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(54) Title: EPITAXY OF HIGH TENSILE SILICON ALLOY FOR TENSILE STRAIN APPLICATIONS

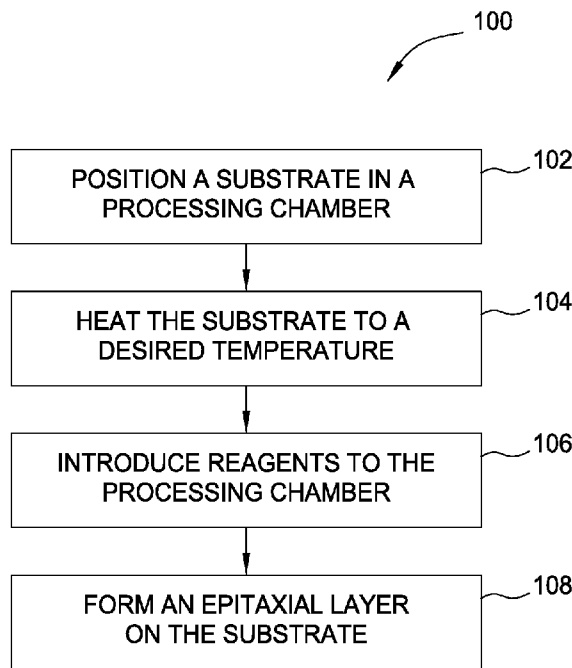


FIG. 1

(57) Abstract: Embodiments of the present invention generally relate to methods for forming silicon epitaxial layers on semiconductor devices. The methods include forming a silicon epitaxial layer on a substrate at increased pressure and reduced temperature. The silicon epitaxial layer has a phosphorus concentration of about 1×10^{21} atoms per cubic centimeter or greater, and is formed without the addition of carbon. A phosphorus concentration of about 1×10^{21} atoms per cubic centimeter or greater increases the tensile strain of the deposited layer, and thus, improves channel mobility. Since the epitaxial layer is substantially free of carbon, the epitaxial layer does not suffer from film formation and quality issues commonly associated with carbon-containing epitaxial layers.

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- *with amended claims (Art. 19(1))*

EPITAXY OF HIGH TENSILE SILICON ALLOY FOR TENSILE STRAIN APPLICATIONS

BACKGROUND OF THE INVENTION

Field of the Invention

[0001] Embodiments of the invention generally relate to the field of semiconductor manufacturing processes and devices, more particularly, to methods of depositing silicon-containing films for forming semiconductor devices.

Description of the Related Art

[0002] Size reduction of metal-oxide-semiconductor field-effect transistors (MOSFET) has enabled the continued improvement in speed performance, density, and cost per unit function of integrated circuits. One way to improve transistor performance is through application of stress to the transistor channel region. Stress distorts (*e.g.*, strains) the semiconductor crystal lattice, and the distortion, in turn, affects the band alignment and charge transport properties of the semiconductor. By controlling the magnitude of stress in a finished device, manufacturers can increase carrier mobility and improve device performance. There are several existing approaches of introducing stress into the transistor channel region.

[0003] One such approach of introducing stress into the transistor channel region is to incorporate carbon into the region during the formation of the region. The carbon present in the region affects the semiconductor crystal lattice and thereby induces stress. However, the quality of epitaxially-deposited films decreases as carbon concentration within the film increases. Thus, there is a limit to the amount of tensile stress which can be induced before film quality becomes unacceptable.

[0004] Generally, carbon concentrations above about 1 atomic percent seriously reduce film quality and increase the probability of film growth issues. For example, film growth issues such as undesired polycrystalline or amorphous silicon growth,

instead of epitaxial growth, may occur due to the presence of carbon concentrations greater than 1 atomic percent. Therefore, the benefits that can be gained by increasing the tensile stress of a film through carbon incorporation are limited to films having carbon concentrations of 1 atomic percent or less. Moreover, even films which contain less than 1 atomic percent carbon still experience some film quality issues.

[0005] Therefore, there is a need for producing a high tensile stress epitaxial film which is substantially free of carbon.

SUMMARY OF THE INVENTION

[0006] Embodiments of the present invention generally relate to methods for forming silicon epitaxial layers on semiconductor devices. The methods include forming a silicon epitaxial layer on a substrate at increased pressure and reduced temperature. The silicon epitaxial layer has a phosphorus concentration of about 1×10^{21} atoms per cubic centimeter or greater, and is formed without the addition of carbon. A phosphorus concentration of about 1×10^{21} atoms per cubic centimeter or greater increases the tensile strain of the deposited layer, and thus, improves channel mobility. Since the epitaxial layer is substantially free of carbon, the epitaxial layer does not suffer from film formation and quality issues commonly associated with carbon-containing epitaxial layers.

