

NITROGEN-CONTAINING COMPOUNDS FOR ETCHING SEMICONDUCTOR STRUCTURES

Cross Reference to Related Applications

[0001] The present application claims the benefit of U.S. Application Serial No. 14/841,271 filed August 31, 2015, herein incorporated by reference in its entirety for all purposes.

Technical Field

[0002] A method for etching silicon-containing films is disclosed. The method includes the steps of introducing a vapor of a nitrogen containing etching compound into a reaction chamber containing a silicon-containing film on a substrate, wherein the nitrogen containing etching compound is an organofluorine compound containing at least one $C\equiv N$ or $C=N$ functional group; introducing an inert gas into the reaction chamber; and activating a plasma to produce an activated nitrogen containing etching compound capable of etching the silicon-containing film from the substrate.

Background

[0003] In memory applications in the semiconductor industries, such as DRAM and 2D NAND, plasma etching removes silicon-containing films, such as SiO or SiN layers, from semiconductor substrates. For 3D NAND (e.g., see US 2011/0180941 to Samsung Electronics Co., Ltd.), etching of stacks of multiple SiO/SiN or SiO/poly-Silicon (p-Si) layers is critical. An etchant having high selectivity between the mask and layers being etched is essential. Furthermore, the etched structure should have a straight vertical profile without bowing and low LER (line etch roughness).

[0004] Traditional etch gases include octafluorocyclobutane (C_4F_8), hexafluoro-1,3-butadiene (C_4F_6), CF_4 , CH_2F_2 , CH_3F , and/or CHF_3 . It is well known that selectivity and polymer deposition rate increase as the ratio of C:F increases (i.e., $C_4F_6 > C_4F_8 > CF_4$). See, e.g., US6387287 to Hung *et al.*

[0005] However, traditional etch chemistries may not be able to provide an aspect ratio higher than 20:1, which is necessary in the newer applications (e.g., 3D NAND), at least due to insufficient etch resistant polymer deposition on sidewalls during the plasma etching processes. The $-C_xF_y-$, wherein x ranges from 0.01 to 1 and y ranges from 0.01 to 4, polymers on sidewalls may be susceptible to etching. As a

result, the etched patterns may not be vertical and the etch structures may show bowing, change in dimensions, pattern collapse and/or increased roughness.

[0006] Bowing may result from sidewall etching of the mask layer, which may often be an amorphous carbon (a-C) material. The a-C materials may be etched by oxygen radicals in the plasma which may cause increased opening of the mask and result in bow-like, or angled/curved, etch structures.

[0007] Ji *et al.* (US 5,814,563) disclose using a mixture of a fluorohydrocarbon, carbon-oxygen, and NH₃-generating gases to achieve high selectivity of dielectric (such as SiO and SiN) to p-Si layers. Shane (US 2003/0162395) discloses addition of nitrogen-comprising gas to fluorocarbon to deposit polymer on mask to improve selectivity while etching silicon dioxide layer. Nemani *et al.* (US 2014/0199851) disclose using a plasma process performed by flowing NF₃ and NH₃ to remove the modified portion of silicon nitride layer to pattern a silicon nitride dielectric film. Hamrah *et al.* (US 5,242,538) discloses using CF₄ and NH₃ etching gases and selectivity of silicon oxide to polysilicon selectivity of up to 100:1 was observed. Pu *et al.* (US 5,843,847) also discloses adding additional nitrogen gas to fluorinated etching gas to assist in feature dimensional control.

[0008] Nitrogen containing compounds have been used as etching gases. For example, Khandelwal, *et al.* ("Dry removal technology for advanced CMOS devices", Nanochip Tech. J., vol. 11, issue 2, 2013, p17-19) disclose an *in-situ* dry removal process using NH₄F as etchant. Garg *et al.* (US 2006/0062914) disclose an activated reactive gas to treat the surface of a substrate. Garg *et al.* describe at paragraph [0019] that the activated reactive gas may include a large variety of fluorine-containing gases, including C₃F₃N₃, fluoroamines such as CF₅N, fluoronitriles such as C₂F₃N, C₃F₆N, and CF₃NO. Felker *et al.* (US 6,508,948) disclose perfluorinated heteroaromatic amine etching compounds, including cyanuric fluoride compounds. One disclosed cyanuric fluoride compound is pentafluoropyridine C₅F₅N.

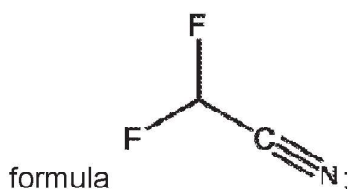
[0009] It is important to minimize bowing and to achieve high aspect ratio (*i.e.*, up to 200:1) needed for current applications (*e.g.*, contact etch or 3D NAND). Additionally, etching today has not been limited to selectivity to the photoresist mask. It is equally important to get high selectivity among other materials such as a-C, SiN, p-Si, SiC or other forms of Si_aC_bO_cH_dN_e materials (where a>0; b, c, d and e ≥0).

[0010] Thus, a need remains for new etch gas compositions for use in plasma etching applications, which may maintain selectivity and high aspect ratio for a wide range of process conditions.

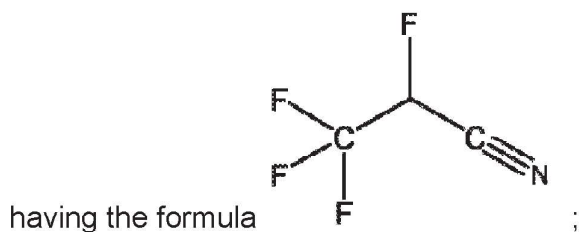
Summary

[0011] Disclosed are methods for plasma etching silicon-containing films. The methods include the steps of: introducing a vapor of a nitrogen containing etching compound into a reaction chamber containing a silicon-containing film on a substrate, wherein the nitrogen containing etching compound comprises an organofluorine compound containing a $C\equiv N$ or $C=N$ functional group; introducing an inert gas into the reaction chamber; and activating a plasma to produce an activated nitrogen containing etching compound capable of etching the silicon-containing film from the substrate. The disclosed methods may include one or more of the following aspects:

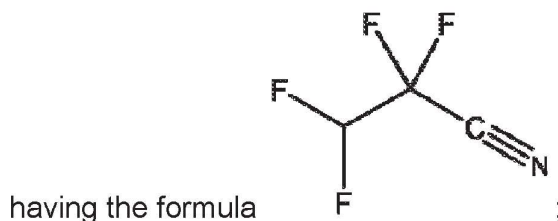
- the organofluorine compound containing a $C\equiv N$ functional group ;
- the organofluorine compound having the formula $N\equiv C-R^1$, wherein R^1 has the formula $H_aF_bC_c$ and $a=1-11$, $b=1-11$, and $c=1-5$;
- the organofluorine compound being difluoroacetonitrile (C_2HF_2N) having the



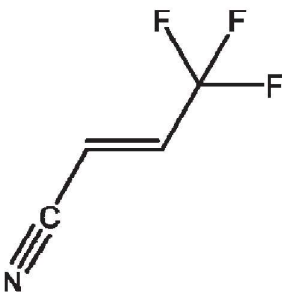
- the organofluorine compound being 2,3,3,3-tetrafluoropropionitrile (C_3HF_4N)



- the organofluorine compound being 2,2,3,3-tetrafluoropropionitrile (C_3HF_4N)

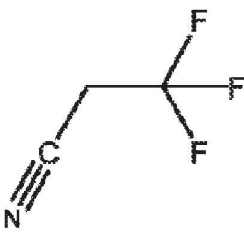


- the organofluorine compound being 4,4,4-Trifluorocrotono-nitrile ($C_4H_2F_3N$)



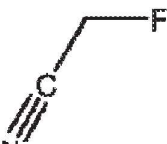
having the formula $N \equiv C - CH = CH - CF_3$;

- the organofluorine compound being 3,3,3-Trifluoropropionitrile ($C_3H_2F_3N$)



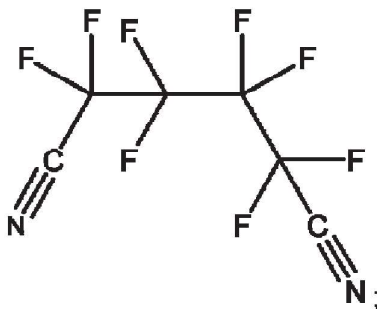
having the formula $N \equiv C - CH_2 - CF_3$;

- the organofluorine compound being fluoroacetonitrile (C_2H_2FN) having the



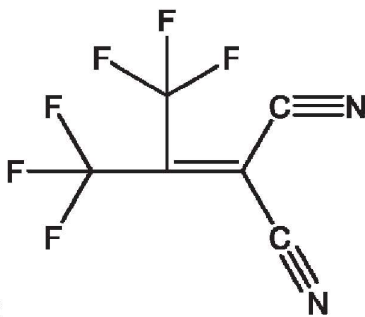
formula $N \equiv C - CH_2 - F$;

- the organofluorine compound having the formula $(N \equiv C-) - (R^2) - (-C \equiv N)$, wherein R^2 independently has the formula $H_aF_bC_c$ with $a=0$, $b=1-11$, and $c=1-5$;
- the organofluorine compound being octafluorohexane-1,6-dinitrile ($C_6F_8N_2$)



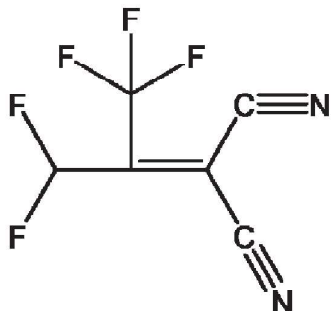
having the formula $N \equiv C - CF_2 - CF_2 - CF_2 - CF_2 - C \equiv N$;

- the organofluorine compound being 1,1-bis(trifluoromethyl)-2,2-dicyanoethylene

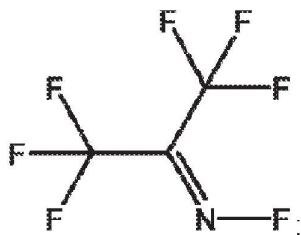


($C_6F_6N_2$) having the formula $(CF_3)_2C = C(C \equiv N)_2$;

- the organofluorine compound having the formula $(\text{N}\equiv\text{C})-(\text{R}^2)-(\text{C}\equiv\text{N})$, wherein R^2 has the formula $\text{H}_a\text{F}_b\text{C}_c$ with $a=1-11$, $b=1-11$, and $c=1-5$;
- the organofluorine compound being 2-[1-(difluoromethyl)-2,2,2-trifluoroethylidene]-propanedinitrile ($\text{C}_6\text{HF}_5\text{N}_2$) having the formula

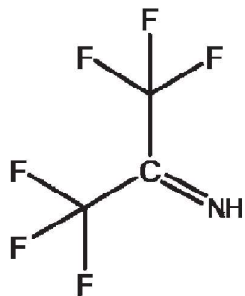


- the organofluorine compound containing a $\text{C}=\text{N}$ functional group;
- the organofluorine compound having the formula $\text{R}^1_x[\text{C}=\text{N}(\text{R}^2_z)]_y$, wherein $x=1-2$, $y=1-2$, $z=0-1$, $x+z=1-3$, and each R^1 and R^2 independently has the formula $\text{H}_a\text{F}_b\text{C}_c$ with $a=0$, $b=1-11$, and $c=0-5$;
- the organofluorine compound being N,1,1,1,3,3,3-heptafluoro-propanamine



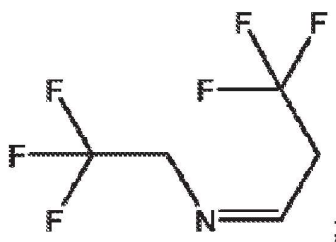
($\text{C}_3\text{F}_7\text{N}$) having the formula

- the organofluorine compound having the formula $\text{R}^1_x[\text{C}=\text{N}(\text{R}^3_z)]_y$, wherein $x=1-2$, $y=1-2$, $z=0-1$, $x+z=1-3$, and each R^1 and R^3 independently has the formula $\text{H}_a\text{F}_b\text{C}_c$ with $a=1-11$, $b=0-11$, and $c=0-5$;
- the organofluorine compound being hexafluoroacetone imine ($\text{C}_3\text{HF}_6\text{N}$) having



the formula

- the organofluorine compound being 1,1,1,6,6,6-Hexafluoro-3-azahex-3-ene



(C₅H₅F₆N) having the formula

- the activated nitrogen containing etching compound reacting with the silicon-containing film to form volatile by-products;
- the volatile by-products being removed from the reaction chamber;
- the inert gas being selected from the group consisting of He, Ar, Xe, Kr, and Ne;
- the inert gas being Ar;
- mixing the vapor of the nitrogen containing etching compound and the inert gas prior to introduction to the reaction chamber to produce a mixture;
- introducing the vapor of the nitrogen containing etching compound into the reaction chamber separately from the inert gas;
- introducing the inert gas continuously into the reaction chamber and introducing the vapor of the nitrogen containing etching compound into the reaction chamber in pulses;
- the inert gas comprising approximately 0.01% v/v to approximately 99.9% v/v of a total volume of the vapor of the nitrogen containing etching compound and inert gas introduced into the reaction chamber;
- the inert gas comprising approximately 90% v/v to approximately 99% v/v of a total volume of the vapor of the nitrogen containing etching compound and inert gas introduced into the reaction chamber;
- introducing an oxidizer into the reaction chamber;
- not introducing an oxidizer into the reaction chamber;
- the oxidizer being selected from the group consisting of O₂, CO, CO₂, NO, N₂O, and NO₂;
- the oxidizer being O₂;
- mixing the vapor of the nitrogen containing etching compound and the oxidizer prior to introduction into the reaction chamber;
- introducing the vapor of the nitrogen containing etching compound into the reaction chamber separately from the oxidizer;

