  and  $C_p$  are density, thermal conductivity and heat capacity of the material respectively. For SiC, the corresponding values are 3210 kg/m<sup>3</sup>, 370 W/(m·K) and 690 J/(kg·K). At both ends of the heating filament, the temperature is fixed at the ambient temperature ( $T_a$ ), which serves as a boundary condition. An air block is inserted between the heater and substrate. The pressure-dependent thermal conductivity of the air  $\lambda_g(P)$  is determined according to equation (6):

$$\lambda_g(P) = \lambda_{g0} \left[ \frac{P/P_0}{1 + (P/P_0)} \right]; \quad P_0 \approx \frac{\eta \lambda_{g0} w T_a}{(w + z)sv} \quad (6)$$

where  $\lambda_{g0}$  is the thermal conductivity of the air at a standard atmospheric pressure.  $P_0$  is the empirical transition pressure.  $v$  is the average velocity of gas molecules. 20 pressure points from 0.1 Pa to 101 kPa were simulated.

The performance of a Pirani gauge ( $l=250 \mu\text{m}$ ,  $w=8 \mu\text{m}$ ,  $z=2 \mu\text{m}$  and  $s=2 \mu\text{m}$ , with 600  $\mu\text{A}$  biasing current) is predicted by both the theoretical model and the COMSOL model mentioned above. As shown in Fig. 3, the results obtained from FEA simulation overlap exactly to the curve predicted by the analytical model. It can, therefore, be concluded that FEA is in agreement with the analytical model.

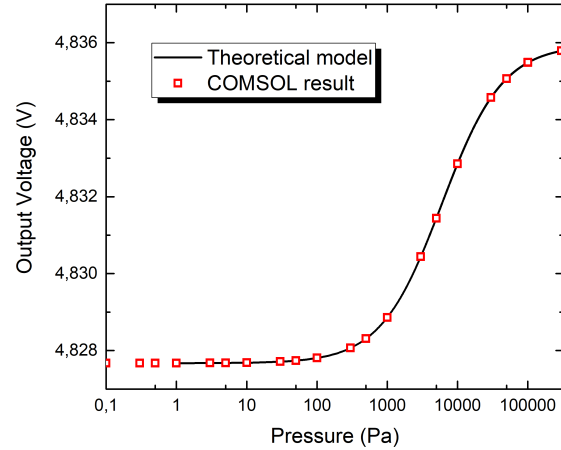


Fig. 3: A comparison between the analytical result and COMSOL simulation.

Tab. 2 summarizes geometries of Pirani gauges that were fabricated. As shown in the table, multiple combinations of length, width, and gap size are taken into consideration. In this study, the thickness of the heating filament is kept constant to 2  $\mu\text{m}$ . As predicted by the analytical model, different geometries would result in different sensor performance such as dynamic range and sensitivity. Therefore, the possibility of integrating two or more Pirani gauges with different geometries into a single chip would allow us to expand the measurement range. The effect brought by different dimensional parameters can be verified at the measurement stage.

TABLE III: Dimensional parameters of the fabricated Pirani gauges

Length ( $\mu\text{m}$ )	Width ( $\mu\text{m}$ )	Gap size ( $\mu\text{m}$ )
250	8	0.5
250	8	1
250	8	2
250	6	2
250	10	2
250	12	2
150	8	2
500	8	2
1000	8	2

#### IV. FABRICATION

The process flow of the Pirani gauge is summarized in Fig. 4. Only three photolithographic steps are needed to fabricate the device. Surface micromachining was used to create the suspended heating filaments. First, a 300 nm  $\text{Si}_3\text{N}_4$  layer was deposited on the silicon substrate by LPCVD with a gas mixture of dichlorosilane ( $\text{SiCl}_2\text{H}_2$ ) and ammonia ( $\text{NH}_3$ ) at 850°C and 20 Pa (Fig. 4a). This layer serves as an additional electrical isolation layer between the metal and the Si substrate, and was patterned with the same mask that will be then used for the metal layer (Fig. 4b). On the front side of the substrate, TEOS (tetraethyl orthosilicate) oxide was coated by means of