[0007] In one embodiment, a method of forming a film on a substrate comprises positioning a substrate within a processing chamber, and heating the substrate to a temperature within a range from about 550 degrees Celsius to about 700 degrees Celsius. One or more process gases are then introduced into the processing chamber. The one or more process gases comprise a silicon source and a phosphorus source. A substantially carbon-free silicon epitaxial layer is then deposited on the substrate. The substantially carbon-free silicon epitaxial layer has a phosphorus concentration of about 1×10^{21} atoms per cubic centimeter or greater. The substantially carbon-free silicon epitaxial layer is deposited at a chamber pressure of about 300 Torr or greater.

[0008] In another embodiment, a method of forming a film on a substrate comprises positioning a substrate within a processing chamber and heating the substrate to a temperature within a range from about 600 degrees Celsius to about 650 degrees Celsius. One or more process gases are then introduced into the processing chamber. The one or more process gases comprise a silicon source and a phosphorus source. A substantially carbon-free silicon epitaxial layer is then deposited on the substrate. The substantially carbon-free silicon epitaxial layer has a phosphorus concentration of about 1×10^{21} atoms per cubic centimeter or greater and is deposited at a chamber pressure of about 300 Torr or greater.

[0009] In another embodiment, a method of forming a film on a substrate comprises positioning a substrate within a processing chamber and heating the substrate to a temperature within a range from about 550 degrees Celsius to about 750 degrees Celsius. Phosphine and at least one silane or disilane are then introduced into the processing chamber and a substantially carbon-free silicon epitaxial layer is deposited on the substrate. The substantially carbon-free silicon epitaxial layer has a phosphorus concentration of about 1×10^{21} atoms per cubic centimeter or greater and is deposited at a chamber pressure of about 150 Torr or greater.

BRIEF DESCRIPTION OF THE DRAWINGS

[0010] So that the manner in which the above recited features of the present invention can be understood in detail, a more particular description of the invention, briefly summarized above, may be had by reference to embodiments, some of which are illustrated in the appended drawings. It is to be noted, however, that the appended drawings illustrate only typical embodiments of this invention and are therefore not to be considered limiting of its scope, for the invention may admit to other equally effective embodiments.

[0011] Figure 1 is a flow chart illustrating a method of forming a phosphorus-containing silicon epitaxial layer.

[0012] Figure 2 is a graph illustrating the dopant profile of a film formed according to embodiments of the invention.

[0013] Figure 3 is a graph illustrating the tensile stress of the film of Figure 2.

[0014] To facilitate understanding, identical reference numerals have been used, where possible, to designate identical elements that are common to the figures. It is contemplated that elements disclosed in one embodiment may be beneficially utilized on other embodiments without specific recitation.

DETAILED DESCRIPTION

[0015] Embodiments of the present invention generally relate to methods for forming silicon epitaxial layers on semiconductor devices. The methods include forming a silicon epitaxial layer on a substrate at increased pressure and reduced temperature. The silicon epitaxial layer has a phosphorus concentration of about 1×10^{21} atoms per cubic centimeter or greater, and is formed without the addition of carbon. A phosphorus concentration of about 1×10^{21} atoms per cubic centimeter or greater increases the tensile strain of the deposited layer, and thus, improves channel mobility. Since the epitaxial layer is substantially free of carbon, the epitaxial layer does not suffer from film formation and quality issues commonly associated with carbon-containing epitaxial layers. Substantially free of carbon as used herein refers to a film which is formed without the use of a carbon-containing precursor; however, it is contemplated that trace amounts of carbon may be present in the film due to contamination.

[0016] Embodiments of the present invention may be practiced in the CENTURA® RP Epi chamber available from Applied Materials, Inc., of Santa Clara, California. It is contemplated that other chambers, including those available from other manufacturers, may be used to practice embodiments of the invention.

[0017] Figure 1 is a flow chart 100 illustrating a method of forming a phosphorus-containing silicon epitaxial layer. In step 102, a monocrystalline silicon substrate is positioned within a processing chamber. In step 104, the substrate is heated to a predetermined temperature. The substrate is generally heated to a temperature within a range from about 550 degrees Celsius to about 700 degrees Celsius. It is desirable to minimize the thermal budget of the final device by heating

the substrate to the lowest temperature sufficient to thermally decompose process reagents and deposit an epitaxial film on the substrate. However, as increased temperatures generally lead to increased throughput, it is contemplated that higher temperatures may be used as dictated by production requirements.