- introducing the oxidizer continuously into the reaction chamber and introducing the vapor of the nitrogen containing etching compound into the reaction chamber in pulses;
- the oxidizer comprising approximately 0.01% v/v to approximately 99.9% v/v of a total volume of the vapor of the nitrogen containing etching compound and oxidizer introduced into the reaction chamber;
- the oxidizer comprising approximately 0.01% v/v to approximately 10% v/v of a total volume of the vapor of the nitrogen containing etching compound and oxidizer introduced into the reaction chamber;
- the silicon-containing film comprising a layer of silicon oxide, silicon nitride, polysilicon, crystalline silicon, SiON, SiOCH, $\text{Si}_a\text{O}_b\text{C}_c\text{N}_d\text{H}_e$ (where $a > 0$; b, c, d and e ≥ 0) or combinations thereof;
- the silicon-containing film comprising oxygen atoms, nitrogen atoms, carbon atoms, hydrogen atoms or combinations thereof;
- the silicon-containing film comprising silicon carbide;
- the silicon-containing film being selectively etched from an a-C layer;
- the silicon-containing film being selectively etched from a photoresist layer;
- the silicon-containing film being selectively etched from a silicon nitride layer;
- the silicon-containing film being selectively etched from a polysilicon layer;
- the silicon-containing film being selectively etched from a crystalline silicon layer;
- the silicon-containing film being selectively etched from a metal contact layer;
- the silicon-containing film being selectively etched from a Titanium Nitride layer;
- the silicon-containing film being selectively etched from a Tantalum layer;
- the silicon-containing film being a silicon oxide layer;
- selectively etching the silicon oxide layer from an a-C layer;
- selectively etching the silicon oxide layer from a photoresist layer;
- selectively etching the silicon oxide layer from a p-Si layer;
- selectively etching the silicon oxide layer from a crystalline silicon layer;
- selectively etching the silicon oxide layer from a metal contact layer;
- selectively etching the silicon oxide layer from a SiN layer;
- the silicon-containing film being a silicon nitride layer;
- selectively etching the silicon nitride layer from an a-C layer;
- selectively etching the silicon nitride layer from a patterned photoresist layer;

- selectively etching the silicon nitride layer from a p-Si layer;
- selectively etching the silicon nitride layer from a crystalline silicon layer;
- selectively etching the silicon nitride layer from a metal contact layer;
- selectively etching the silicon nitride layer from a silicon oxide layer;
- the silicon-containing film being a SiON layer;
- selectively etching the SiON layer from photoresist layer;
- the silicon-containing film being a SiCOH layer;
- selectively etching the SiCOH layer from titanium nitride layer;
- selectively etching the SiCOH layer from a-C layer;
- selective etching the SiCOH layer from photoresist layer;
- the silicon-containing film being alternating silicon oxide and silicon nitride layers;
- etching both silicon oxide and silicon nitride layers at similar etch rates;
- selectively etching both silicon oxide and silicon nitride layers from a silicon layer;
- selectively etching both silicon oxide and silicon nitride layers from a p-Si layer;
- selectively etching both silicon oxide and silicon nitride layers from a crystalline silicon layer;
- selectively etching both silicon oxide and silicon nitride layers from an a-C layer;
- the silicon-containing film being alternating silicon oxide and p-Si layers;
- etching both silicon oxide and p-Si layers at similar etch rates;
- selectively etching both silicon oxide and p-Si layers from an a-C layer;
- selectively etching both silicon oxide and p-Si layers from a silicon nitride layer;
- producing an aperture in the silicon-containing film having an aspect ratio between approximately 10:1 and approximately 200:1;
- producing a gate trench;
- producing a staircase contact;
- producing a channel hole;
- producing a channel hole having an aspect ratio between approximately 60:1 and approximately 100:1;
- producing a channel hole having a diameter ranging from approximately 5 nm to approximately 100 nm;

- producing a channel hole having a diameter ranging from approximately 10 nm to approximately 50 nm;
- improving selectivity by introducing an etch gas into the reaction chamber;
- the etch gas being selected from the group consisting of cC_4F_8 , C_4F_8 , C_4F_6 , CF_4 , CH_3F , CF_3H , CH_2F_2 , COS , CF_3I , $\text{C}_2\text{F}_3\text{I}$, $\text{C}_2\text{F}_5\text{I}$, $\text{F-C}\equiv\text{N}$, CS_2 , SO_2 , *trans*-1,1,1,4,4,4-hexafluoro-2-butene (*trans*- $\text{C}_4\text{H}_2\text{F}_6$), *cis*-1,1,1,4,4,4-hexafluoro-2-butene (*cis*- $\text{C}_4\text{H}_2\text{F}_6$), hexafluoroisobutene ($\text{C}_4\text{H}_2\text{F}_6$), *trans*-1,1,2,2,3,4-hexafluorocyclobutane (*trans*- $\text{C}_4\text{H}_2\text{F}_6$), 1,1,2,2,3-pentafluorocyclobutane ($\text{C}_4\text{H}_3\text{F}_5$), 1,1,2,2-tetrafluorocyclobutane ($\text{C}_4\text{H}_4\text{F}_4$), or *cis*-1,1,2,2,3,4-hexafluorocyclobutane (*cis*- $\text{C}_4\text{H}_2\text{F}_6$);
- the etch gas being cC_5F_8 ;
- the etch gas being cC_4F_8 ;
- the etch gas being C_4F_6 ;
- mixing the vapor of the nitrogen containing etching compound and the etch gas prior to introduction to the reaction chamber;
- introducing the vapor of the nitrogen containing etching compound into the reaction chamber separately from the etch gas;
- introducing approximately 0.01% v/v to approximately 99.99% v/v of the etch gas into the reaction chamber;
- activating the plasma by applying RF power;
- activating the plasma by a RF power ranging from approximately 25 W to approximately 10,000 W;
- the reaction chamber having a pressure ranging from approximately 1 mTorr to approximately 10 Torr;
- introducing the vapor of the nitrogen containing etching compound into the reaction chamber at a flow rate ranging from approximately 0.1 sccm to approximately 1 slm;
- maintaining the substrate at a temperature ranging from approximately -196°C to approximately 500°C ;
- maintaining the substrate at a temperature ranging from approximately -120°C to approximately 300°C ;
- maintaining the substrate at a temperature ranging from approximately -100°C to approximately 50°C ;

- maintaining the substrate at a temperature ranging from approximately -10°C to approximately 40°C; and
- measuring the activated nitrogen containing etching compound by Quadrupole mass spectrometer, optical emission spectrometer, FTIR, or other radical/ion measurement tool.

[0011a] In one aspect, there is provided a method of depositing an etch-resistant polymer layer on a substrate, the method comprising introducing a vapor of an organofluorine compound into a reaction chamber containing a substrate, the organofluorine compound having a formula selected from the group consisting of $N\equiv C-R^1$, wherein R^1 has the formula $H_aF_bC_c$ and $a=1-11$, $b=1-11$, and $c=1-5$; $(N\equiv C)-(R^2)-(-C\equiv N)$, wherein R^2 has the formula $H_aF_bC_c$ with $a=0-11$, $b=1-11$, and $c=1-5$; and $R^1_x[-C=N(R^2_z)]_y$, wherein $x=1-2$, $y=1-2$, $z=0-1$, $x+z=1-3$, and each R^1 and R^2 independently has the formula $H_aF_bC_c$ with $a=0-11$, $b=0-11$, and $c=0-5$; and plasma activating an organofluorine compound to form the etch-resistant polymer layer on the substrate, wherein the organofluorine compound having a purity ranging from 95% to 99.999% by volume and comprising between 10 parts per trillion to 5% by volume water vapor impurities.

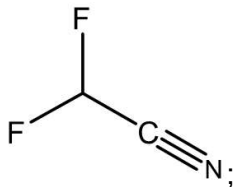
[0011b] In another aspect, there is provided a method of depositing an N-containing polymer passivation layer on a substrate, the method comprising introducing a vapor of an organofluorine compound into a reaction chamber containing a substrate, the organofluorine having a formula selected from the group consisting of $N\equiv C-R^1$, wherein R^1 has the formula $H_aF_bC_c$ and $a=1-11$, $b=1-11$, and $c=1-5$; $(N\equiv C)-(R^2)-(-C\equiv N)$, wherein R^2 has the formula $H_aF_bC_c$ with $a=0-11$, $b=1-11$, and $c=1-5$; and $R^1_x[-C=N(R^2_z)]_y$, wherein $x=1-2$, $y=1-2$, $z=0-1$, $x+z=1-3$, and each R^1 and R^2 independently has the formula $H_aF_bC_c$ with $a=0-11$, $b=0-11$, and $c=0-5$; and producing fragments of the organofluorine compound by activating a plasma to form the N-containing polymer passivation layer on the substrate, wherein the organofluorine compound having a purity ranging from 95% to 99.999% by volume and comprising between 10 parts per trillion to 5% by volume water vapor impurities.

[0012] Also disclosed are nitrogen containing etching compound comprising an organofluorine compound having a $C\equiv N$ or $C=N$ functional group. The disclosed nitrogen containing etching compounds include one or more of the following aspects:

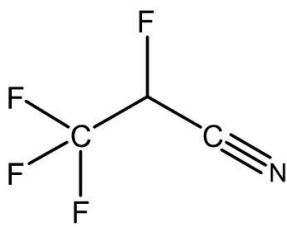
- the organofluorine compound containing a $C\equiv N$ functional group ;

- the organofluorine compound having the formula $\text{N}\equiv\text{C}-\text{R}^1$, wherein R^1 has the formula $\text{H}_a\text{F}_b\text{C}_c$ and $a=1-11$, $b=1-11$, and $c=1-5$;

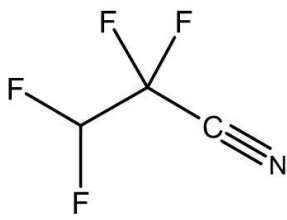
- the organofluorine compound being difluoroacetonitrile ($\text{C}_2\text{HF}_2\text{N}$) having the



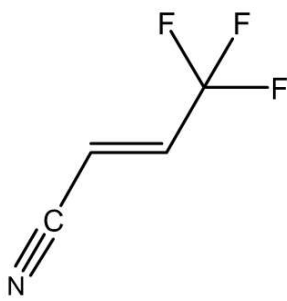
- the organofluorine compound being 2,3,3,3-tetrafluoropropionitrile ($\text{C}_3\text{HF}_4\text{N}$)



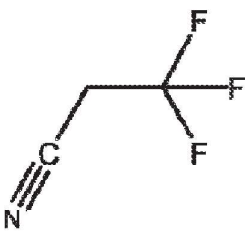
- the organofluorine compound being 2,2,3,3-tetrafluoropropionitrile ($\text{C}_3\text{HF}_4\text{N}$)



- the organofluorine compound being 4,4,4-Trifluorocrotono-nitrile ($\text{C}_4\text{H}_2\text{F}_3\text{N}$)

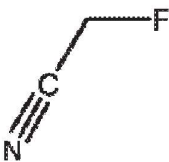


- the organofluorine compound being 3,3,3-Trifluoropropionitrile ($C_3H_2F_3N$)



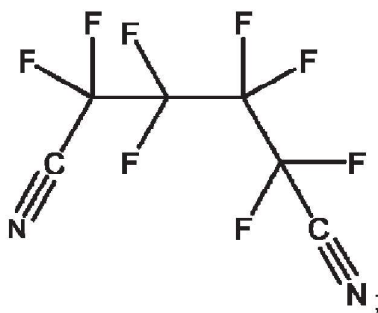
having the formula $N \equiv C - CH_2 - CF_3$;

- the organofluorine compound being fluoroacetonitrile (C_2H_2FN) having the



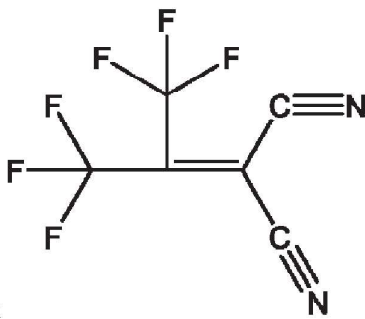
formula $N \equiv C - CH_2 - F$;

- the organofluorine compound having the formula $(N \equiv C -)(R^2) - (C \equiv N)$, wherein R^2 has the formula $H_aF_bC_c$ with $a=0$, $b=1-11$, and $c=1-5$;
- the organofluorine compound being octafluorohexane-1,6-dinitrile ($C_6F_8N_2$)



having the formula $N \equiv C - (CF_2)_4 - C \equiv N$;

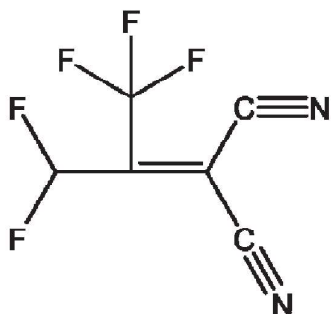
- the organofluorine compound being 1,1-bis(trifluoromethyl)-2,2-dicyanoethylene



