VARYING POTENTIAL SILICON CARBIDE GAS SENSOR

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Field of Search: G01N 27/12; G01N 27/06; HOI1 29/66

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A hydrocarbon gas detection device operates by dissociating or electro-chemically oxidizing hydrocarbons adsorbed to a silicon carbide detection layer. Dissociation or oxidation are driven by a varying potential applied to the detection layer. Different hydrocarbon species undergo reaction at different applied potentials so that the device is able to discriminate among various hydrocarbon species. The device can operate at temperatures between 100° C. and at least 650° C., allowing hydrocarbon detection in hot exhaust gases. The dissociation reaction is detected either as a change in a capacitor or, preferably, as a change of current flow through an FET which incorporates the silicon carbide detection layers. The silicon carbide detection layer can be augmented with a pad of catalytic material which provides a signal without an applied potential. Comparisons between the catalytically produced signal and the varying potential produced signal may further help identify the hydrocarbon present.

27 Claims, 5 Drawing Sheets
**FIG. 6**

![Graph showing the relationship between Relative Current (mA) and Applied Potential (Volts) for Methane and Propene.]

- **Methane** represented by a square symbol.
- **Propene** represented by a circle symbol.
FIG. 8

80. PLACE SENSOR

84. HEAT SENSOR

86. PROVIDE SAMPLE

88. APPLY VARYING POTENTIAL

90. MEASURE CURRENT AS QUANTITY OF HYDROCARBON

92. DETERMINE HYDROCARBON AS FUNCTION OF POTENTIAL

94. REVERSE POTENTIAL TO RENEW SENSOR
VARYING POTENTIAL SILICON CARBIDE GAS SENSOR

ORIGIN OF THE INVENTION

The invention described herein was made in the performance of work under a NASA contract, and is subject to the provisions of Public Law 96-517 (35 U.S.C. §202) in which the Contractor has elected not to retain title.

TECHNICAL FIELD

This invention is in the field of semiconductor sensors and, more specifically, involves the detection of hydrocarbon gases by a semiconductor sensor.

BACKGROUND ART

There is a significant need for a compact, rugged component capable of detecting hydrocarbon gases at relatively high temperatures. Such a device would be useful for monitoring the presence of unburned hydrocarbons in the exhaust gases of furnaces, turbines, or internal combustion engines that operate by burning hydrocarbon fuels. When a combustion device fails to completely burn its fuel, there is, of course, significant economic waste as purchased fuel is discarded into the exhaust stream without yielding usable energy. Furthermore, there is often a deposition of soot and other products of partial combustion that may contribute to mechanical wear and failure.

An even more pressing problem posed by incomplete combustion is the significant contribution of unburned hydrocarbons to air pollution. The hydrocarbons expelled with an engine’s exhaust react with ozone and oxides of nitrogen to create photochemical smog, that nostrous brown pall that is the bane of many highly populated urban areas. Not only does smog have a loathsome appearance, obscuring distant vistas, but it also poses significant health hazards. Children growing up in a polluted atmosphere show distinct respiratory impairment. Adults exposed to smog experience a significantly greater incidence of asthma and other respiratory diseases as well.

It is well recognized that automobiles and other vehicles with internal combustion engines are significant contributors to smog. Consequently, modern autos are equipped with catalytic converters to remove hydrocarbons in the exhaust through oxidation. In addition, modern automobiles employ complex microprocessor-controlled systems to ensure that fuel mixtures burn completely so as to avoid overloading the catalytic converter. However, the absence of an effective hydrocarbon sensor capable of operating in the hot exhaust stream means that the automobile control systems are unable to directly sense the presence of hydrocarbons; therefore, they often err on the side of lean mixtures to avoid excess hydrocarbon emissions, frequently compromising performance in the process.

When a mechanical defect results in excessive hydrocarbon emission, the problem is often not detected until after the catalytic converter is overloaded and damaged. If the control system could directly sense the presence of hydrocarbons, the fuel mixture could be constantly adjusted to provide maximum performance without excess hydrocarbon emission. If hydrocarbon emission did become excessive through a mechanical failure, an alarm could be sounded so the car owner could immediately obtain necessary repairs.