[0018] In step 106, process gases containing one or more processing reagents are introduced into the processing chamber. The process gases include a silicon source and phosphorus source for depositing a phosphorus-containing silicon epitaxial layer on the substrate. Optionally, the one or more process gases may include a carrier gas for delivering the silicon source and the phosphorus source to the processing chamber, as well as an etchant when performing selective deposition processes.

[0019] An exemplary phosphorus source includes phosphine, which may be delivered to the processing chamber at a rate of about 2 sccm to about 30 sccm or greater. For example, the flow rate of phosphine may be about 12 sccm to about 15 sccm. Suitable carrier gases include nitrogen, hydrogen, or other gases which are inert with respect to the deposition process. The carrier gas may be provided to the processing chamber at a flow rate within a range from about 3 SLM to about 30 SLM. Suitable silicon sources include dichlorosilane, silane, and disilane. The silicon source may be delivered to the processing chamber at a flow rate between about 300 sccm and 400 sccm. While other silicon and phosphorus sources are contemplated, it is generally desirable that carbon addition to the processing atmosphere is minimized, thus, carbon-containing precursors should be avoided.

[0020] In step 108, the mixture of reagents is thermally driven to react and deposit a phosphorus-containing silicon epitaxial layer on the substrate surface. During the deposition process, the pressure within the processing chamber is maintained at about 150 Torr or greater, for example, about 300 Torr to about 600 Torr. It is contemplated that pressures greater than about 600 Torr may be utilized when low pressure deposition chambers are not employed. In contrast, typical epitaxial growth processes in low pressure deposition chambers maintain a processing pressure of about 10 Torr to about 100 Torr and a processing

temperature greater than 700 degrees Celsius. However, by increasing the pressure to about 150 Torr or greater, the deposited epitaxial film is formed having a greater phosphorus concentration (e.g., about 1×10^{21} atoms per cubic centimeter to about 5×10^{21} atoms per cubic centimeter) compared to lower pressure epitaxial growth processes. Furthermore, high flow rates of phosphorus source gas provided during low pressure depositions often result in "surface poisoning" of the substrate, which suppresses epitaxial formation. Surface poisoning is typically not experienced when processing at pressures above 300 Torr, due to the silicon source flux overcoming the poisoning effect. Thus, increased processing pressures are desirable for epitaxial processes utilizing high dopant flow rates.

[0021] The phosphorus concentration of an epitaxial film formed at a pressure less than 100 Torr is approximately 3×10^{20} atoms per cubic centimeter when providing a phosphine flow rate of about 3 sccm to about 5 sccm. Thus, epitaxial layers formed at higher pressures (e.g., 300 Torr or greater) experience approximately a tenfold increase in phosphorus concentration compared to epitaxial films formed at pressures below about 100 Torr or less. It is believed that at a phosphorus concentration of about 1×10^{21} atoms per cubic centimeter or greater, the deposited epitaxial film is not purely a silicon film doped with phosphorus, but rather, that the film is an alloy between silicon and silicon phosphide (e.g., pseudocubic Si_3P_4). It is believed that the silicon/silicon phosphide alloy attributes to the increased tensile stress of the epitaxial film. The likelihood of forming the silicon/silicon phosphide alloy increases with greater phosphorus concentrations, since the probability of adjacent phosphorus atoms interacting is increased.

[0022] Epitaxial films which are formed at process temperatures between about 550 degrees Celsius and about 750 degrees Celsius and at pressures above 300 Torr experience increased tensile stress when doped to a sufficient phosphorus concentration (e.g., about 1×10^{21} atoms per cubic centimeter or greater). Carbon-free epitaxial films formed under such conditions experience approximately 1 gigapascal to about 1.5 gigapascals of tensile stress, which is equivalent to a low pressure silicon epitaxial film containing about 1.5 percent carbon. However, as

noted above, epitaxial films containing greater than about 1 percent carbon suffer from decreased film quality, and are thus undesirable. Furthermore, carbon-doped silicon epitaxy processes typically utilize cyclical deposition-etch processes which increase process complexity and cost. Producing an epitaxial film according to embodiments herein not only results in a film having a tensile stress equal to or greater than a 1.5 percent carbon-containing epitaxial film, but the resistivity of the carbon-free film is also lower (e.g., about 0.6 milliohm-centimeters compared to about 0.9 milliohm-centimeters). Thus, the substantially carbon-free epitaxial film exhibits higher film quality, lower resistivity, and equivalent tensile stress when compared to carbon-containing epitaxial films.