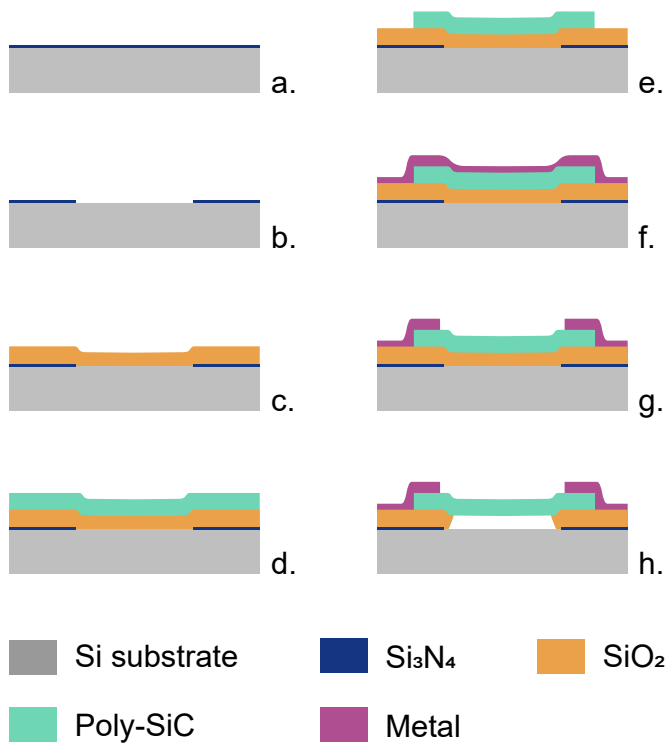


Fig. 4: Schematic illustration of the process steps for the fabrication of the SiC-based Pirani gauge.

plasma enhanced chemical vapor deposition (PECVD) (Fig. 4c). This oxide layer is not only a sacrificial layer for the release of the suspended heating filament (micro-bridge) but also an electrical and thermal isolation layer. The thickness of the oxide defines the gap size of the Pirani gauge. 0.5  $\mu\text{m}$ , 1  $\mu\text{m}$ , and 2  $\mu\text{m}$  oxide layer were deposited on separate batches. After the oxide deposition, a poly-SiC layer of 2000 nm was deposited with  $\text{SiCl}_2\text{H}_2$  (dichlorosilane),  $\text{C}_2\text{H}_2$  (acetylene) as precursor gases, and was in-situ doped with  $\text{NH}_3$  at 860°C and 80 Pa (Fig. 4d). More information on the poly-SiC deposition process can be found in [25]. The deposited carbide layer has a tensile stress of 403.9 MPa, which is within the specification comparing to the result obtained by Bruno *et al.* Subsequently, a patterning step was conducted to define the suspended micro-bridge structure and the anchor area (Fig. 4e).

Then, a metal layer was sputtered (Fig. 4f) and patterned (Fig. 4g) to form the metal lines and pads for electrical measurement. Aluminum (Al) and titanium (Ti) were used as metallization materials. Earlier research showed that these metals will form an ohmic contact to the doped poly-SiC layers employed in this work [5], [26], [27]. A post-deposition annealing process in a nitrogen environment was followed to relieve the strain and improve the contact property at the metal-semiconductor interface. Finally, the sacrificial oxide layer was etched away to release the micro-bridge (Fig. 4h). In order to avoid any stiction caused by the capillary forces during a wet releasing step, vapor HF was used.

The longest Pirani gauge that was fabricated had a length of 1000  $\mu\text{m}$ . Although this geometry is in principle most likely to show the highest deflection between the different designs,

it can be seen from Fig. 5 that the realized 1000  $\mu\text{m}$  does not show any deflection after realization.