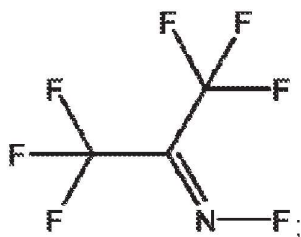
($C_6F_8N_2$) having the formula $(CF_3)_2C = C(C \equiv N)_2$;

- the organofluorine compound having the formula $(N \equiv C -)(R^2) - (C \equiv N)$, wherein R^2 has the formula $H_aF_bC_c$ with $a=1-11$, $b=1-11$, and $c=1-5$;

- the organofluorine compound being 2-[1-(difluoromethyl)-2,2,2-trifluoroethylidene]-propanedinitrile ($C_6HF_5N_2$) having the formula

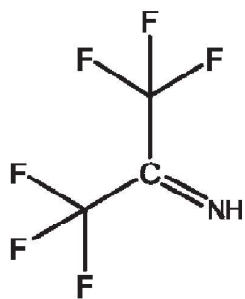


- the organofluorine compound containing a C=N functional group;
- the organofluorine compound having the formula $R^1_x[-C=N(R^2_z)]_y$, wherein $x=1-2$, $y=1-2$, $z=0-1$, $x+z=1-3$, and each R^1 and R^2 independently has the formula $H_aF_bC_c$ with $a=0$, $b=0-11$, and $c=0-5$;
- the organofluorine compound being N,1,1,1,3,3,3-heptafluoro-propanamine



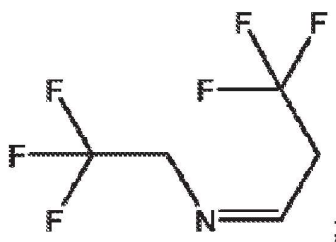
(C_3F_7N) having the formula

- the organofluorine compound having the formula $R^1_x[-C=N(R^2_z)]_y$, wherein $x=1-2$, $y=1-2$, $z=0-1$, $x+z=1-3$, and each R^1 and R^2 independently has the formula $H_aF_bC_c$ with $a=1-11$, $b=0-11$, and $c=0-5$;
- the organofluorine compound being hexafluoroacetone imine (C_3HF_6N) having



the formula

- the organofluorine compound being 1,1,1,6,6,6-Hexafluoro-3-azahex-3-ene



(C₅H₅F₆N) having the formula

- having a purity ranging from approximately 95% to approximately 99.999% by volume;
- comprising between approximately 10 parts per trillion to approximately 5 % by volume trace gas impurities;
- the trace gas impurities comprising water;
- the trace gas impurities comprising CO₂;
- the trace gas impurities comprising N₂; and
- the nitrogen containing etching compound having a water content of less than 20 ppmw.

Notation and Nomenclature

[0013] The following detailed description and claims utilize a number of abbreviations, symbols, and terms, which are generally well known in the art. While definitions are typically provided with the first instance of each acronym, for convenience, Table 1 provides a list of the abbreviations, symbols, and terms used along with their respective definitions.

Table 1

a or an	one or more than one
approximately or about	±10% of the value stated
LCD-TFT	liquid-crystal display-thin-film transistor
DRAM	dynamic random-access memory
FINFET	fin-shaped field-effect transistor
Bulk CMOS	Bulk complementary metal-oxide-semiconductor
FD-SOI	fully depleted silicon-on-insulator
FEOL	front end of the line
BEOL	back end of the line
TSV	through silicon via
OLED	organic light-emitting diode
RIE	reactive ion etching

CCP	capacitively coupled plasma
ICP	inductively coupled plasma
sccm	standard cubic centimeters per minute
slm	standard liter per minute
ONON	silicon oxide and silicon nitride alternating layers
TCAT	terabit cell array transistor
P-BICS	pipe-shaped bit cost scalable
QMS	Quadrupole mass spectrometer
ARC or BARC	Anti-reflecting coating or Bottom anti-reflecting coating
APF or a-C	Amorphous carbon. Trademark of Applied Materials
CAS	chemical abstract service
PCTFE	Polychlorotrifluoroethene
PVDF	polyvinylidene fluoride
PTFE	Polytetrafluoroethylene
SEM	scanning electron microscope
NAND	Not AND or Negative-NAND or Negated AND
TEOS	Tetraethylorthosilicate
LDTEOS	low deposition rate TEOS
BPSG	Borophosphosilicateglass
p-Si	poly-silicon
XPS	X-ray photoelectron spectroscopy
SiN	Silicon nitride
SiO	Silicon oxide

[0014] As used herein, the term "etch" or "etching" refers to a plasma etch process (*i.e.*, a dry etch process) in which ion bombardment accelerates the chemical reaction in the vertical direction so that vertical sidewalls are formed along the edges of the masked features at right angles to the substrate (Manos and Flamm, Plasma Etching An Introduction, Academic Press, Inc. 1989 pp.12-13). The etching process produces apertures, such as vias, trenches, channel holes, gate trenches, staircase contacts, capacitor holes, contact holes, *etc.*, in the substrate.

[0015] The term "pattern etch" or "patterned etch" refers to etching a non-planar structure, such as a patterned mask layer on a stack of silicon-containing films.

[0016] The term "mask" refers to a layer that resists etching. The mask layer may be located above the layer to be etched.

[0017] The term “etch stop” refers to a layer below the layer to be etched that protects layers underneath.

[0018] The term “device channel” refers to layers that are part of actual device and any damage to it will affect device performance.

[0019] The term “aspect ratio” refers to a ratio of the height of a trench (or via) to the width of the trench (or the diameter of the via).

[0020] The term "selectivity" means the ratio of the etch rate of one material to the etch rate of another material. The term "selective etch" or "selectively etch" means to etch one material more than another material, or in other words to have a greater or less than 1:1 etch selectivity between two materials.

[0021] The term “independently” when used in the context of describing R groups should be understood to denote that the subject R group is not only independently selected relative to other R groups bearing the same or different subscripts or superscripts, but is also independently selected relative to any additional species of that same R group. For example in the formula $MR^1_x (NR^2R^3)_{(4-x)}$, where M is an atom, x is 2 or 3, the two or three R^1 groups may, but need not be identical to each other or to R^2 or to R^3 . Further, it should be understood that unless specifically stated otherwise, values of R groups are independent of each other when used in different formulas.

[0022] Note that herein, the terms “film” and “layer” may be used interchangeably. It is understood that a film may correspond to, or related to a layer, and that the layer may refer to the film. Furthermore, one of ordinary skill in the art will recognize that the terms “film” or “layer” used herein refer to a thickness of some material laid on or spread over a surface and that the surface may range from as large as the entire wafer to as small as a trench or a line.

[0023] Note that herein, the terms “etching compound” and “etching gas” may be used interchangeably. It is understood that an etching compound may correspond to, or related to an etching gas, and that the etching gas may refer to the etching compound.

[0024] As used herein, the abbreviation "NAND" refers to a "Negated AND" or "Not AND" gate; the abbreviation "2D" refers to 2 dimensional gate structures on a planar substrate; the abbreviation "3D" refers to 3 dimensional or vertical gate structures, wherein the gate structures are stacked in the vertical direction.

[0025] The standard abbreviations of the elements from the periodic table of elements are used herein. It should be understood that elements may be referred to by these abbreviations (e.g., Si refers to silicon, N refers to nitrogen, O refers to oxygen, C refers to carbon, H refers to hydrogen, F refers to fluorine, etc.).

[0026] The unique CAS registry numbers (*i.e.*, "CAS") assigned by the Chemical Abstract Service are provided to help better identify the molecules disclosed.

[0027] Please note that the silicon-containing films, such as SiN and SiO, are listed throughout the specification and claims without reference to their proper stoichiometry. The silicon-containing films may include pure silicon (Si) layers, such as crystalline Si, poly-silicon (p-Si or polycrystalline Si), or amorphous silicon; silicon nitride (Si_kN_l) layers; or silicon oxide (Si_nO_m) layers; or mixtures thereof, wherein k, l, m, and n, inclusively range from 0.1 to 6. Preferably, silicon nitride is Si_kN_l , where k and l each range from 0.5 to 1.5. More preferably silicon nitride is Si_3N_4 . Preferably silicon oxide is Si_nO_m , where n ranges from 0.5 to 1.5 and m ranges from 1.5 to 3.5. More preferably, silicon oxide is SiO_2 . Herein, SiO in the following description may be used to represent Si_nO_m containing layers. The silicon-containing film could also be a silicon oxide based dielectric material such as organic based or silicon oxide based low-*k* dielectric materials such as the Black Diamond II or III material by Applied Materials, Inc. with a formula of SiOCH . Silicon-containing film may also include $\text{Si}_a\text{O}_b\text{N}_c$ where a, b, c range from 0.1 to 6. The silicon-containing films may also include dopants, such as B, C, P, As and/or Ge.

Brief Description of the Drawings

[0028] For a further understanding of the nature and objects of the present invention, reference should be made to the following detailed description, taken in conjunction with the accompanying drawings, in which like elements are given the same or analogous reference numbers and wherein:

FIG. 1a is a cross-sectional side view of exemplary layers in a 3D NAND stack;

FIG. 1b is a cross-sectional side view of the exemplary layers in the 3D NAND stack showing polymer deposited on a sidewall during etching of the 3D NAND stack;

FIG. 1c is a cross-sectional side view of the exemplary layers in the 3D NAND stack showing particles generated during alternating SiO/SiN layer etching of a 3D NAND stack;

FIG. 1d is a cross-sectional side view of the exemplary layers in the 3D NAND stack showing selective etching of SiN exposed on the sidewall in the 3D NAND stack;

FIG. 2 is a cross-sectional side view of exemplary layers of a DRAM stack;

FIG. 3a is a cross-sectional side view of exemplary layers showing photoresist pattern over SiO insulation layer surrounding typical transistor device region to produce a transistor structure;

FIG. 3b is a cross-sectional side view of the exemplary layers of **FIG. 3a** after etching the SiO insulation layer;

Fig. 4 is a cross-sectional side view of an exemplary reactor system applied in the exemplary deposition and etching tests;

FIG. 5 is a graph demonstrating the electron impact ionization energy versus species concentration of C_3HF_4N ;

FIG. 6 is a graph demonstrating the electron impact ionization energy versus species concentration of C_2HF_2N ;

FIG. 7 is a X-ray Photoelectron Spectrometric graph demonstrating the element content of a polymerizing film deposited from C_3HF_4N ;

FIG. 8 is a graph demonstrating the deposition or etch rates of SiO, SiN, p-Si and a-C films as a function of oxygen flow rate using C_2HF_2N and O_2 ;

FIG. 9 is a graph demonstrating the deposition or etch rates of SiO, SiN, p-Si and a-C films as a function of oxygen flow rate using C_3HF_4N and O_2 ;

FIG. 10 is a graph demonstrating etch rates of SiO, SiN, p-Si and a-C films as a function of oxygen flow rate using C_3HF_4N and CF_4 ;

FIG. 11 is a graph comparing the silicon oxide deposition and etch rates without oxygen addition of C_3HF_4N , C_2HF_2N , C_5F_5N , C_2F_3N , cC_4F_8 , and C_4F_6 ;

FIG. 12 is a graph comparing the selectivity of silicon oxide to silicon nitride without oxygen addition of C_3HF_4N , C_2HF_2N , C_5F_5N , C_2F_3N , cC_4F_8 , and C_4F_6 ;

FIG. 13 is a graph comparing the silicon oxide etch rates with 5 sccm oxygen addition of C_3HF_4N , C_2HF_2N , C_5F_5N , C_2F_3N , cC_4F_8 , and C_4F_6 ;

FIG. 14 is a graph comparing the selectivity of silicon oxide to silicon nitride with 5 sccm oxygen addition of C_3HF_4N , C_2HF_2N , C_5F_5N , C_2F_3N , cC_4F_8 , and C_4F_6 ;

FIG. 15 is a graph comparing the silicon oxide etch rates with 10 sccm oxygen addition of C_3HF_4N , C_2HF_2N , C_5F_5N , C_2F_3N , cC_4F_8 , and C_4F_6 ;

FIG. 16 is a graph comparing the selectivity of silicon oxide to silicon nitride with 10 sccm oxygen addition of C_3HF_4N , C_2HF_2N , C_5F_5N , C_2F_3N , cC_4F_8 , and C_4F_6 ;

FIG. 17 is a graph comparing the silicon oxide etch rates with 15 sccm oxygen addition of C_3HF_4N , C_2HF_2N , C_2F_3N , cC_4F_8 , and C_4F_6 ;

FIG. 18 is a graph comparing the selectivity of silicon oxide to silicon nitride with 15 sccm oxygen addition of C_3HF_4N , C_2HF_2N , C_2F_3N , cC_4F_8 , and C_4F_6 ;

FIG. 19 is a graph demonstrating the effect of adding N_2 on cC_4F_8 and O_2 etch rates of different substrate materials;

FIG. 20 is a graph demonstrating the effect of adding O_2 to cC_4F_8 and NH_3 on the deposition and etch rates of different substrate materials;

FIG. 21 is a graph demonstrating the electron impact ionization energy versus species concentration of C_2F_3N ;

FIG. 22 is a graph demonstrating etch rates of SiO , SiN , p-Si and a-C films as a function of oxygen flow rates using C_2F_3N and O_2 ;

FIG. 23 is a graph demonstrating the electron impact ionization data versus energy for $C_3H_3F_6N$; and

FIG. 24 is a graph demonstrating the deposition and etch rates without oxygen addition of $C_3H_3F_6N$ on different substrate materials.