[0023] The tensile strain of the epitaxially-grown film can further be increased by reducing the deposition temperature during the epitaxial growth process. In a first example, a phosphorus-doped silicon epitaxial film is deposited at a chamber pressure of 700 Torr and a temperature of about 750 degrees Celsius. Process gases containing 300 sccm of dichlorosilane and 5 sccm of phosphine were provided to a process chamber during the growth process. The deposited film contained a phosphorus concentration of about 3×10^{20} atoms per cubic centimeter, and exhibited a tensile strain equal to a silicon epitaxial film having a carbon concentration of about 0.5 atomic percent. In a second example, a phosphorus-doped silicon epitaxial film was deposited on another substrate under similar process conditions; however, the process temperature was reduced to about 650 degrees Celsius, and the flow rate of phosphine was increased to 20 sccm. The phosphorus-doped silicon epitaxial film had a tensile strain equivalent to a film containing 1.8 atomic percent carbon. Thus, as process temperature is reduced and dopant concentration is increased, the tensile strain within the deposited epitaxial film increases. It is to be noted, however, that the tensile strain benefits due to decreased temperature may be limited, since there is minimum temperature which is required to react and deposit the process reagents.

[0024] In a third example, a phosphorus-doped silicon epitaxial film was formed under similar process conditions as the first example; however, the flow rate of phosphine during processing was reduced to about 2 sccm. The resultant

phosphorus-doped silicon epitaxial film had a tensile strain equivalent to a film having about 0.2 percent carbon. Additionally, the resultant film had a resistivity of about 0.45 milliohm-centimeters compared to 0.60 milliohm-centimeters for the film of the first example. Thus, not only can the tensile strain of an epitaxial film be adjusted by varying temperature and or pressure during the deposition process, but the resistivity can also be adjusted by varying the amount of dopant provided to the processing chamber.

[0025] Figure 2 is a graph illustrating the dopant profile of a film formed according to embodiments of the invention. The analyzed film of Figure 2 was formed by heating a silicon substrate having a silicon-germanium layer thereon to a temperature of about 650 degrees Celsius. Approximately 300 sccm of dichlorosilane and 30 sccm of phosphine were delivered to a processing chamber maintained at a pressure of about 600 Torr. A 450 angstrom silicon epitaxial film was formed on the silicon-germanium layer. As determined by secondary ion mass spectroscopy, the phosphorus-doped epitaxial film had a uniform phosphorus concentration of about 3×10^{21} atoms per cubic centimeter, and was substantially free of carbon. In contrast to the film analyzed in Figure 2, epitaxial films formed at lower pressures, such as less than 300 Torr, have a phosphorus concentration of about 3×10^{20} atoms per cubic centimeter. Thus, the epitaxial film formed according to embodiments described herein exhibited a tenfold increase in phosphorus concentration as compared to epitaxial films formed at lower pressures.

[0026] Figure 3 is a graph illustrating the tensile stress of the film of Figure 2 as determined by high resolution X-ray diffraction. The peak A corresponds to the tensile stress of the monocrystalline silicon substrate, while the peak B corresponds to the tensile stress of the silicon-germanium layer. The peak C corresponds to the tensile stress of the phosphorus-containing epitaxial layer. The well defined edges of the peak B and the peak C are indicative of high quality epitaxial films having uniform composition. The peak B corresponds to a silicon-germanium epitaxial layer containing about 12.3 percent germanium. The peak B has a shift between about -1000 arc seconds and about -1500 arc seconds (e.g., compressed stress), and an intensity of about 1000 a.u. The peak C has a peak shift of about 1700 arc

seconds to about 2400 arc seconds (e.g., tensile stress), and an intensity of about 800 a.u. The stress corresponding to peak C is similar to that of an epitaxial film having a carbon concentration of about 1.8 atomic percent. As discussed above, epitaxial films containing greater than about 1 atomic percent carbon have unacceptable film quality. Thus, while the tensile strength of highly phosphorus-doped epitaxial films is about equal to an epitaxial film containing 1.8 atomic percent carbon, the highly phosphorus-doped epitaxial films exhibit a higher film quality than the carbon-doped epitaxial films of comparable tensile strain.

[0027] Benefits of the invention include high quality silicon epitaxial films exhibiting high tensile strain. Increased process pressures combined with reduced process temperatures allow for formation of a silicon epitaxial film having a phosphorus concentration of 3×10^{21} atoms per cubic centimeter or greater, without experiencing surface poisoning. The high phosphorus concentration induces stress within the deposited epitaxial film, thereby increasing tensile strain, leading to increased carrier mobility and improved device performance. The tensile strain obtained by highly phosphorus-doped epitaxial silicon is comparable to epitaxial films containing up to 1.8 atomic percent carbon. However, highly phosphorus-doped epitaxial silicon of the present invention avoids the quality issues associated with carbon-doped films.