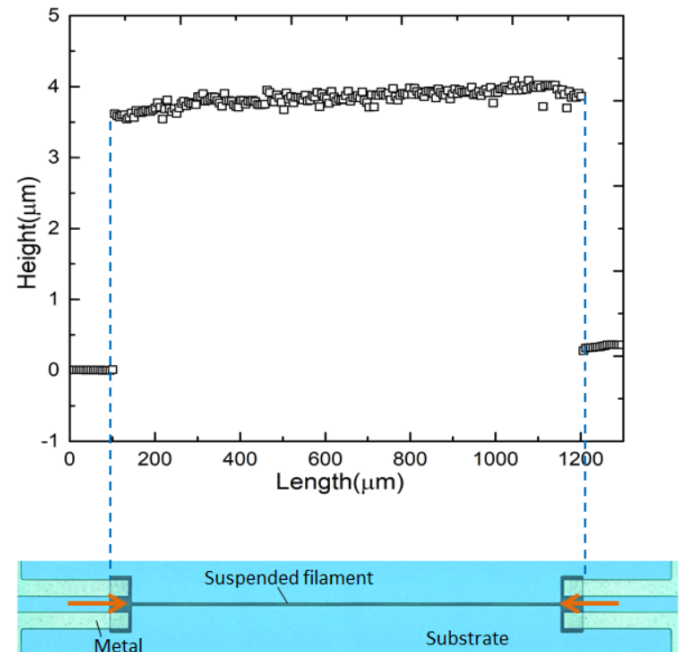


Fig. 5: Distance between the top surface of the micro-bridge and Si substrate along the beam. The orange arrows indicate the position of the cross-section.

## V. RESULTS AND DISCUSSION

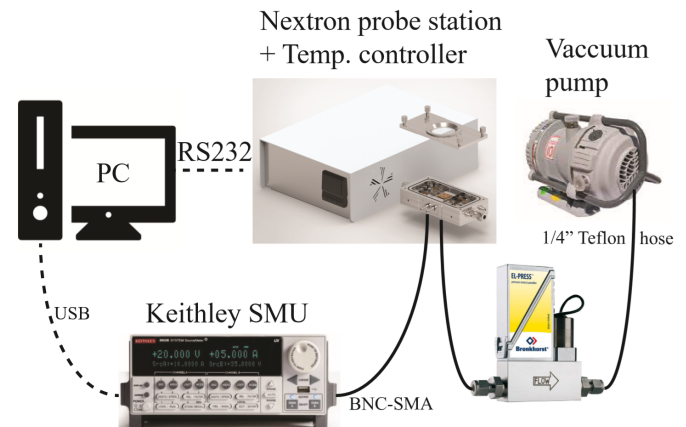


Fig. 6: Measurement setup used for calibration [28]. The setup contains a MPS, a mass flow controller, a temperature controller and a vacuum pump.

### A. Measurement Setup

A dedicated measurement setup for calibrating Pirani gauges is built as shown in Fig. 6. The setup is able to provide a low-pressure environment with values ranging from 10 Pa to atmospheric pressure. The controller of this setup can change the pressure in steps of 500 Pa while the internal pressure sensor has a resolution of 10 Pa. The micro-fabricated Pirani

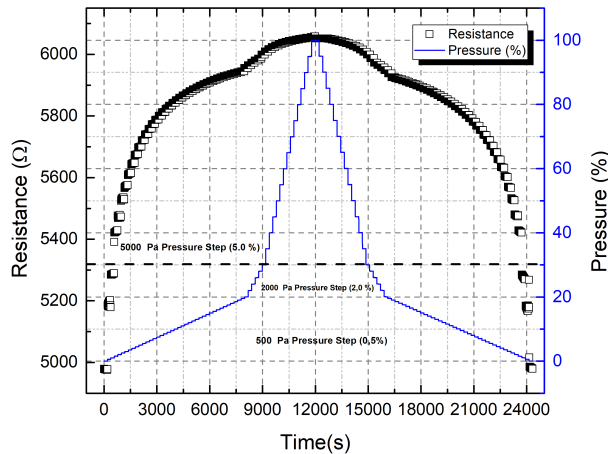


Fig. 7: The pressure profile for the calibration (blue curve) and the output voltage of the reference Pirani gauge as a function of pressure (black dots).

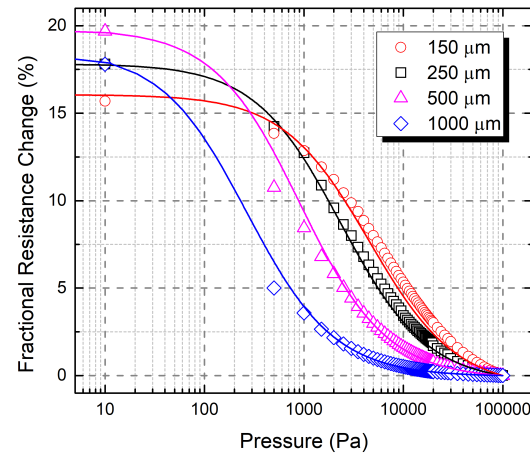
gauges are calibrated by using the pressure values from the internal pressure gauge.