Description of Preferred Embodiments

[0029] Disclosed are methods for plasma etching semiconductor structures, such as, channel holes, gate trenches, staircase contacts, capacitor holes, contact holes, *etc.*, in silicon-containing films. The disclosed methods include the steps of i) introducing a vapor of a nitrogen containing etching compound into a reaction chamber containing a silicon-containing film on a substrate, wherein the nitrogen containing etching compound is an organofluorine compound containing at least one $C\equiv N$ or $C=N$ functional group; ii) introducing an inert gas into the reaction chamber; and iii) activating a plasma to produce an activated nitrogen containing etching compound capable of etching the silicon-containing film from the substrate. Herein the organofluorine compounds are organic compounds that contain a carbon-fluorine (C-F) bonds

[0030] The disclosed organofluorine compounds comprising at least one $C\equiv N$ functional group have the general formulae:



wherein R^1 is a functional group having the formula $H_aF_bC_c$ with $a=1-11$, $b=1-11$, and $c=0-5$ and R^2 is a functional group having the formula $H_aF_bC_c$ with $a=0-11$, $b=1-11$, and $c=0-5$. The $H_aF_bC_c$ functional group may be linear, branched, or cyclic and saturated or unsaturated. Applicants believe that the $C\equiv N$ functionality may provide improved etch resistant polymer deposition benefits compared to C_xF_y polymers, wherein x ranges from 0.01 to 1 and y ranges from 0.01 to 4. When the organofluorine compound includes at least one H, the etch rate of SiN may be enhanced while maintaining high selectivity to the mask layer.

[0031] The disclosed organofluorine compounds comprising at least one $C=N$ functional group have the general formula:



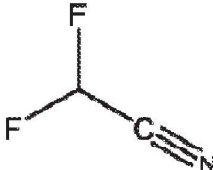
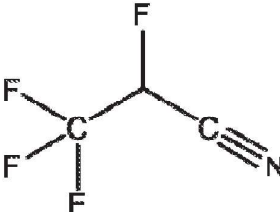
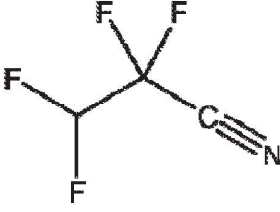
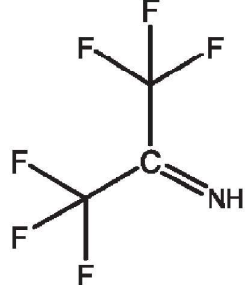
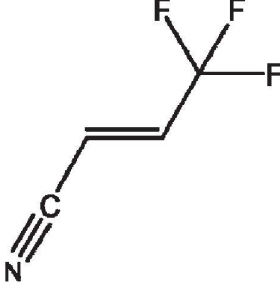
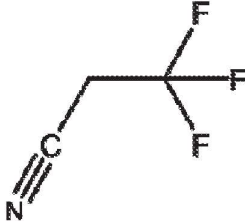
wherein $x=1-2$, $y=1-2$, $z=0-1$, $x+z=1-3$, and each R^1 and R^2 independently has the formula $H_aF_bC_c$ with $a=0-11$, $b=0-11$, and $c=0-5$. The $H_aF_bC_c$ functional group may be linear, branched, or cyclic and saturated or unsaturated. Applicants believe that the $C=N$ functionality may provide improved selectivity among silicon layers compared to C_xF_y polymers, wherein x ranges from 0.01 to 1 and y ranges from 0.01 to 4,. When the organofluorine compound includes at least one H, and preferably when R^2 is H, the etch rate of SiN may be enhanced while maintaining high selectivity to the mask layer.

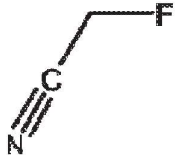
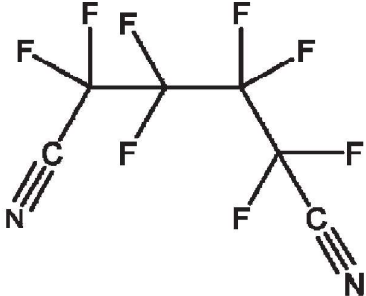
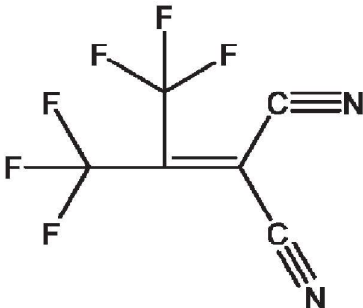
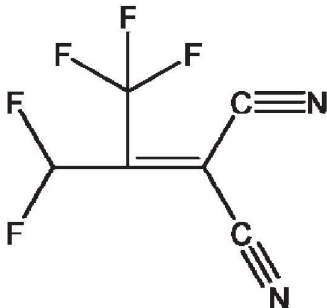
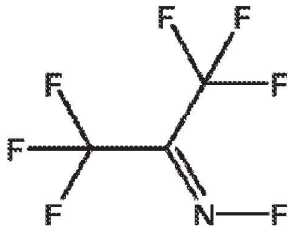
[0032] The disclosed organofluorine compounds may be a hydrofluorocarbon ($C_xH_yF_z$) or fluorocarbon (C_mF_n) compound containing at least one $C\equiv N$ or $C=N$ functional group, or a hetero-hydrofluorocarbon ($C_xH_yF_z$) or hetero-fluorocarbon (C_mF_n) compound containing at least one $C\equiv N$ or $C=N$ functional group, where m , n , x , y and z are integers.

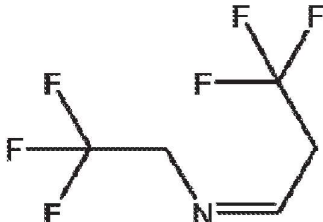
[0033] The disclosed nitrogen containing etching compounds include: difluoroacetonitrile (C_2HF_2N), 2,3,3,3-tetrafluoropropionitrile (C_3HF_4N), 2,2,3,3-tetrafluoropropionitrile (C_3HF_4N), or Hexafluoroacetone imine (C_3HF_6N), which are listed in Table 2. These molecules are commercially available and their structure formula, CAS numbers and boiling points are also included in Table 2.

Table 2. Commercially available etching compounds

Etching compounds	Formula	Structure	CAS number	Boiling point ($^{\circ}C$)
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difluoroacetonitrile	C_2HF_2N		359-12-6	23
2,3,3,3-tetrafluoropropionitrile	C_3HF_4N		431-32-3	40
2,2,3,3-tetrafluoropropionitrile	C_3HF_4N		425-85-4	27
Hexafluoroacetone imine	C_3HF_6N		1645-75-6	16.5
4,4,4-Trifluorocrotononitrile	$C_4H_2F_3N$		406-86-0	84-85
3,3,3-Trifluoropropionitrile	$C_3H_2F_3N$		20530-38-5	92

fluoroacetonitrile	C_2H_2FN		503-20-8	79-80
octafluorohexane-1,6-dinitrile	$C_6F_8N_2$		376-53-4	63-64
1,1-bis(trifluoromethyl)-2,2-dicyanoethylene	$C_6F_6N_2$		1113-69-5	226.2
1-trifluoromethyl, 1-difluoromethyl, -2,2-dicyanoethylene	$C_6HF_5N_2$		439913-54-9	255.9
N,1,1,1,3,3,3-heptafluoropropanamine	C_3F_7N		2802-70-2	-25

1,1,1,6,6,6-Hexafluoro-3-azahex-3-ene	$C_5H_5F_6N$		885275-84-3	87
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[0034] The disclosed nitrogen containing etching compounds may also include 4,4,4-Trifluorocrotononitrile ($C_4H_2F_3N$); 3,3,3-Trifluoropropionitrile ($C_3H_2F_3N$); fluoroacetonitrile (C_2H_2FN); octafluorohexane-1,6-dinitrile ($C_6F_8N_2$); 1,1-bis(trifluoromethyl)-2,2-dicyanoethylene ($C_6F_8N_2$); N,1,1,1,3,3,3-heptafluoropropanamine (C_3F_7N); 1,1,1,6,6,6-Hexafluoro-3-azahex-3-ene ($C_5H_5F_6N$). Their structure formula, CAS numbers and boiling points or predicted boiling points are also included in Table 2. One of ordinary skill in the art will recognize that the synthesis methods for these compounds may be obtained using the CAS numbers provided.

[0035] The disclosed nitrogen-containing etching compounds may provide high selectivity to mask layers, etch stop layers and device channel materials and no profile distortion in high aspect ratio structures, such as those having an aspect ratio ranging from 10:1 to 200:1 such as DRAM and 3D NAND applications. Alternatively, the disclosed nitrogen-containing etching compounds may also provide high selectivity to mask layers or silicon nitride, such as those having an aspect ratio ranging from 1:1 to 50:1 in contact etch applications.

[0036] The disclosed nitrogen-containing etching compounds may provide infinite selectivity for wide process conditions of etching. Herein the selectivity refers to the etching rate ratio of two different layers. For example, the selectivity for SiO layer vs. a-C layer is the etch rate of the SiO divided by the etching rate of the a-C layer. The disclosed nitrogen-containing etching compounds may provide improved selectivity between the silicon-containing films and mask materials, less damage to channel region, and reduced bowing in pattern high aspect ratio structures. The disclosed nitrogen-containing etching compounds may also etch through alternating layers of p-Si, SiO, and/or SiN, resulting in a vertical etch profile (i.e., demonstrating selectivity ranging from 2:1 to 1:2 between the alternating layers).

[0037] The disclosed nitrogen-containing etching compounds are provided at greater than 95% v/v purity, preferably at greater than 99.99% v/v purity, and more

preferably at greater than 99.999% v/v purity. The disclosed nitrogen-containing etching compounds contain less than 5% by volume trace gas impurities, with less than 150 ppm by volume of impurity gases, such as N₂ and/or H₂O and/or CO₂, contained in said trace gaseous impurities. Preferably, the water content in the plasma etching gas is less than 20 ppmw by weight. The purified product may be produced by distillation and/or passing the gas or liquid through a suitable adsorbent, such as a 4A molecular sieve.

[0038] The disclosed nitrogen-containing etching compounds contain less than 10% v/v, preferably less than 1% v/v, more preferably less than 0.1% v/v, and even more preferably less than 0.01% v/v of any of its isomers, which may be purified by distillation of the gas or liquid to remove isomers and may provide better process repeatability.

[0039] Alternatively, the disclosed nitrogen-containing etching compounds may contain between 5% v/v and 50% v/v of its isomers, particularly when the isomer mixture provides improved process parameters or if isolation of the target isomer is too difficult or expensive. For example, the disclosed nitrogen-containing etching compounds may comprise between approximately 50% v/v and approximately 75% v/v 2,3,3,3-tetrafluoropropionitrile and between approximately 25% v/v and approximately 50% v/v 2,2,3,3-tetrafluoropropionitrile. The mixture of isomers may reduce the need for two or more gas lines to the reaction chamber.

[0040] Some of the disclosed nitrogen-containing etching compounds are gaseous at room temperature and atmospheric pressure. For the non-gaseous (*i.e.*, liquid or solid) compounds, their gas form may be produced by vaporizing the compounds through a conventional vaporization step, such as direct vaporization or by bubbling with inert gas (N₂, Ar, He). The non-gaseous compounds may be fed in liquid state to a vaporizer where it is vaporized before it is introduced into a reactor.

[0041] The disclosed nitrogen-containing etching compounds are suitable for plasma etching semiconductor structures, such as, channel holes, gate trenches, staircase contacts, capacitor holes, contact holes, *etc.*, in the silicon-containing films. The disclosed nitrogen-containing etching compounds are not only compatible with currently available mask materials but also compatible with the future generations of mask materials because the disclosed nitrogen-containing etching compounds induce little to no damage on the mask along with good profile of high aspect ratio structures. In other words, the disclosed nitrogen-containing etching compounds may

produce vertical etched patterns having minimal to no bowing, pattern collapse, or roughness. In order to achieve these properties, the disclosed nitrogen-containing etching compounds may deposit an etch-resistant polymer layer during etching to help reduce the direct impact of the oxygen and fluorine radicals during the etching process. The disclosed nitrogen-containing etching compounds may also reduce damage to p-Si or crystalline Si channel structure during etching. Preferably, the disclosed nitrogen-containing etching compounds are suitably volatile and stable during the etching process for delivery into the reactor/chamber.