[0028] While the foregoing is directed to embodiments of the present invention, other and further embodiments of the invention may be devised without departing from the basic scope thereof, and the scope thereof is determined by the claims that follow.

We claim:

1. A method of forming a film on a substrate, comprising:
positioning a substrate within a processing chamber;
heating the substrate to a temperature within a range from about 550 degrees Celsius to about 750 degrees Celsius;
introducing one or more process gases into the processing chamber, the one or more process gases comprising a silicon source and a phosphorus source; and
depositing a substantially carbon-free silicon epitaxial layer on the substrate, the substantially carbon-free silicon epitaxial layer having a phosphorus concentration of about 1×10^{21} atoms per cubic centimeter or greater, wherein the substantially carbon-free silicon epitaxial layer is deposited at a chamber pressure of about 150 Torr or greater.
2. The method of claim 1, wherein the chamber pressure is about 300 Torr or greater.
3. The method of claim 1, wherein the silicon precursor is dichlorosilane.
4. The method of claim 3, wherein the phosphorus precursor is phosphine.
5. The method of claim 1, wherein the temperature is within a range from about 600 degrees Celsius to about 650 degrees Celsius.
6. The method of claim 5, wherein the silicon precursor is silane or disilane.
7. The method of claim 1, wherein the silicon epitaxial layer has a tensile strain of about 1 gigapascal to about 1.5 gigapascals.

8. A method of forming a film on a substrate, comprising:
 - positioning a substrate within a processing chamber;
 - heating the substrate to a temperature within a range from about 600 degrees Celsius to about 650 degrees Celsius;
 - introducing one or more process gases into the processing chamber, the one or more process gases comprising a silicon source and a phosphorus source; and
 - depositing a substantially carbon-free silicon epitaxial layer on the substrate, the substantially carbon-free silicon epitaxial layer having a phosphorus concentration of about 1×10^{21} atoms per cubic centimeter or greater, wherein the substantially carbon-free silicon epitaxial layer is deposited at a chamber pressure of about 300 Torr or greater.
9. The method of claim 8, wherein the silicon epitaxial layer has a tensile strain of about 1 gigapascal to about 1.5 gigapascals.
10. The method of claim 9, wherein the silicon precursor is silane or disilane.
11. The method of claim 8, wherein the phosphorus precursor is phosphine.
12. The method of claim 1, wherein the silicon precursor is dichlorosilane.
13. A method of forming a film on a substrate, comprising:
 - positioning a substrate within a processing chamber;
 - heating the substrate to a temperature within a range from about 550 degrees Celsius to about 750 degrees Celsius;
 - introducing phosphine and at least one silane or disilane into the processing chamber; and
 - depositing a substantially carbon-free silicon epitaxial layer on the substrate, the substantially carbon-free silicon epitaxial layer having a phosphorus concentration of about 1×10^{21} atoms per cubic centimeter or greater, wherein the substantially carbon-free silicon epitaxial layer is deposited at a chamber pressure of about 150 Torr or greater.

14. The method of claim 13, wherein the chamber pressure is about 300 Torr or greater.

15. The method of claim 4 wherein the temperature is within a range from about 600 degrees Celsius to about 650 degrees Celsius, and wherein the silicon epitaxial layer has a tensile strain of about 1 gigapascal to about 1.5 gigapascals.

AMENDED CLAIMS

received by the International Bureau on 06 June 2012 (06.06.2012)

1. A method of forming a film on a substrate, comprising:
positioning a substrate within a processing chamber;
heating the substrate to a temperature within a range from about 550 degrees Celsius to about 750 degrees Celsius;
introducing one or more process gases into the processing chamber, the one or more process gases comprising a silicon source and a phosphorus source; and
depositing a substantially carbon-free epitaxial layer comprising Si_3P_4 on the substrate, the substantially carbon-free epitaxial layer having a phosphorus concentration of 1×10^{21} atoms per cubic centimeter or greater, wherein the substantially carbon-free epitaxial layer is deposited at a chamber pressure of about 150 Torr or greater.
2. The method of claim 1, wherein the chamber pressure is about 300 Torr or greater.
3. The method of claim 1, wherein the silicon precursor is dichlorosilane.
4. The method of claim 3, wherein the phosphorus precursor is phosphine.
5. The method of claim 1, wherein the temperature is within a range from about 600 degrees Celsius to about 650 degrees Celsius.
6. The method of claim 5, wherein the silicon precursor is silane or disilane.
7. The method of claim 1, wherein the substantially carbon-free epitaxial layer has a tensile strain of about 1 gigapascal to about 1.5 gigapascals.
8. A method of forming a film on a substrate, comprising:
positioning a substrate within a processing chamber;