Semiconductor gas sensors are already known. There are sensors capable of detecting, for example, carbon monoxide, hydrogen, and oxygen. Many of these sensors are based on semiconductor materials, such as tin dioxide or titanium dioxide, that change electrically resistive properties upon absorbing a gas. Thus, the sensors are conductometric—that is, changing their resistance in a predictable manner upon absorption of a given gas. Generally, these sensors can operate with only one species of adsorbable gas present since, with more than one gas capable of being adsorbed by the sensor, there is no way of knowing which of the several gases is actually present and causing a change in resistance. In theory, a conductometric hydrocarbon sensor might be created, but discrimination among the large number of different hydrocarbons would pose a significant problem.

It is also known in the art that properly prepared crystalline silicon carbide is useful in the formation of semiconductor devices (see J. H. Edgar, J. Mater. Res. 7:235-52 (1992)). Silicon carbide is known to have a wider energy gap band than more traditional semiconductors such as silicon. This allows the construction of silicon carbide semiconductor devices which are capable of operating at higher temperatures, i.e., well above 280°C, than are silicon semiconductors. The favorable properties of silicon carbide make it possible to prepare semiconductors such as field effect transistors (FET) from this material. In fact, metal oxide field effect transistors (MOSFET), as well as other devices, have already been prepared using silicon carbide.

Although, prior to the instant invention, no hydrocarbon-sensitive silicon carbide FETs had been developed, there has been some work using metal oxide-silicon carbide capacitors to detect hydrocarbon gases. These capacitors consist of a layer of catalytic metal deposited on an insulator which, in turn, is disposed on a layer of silicon carbide. Some of these experiments are reported in A. Arbab, S. Spetz, Q. Wahab, M. Willander, and I. Lundstrom, Sensors and Materials 4:173-85 (1993), and A. Arbab, S. Spetz, and I. Lundstrom, Sensors and Actuators B 15/16:19-23 (1993). The devices reported produced only low-level signals, required very high operating temperatures (generally >500°C), and did not appear capable of discriminating among various hydrocarbons.

STATEMENT OF THE INVENTION

A hydrocarbon gas detection device operates by dissociating or electrochemically oxidizing hydrocarbons adsorbed to a silicon carbide detection layer. Dissociation or oxidation are driven by a varying potential applied to the detection layer. Different hydrocarbon species undergo reaction at different applied potentials so that the device is able to discriminate between various hydrocarbon species. The device can operate at temperatures from 100°C to over 50°C, allowing hydrocarbon detection in hot exhaust gases. The dissociation reaction is detected either as an electrical change in a metalinsulator silicon carbide (MIS) capacitor or, preferably, as a change of current flow through a metalinsulator silicon carbide field effect transistor (MISFET) of which the silicon carbide detection layer forms the gate. The silicon carbide detection layer can be augmented with a pad of catalytic material which can provide a signal without an applied potential; in addition, potential can be applied to the catalytic pad to enhance the catalytic reaction. Comparisons between the catalytically-produced signal and the varying potential produced signal further helps identify the hydrocarbon present.

BRIEF DESCRIPTION OF THE DRAWINGS

The objects and features of the present invention, which are believed to be novel, are set forth with particularity in the
The invention is based on the discovery that a process, having opposite electrical effects, may have some similarity to the electronic attraction of a catalytic pad; it may represent an electrochemical attraction. If an electrical potential at which a sensor operates is varied continuously, the net change in charge caused by the varying potential-induced processes can control the flow of current through the channel of the silicon carbide MISFET either through enhancement or depletion, depending on the precise configuration of the FET design chosen.