[0042] Material compatibility tests are important to determine if any of the disclosed nitrogen-containing etching compounds will react with chamber materials and degrade the performance of the chamber with short term or long term use. Key materials involved in parts of the chamber, valves, *etc.* include stainless steel, aluminum, nickel, PCTFE, PVDF, PTFE and other metals and polymers. At times these materials are exposed to high temperatures, for example, higher than 20°C, and high pressures, for example, higher than 1 atm, which may enhance their degradation. The metrology methods may include visual inspection, weight measurement, measuring nanometer scale changes in scanning electron microscopy (SEM), tensile strength, hardness, *etc.*

[0043] The disclosed nitrogen-containing etching compounds may be used to plasma etch silicon-containing films on a substrate. The disclosed plasma etching method may be useful in the manufacture of semiconductor devices such as NAND or 3D NAND gates or Flash or DRAM memory or transistors such as fin-shaped field-effect transistor (FinFET), Bulk complementary metal-oxide-semiconductor (Bulk CMOS), fully depleted silicon-on-insulator (FD-SOI) structures. The disclosed nitrogen-containing etching compounds may be used in other areas of applications, such as different front end of the line (FEOL) and back end of the line (BEOL) etch applications. Additionally, the disclosed nitrogen-containing etching compounds may also be used for etching Si in 3D through silicon via (TSV) etch applications for interconnecting memory to logic on a substrate.

[0044] The plasma etching method includes providing a reaction chamber having a substrate disposed therein. The reaction chamber may be any enclosure or chamber within a device in which etching methods take place such as, and without limitation, reactive ion etching (RIE), capacitively coupled plasma (CCP) with single or multiple frequency RF sources, inductively coupled plasma (ICP), or microwave

plasma reactors, or other types of etching systems capable of selectively removing a portion of the silicon-containing film or generating active species. One of ordinary skill in the art will recognize that the different plasma reaction chamber designs provide different electron temperature control. Suitable commercially available plasma reaction chambers include but are not limited to the Applied Materials magnetically enhanced reactive ion etcher sold under the trademark eMAX™ or the Lam Research Dual CCP reactive ion etcher dielectric etch product family sold under the trademark 2300® Flex™. The RF power in such may be pulsed to control plasma properties and thereby improving the etch performance (selectivity and damage) further.

[0045] Alternatively, the plasma-treated reactant may be produced outside of the reaction chamber. The MKS Instruments' ASTRONi® reactive gas generator may be used to treat the reactant prior to passage into the reaction chamber. Operated at 2.45 GHz, 7kW plasma power, and a pressure ranging from approximately 0.5 Torr to approximately 10 Torr, the reactant O₂ may be decomposed into two O· radicals. Preferably, the remote plasma may be generated with a power ranging from about 1 kW to about 10 kW, more preferably from about 2.5 kW to about 7.5 kW.

[0046] The reaction chamber may contain one or more than one substrate. For example, the reaction chamber may contain from 1 to 200 silicon wafers having from 25.4 mm to 450 mm diameters. The substrates may be any suitable substrates used in semiconductor, photovoltaic, flat panel or LCD-TFT device manufacturing. Examples of suitable substrates include wafers, such as silicon, silica, glass, or GaAs wafers. The wafer will have multiple films or layers on it from previous manufacturing steps, including silicon-containing films or layers. The layers may or may not be patterned. Examples of suitable layers include without limitation silicon (such as amorphous silicon, p-Si, crystalline silicon, any of which may further be p-doped or n-doped with B, C, P, As, and/or Ge), silica, silicon nitride, silicon oxide, silicon oxynitride, Si_aO_bH_cC_dN_e, (wherein a>0; b, c, d, e ≥ 0), mask layer materials such as amorphous carbon, antireflective coatings, photoresist materials, tungsten, titanium nitride, tantalum nitride or combinations thereof, etch stop layer materials such as silicon nitride, polysilicon, crystalline silicon, silicon carbide, SiCN or combinations thereof, device channel materials such crystalline silicon, epitaxial silicon, doped silicon, Si_aO_bH_cC_dN_e, (wherein a>0; b, c, d, e ≥ 0) or combinations thereof. The silicon oxide layer may form a dielectric material, such as an organic

based or silicon oxide based low- k dielectric material (e.g., a porous SiCOH film). An exemplary low- k dielectric material is sold by Applied Materials under the trade name Black Diamond II or III. Additionally, layers comprising tungsten or noble metals (e.g. platinum, palladium, rhodium or gold) may be used. Furthermore, examples of the silicon-containing films may be $\text{Si}_a\text{O}_b\text{H}_c\text{C}_d\text{N}_e$, (wherein $a > 0$; $b, c, d, e \geq 0$). Throughout the specification and claims, the wafer and any associated layers thereon are referred to as substrates.

[0047] The following are exemplary embodiments of the substrates on which the disclosed nitrogen-containing etching compounds may be applied to etch.

[0048] In one embodiment, a substrate **100** may include a stack of multiple layers as shown in **FIG. 1a**. **FIG. 1a** is a cross-sectional side view of exemplary layers in a 3D NAND stack to produce a 3D NAND gate. In **FIG. 1a**, a stack of seven alternative SiO/SiN (*i.e.*, **104a/104b**) layers **104** is located on top of a silicon wafer **102** (*i.e.*, ONON or TCAT technology). One of ordinary skill in the art will recognize that some technologies replace the SiN layers **104a** with p-Si layers (e.g., SiO/p-Si or P-BICS technology). An amorphous carbon (a-C) mask layer **106** is located on the top of the seven SiO/SiN layers **104**. The a-C mask layer **106** may contain C and H, as well as other elements, such as boron, nitrogen, *etc.*, to improve etch resistance during SiO/SiN layer etch. An antireflective coating layer **108** is located on top of the a-C mask layer **106**. A patterned photoresist layer **110** is located on top of the antireflective coating layer **108**. Herein, a SiON layer (not shown) may be present between the antireflective coating layer **108** and the a-C mask layer **106** to transfer pattern in photoresist layer **110** to the a-C layer **106**. One of ordinary skill in the art will recognize that the stack of layers in the substrate **100** in **FIG. 1a** is provided for exemplary purposes only and that the disclosed nitrogen-containing etching compounds may be used to etch other types of stacks of layers. Furthermore, one of ordinary skill in the art will recognize that the number alternating SiO/SiN or SiO/p-Si layers **104** in the stack of the substrate **100** may vary (*i.e.*, may include more or less than the seven SiO/SiN (**104a/104b**) layers depicted).

[0049] **FIG. 1b** is a cross-sectional side view of the exemplary layers in the 3D NAND stack showing polymer deposited on a sidewall during etching. The disclosed nitrogen containing compounds may produce fragments during the plasma process that are suitable for both anisotropically etching the silicon-containing films **104** and depositing a N-containing polymer passivation layer **212** on sidewalls of a structure

being etched as shown in **FIG. 1b**. The difference between **FIG. 1b** and **FIG. 1a** is in **FIG. 1b** via **214** is formed in substrate **100** by plasma etching using the disclosed nitrogen-containing etch compounds, which also deposit the polymer passivation layer **212** on the sidewalls of the via **214**. The polymer passivation layer **212** also provides smoother sidewall, less bowing and less deformation at the bottom of the via **214**. The polymer passivation layer **212** may however be easily removed or cleaned by dry or wet etch chemistries known in the art.

[0050] **FIG. 1c** is a cross-sectional side view of the exemplary layers in the 3D NAND stack showing particles **316** generated during alternating SiO/SiN layer etching in a 3D NAND stack. The particles **316** generated on the sidewalls of the alternative SiO/SiN (*i.e.*, **104a/104b**) layers **104** as shown in **FIG. 1c** may be minimized by using the disclosed nitrogen-containing compounds. The difference between **FIG. 1c** and **FIG. 1b** is in **FIG. 1c** the alternative SiO/SiN exposed sidewall has particle **316** generated during plasma etching. Applicants do not believe that the disclosed nitrogen-containing compounds will generate the particles **316** shown in **Fig. 1c**.

[0051] **FIG. 1d** is a cross-sectional side view of the exemplary layers in the 3D NAND stack showing selective etching of SiN exposed on the sidewall in the 3D NAND stack after etching. The SiN exposed sidewall in stack **100** may be etched selectively as shown in **FIG. 1d** by using the disclosed nitrogen containing compounds to selectively break Si-N bond in SiN layers **104b** over Si-O bond in SiO layers **104a** forming a selective sidewall SiN etch **418** on the stack of SiO/SiN layers **104** in the via **214**. The difference between **FIG. 1d** and **FIG. 1b** is in **FIG. 1d** SiN exposed on the alternative SiO/SiN sidewall is selectively etched by the disclosed nitrogen containing compounds forming the selective sidewall SiN etch **418**.

Typically, the selective sidewall SiN etch **418** is performed by wet etching using mixtures with phosphoric acid. Replacing wet etch process with dry plasma etch processes is known to greatly improve the economics of semiconductor device fabrication process since wet etching requires moving the substrate to different wet etching equipments. With the disclosed methods, all etching including selective sidewall SiN etch may be performed in one etch equipment, which may reduce the cost of the semiconductor fabrication.

[0052] In alternative embodiment, substrate **100** may include a stack of multiple layers thereon as shown in **FIG. 2**. **FIG. 2** is a cross-sectional side view of

exemplary layers in a DRAM stack to produce a DRAM memory. In **FIG. 2**, a stack of four layers is located on top of a silicon wafer **102**. An a-C mask layer **106** is located on top of a large SiO layer **104a**. An antireflective coating layer **108** is located on top of the a-C mask **106**. A pattern photoresist layer **110** is located on top of the antireflective coating **108**. Herein, a SiON layer (not shown) may be present between the antireflective coating layer **108** and the a-C mask layer **106** to transfer pattern in photoresist layer **110** to the a-C layer **106**. One of ordinary skill in the art will recognize that the stack of layers in **FIG. 2** is provided for exemplary purposes only and that the disclosed nitrogen-containing etching compounds may be used to etch other stacks of layers, for example, for a stack where the a-C mask layer **106** is replaced with a TiN layer. Furthermore, one of ordinary skill in the art will recognize that the number of layers in the stack may vary (*i.e.*, may include more or less than the layers depicted).

[0053] **FIG. 3a** is a cross-sectional side view of exemplary layers showing photoresist pattern over SiO insulation layer surrounding a typical transistor device region to produce a transistor structure. Substrate **600** may include a stack of four layers surrounding a transistor gate electrode region supported on the Silicon wafer **602** as shown in **FIG. 3a**. The transistor region shown in **FIG. 3a** includes two doped silicon regions **606** acting as source and drain. Transistor gate dielectric **614** is present underneath gate electrode **616**. The whole transistor, *i.e.*, transistor gate dielectric **614** and gate electrode **616**, is surrounded by a thin SiN layer **608** which may later act as etch stop layer during contact etch. Each transistor device region **616/606** is separated by SiO isolation regions **604** in the silicon wafer **602** to minimize electrical interference. One of ordinary skill in the art will recognize that layer **602** may be located on top of a silicon oxide layer of the Silicon on Insulator (SOI) wafer. Another SiO layer **610** is deposited on the transistor and used to insulate metal contact to the transistor device regions **606**. Photoresist mask **612** is used to pattern the SiO layer **610**. Etching is performed using the disclosed nitrogen-containing etching compounds in plasma environment. Herein, photoresist serves as a mask to etch the SiO layer **610** and etching is stopped on the SiN layer **608** as shown in **FIG. 3b**.

[0054] **FIG. 3b** is a cross-sectional side view of the exemplary layers of **FIG. 3a** after etching the SiO insulation layer. The difference between **FIG. 3b** and **FIG. 3a** is in **FIG. 3b** vias **718** are formed in the SiO layer **610** through etching by the disclosed

nitrogen-containing compounds. The SiO layer **610** may be etched with photoresist layer **612** as a mask layer. The mask layer may be any suitable photoresist mask materials, such as TiN, a-C and so on. The etching may be stopped at the underlying SiN layer **608**.

[0055] The disclosed nitrogen-containing etching compounds may also be used to etch the SiN layer **608** with different plasma conditions and different mixtures. One of ordinary skill in the art will recognize that the stack and geometry of layers in **FIG. 3a** and **FIG. 3b** is provided for exemplary purposes only and that the disclosed nitrogen-containing etching compounds may be used to etch other types of stacks of layers. Furthermore, one of ordinary skill in the art will recognize that the number of layers in the stack may vary (*i.e.*, may include more or less than the four layers depicted).

[0056] The vapor of the disclosed nitrogen-containing etching compounds is introduced into the reaction chamber containing the substrate and silicon-containing films. The vapor may be introduced to the chamber at a flow rate ranging from approximately 0.1 sccm to approximately 1 slm. For example, for a 200 mm wafer size, the vapor may be introduced to the chamber at a flow rate ranging from approximately 5 sccm to approximately 50 sccm. Alternatively, for a 450 mm wafer size, the vapor may be introduced to the chamber at a flow rate ranging from approximately 25 sccm to approximately 250 sccm. One of ordinary skill in the art will recognize that the flow rate may vary from tool to tool.

[0057] The disclosed nitrogen-containing etching compounds may be supplied either in neat form or in a blend with an inert gas, such as N₂, Ar, He, Xe, etc, or solvent. The disclosed nitrogen-containing etching compounds may be present in varying concentrations in the blend. For liquid nitrogen-containing compounds, the vapor form of the nitrogen-containing etching compounds may be produced by vaporizing the neat or blended nitrogen-containing etching compound solution through a conventional vaporization step such as direct vaporization or by bubbling. The neat or blended nitrogen-containing etching compounds may be fed in liquid state to a vaporizer where it is vaporized before it is introduced into the reactor.