heating the substrate to a temperature within a range from about 600 degrees Celsius to about 650 degrees Celsius;

introducing one or more process gases into the processing chamber, the one or more process gases comprising a silicon source and a phosphorus source; and

depositing a substantially carbon-free epitaxial layer comprising Si_3P_4 on the substrate, the substantially carbon-free epitaxial layer having a phosphorus concentration of 1×10^{21} atoms per cubic centimeter or greater, wherein the substantially carbon-free epitaxial layer is deposited at a chamber pressure of about 300 Torr or greater.

9. The method of claim 8, wherein the substantially carbon-free epitaxial layer has a tensile strain of about 1 gigapascal to about 1.5 gigapascals.

10. The method of claim 9, wherein the silicon precursor is silane or disilane.

11. The method of claim 8, wherein the phosphorus precursor is phosphine.

12. The method of claim 1, wherein the silicon precursor is dichlorosilane.

13. A method of forming a film on a substrate, comprising:

positioning a substrate within a processing chamber;

heating the substrate to a temperature within a range from about 550 degrees Celsius to about 750 degrees Celsius;

introducing phosphine and at least one silane or disilane into the processing chamber; and

depositing a substantially carbon-free epitaxial layer comprising Si_3P_4 on the substrate, the substantially carbon-free epitaxial layer having a phosphorus concentration of 1×10^{21} atoms per cubic centimeter or greater, wherein the substantially carbon-free epitaxial layer is deposited at a chamber pressure of about 150 Torr or greater.

14. The method of claim 13, wherein the chamber pressure is about 300 Torr or greater.

15. The method of claim 4 wherein the temperature is within a range from about 600 degrees Celsius to about 650 degrees Celsius, and wherein the silicon epitaxial layer has a tensile strain of about 1 gigapascal to about 1.5 gigapascals.