Depending on the exact characteristics of the measured gas (e.g., the presence of oxygen or water) and the magnitude of the applied varying potential (0 to about 2 V versus 2 to about 10 V), either or both electrochemical oxidation and bond dissociation can occur on the same sensor. In practice, the varying potential is varied continuously (swept) at a constant rate from 0 to over 10 V to cover the entire range. The actual potential at which a signal is observed will indicate which process is occurring and which type of hydrocarbon is involved. Further, depending on the configuration of the MISFET (i.e., p-type or n-type, depletion mode, or enhancement mode) each process will either increase or decrease current flow through the transistor, with the two processes, having opposite effects because electrochemistry yields negative ions, while dissociation yields positive ions. FIG. 1 shows a cross-section of a diagram of one configuration of a sensor 10 of the present invention; FIG. 2 shows a top view of the device 11; as will be explained below, various fabrication constraints result in other configurations being preferred. In the basic design of FIG. 1, the actual potential at which a sensor operates is varied continuously (swept) at a constant rate from 0 to over 10 V to cover the entire range. The actual potential at which a signal is observed will indicate which process is occurring and which type of hydrocarbon is involved. Further, depending on the configuration of the MISFET (i.e., p-type or n-type, depletion mode, or enhancement mode) each process will either increase or decrease current flow through the transistor, with the two processes, having opposite effects because electrochemistry yields negative ions, while dissociation yields positive ions. 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structing a multilayered semiconductor device, attention must be paid to crystal structures created to ensure that the crystal lattices of the various layers align properly. Two of the silicon carbide crystal types are of particular interest: the cubic (zincblende) or 3C—SiC form and the hexagonal or 6H—SiC form. A MISFET-based sensor 10 is constructed by forming a region of doped (with added impurities) silicon carbide called a current channel 14 and a gate 18 of undoped polycrystalline silicon carbide separated by a thin insulator layer 17. Figure 17 shows a cross-section of a sensor 30 constructed to permit epitaxial growth. A typical epilayer can be grown on the substrate following methods well known in the art (see, for example, R. Davis, Physica B 185:1-15 (1993)). The first epilayer forms the current channel 14 of the device 10. An epilayer of aluminum nitride (see L. B. Rowland, R. S. Kern, S. Tanaka, R. F. Davis, Appl. Phys. Lett. 62:3333-35 (1993)) is then grown on top of the silicon carbide channel 14 followed by a polycrystalline layer 17 of silicon carbide grown in the same reaction chamber to prevent contamination. The exposed surfaces of the device 10 are protected by passivation using thermal oxidation in dry oxygen for about 30 minutes at 1100°C. The oxide is removed from the polycrystalline silicon carbide layer 17 and from the contact areas for the source electrode 22 and the drain electrode 24 using hydrofluoric acid. Chromium to form the metal grid 26 and the varying potential 28, the source 22 and the drain 24 electrodes is then vacuum evaporated onto the device 10 typically at 1×10^-6 to 1×10^-7 torr.

As reported in J. S. Shor, I. Gringberg, B. Weiss and A. D. Kurtz, Appl. Phys. Lett. 62:2836-2838 (1993), it is possible to create porous silicon carbide by an etching process which creates pores oriented along the C-axis of hexagonal or 6H—SiC. Porous silicon carbide is ideal if the pores are oriented perpendicularly to the top surface of the structure so that a large number of pore openings are exposed for the hydrocarbon gases to enter, thereby providing a large surface area for gas adsorption. However, the process of etching silicon carbide imposes certain limitations on the configuration of the resulting device.

The silicon carbide is etched through a photoelectrochemical anodization process. The silicon carbide must be attached to an electrical conductor on one surface, while the opposite surface is exposed to a strong (about 300 mW/cm²) beam of ultraviolet (230-400 nm) light in a dilute hydrofluoric acid bath. It is not possible to readily apply this process to the silicon carbide gate 18 as configured in FIGS. 1 and 2. Therefore, one must employ a different fabrication scheme which starts with a silicon carbide wafer and converts that wafer into a porous gate.

FIG. 3 shows a cross-section of a sensor 30 constructed to permit the use a porous etched silicon carbide current gate 48. The porous gate 48 is topped by a metal grid 26 and varying potential electrode 28 just as in the sensor 10 shown in FIGS. 1 and 2. A porous layer 47 is opposite the aluminum nitride insulator 16 so that the porous silicon carbide layer 47 is sandwiched between the grid 26 and the insulator 16. A doped silicon carbide channel 44 covers the insulator 16. Source 22 and drain 24 electrodes are deposited onto the channel 44.