[0058] Alternatively, the neat or blended nitrogen-containing etching compounds may be vaporized by passing a carrier gas into a container containing the disclosed nitrogen-containing etching compounds or by bubbling the carrier gas into the disclosed nitrogen-containing compounds. The carrier gas may include, but is not

limited to, Ar, He, N₂, and mixtures thereof. Bubbling with a carrier gas may also remove any dissolved oxygen present in the neat or blended nitrogen-containing etching compounds solution. The carrier gas and disclosed nitrogen-containing etching compounds are then introduced into the reactor as a vapor.

[0059] If necessary, the container containing the disclosed nitrogen-containing etching compounds may be heated to a temperature that permits the nitrogen-containing etching compounds to be in liquid phase and to have a sufficient vapor pressure for delivery into an etching tool. The container may be maintained at temperatures in the range of, for example, approximately 0°C to approximately 150°C, preferably from approximately 25°C to approximately 100°C, more preferably from approximately 25°C to approximately 50°C. More preferably, the container is maintained at room temperature (approximately 25°C) in order to avoid heating lines to the etch tool. Those skilled in the art recognize that the temperature of the container may be adjusted in a known manner to control the amount of nitrogen-containing compound vaporized.

[0060] Additionally, the nitrogen-containing etching compounds are delivered in purity ranging from 95% to 99.999% by volume and may be purified with known standard purification techniques for removal of CO, CO₂, N₂, H₂O, HF, H₂S, SO₂, halides, and other hydrocarbons or hydrohalocarbons.

[0061] An inert gas is also introduced into the reaction chamber in order to sustain the plasma. The inert gas may be He, Ar, Xe, Kr, Ne, N₂ or combinations thereof. The etching gas and the inert gas may be mixed prior to introduction to the chamber, with the inert gas comprising between approximately 0.01% v/v and approximately 99.9% v/v of the resulting mixture. Alternatively, the inert gas may be introduced to the chamber continuously while the etching gas is introduced to the chamber in pulses.

[0062] The vapor of the disclosed etching gas and inert gas are activated by plasma to produce an activated etching gas. The plasma decomposes the etching gas into radical form (*i.e.*, the activated etching gas). The plasma may be generated by applying RF or DC power. The plasma may be generated with a RF power ranging from about 25W to about 10,000W. The plasma may be generated remotely or within the reactor itself. The plasma may be generated in dual CCP or ICP mode with RF applied at both electrodes. RF frequency of plasma may range from 200 KHz to 1GHz. Different RF sources at different frequency may be coupled and

applied at same electrode. Plasma RF pulsing may be further used to control molecule fragmentation and reaction at substrate. One of skill in the art will recognize methods and apparatus suitable for such plasma treatment.

[0063] A quadrupole mass spectrometer (QMS), optical emission spectrometer, FTIR, or other radical/ion measurement tools may measure the activated etching gas from the chamber exhaust to determine the types and numbers of species produced. If necessary, the flow rate of the etching gas and/or the inert gas may be adjusted to increase or decrease the number of radical species produced.

[0064] The disclosed etching gases may be mixed with other gases either prior to introduction into the reaction chamber or inside the reaction chamber. Preferably, the gases may be mixed prior to introduction to the chamber in order to provide a uniform concentration of the entering gas.

[0065] In another alternative, the vapor of the nitrogen-containing compound may be introduced into the chamber independently of the other gases such as when two or more of the gases react.

[0066] In another alternative, the etching gas and the inert gas are the only two gases that are used during the etching process.

[0067] Exemplary other gases include, without limitation, oxidizers such as O₂, O₃, CO, CO₂, NO, N₂O, NO₂, and combinations thereof. The disclosed etching gases and the oxidizer may be mixed together prior to introduction into the reaction chamber.

[0068] Alternatively, the oxidizer may be introduced continuously into the chamber and the etching gas introduced into the chamber in pulses. The oxidizer may comprise between approximately 0.01% v/v to approximately 99.99% v/v of the mixture introduced into the chamber (with 99.99% v/v representing introduction of almost pure oxidizer for the continuous introduction alternative).

[0069] Other exemplary gases with which the etching gas may be mixed include additional etching gases, such as cC₄F₈, C₄F₈, C₄F₆, CF₄, CH₃F, CF₃H, CH₂F₂, COS, CS₂, CF₃I, C₂F₃I, C₂F₅I, SO₂, trans-1,1,1,4,4,4-hexafluoro-2-butene (*trans*-C₄H₂F₆), cis-1,1,1,4,4,4-hexafluoro-2-butene (*cis*-C₄H₂F₆), hexafluoroisobutene (C₄H₂F₆), trans-1,1,2,2,3,4-hexafluorocyclobutane (*trans*-C₄H₂F₆), 1,1,2,2,3-pentafluorocyclobutane (C₄H₃F₅), 1,1,2,2-tetrafluorocyclobutane (C₄H₄F₄), or cis-1,1,2,2,3,4-hexafluorocyclobutane (*cis*-C₄H₂F₆).

[0070] The vapor of the etching gas and the additional gas may be mixed prior to introduction to the reaction chamber. The additional etching gas may comprise between approximately 0.01% v/v to approximately 99.99% v/v of the mixture introduced into the chamber.

[0071] In one non-limiting exemplary plasma etch process, the vapor of 2,3,3,3-Tetrafluoropropionitrile is introduced into a 200 mm Dual CCP plasma etch tool using a controlled gas flow device. The controlled gas flow device may be a mass flow controller or a bubbler design with inert gas flow to deliver the vapor of the desired molecule. In case of high boiling point molecules, a special low pressure drop mass flow controller from Brooks Automation (No. GF120XSD), MKS Instruments, *etc.*, may be used. The pressure of the reaction chamber is set at approximately 30 mTorr. No gas source heating is necessary, as the vapor pressure of 2,3,3,3-Tetrafluoropropionitrile is approximately 498 torr at room temperature. The distance between the two CCP electrodes is kept at 1.35 cm and the top electrode RF power is fixed at 750 W. The bottom electrode RF power is varied to analyze the performance of the molecule. The reaction chamber contains a substrate having silicon-containing films thereon, similar to those shown in **FIG. 1a**. The antireflective coating layer **108** is patterned/etched by a fluorocarbon (*e.g.*, CF_4 and CH_2F_2) and oxygen-containing gas (*e.g.*, O_2). The amorphous carbon mask layer is patterned/etched by an oxygen-containing gas. The SiO and SiN layers **104** are patterned by the plasma of the disclosed nitrogen containing organofluorine compounds (*e.g.*, 2,3,3,3-Tetrafluoropropionitrile) and Argon. Argon is independently introduced into the chamber at a 250 sccm flow rate. 2,3,3,3-Tetrafluoropropionitrile is independently introduced into the chamber at 15 sccm. O_2 is independently introduced into the chamber and varied from 0 sccm to 20 sccm to determine optimum etching conditions. An aperture having an aspect ratio equal to or greater than 30:1 is produced, which may be used as a channel hole in vertical NAND. Similar example may be used for other stack layers as shown in **Figs 2 and 3a**.

[0072] In another non-limiting exemplary plasma etch process, difluoroacetonitrile is introduced into a 200 mm Dual CCP plasma etch tool using a controlled gas flow device. The controlled gas flow device may be a mass flow controller. In case of high boiling point molecules, a special low pressure drop mass flow controller from BrooksAutomation (No. GF120XSD), MKS Instruments, *etc.*, may be used. The pressure of the reaction chamber is set at approximately 30 mTorr. No gas source

heating is necessary, as the vapor pressure of difluoroacetonitrile is approximately 900 Torr at 20°C. The distance between the two CCP electrodes is kept at 1.35 cm and the top electrode RF power is fixed at 750 W. The bottom electrode RF power is varied to analyze the performance of difluoroacetonitrile. The reaction chamber contains a substrate **100** having a thick SiO layer **104a** thereon, similar to the layer shown in **FIG. 2**. Prior to this process, the antireflective coating layer **108** is removed by a fluorocarbon (e.g., CF₄) and oxygen-containing gas (e.g., O₂) and the A-c mask layer **106** is removed by an oxygen-containing gas. Argon is independently introduced into the chamber at a 250 sccm flow rate. Difluoroacetonitrile is independently introduced into the chamber at 15 sccm. O₂ is independently introduced into the chamber at 0-20 sccm to determine optimum etching conditions. An aperture having an aspect ratio equal to or greater than 10:1 is produced, which may be used as a contact hole in DRAM. Similar examples may be used for other stack layers as shown in **Figs 1a** and **3a**.

[0073] The silicon-containing films and the activated etching gas react to form volatile by-products that are removed from the reaction chamber. The a-C mask, antireflective coating, and photoresist layer are less reactive with the activated etching gas. Thus, the activated etching gas selectively reacts with the silicon-containing films to form volatile by-products.

[0074] The temperature and the pressure within the reaction chamber are held at conditions suitable for the silicon-containing film to react with the activated etching gas. For instance, the pressure in the chamber may be held between approximately 0.1 mTorr and approximately 1000 Torr, preferably between approximately 1 mTorr and approximately 10 Torr, more preferably between approximately 10 mTorr and approximately 1 Torr, and more preferably between approximately 10 mTorr and approximately 100 mTorr, as required by the etching parameters. Likewise, the substrate temperature in the chamber may range between about approximately -196°C to approximately 500°C, preferably between approximately -120°C to approximately 300°C, more preferably between approximately -100°C to approximately 50°C; and more preferably between approximately -10°C to approximately 40°C. Chamber wall temperatures may range from approximately -196°C to approximately 300°C depending on the process requirements.

[0075] The reactions between the silicon-containing film and the activated etching gas result in anisotropic removal of the silicon-containing films from the substrate.

Atoms of nitrogen, oxygen, and/or carbon may also be present in the silicon-containing film. The removal is due to a physical sputtering of silicon-containing film from plasma ions (accelerated by the plasma) and/or by chemical reaction of plasma species to convert Si to volatile species, such as SiF_x , wherein x ranges from 1-4.

[0076] The plasma activated vapor of the disclosed nitrogen-containing etching compounds preferably exhibits high selectivity toward the mask and etches through the alternating layers of SiO and SiN resulting in a vertical etch profile with no bowing or roughness, which is important for 3D NAND applications. Additionally, plasma activated vapor deposits polymer on sidewall to minimize feature profile deformation. For other applications, such as DRAM and 2D NAND, for example, the plasma activated etching gas under different process conditions may selectively etch SiO from SiN. The plasma activated etching gas may selectively etch SiO and/or SiN from mask layers, such as a-C, photoresist, p-Si, or silicon carbide; or from metal contact layers, such as Cu; or from channel regions consisting of SiGe or polysilicon regions.

[0077] The disclosed etch processes using the disclosed nitrogen-containing etching compounds as the etching gases produce channel holes, gate trenches, staircase contacts, capacitor holes, contact holes, *etc.*, in the silicon-containing films. The resulting aperture may have an aspect ratio ranging from approximately 10:1 to approximately 200:1 and a diameter ranging from approximately 5 nm to approximately 50 nm. For example, one of ordinary skill in the art will recognize that a channel hole etch produces apertures in the silicon-containing films having an aspect ratio greater than 60:1.

[0078] Typical materials that need to be etched may be SiO. A process of etching SiO may be relevant to etching trenches in Borophosphosilicateglass (BPSG), Tetraethylorthosilicate (TEOS), or low deposition rate TEOS (LDTEOS). An etch stop layer may be silicon nitride or silicon oxygen nitride (SiON) or poly silicon. A mask material used may be a-C, p-Si, or photo resist materials. Herein, the disclosed nitrogen-containing etching compounds are applied to etch SiO, SiN, p-Si and/or a-C substrate films.

Examples

[0079] The following non-limiting examples are provided to further illustrate embodiments of the invention. However, the examples are not intended to be all inclusive and are not intended to limit the scope of the inventions described herein.

[0080] In the following examples, the etch performance of C_2HF_2N and C_3HF_4N nitrogen containing compounds are evaluated and compared against C_5F_5N (CAS 700-16-3), C_2F_3N , and standard gases like cC_4F_8 and C_4F_6 . The results show that C_2HF_2N and C_3HF_4N nitrogen-containing etching compounds offer sidewall protection and could be used for etching semiconductor structures, such as contact etch.

[0081] Herein, the target etching requirements for etchants or etching gases for contact etch are:

1. Etch rates of oxide (*i.e.*, SiO) have to be high;
2. Selectivity of SiO to other materials, typically SiN, should be high;
3. Selectivity of SiO to other materials, typically p-Si or a-C, should be high.

[0082] As will be shown, the results of the compounds C_3HF_4N and C_2HF_2N show greater promise because they meet all the required etching targets for contact etching applications.

[0083] Etching experiments were performed on four $1 \times 1 \text{ cm}^2$ coupons having four different substrate materials including SiO, SiN, p-Si, and a-C. Deposition and/or etch rates are measured using an ellipsometer and/or SEM by measuring the change in etch thickness as a function of etching time. The coupons are placed on 200 mm diameter carrier wafer and held in contact by using double sided carbon tape obtained from 2spi manufacturer. Alternatively, thermal paste could be used to stick coupons on carrier wafer.

[0084] Deposition tests are performed on $1 \times 1 \text{ cm}^2$ Si coupon at 30 mTorr, and source power of 750 W (27 MHz), with no bias power at the substrate. The process feed mixture contains 250 sccm of Ar and 15 sccm of etch gas. The deposition test sample is then sent for X-ray Photoelectron spectrometry (XPS) analysis to study the type of polymerizing film formed on the substrate.