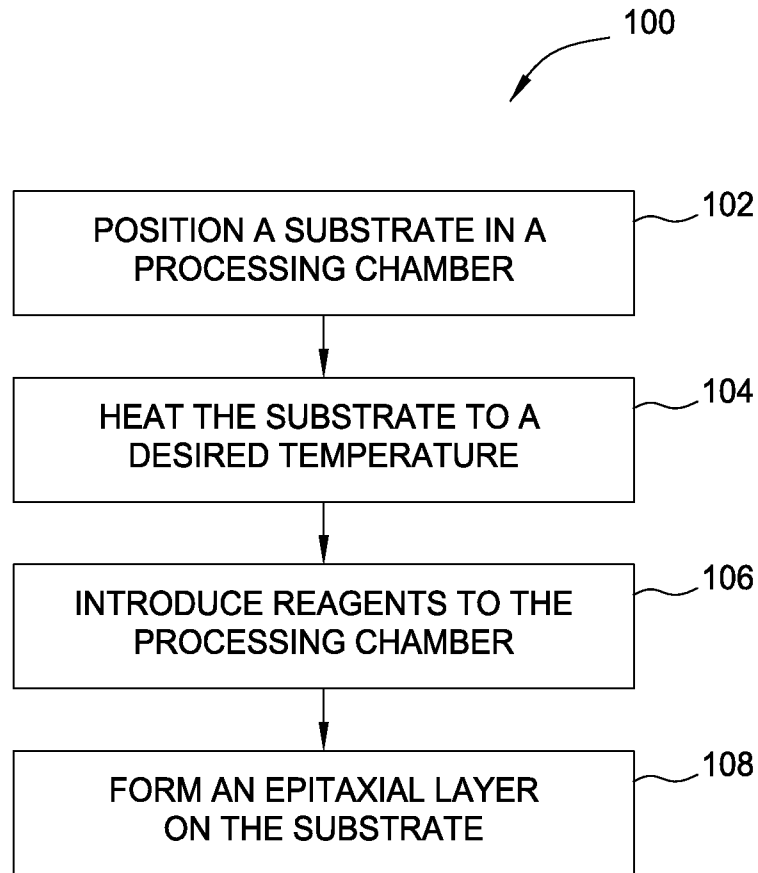


FIG. 1

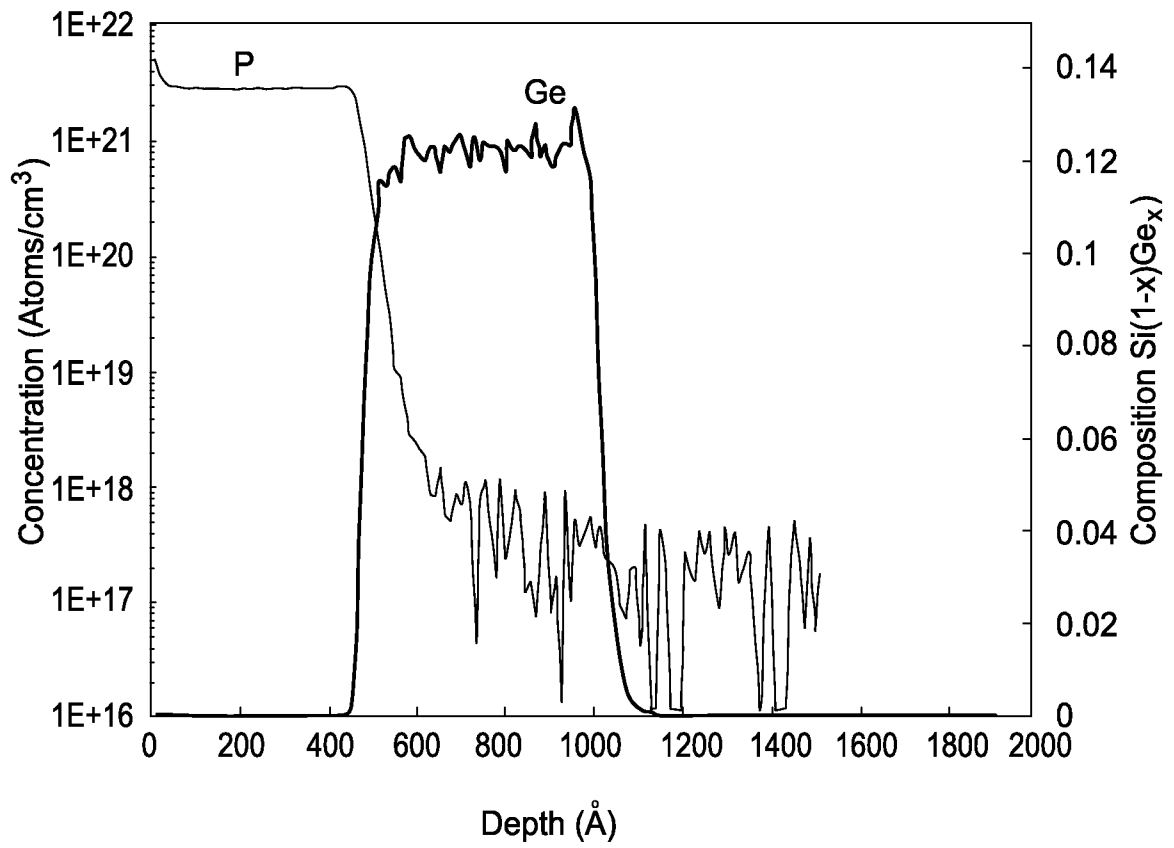


FIG. 2

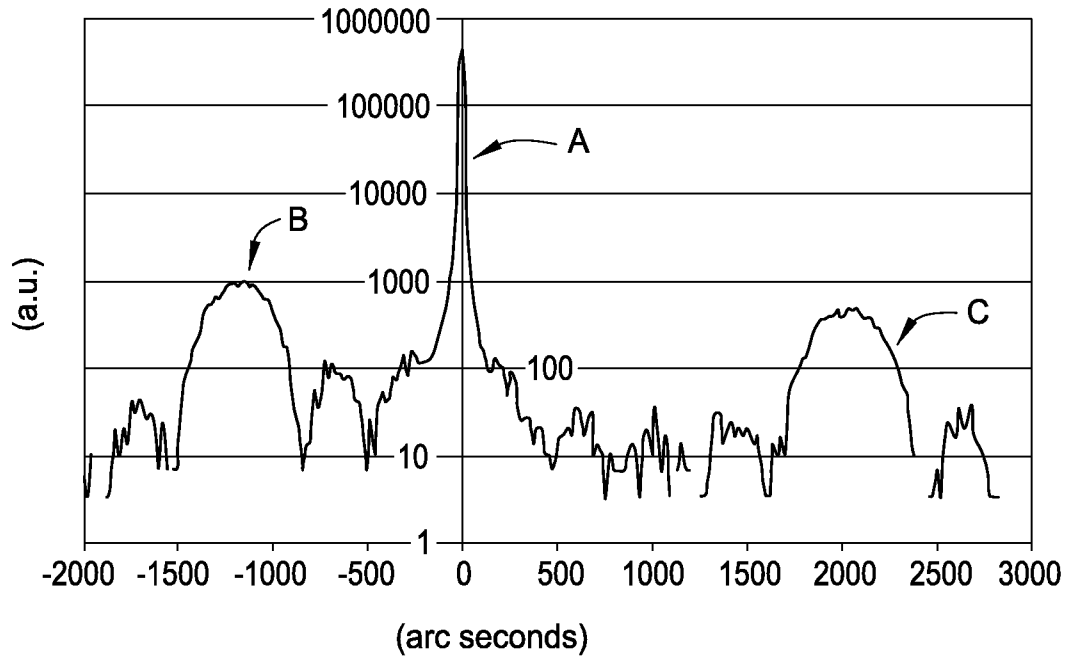


FIG. 3

A. CLASSIFICATION OF SUBJECT MATTER*H01L 21/20(2006.01)i, H01L 21/205(2006.01)i, H01L 21/336(2006.01)i, H01L 29/78(2006.01)i*

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

H01L 21/20; H01L 29/167; C30B 25/04; H01L 21/00

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Korean utility models and applications for utility models

Japanese utility models and applications for utility models

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

eKOMPASS(KIPO internal) & Keywords:epitaxial,carbon-free,silicon,phosphorus,strain

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 2005-0079691 A1 (YIHWAN KIM et al.) 14 April 2005 See abstract and paragraphs [0009]-[0013].	1-15
A	US 05731626 A (EAGLESHAM; DAVID JAMES et al.) 24 March 1998 See column 3, line 17 - column 4, line 67.	1-15
A	US 2008-0138939 A1 (KIM YIHWAN) 12 June 2008 See abstract and paragraphs [0021],[0024],[0031],[0036]-[0038].	1-15
A	US 06107197 A (SUZUKI; TATSUYA) 22 August 2000 See column 3, line 25-column 4, line 33; and column 6, line 47-49.	1-15

 Further documents are listed in the continuation of Box C. See patent family annex.

* Special categories of cited documents:

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Date of the actual completion of the international search

16 MARCH 2012 (16.03.2012)

Date of mailing of the international search report

16 MARCH 2012 (16.03.2012)

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Bang Kee In