The structural differences between the polycrystalline sensor 10 and the porous layer gate sensor 30 are a result of the requirement for a backing conductor during the photoelectrochemical etching process. Manufacture of the porous gate sensor 30 starts with a thin wafer 32 of hexagonal silicon carbide cut so that the C-axis is normal to the wafers planar surfaces. A metal electrode (not illustrated) is deposited on one surface of the wafer and the wafer is photoelectrochemically etched so that 10-100 nm pores are formed on the planar surface opposite from the electrode. This leaves a top surface of the wafer 32 transformed into the porous silicon layer 47 which forms the porous layer gate 48.

Generally an undoped wafer 32 about 200-500 μm thickness is used and a porous layer 47 one to a few micrometers thick is etched into the wafer 32. Thus, much of the unetched wafer 32 remains. While the undoped wafer acts as an n-type...
A semiconductor due to atmospheric nitrogen contaminants incorporated during manufacture, the porous etched silicon carbide layer has very high resistivity. The backing metal electrode is then removed chemically and the aluminum nitride insulator 16 is deposited in its place. A layer of n-type doped silicon carbide is then grown on the surface of the insulator 16 to constitute the channel 44. The metal grid 26 and the varying potential electrode 28 are deposited onto the porous surface 47 of the porous silicon carbide gate 48. The sensor 30 is completed by depositing the source 22 and drain 24 electrodes onto the channel 44. These fabrication steps are the same as those explained above for FIGS. 1 and 2. Those of ordinary skill in the art will appreciate that the passivation step is important to protect the device.

Thus, the porous gate sensor 30 is not formed by growing various layers onto a silicon carbide wafer 12 as is the case with the polycrystalline gate sensor 10. Rather, a silicon carbide wafer 32 is etched to form the porous silicon gate 48 and the remaining layers of the device are then grown onto the gate 48. The major difference visible in the final device is the absence of the silicon carbide wafer 12 backing the channel 44. The polycrystalline silicon carbide gate device 10 shows the metal grid 26 on one surface and the silicon wafer 12 on the opposite surface. The porous gate device 30 grown according to this method shows the metal grid 26 on one surface and the channel 44 with electrodes 22 and 24 on the opposite surface.

Both the polycrystalline gate MISFET 10 and the porous gate MISFET 30 have been illustrated as depletion mode devices. As is well known in the art, these devices can readily be constructed as enhancement mode MISFETs. The invention lies in the employment of a porous or polycrystalline silicon carbide gate to which a varying potential is applied. The gate can be electrochemical oxidation or dissociation of adsorbed hydrocarbons.

In carrying out experiments to test various parameters of the polycrystalline or porous silicon carbide sensor 10, 30, there are considerable advantages in cost and time to testing the porous silicon carbide or polycrystalline silicon carbide alone as opposed to constructing entire MISFETs. FIG. 5a shows a device 60 fabricated to test an individual polycrystalline layer 17; the polycrystalline silicon carbide layer 17 is deposited on the insulator 16 which, in turn, has been deposited directly onto the silicon carbide wafer 12, the metal grid 26, and the varying potential electrode 28 are then deposited onto the polycrystalline silicon carbide 17. The back electrode 46 is added to complete the device 60. It has been found that polycrystalline silicon carbide may be excessively conductive due to grain boundary conduction. This problem can be controlled by slightly doping the polycrystalline silicon carbide with an electron acceptor such as boron.

FIG. 5b shows a device 50 fabricated to test an individual porous silicon carbide layer 47; the silicon carbide wafer 32 is equipped with a back electrode 46 for the photoelectrochemical etching process as explained above. After the etching is complete, the open metal grid 26 and the varying potential electrode 28 are deposited on the porous surface 47.

The described test structures, FIGS. 5a, 5b, comprise sandwiches of two electrodes (the back electrode 46 and the metal grid 26) with other layers between. These devices act as MIS capacitors 60. Those of ordinary skill in the art will appreciate that the presence of hydrocarbons adsorbed to the polycrystalline silicon carbide layer 17 and the porous silicon carbide layer 17, respectively, when adsorbed hydrocarbons are dissociated by the applied varying potential.