[0085] Etching tests are also performed at 30 mTorr, source power of 750 W (27 MHz), and bias power of 1500 W (2 MHz). The feed mixture contains 250 sccm of Ar, 15 sccm of etch gas, while O_2 is varied in the range 0 to 15 sccm.

[0086] **Fig. 4** is an exemplary cross-sectional side view of exemplary reactor system applied in deposition and etching tests. As shown, the reactor **800** includes a reactor chamber **802**. Inside of the reactor chamber **802**, a wafer **806** attached on the top of a bottom electrode **804** may be placed in the bottom portion of the reactor chamber **802**, and a silicon top electrode showerhead **808** may be placed on the top portion of the reactor chamber **802**. The bottom electrode **804** may be an electrostatic chuck having bias power applied thereto. For example, 2 MHz RF bias power may be applied to the bottom electrode **804**. The wafer **806** may have multiple layers that need to be etched. The silicon top electrode showerhead **808** has a plurality of holes **810** in the showerhead through which the gases pass. The gases may be introduced into the reactor chamber **802** through gas inlet **812** and then pass through holes **810** in the showerhead **808** for uniform gas distribution. Source power may be applied to the silicon top electrode showerhead **808**. For example, 27 MHz RF source power may be applied to the silicon top electrode showerhead **808**. Between the silicon top electrode showerhead **808** and the bottom electrode **804** is the plasma region. The gases passing through the holes **810** in the showerhead **808** may be ionized in the plasma region and then perform etching on the wafer **806**. The gases may be removed by pumping the gases out of the reactor chamber **802** from outlet **814**.

[0087] Additionally, a mass spectrometer may be used to study electron impact ionizations of the etch gases. For this test, the etch gases are allowed to flow through the mass spectrometer chamber and a Quadrupole mass spectrometer (Hiden Analytical Inc.) detector is used to study the fragments from the etch gas as a function of electron energy.

Example 1

[0088] **FIG. 5** is a graph demonstrating electron impact ionization data for C_3HF_4N . In **FIG. 5**, the x-axis represents electron energy and y-axis represents the partial pressure of the fragment species. **FIG. 5** shows that the major fragments for C_3HF_4N are CF_3 and C_2HFN . The C_2HFN fragment has 1:2 F/C ratios and may readily polymerize upon reaching the substrate.

Example 2

[0089] **FIG. 6** is a graph demonstrating electron impact ionization data for C_2HF_2N . In **FIG. 6**, the x-axis represents electron energy and y-axis represents the partial pressure of the fragment species. **FIG. 6** shows that the major fragments for C_2HF_2N are CF_2 and C_2F_2N . The C_2F_2N fragment has a low F/C ratio and will readily polymerize reaching the substrate.

Example 3

[0090] The deposition test is an experiment conducted on a blanket $1 \times 1 \text{ cm}^2$ Si coupon, where the source power (750 W @ 27 MHz) is applied with no bias power. Because of the absence of bias power, the ions reaching the substrate may not have sufficient energy to etch. In addition, the neutrals and active species reaching the surface stick to the surface, based on their sticking coefficient, and deposit a thin polymer layer. This thin polymer layer may be responsible for sidewall passivation that often provides the selectivity. The deposition test experimental condition helps simulate the polymer layer formed during plasma processing of patterns either on surface or on sidewalls.

[0091] When the deposition test has been performed, for 60 sec, at 30 mTorr pressure with process gas mixture containing 250 sccm of Ar and 15 sccm of C_2F_3N , a 260 nm polymer film is deposited on Si. The deposition rate is thus 260 nm/min for C_2F_3N .

[0092] For the same deposition test conditions, it was found that C_2HF_2N yielded a deposition rate of 150 nm/min and C_3HF_4N yielded a deposition rate of 190 nm/min. For C_5F_5N , the deposition test has been performed, for 60 sec, at 30 mTorr pressure with process gas mixture containing 250 sccm of Ar and 5 sccm of C_5F_5N , and it yields deposition rate of 120 nm/min.

[0093] The sample is further sent for XPS analysis where the nature of the polymerizing layer is investigated. **FIG. 7** is a graph providing XPS data of a polymerizing film formed with C_3HF_4N during deposition test. The wide elemental scan from XPS analysis, from **FIG. 7**, clearly shows the presence of C, F, O, and N peaks, which shows the evidence of Nitrogen present in the polymerizing film. Thus, nitrogen is present in the polymerizing film and it may act as a better sidewall passivation layer and helps create vertical profiles during high aspect ratio etching.

Example 4

[0094] **FIG. 8** is a graph demonstrating etch rates of SiO, SiN, p-Si and a-C with C₂HF₂N and O₂. In **FIG. 8**, the positive y-axis represents etch rates while the negative y-axis represents deposition rates; the x-axis is O₂ flow rate in sccm; the C₂HF₂N flow rate is fixed at 15 sccm while the O₂ flow rate is varied from 0 to 15 sccm.

[0095] As shown, when no oxygen is added (0 sccm O₂ condition), C₂HF₂N readily etches silicon oxide but does not etch other substrates. This is a very important result as C₂HF₂N etches silicon oxide and protects the other substrate materials offering infinite selectivity of oxide to other substrate materials. Small damage of few nanometer due to plasma ion bombardment is expected before the etch gas starts protecting the other substrate materials. This damage is not characterized and not accounted for in the selectivity measurements. When 5 sccm of oxygen is added to the mixture, the etch rate of silicon oxide decreases compared to the 0 sccm oxygen condition, but C₂HF₂N still deposits on all other substrates preserving the selectivity of Oxide to other substrate materials. When 10 sccm of oxygen is added to the mixture, the etch rate of silicon oxide continues to decrease, but, C₂HF₂N still deposits on all other substrates preserving the selectivity of silicon oxide to other substrate materials. When 15 sccm of oxygen is added to the mixture, while the etch rate of silicon oxide once again decreases, C₂HF₂N still deposits on Nitride and p-Si, and preserves selectivity of silicon oxide to silicon nitride and p-Si. At 15 sccm O₂ test condition, etching is observed on a-C and the selectivity of silicon oxide to a-C decreases drastically from infinity to ~2 (*i.e.*, etch rate of SiO/ Etch rate of a-C). Overall, C₂HF₂N offers the widest possible range of process conditions that give infinite selectivity to silicon nitride and p-Si substrates. The etch rates of silicon oxide are lower than the standard cC₄F₈ gas (which is above 550 nm/min) but may be readily increased by adding an additive gas like CF₄, C₃F₈, cC₄F₈ or C₄F₆.

Example 5

[0096] **FIG. 9** is a graph demonstrating etch rates of SiO, SiN, p-Si and a-C with C₃HF₄N and O₂. In **FIG. 9**, the positive y-axis represents etch rates while the negative y-axis represents deposition rates; the x-axis is O₂ flow rate in sccm; the C₃HF₄N flow rate is fixed at 15 sccm while the O₂ flow rate is varied from 0 to 15 sccm.

[0097] As shown, when no oxygen is added (0 sccm O₂ condition), C₃HF₄N deposits on all substrates. As 5 sccm of oxygen is added to the mixture, C₃HF₄N starts etching silicon oxide but deposits on all the other substrate materials offering infinite selectivity of silicon oxide to the other substrate materials. Similar behavior is seen at 10 sccm of O₂. When 15 sccm of oxygen is added, C₃HF₄N starts etching silicon nitride and the selectivity to silicon nitride drops to ~3 (*i.e.*, etch rate of SiO₂/Etch rate of SiN) while still maintaining high selectivity of ~40 to a-C.

Example 6

[0098] **FIG. 10** is a graph demonstrating etch rates of SiO₂, SiN, p-Si and a-C with C₃HF₄N and CF₄. In **FIG. 10**, the y-axis represents etch rates and the x-axis is CF₄ flow in sccm. The C₃HF₄N flow rate is fixed at 15 sccm, the O₂ flow rate is fixed at 5 sccm, while the CF₄ flow rate is varied from 10 to 15 sccm.

[0099] In order to enhance the etch rates of SiO₂, comparable to the performance of cC₄F₈ (see comparative examples below), CF₄ is added to the etch gas mixture of 250 sccm Ar and 15 sccm C₃HF₄N. **FIG. 10** is a graph demonstrating that by adding CF₄ to the process gas mixture, the etch rates of SiO₂ have increased up to 500 nm/min while maintaining good selectivity to p-Si and a-C. Additives like C_xF_{2x+2} (x=1 to 5), C_xF_{2x} (x= 3 to 5), C_xF_{2x-2} (x=4 to 5) may be added to the mixture to enhance Oxide etch rate.

Comparative example 1

[00100] **FIG. 11** is a comparative graph demonstrating silicon oxide etch rates of C₃HF₄N, C₂HF₂N, C₅F₅N, C₂F₃N, cC₄F₈, and C₄F₆ etch gases. In **FIG. 11**, the positive y-axis represents etch rates while the negative y-axis represents deposition rates; the x-axis represents the compounds to be compared. **FIG. 11** shows that the nitrogen containing compounds C₂HF₂N and C₂F₃N have higher etch rates than those of C₃HF₄N and C₅F₅N.

Comparative example 2

[00101] **FIG. 12** is a comparative graph demonstrating selectivity of silicon oxide to silicon nitride with C₃HF₄N, C₂HF₂N, C₅F₅N, C₂F₃N, cC₄F₈, and C₄F₆ etch gases with no oxygen addition. In **FIG. 12**, the y-axis represents SiO₂:SiN selectivity; the x-axis represents the compounds compared. **FIG. 12** shows that C₂HF₂N and C₅F₅N have

infinite selectivity (represented as 150) of silicon oxide to silicon nitride with 0 sccm oxygen addition.

Comparative example 3

[00102] **FIG. 13** is a comparative graph demonstrating silicon oxide etch rates of C_3HF_4N , C_2HF_2N , C_5F_5N , C_2F_3N , cC_4F_8 , and C_4F_6 etch gases including a 5 sccm oxygen addition. In **FIG. 13**, the y-axis represents etch rates and the x-axis represents the compounds compared. As shown in **FIG. 13**, the rank of the etching rates with 5 sccm oxygen addition are $cC_4F_8 > C_2F_3N > C_3HF_4N > C_2HF_2N > C_4F_6 > C_5F_5N$.

Comparative example 4

[00103] **FIG. 14** is a comparative graph demonstrating selectivity (SiO/SiN) of C_3HF_4N , C_2HF_2N , C_5F_5N , C_2F_3N , cC_4F_8 , and C_4F_6 etch gases with 5 sccm oxygen addition. In **FIG. 14**, the y-axis represents SiO:SiN selectivity and the x-axis represents the compounds compared. **FIG. 14** shows that C_3HF_4N , C_2HF_2N and C_5F_5N have infinite selectivity of oxide to nitride with 5 sccm oxygen addition represented as 150.

Comparative example 5

[00104] **FIG. 15** is a comparative graph demonstrating silicon oxide etch rates of C_3HF_4N , C_2HF_2N , C_5F_5N , C_2F_3N , cC_4F_8 , and C_4F_6 etch gases with 10 sccm oxygen addition. In **FIG. 15**, the y-axis represents etch rates and the x-axis represents the compounds compared. As shown in **FIG. 15**, the rank of the etching rates with 10 sccm oxygen addition are $cC_4F_8 > C_4F_6 > C_3HF_4N > C_2F_3N > C_2HF_2N > C_5F_5N$.

Comparative example 6

[00105] **FIG. 16** is a comparative graph demonstrating selectivity (SiO/SiN) of C_3HF_4N , C_2HF_2N , C_5F_5N , C_2F_3N , cC_4F_8 , and C_4F_6 etch gases with 10 sccm oxygen addition. In **FIG. 16**, the y-axis represents SiO:SiN selectivity and the x-axis represents the compounds compared. **FIG. 16** shows that C_3HF_4N , C_2HF_2N and C_5F_5N have the infinite selectivity of oxide to nitride with 10 sccm oxygen addition represented as 150.

Comparative example 7

[00106] **FIG. 17** is a comparative graph demonstrating silicon oxide etch rates of C_3HF_4N , C_2HF_2N , C_2F_3N , cC_4F_8 , and C_4F_6 etch gases with 15 sccm oxygen addition. In **FIG. 17**, the y-axis represents etch rates and the x-axis represents the compounds compared. As shown in **FIG. 17**, the rank of the etching rates with 15 sccm oxygen addition are $C_4F_6 > cC_4F_8 > C_3HF_4N > C_2F_3N > C_2HF_2N$.

Comparative example 8

[00107] **FIG. 18** is a comparative graph demonstrating selectivity (SiO/SiN) of C_3HF_4N , C_2HF_2N , C_2F_3N , cC_4F_8 , and C_4F_6 etch gases with 15 sccm oxygen addition. In **FIG. 18**, the y-axis represents etch rates and the x-axis represents the compounds compared. **FIG. 18** shows that only C_2HF_2N has the infinite selectivity of oxide to nitride with 15 sccm oxygen addition represented as 150.

Comparative example 9

[00108] N_2 was added to the etch gas mixture (including cC_4F_8) to see the effect of N_2 on etch rate and selectivity. The etching tests were performed at 30 mTorr, source power of 750 W (27 MHz), and bias power of 1500 W (2 MHz). The feed mixture contained 250 sccm of Ar, 15 sccm of cC_4F_8 , 10 sccm of O_2 , while N_2 has been varied in the amounts of 0 sccm to 20 sccm. **FIG. 19** is a graph demonstrating the effect of adding N_2 on etch rates of different substrate materials. In **FIG. 19**, the y-axis represents etch rates and the x-axis represents the N_2 flow rate.