The setup also contains a microprobe station (MPS) which is equipped with four-probe needles. These probe needles are connected to a source meter (Keithley 2612B) to measure the resistance of Pirani gauges. A hotplate is also integrated with the probe station and is able to create a hot surface up to 750°C.

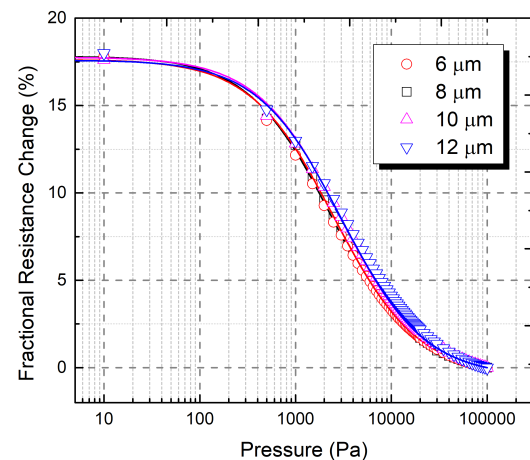
When calibrating a Pirani gauge, the device is put in the MPS where a certain vacuum is created. Below 20 kPa, the pressure increment was 500 Pa, because the sensitivity of the realized gauges is larger in this pressure regime. As for 20-30 kPa and 30-100 kPa, the pressure step was set as 2000 Pa and 5000 Pa respectively. In total, there were 60 pressure set-points used for calibration. For each of the points, the measurement was paused for 200 s so that the pressure in the micro-chamber can fully stabilize whereafter the electrical measurement is performed. The pressure profile for calibration is shown by the blue curve in Fig. 7.

### B. Dimensioning of the Pirani gauge

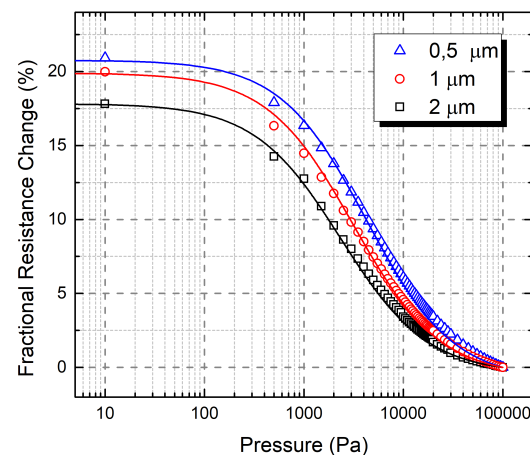
As stated in Mastrangelo's model, dimensional parameters play a significant role in the performance of the sensor. To study the effect of the variation in the geometry on the performance, a Pirani gauge with a length of 250  $\mu\text{m}$ , a width of 8  $\mu\text{m}$ , a thickness of 2  $\mu\text{m}$  and a gap size of 2  $\mu\text{m}$  is taken as a reference device. The performance of the reference gauge was measured and was compared to the performance of gauges with other dimensions. The reference device was biased with a constant current of 600  $\mu\text{A}$ . The calibration curve of the reference device is plotted in Fig. 7. The gauge resistance ranges from 4978.3  $\Omega$  to 6060.0  $\Omega$  from the lowest pressure set-point of 10 Pa up to 100 kPa, which corresponds to a maximum resistance deviation of 17.8 %. The Pirani gauge shows a response to the pressure change without hysteresis, as



(a)



(b)



(c)

Fig. 8: The calibration curves for device with different (a) lengths; (b) widths; (c) gap sizes comparing to the reference device. The scattered points represent experimental data, while continuous curves represent fitted data.