FIG. 6 shows typical results of testing the pores silicon carbide capacitor 50. A flow of current (Relative Current) is measured across the electrodes 28 and 46 as the applied potential is varied between -8 V and +8 V at a temperature of 223°C. This temperature is considerably lower than that reported for gas detecting capacitors based on catalytic materials. A first trace represents the current produced by methane, the smallest alkane. A second trace shows the current produced by propane, a three carbon alkene. Both traces are shown relative to the current produced by the sensor 50 in an atmosphere of inert argon, to zero out any current leakage in the capacitor. The difference in shape between the traces produced by the two different hydrocarbons reflects the differences in bond energy between propane and methane showing that the device can discriminate between different hydrocarbons.

One problem that may affect the performance of the sensors of the present invention is excess accumulation of reaction products or "foiling" of the detection layer by adsorbed hydrocarbons or impurities. It has been found that reversing the polarity of the varying potential and increasing its magnitude often results in expulsion of at least some of the excess accumulation, thus regenerating the detection layer. Care must be taken not to use so high a potential as to puncture the thin insulator 16. In most cases a potential of 100 V or so can be safely, tolerated. In many cases the devices of the present invention will be fabricated with integral zener diodes to protect the insulator 16 as is common in many MOSFETs. If potential reversal is not adequate to regenerate the device, mechanical cleaning such as by ultrasound may be necessary.

FIGS. 7a–7b show additional modifications that can be made to the porous silicon carbide layer 47; the modification of FIG. 7a can be made to the polycrystalline silicon carbide layer 17 as well. The modifications can be applied to detection layers of either MISFETs or to MIS capacitors. The figures illustrate only a portion of the detection layer. The variable potential electrode 28 as well as other layers of a functioning device are not illustrated.

In FIG. 7a a pad 52 (or pads) of a catalytic metal or a catalytic metal compound is deposited over a portion of the polycrystalline or porous silicon carbide layers 47. The pad 52 can be deposited by conventional evaporation techniques in the case of a pure metal or by CVD in the case of metal compounds. The list of usable metals and compounds is long; virtually any material that shows catalytic activities towards hydrocarbons is useful. This includes platinum, vanadium, molybdenum carbide, tungsten carbide, titanium carbide, niobium carbide, and palladium. The pad 52 can be in contact with the metal grid 26 (not illustrated), in which case the applied potential may also augment the catalytic reaction. Alternatively, the pad 52 may be isolated from the grid 26 so that there is no interaction between the applied potential and the pad 52.

The catalytic material is able to dissociate hydrocarbons without an applied varying potential, although it requires a higher temperature than is needed with porous silicon carbide. By covering only part of the porous surface 47 with the pad 52, there is still plenty of area of porous silicon carbide to adsorb hydrocarbon gases. If the device is operated at a low temperature (i.e., 100°C–400°C), the signal developed will be primarily due to dissociation of hydrocarbons in the porous silicon carbide 48 and will be dependent upon the applied varying potential. At a higher temperature (i.e.,
catalytic pads 52, it may be advantageous to include time periods where the varying potential is entirely switched off so that the output signal is indicative of the catalytic reaction only.

Step 90 comprises measuring an electrical characteristic that reflects binding and dissociation of hydrocarbons in the silicon carbide detection layer. In the case of transistor devices this would represent a change in the amount of current passing through the device from the source electrode 22 to the drain electrode 24. In the case of capacitor devices this would represent either a change in capacitor voltage measured by subjecting the capacitor to a higher frequency signal supplied in addition to the more slowly changing varying potential or a change in a current measured between the variable potential electrode 28 and the back electrode 46.

In step 92 the electrical characteristic signal is correlated with the potential at which it occurs to identify the type of hydrocarbon present. As explained above, electrochemical reactions occur mostly between 0.3 V and 2–3 V, while dissociations occur mostly between 2 V and about 10 V. Individual chemical compounds tend to fall at slightly different potentials. Where both dissociation and electrochemical oxidation occurs, comparison of the potential for each type of reaction may identify a given chemical species. In other cases, comparison of a catalytic signal with a varying potential signal may identify the chemical species being measured.