[00109] As shown, for each substrate (SiO, SiN, p-Si, or a-C), the etch rates varied less than 10% compared to the etch rate when no nitrogen (0 sccm) is added. Thus, nitrogen addition has minimal effect on the etch rates of different substrate materials based on the results shown in **FIG. 19**.

Comparative example 10

[00110] NH_3 was added to the etch gas mixture to see the effect of nitrogen on etch rate and selectivity. The etching tests were performed at 30 mTorr, source power of 750 W (27 MHz), and bias power of 1500 W (2 MHz). The feed mixture contained 250 sccm of Ar, 15 sccm of cC_4F_8 , 15 sccm of NH_3 , while O_2 was varied between 0 sccm to 15 sccm. **FIG. 20** is a graph demonstrating the effect of adding O_2 on etch rates of different substrate materials. In **FIG. 20**, the positive y-axis

represents etch rates while the negative y-axis represents deposition rates; the x-axis represents O₂ flow.

[00111] As shown, infinite selectivity of silicon oxide to silicon nitride is obtained when no oxygen is added. However, this condition results in excess polymerization leading to etch stop.

[00112] As described above in **FIG. 1b**, polymer may be deposited on a sidewall during etching. In order to obtain straight etch profiles that are deep, the amount of polymer deposition should be very well controlled. If there is excessive polymerization, it may result in a condition called etch stop phenomenon. To prevent the etch stop, oxygen is often added to the etch gas mixture. However, excess oxygen may result in loss of selectivity. Thus, there is a trade-off between etch stop and selectivity.

[00113] As shown in **FIG. 20**, as oxygen is added, the selectivity decreases. The selectivity of SiO to other materials is significantly better for nitrogen containing organofluorine compounds than by adding Ammonia to cC₄F₈ and varying oxygen flow rates between 5 sccm to 15 sccm (*cf.* **Figs 8 and 9**). Moreover, the need for another primary organofluorine compound to ammonia in the mixture makes its use difficult for commercial applications.

[00114] Deposition tests were performed on 1 x 1 cm² Si coupon at 30 mTorr, and source power of 750 W (27 MHz), with no bias power. The process feed mixture contained 250 sccm of Ar, 15 sccm of cC₄F₈, and 15 sccm of NH₃. For these conditions, the deposition rate is found to be 180 nm/min.

Comparative Example 11

[00115] **FIG. 21** is a graph demonstrating electron impact ionization data for C₂F₃N. In **FIG. 21**, the x-axis represents electron energy and y-axis represents the partial pressure of the fragment species. **FIG. 21** shows that the major fragments for C₂F₃N are CF₃ and C₂F₂N. The C₂F₂N fragment has a low F/C ratio and will readily polymerize reaching the substrate.

Comparative Example 12

[00116] **FIG. 22** is a graph demonstrating etch rates of SiO, SiN, p-Si and a-C with C₂F₃N and O₂. In **FIG. 22**, the positive y-axis represents etch rates while the

negative y-axis represents deposition rates; the x-axis is O_2 flow in sccm. The C_2F_3N flow is fixed at 15 sccm while O_2 flow is varied from 0 to 15 sccm.

[00117] As shown, when no oxygen is added (0 sccm O_2 condition), C_2F_3N readily etches silicon oxide but etches other substrates at a very slow rate. This results in a very high selectivity of ~40 for silicon oxide to silicon nitride and selectivity of ~30 for silicon oxide to a-C and infinite selectivity to p-Si. However, as the oxygen flow rate increased to 5 sccm and beyond, the selectivity slows drops.

Comparative example 13

[00118] FIG. 23 is a graph demonstrating electron impact ionization data for



$C_3H_3F_6N$ (1,1,1,3,3,3-hexafluoroisopropylamine, $C_3H_3F_6N$). In FIG. 23, the x-axis represents electron energy and y-axis represents the partial pressure of the fragment species. FIG. 23 shows that the major fragments for $C_3H_3F_6N$ is $C_2H_3F_3N$. The $C_2H_3F_3N$ fragment has high F/C ratio and may not provide etch resistance polymers on the substrate.

Comparative example 14

[00119] FIG. 24 is a graph of the results demonstrating etch rates without oxygen addition of $C_3H_3F_6N$ on different substrate materials. For the deposition test conditions, it deposits at 210 nm/min and so may offer sidewall protection. This molecule offers good selectivity to p-Si and a-C but loses selectivity to SiN film even without any oxygen addition.

[00120] In summary, the evaluation of the dry etching of SiO, SiN, p-Si and a-C films with nitrogen containing organofluorine compound plasmas shows that the nitrogen containing HFCs yield highest (up to infinite) selectivity of silicon oxide to silicon nitride and p-Si than the prior art fluorocarbons. The reason for high selectivity may be attributed to the formation of low F/C nitrogen containing fragments during plasma dissociation of the etch gases, which results in the formation of a protective polymer film on the substrates. The polymer film, when analyzed with XPS, shows the evidence of C, F, and N. The resulting film has the formula CF_xN_y , wherein x, y=0.01 to 5. The presence of nitrogen in the polymerizing film offers an extra layer of

protection in comparison to the standard cC_4F_8 gas where only a CF_x type polymer is formed, wherein $x=0.01$ to 5. The etching gas results presented herein show they are not only ready for contact etch process, but also may be beneficial for other etching processes on silicon or metal containing substrates.

[00121] While embodiments of this invention have been shown and described, modifications thereof may be made by one skilled in the art without departing from the spirit or teaching of this invention. The embodiments described herein are exemplary only and not limiting. Many variations and modifications of the composition and method are possible and within the scope of the invention. Accordingly the scope of protection is not limited to the embodiments described herein, but is only limited by the claims which follow, the scope of which shall include all equivalents of the subject matter of the claims.

What is claimed is:

1. A method of depositing an etch-resistant polymer layer on a substrate, the method comprising:

introducing a vapor of an organofluorine compound into a reaction chamber containing a substrate, the organofluorine compound having a formula selected from the group consisting of $\text{N}\equiv\text{C}-\text{R}^1$, wherein R^1 has the formula $\text{H}_a\text{F}_b\text{C}_c$ and $a=1-11$, $b=1-11$, and $c=1-5$; $(\text{N}\equiv\text{C})-(\text{R}^2)-(\text{C}\equiv\text{N})$, wherein R^2 has the formula $\text{H}_a\text{F}_b\text{C}_c$ with $a=0-11$, $b=1-11$, and $c=1-5$; and $\text{R}^1_x[-\text{C}=\text{N}(\text{R}^2_z)]_y$, wherein $x=1-2$, $y=1-2$, $z=0-1$, $x+z=1-3$, and each R^1 and R^2 independently has the formula $\text{H}_a\text{F}_b\text{C}_c$ with $a=0-11$, $b=0-11$, and $c=0-5$; and

plasma activating an organofluorine compound to form the etch-resistant polymer layer on the substrate,

wherein the organofluorine compound having a purity ranging from 95% to 99.999% by volume and comprising between 10 parts per trillion to 5% by volume water vapor impurities.

2. The method of claim 1, wherein the organofluorine compound has the formula: $\text{N}\equiv\text{C}-\text{R}^1$, wherein R^1 has the formula $\text{H}_a\text{F}_b\text{C}_c$ and $a=1-11$, $b=1-11$ and $c=1-5$, or $(\text{N}\equiv\text{C})-(\text{R}^2)-(\text{C}\equiv\text{N})$, wherein R^2 is a functional group having the formula $\text{H}_a\text{F}_b\text{C}_c$ with $a=0-11$, $b=1-11$, and $c=1-5$.

3. The method of claim 1, wherein the organofluorine compound has the formula: $\text{R}^1_x[-\text{C}=\text{N}(\text{R}^2_z)]_y$, wherein $x=1-2$, $y=1-2$, $z=0-1$, $x+z=1-3$, and each R^1 and R^2 independently has the formula $\text{H}_a\text{F}_b\text{C}_c$ with $a=0-11$, $b=0-11$, and $c=0-5$.

4. The method of claim 1, wherein the organofluorine compound is selected from the group consisting of 2,2,3,3-tetrafluoropropionitrile, 2,3,3,3-tetrafluoropropionitrile ($\text{C}_3\text{HF}_4\text{N}$), difluoroacetonitrile ($\text{C}_2\text{HF}_2\text{N}$), Hexafluoroacetone imine ($\text{C}_3\text{HF}_6\text{N}$); 4,4,4-Trifluorocrotono-nitrile, 3,3,3-Trifluoropropionitrile, fluoroacetonitrile, octafluorohexane-1,6-dinitrile, 1,1-bis(trifluoromethyl)-2,2-dicyanoethylene, N,1,1,1,3,3,3-heptafluoro-propanamine, and 1,1,1,6,6,6-Hexafluoro-3-azahex-3-ene.

5. The method of claim 1, wherein the organofluorine compound is 2,3,3,3-tetrafluoropropionitrile (C_3HF_4N).
6. The method of claim 1, wherein the organofluorine compound is difluoroacetonitrile (C_2HF_2N).
7. The method of claim 1, wherein the etch-resistant polymer layer forms a N-containing polymer passivation layer on sidewalls of a pattern etch structure.
8. The method of claim 7, wherein the pattern etch structure has an aspect ratio ranging from 1:1 to 50:1.
9. The method of claim 7, wherein the pattern etch structure has an aspect ratio ranging from 10:1 to 200:1.
10. The method of claim 7, wherein the N-containing polymer passivation layer prevents ions and radicals from etching the sidewalls.
11. The method of claim 7, wherein the etch-resistant polymer layer results in the pattern etch structure having a vertical profile that is straight with no bowing.
12. The method of any one of claims 1 to 11, further comprising removing the etch-resistant polymer layer using dry or wet etch chemistries.
13. The method of any one of claims 1 to 11, further comprising introducing an inert gas into the reaction chamber.
14. The method of claim 13, wherein the inert gas is selected from the group consisting of He, Ar, Xe, Kr, Ne, N_2 , and combinations thereof.
15. A method of depositing an N-containing polymer passivation layer on a substrate, the method comprising:
 - introducing a vapor of an organofluorine compound into a reaction chamber containing a substrate, the organofluorine having a formula selected from the group

consisting of $\text{N}\equiv\text{C}-\text{R}^1$, wherein R^1 has the formula $\text{H}_a\text{F}_b\text{C}_c$ and $a=1-11$, $b=1-11$, and $c=1-5$; $(\text{N}\equiv\text{C})-(\text{R}^2)-(\text{C}\equiv\text{N})$, wherein R^2 has the formula $\text{H}_a\text{F}_b\text{C}_c$ with $a=0-11$, $b=1-11$, and $c=1-5$; and $\text{R}^1_x[-\text{C}=\text{N}(\text{R}^2_z)]_y$, wherein $x=1-2$, $y=1-2$, $z=0-1$, $x+z=1-3$, and each R^1 and R^2 independently has the formula $\text{H}_a\text{F}_b\text{C}_c$ with $a=0-11$, $b=0-11$, and $c=0-5$; and producing fragments of the organofluorine compound by activating a plasma to form the N-containing polymer passivation layer on the substrate, wherein the organofluorine compound having a purity ranging from 95% to 99.999% by volume and comprising between 10 parts per trillion to 5% by volume water vapor impurities.

16. The method of claim 15, wherein the organofluorine compound is selected from the group consisting of 2,2,3,3-tetrafluoropropionitrile, 2,3,3,3-tetrafluoropropionitrile ($\text{C}_3\text{HF}_4\text{N}$), difluoroacetonitrile ($\text{C}_2\text{HF}_2\text{N}$), trifluoroacetonitrile ($\text{C}_2\text{F}_3\text{N}$); Hexafluoroacetone imine ($\text{C}_3\text{HF}_6\text{N}$); 4,4,4-Trifluorocrotono-nitrile, 3,3,3-Trifluoropropionitrile, fluoroacetonitrile, octafluorohexane-1,6-dinitrile, 1,1-bis(trifluoromethyl)-2,2-dicyanoethylene, N,1,1,1,3,3,3-heptafluoro-propanamine, and 1,1,1,6,6,6-Hexafluoro-3-azahex-3-ene.

17. The method of claim 15 or 16, further comprising removing the N-containing polymer passivation layer using dry or wet etch chemistries.

18. The method of claim 15 or 16, further comprising introducing an inert gas into the reaction chamber.

19. The method of claim 18, wherein the inert gas is selected from the group consisting of He, Ar, Xe, Kr, Ne, N_2 , and combinations thereof.

NITROGEN-CONTAINING COMPOUNDS FOR ETCHING SEMICONDUCTOR STRUCTURES

ABSTRACT

A method for etching silicon-containing films is disclosed. The method includes the steps of introducing a vapor of a nitrogen containing etching compound into a reaction chamber containing a silicon-containing film on a substrate, wherein the nitrogen containing etching compound is an organofluorine compound containing at least one $\text{C}\equiv\text{N}$ or $\text{C}=\text{N}$ functional group; introducing an inert gas into the reaction chamber; and activating a plasma to produce an activated nitrogen containing etching compound capable of etching the silicon-containing film from the substrate.

[No Figure]