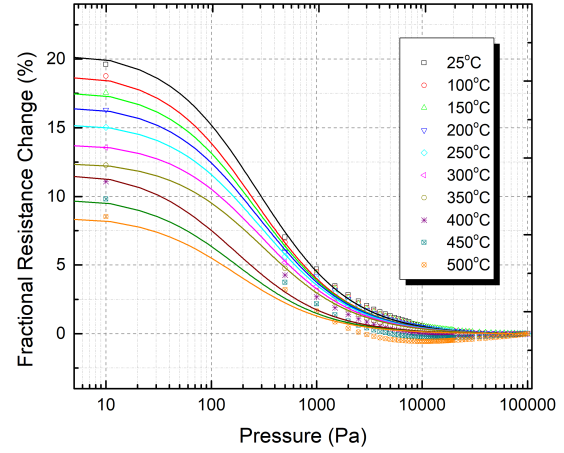
indicated by the symmetrical shape of the calibration curve in Fig. 7. The performances of the gauges with other dimensions are demonstrated in Fig. 8. In this figure, the scattered data points are obtained with the measurement setup, and the continuous curves are fitting curves based on equation (3) according to the experimental data. From the fitting data, the reference gauge has the highest sensitivity of  $0.47 \text{ } \Omega/\text{Pa}$  at around  $10 \text{ Pa}$ .

Fig. 8a shows the performance of gauges with different lengths. Since Pirani gauges with various dimensions have different initial resistance, the output resistance is expressed with the fractional resistance change  $R_{fractional}(P)$  as shown in equation (7). With increasing length, the calibration curve shifts towards the left because  $Q_{gas}$  takes a larger proportion in the total heat transfer. As a result, the longer gauge has a smaller  $P_L$ . From  $150 \text{ } \mu\text{m}$  to  $1000 \text{ } \mu\text{m}$ , the maximum sensitivity increases from  $0.13 \text{ } \Omega/\text{Pa}$  to  $16.76 \text{ } \Omega/\text{Pa}$ . It is worth noting that the maximum sensitivity of each gauge locates at  $10 \text{ Pa}$ , which implies that the gauge is still sensitive below  $10 \text{ Pa}$ .

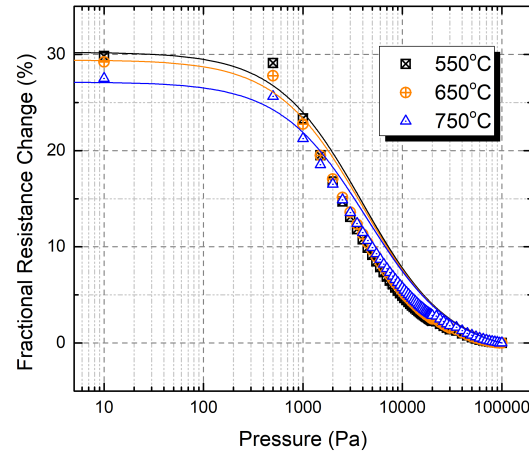
$$R_{fractional}(P) = \frac{R_0 - R_{filament}(P)}{R_0} \times 100\% \quad (7)$$

Fig. 8b shows the impact of the width variations. The reduction of the heater width only causes a slight left shift of the calibration curve. Despite a larger width increases  $Q_{gas}$ , the solid contact between the filament and the anchor region also increases with the width, causing  $Q_{solid}$  to increase with the same order. Therefore, the dynamic range of gauges with different width does not vary too much. However, due to the increase in the heat exchange surface, the wider gauge is less power-efficient. For example, the narrowest gauge ( $w=6 \text{ } \mu\text{m}$ ) and the widest gauge ( $w=12 \text{ } \mu\text{m}$ ) have initial resistances of  $8591.5 \text{ } \Omega$  and  $3771.1 \text{ } \Omega$ , and are biased with a current of  $424 \text{ } \mu\text{A}$  and  $965 \text{ } \mu\text{A}$  respectively. As shown in Fig. 8b, both of them have a similar fractional resistance change of around  $17.8 \%$ , while the narrowest gauge consumes  $56.2 \%$  less power than the widest gauge.

Devices with different gap sizes ( $0.5 \text{ } \mu\text{m}$ ,  $1 \text{ } \mu\text{m}$ , and  $2 \text{ } \mu\text{m}$ ) were measured. As can be seen from Fig. 8c, the device with a smaller gap (blue curve) tends to saturate at a higher pressure. In general, the experimental results align with the prediction of the analytical model qualitatively: longer beam will have larger sensitivity in low pressure range, and the gauge with smaller air gap has larger sensitivity near the standard atmospheric pressure. However, numerically, the experimental results do not match the analytical results. From the measurement, the reference gauge shows a  $17.8 \%$  fractional resistance change, while the analytical model and COMSOL simulation both predict only  $0.16 \%$  of fractional resistance change. The reason for the discrepancy could be that the material properties of poly-SiC reported in the literature which used as inputs of the analytical model, such as thermal conductivity, are different from the material properties of SiC we deposited.