The final optional step 94 is the application of a reverse potential to regenerate the sensor by helping to drive off reaction products and attracted impurities. Generally, the regenerating potential is about equal to the maximum dissociation potential (10 V) but is of an opposite polarity. In some cases a higher regeneration potential (i.e., up to about 100 V) gives improved regeneration. There is also some advantage to applying the dissociating potential as a continuously varying potential that sweeps from one extreme (say 10 V) to the opposite extreme (–10 V). In this way the device is regenerated constantly.

Those skilled in the art will appreciate that various adaptations and modifications of the just-described preferred embodiment can be configured without departing from the scope and spirit of the invention. Therefore, it is to be understood that, within the scope of the appended claims, the invention may be practiced other than as specifically described herein.

We claim:

1. An electronic device of semiconductor design for detecting hydrocarbon gases comprising:
   a detection layer having first and second surfaces, and forming a gate of a metal insulator field effect transistor, the detection layer comprising:
   a wafer of silicon carbide;
   a layer of porous silicon carbide formed at a first surface of said silicon carbide wafer by photoelectrochemical etching, the layer of porous silicon carbide for adsorbing hydrocarbon gases;
   a metal grid deposited onto the porous silicon carbide layer; and
   an electrode in electrical communication with the metal grid for applying a varying dissociating potential to said porous silicon carbide layer for dissociating hydrocarbon molecules adsorbed to said porous silicon carbide layer;
   an insulation layer formed on a second surface of the detection layer;
   a channel layer of doped silicon carbide formed on an exposed surface of the insulation layer; and
11. Drain and source electrodes disposed on the channel layer at opposite sides thereof so that dissociation of hydrocarbons in the detection layer causes an alteration in a current flow between the drain electrode and the source electrode, a magnitude of said alteration indicating a quantity of hydrocarbon gas adsorbed to the detection layer and a magnitude of the dissociating potential indicating a type of hydrocarbon gas adsorbed to the detection layer.

2. The electronic device of claim 1, wherein the wafer of silicon carbide forming the detection layer is hexagonal form silicon carbide fabricated with its C-axis normal to the first surface of said silicon carbide wafer.

3. The electronic device of claim 1, wherein the dissociating potential varies between 0 V and about 10 V.

4. The electronic device of claim 1, wherein the insulation layer comprises a layer of single crystal aluminum nitride.

5. The electronic device of claim 1, wherein the detection layer further comprises a layer of catalytic material deposited over an portion of the first surface of the porous silicon carbide layer.

6. The electronic device of claim 5, wherein the catalytic material is selected from the group consisting of platinum, vanadium, molybdenum carbide, tungsten carbide, titanium carbide, niobium carbide and palladium.

7. The electronic device of claim 5, wherein a region of the porous silicon layer is removed beneath the catalytic material so that the layer of catalytic material contacts the silicon carbide wafer.

8. An electronic device of semiconductor design for detecting hydrocarbon gases comprising:

   a detection layer having a first and second surfaces, comprising:
   a region of porous silicon carbide formed at a first surface of said silicon carbide wafer by photoelectrochemical etching, the region of porous silicon carbide for adsorbing hydrocarbon gases;
   a metal grid deposited on the porous silicon carbide region; and
   a first electrode in electrical communication with the metal grid for applying a varying dissociating potential to said porous silicon carbide region and a magnitude of the dissociating potential indicating a type of hydrocarbon adsorbed to said porous silicon region.

9. The electronic device of claim 8, wherein the dissociating potential varies between 0 V and about 10 V.

10. The electronic device of claim 8, wherein the silicon carbide wafer forming the detection layer is hexagonal form silicon carbide with the C-axis normal to the first surface.

11. The electronic device of claim 8, wherein the detection layer further comprises a layer of catalytic material deposited over a portion of the porous silicon carbide region.

12. The electronic device of claim 11, wherein the catalytic material is selected from the group consisting of platinum, vanadium, molybdenum carbide, tungsten carbide, titanium carbide, niobium carbide and palladium.

13. The electronic device of claim 11, wherein an area of the porous silicon layer is removed beneath the layer of catalytic material so that the layer of catalytic material contacts the silicon carbide wafer.