Telephone No. 82-42-481-8684



INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

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Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 2005-0079691 A1	14.04.2005	CN 100468625 C0	11.03.2009
		CN 100511587 C	08.07.2009
		CN 101483136 A	15.07.2009
		CN 101593680 A	02.12.2009
		CN 101593680 B	23.02.2011
		CN 1875461 A	06.12.2006
		CN 1926664 A	07.03.2007
		CN 1926664 C0	07.03.2007
		EP 1680808 A1	19.07.2006
		EP 1745503 A2	24.01.2007
		JP 2007-514294 A	31.05.2007
		JP 2007-537601 A	20.12.2007
		KR 10-2006-0110291 A	24.10.2006
		KR 10-2007-0022046 A	23.02.2007
		US 2005-0079692 A1	14.04.2005
		US 2006-0234488 A1	19.10.2006
		US 2007-0082451 A1	12.04.2007
		US 2009-0011578 A1	08.01.2009
		US 2010-0317177 A1	16.12.2010
		US 7132338 B2	07.11.2006
		US 7166528 B2	23.01.2007
		US 7439142 B2	21.10.2008
		US 7517775 B2	14.04.2009
		US 7737007 B2	15.06.2010
		WO 2005-038890 A1	28.04.2005
		WO 2005-112577 A2	01.12.2005
		WO 2005-112577 A3	01.12.2005
US 05731626 A	24.03.1998	EP 0717435 A1	19.06.1996
		JP 04-117914 B2	16.07.2008
		JP 08-227863 A	03.09.1996
		JP 2008-211222 A	11.09.2008
		KR 10-1996-0026521 A	22.07.1996
		US 06043139 A	28.03.2000
		US 06153920 A	28.11.2000
US 2008-0138939 A1	12.06.2008	JP 2010-512668 A	22.04.2010
		TW 200832529 A	01.08.2008
		US 2008-0182075 A1	31.07.2008
		US 7960236 B2	14.06.2011
		WO 2008-073894 A1	19.06.2008
		WO 2009-079485 A1	25.06.2009
US 06107197 A	22.08.2000	EP 0784337 A2	16.07.1997
		EP 0784337 A3	03.06.1998
		EP 1045432 A2	18.10.2000
		EP 1045432 A3	13.12.2000
		JP 09-190979 A	22.07.1997