(a)



(b)

Fig. 9: The calibration curve of devices ( $l=1000 \text{ } \mu\text{m}$ ,  $w=8 \text{ } \mu\text{m}$ ,  $z=2 \text{ } \mu\text{m}$  and  $s=2 \text{ } \mu\text{m}$ ) (a) with Al pads from  $25^\circ\text{C}$  to  $500^\circ\text{C}$ ; (b) with Ti pads from  $550^\circ\text{C}$ ,  $650^\circ\text{C}$  and  $750^\circ\text{C}$ . The sensitivity decreases as temperature rises. The scattered points represent experimental data, while continuous curves represent fitted data.

### C. High-temperature Characterization

The micro-fabricated Pirani gauges were characterized at different temperatures to examine the high-temperature performance. The longest Pirani gauge ( $1000 \text{ } \mu\text{m}$ ) was used for the test because it is most likely to experience a buckling among all the geometries. First, the Pirani gauges with conventional Al metallization was measured. The measurement was carried out from  $25^\circ\text{C}$  to  $500^\circ\text{C}$  in steps of  $50^\circ\text{C}$ . The calibration curves are plotted in Fig. 9a. As the temperature increases, the fractional resistance change reduces and the maximum sensitivity decreases from  $17.63 \text{ } \Omega/\text{Pa}$  at  $25^\circ\text{C}$  to  $6.15 \text{ } \Omega/\text{Pa}$  at  $500^\circ\text{C}$ , which results from the decrease in the material TCR at an elevated temperature.



For characterization beyond 500°C, the realized devices with Ti metallization were measured. The reason for this is the higher melting point of Ti of 1668°C compared to 575°C for Al with 1 % Si. Additionally, Ti has a work function of 4.33 eV, which is similar to Al (4.06-4.26 eV). As reported in [26], [27], Ti is able to form an Ohmic contact with SiC and is less reactive and more compatible with high-temperature environments. The measurement results at 550, 650, and 750°C are given in Fig. 9b. Even at these temperatures, the device demonstrates a repeatable response to pressure change. The maximum sensitivity can still be obtained at 10 Pa, which is 1.35  $\Omega$ /Pa at 550°C, 1.23  $\Omega$ /Pa at 650°C, and 0.81  $\Omega$ /Pa at 750°C. To the best of authors' knowledge, this is the first time that the operating temperature of the Pirani gauge has been pushed to such a high temperature.

## VI. CONCLUSION

This work reports on the realization of a CMOS compatible vacuum gauge operating at elevated temperatures. A MEMS Pirani gauge in SiC is fabricated with surface micro-machining technique by means of a three-mask process. The use of SiC combines favorable properties such as a useful TCR, high mechanical stiffness, and compatibility with high temperatures and harsh environments. The realized Pirani gauges can achieve a length of 1000  $\mu$ m without any mechanical supporting structure to prevent buckling. At room temperature, the gauge with reference geometry showed a maximum sensitivity of 0.47  $\Omega$ /Pa and a fractional resistance change of 17.8 %. High-temperature characterization on the 1000  $\mu$ m long beam revealed the desired operation even up to a maximum temperature of 750°C, with a maximum sensitivity of 0.81  $\Omega$ /Pa. These results further illustrate the potential of poly-SiC Pirani gauges for applications at high temperatures and in harsh environments. However, at 750°C, the filament resistance of some devices increased while the pressure was decreasing. The future work will include a closer investigation into the undesired device performance beyond 750°C. In such an analysis the temperature dependency of the TCR, the possible buckling and radiative losses at high temperatures need carefully to be taken into account.

The future outlook of this work is to be integrated with state of the art SiC CMOS technologies to achieve an all-SiC monolithically integrated MEMS + ASIC platform. In order to achieve this, the process flow of this poly-SiC MEMS sensor should be integrated into the back-end of line processing of the SiC CMOS process, where thermal budget, metallization, contamination and passivation aspects need to be considered.

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