14. An electronic device of semiconductor design for detecting hydrocarbon gases comprising:

   a wafer of silicon carbide;
   a channel layer of doped silicon carbide formed on an exposed surface of the wafer with metallic drain and source electrodes disposed on the channel layer at opposite sides thereof; and
   an insulation layer formed on an exposed surface of the channel layer;

    a detection layer having a first and second surface, and forming a gate of a field effect transistor, the detection layer comprising:
    a layer of polycrystalline silicon carbide formed on an exposed surface of said insulation layer for adsorbing hydrocarbon gases;
    a metal grid deposited on the polycrystalline silicon carbide layer; and
    an electrode in electrical communication with the metal grid for applying a varying dissociating potential to said polycrystalline silicon carbide layer for dissociating hydrocarbon gas molecules adsorbed to the polycrystalline silicon carbide layer so that dissociation of hydrocarbon gas molecules causes an alteration in a current flow between the drain electrode and the source electrode, a magnitude of said alteration indicating a quantity of hydrocarbon gas adsorbed to the detection layer and a magnitude of the dissociating potential indicating a type of hydrocarbon gas adsorbed to the detection layer.

15. The electronic device of claim 14, wherein the dissociating potential varies between 0 V and about 10 V.

16. The electronic device of claim 14, wherein the silicon carbide wafer is hexagonal form silicon carbide.

17. The electronic device of claim 14, wherein the silicon carbide wafer is cubic form silicon carbide.

18. The electronic device of claim 14, wherein the insulation layer comprises a layer of single crystal aluminum nitride.

19. The electronic device of claim 14, wherein the detection layer further comprises a layer of catalytic material deposited over a portion of the polycrystalline silicon carbide layer.

20. An electronic device of semiconductor design for detecting hydrocarbon gases comprising:

   a detection layer with a first surface and a second surface, comprising:
   a wafer of silicon carbide;
   a layer of polycrystalline silicon carbide formed on a first surface of said wafer for adsorbing hydrocarbon gases;
   a metal grid deposited on the polycrystalline silicon carbide layer; and
   an electrode in electrical communication with the metal grid for applying a varying dissociating potential to said polycrystalline silicon carbide layer, said dissociating potential for dissociating hydrocarbon gas molecules adsorbed to the polycrystalline silicon carbide layer so that dissociation of hydrocarbon gas molecules causes an alteration in a current flow between the drain electrode and the source electrode, a magnitude of said alteration indicating a quantity of hydrocarbon gas adsorbed to the detection layer and a magnitude of the dissociating potential indicating a type of hydrocarbon gas adsorbed to the detection layer.
indicating a quantity of hydrocarbon adsorbed to said polycrystalline silicon carbide layer and a magnitude of the dissociating potential indicating a type of hydrocarbon adsorbed to said polycrystalline silicon carbide layer.

21. The electronic device of claim 20, wherein the dissociating potential varies between 0 V and about 10 V.

22. The electronic device of claim 20, wherein the detection layer further comprises a layer of catalytic material deposited over an area of the poly-crystalline silicon carbide layer.

23. A method of employing a semiconductor device to detect hydrocarbon gases comprising the steps of:
   providing a semiconductor device with a detection layer of silicon carbide to interact with a gas sample;
   operating said semiconductor device at temperatures between about 100° C. and 650° C.;
   introducing a gas sample to be tested by exposure to a surface of the detection layer so that the gas sample adsorbs to the detection layer;
   varying a dissociating potential applied to the detection layer;
   measuring a change in an electrical characteristic of the semiconductor device as indicative of a quantity of hydrocarbon gas adsorbed to the detection layer; and
   measuring the potential at which the change in an electrical characteristic occurs as indicative of a type of hydrocarbon gas present in the detection layer.

24. The method of claim 23, wherein the detection layer further comprises a pad of catalytic material covering a portion of the silicon carbide of the detection layer.

25. The method of claim 24, wherein the catalytic material is selected from the group consisting of platinum, vanadium, molybdenum carbide, tungsten carbide, titanium carbide, niobium carbide and palladium.

26. The method of claim 23, wherein the dissociating potential is varied between about 0 V and 10 V.

27. The method of claim 23 further comprising a step of reversing polarity of the dissociating potential to regenerate the semiconductor device.

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