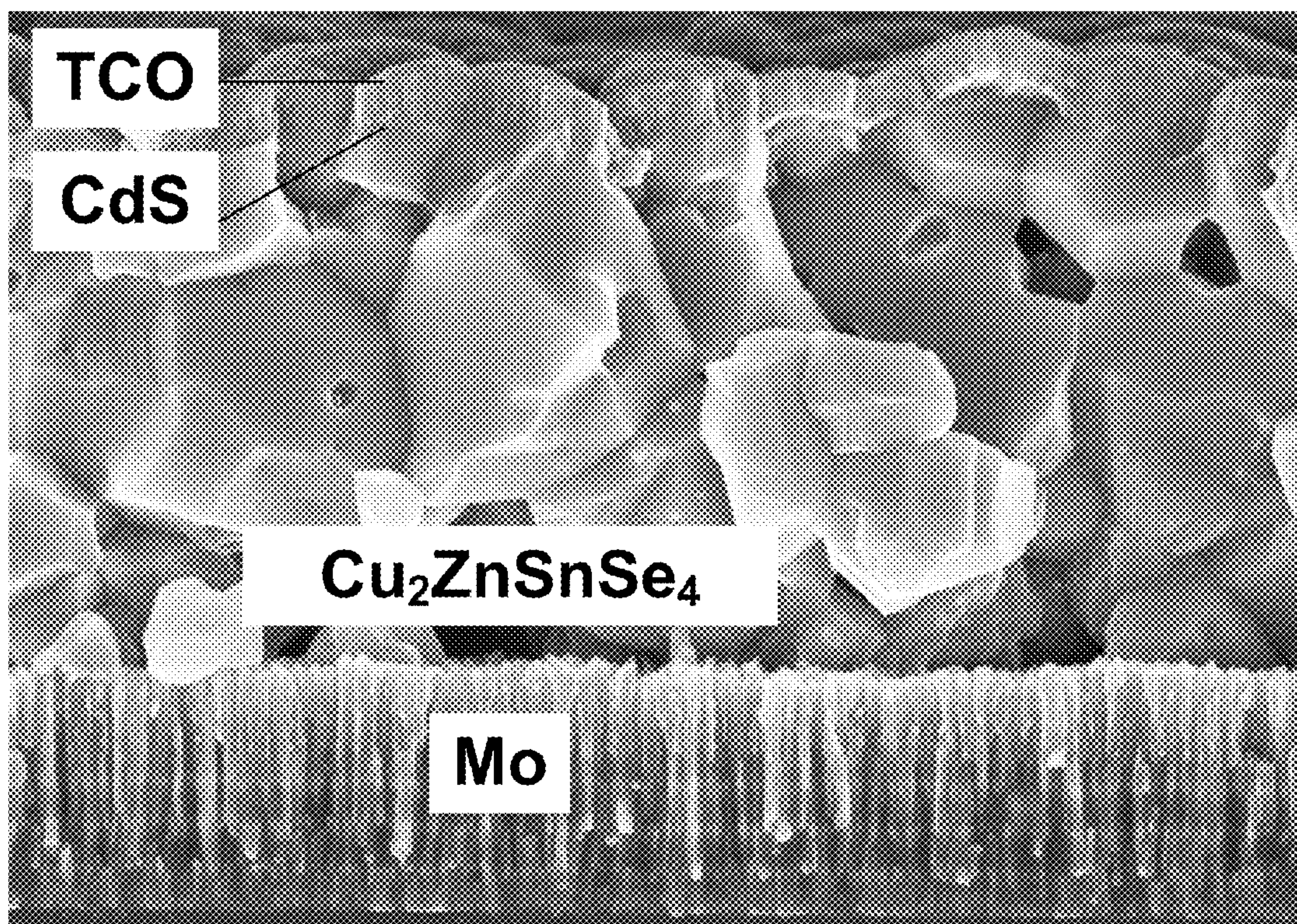


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(19) **United States**(12) **Patent Application Publication**
Mitzi et al.(10) **Pub. No.: US 2011/0094557 A1**(43) **Pub. Date: Apr. 28, 2011**(54) **METHOD OF FORMING SEMICONDUCTOR
FILM AND PHOTOVOLTAIC DEVICE
INCLUDING THE FILM****B05D 5/12** (2006.01)**H01B 1/02** (2006.01)(52) **U.S. Cl. 136/244; 136/252; 427/76; 427/75;
252/519.14**(75) **Inventors:** **David B. Mitzi**, Mahopac, NY
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Armonk, NY (US)(21) **Appl. No.: 12/606,210**(22) **Filed: Oct. 27, 2009****Publication Classification**(51) **Int. Cl.**
H01L 31/042 (2006.01)
H01L 31/00 (2006.01)(57) **ABSTRACT**

A method of depositing a kesterite film which includes a compound of the formula: $\text{Cu}_{2-x}\text{Zn}_{1+y}\text{Sn}(\text{S}_{1-z}\text{Se}_z)_{4+q}$, wherein $0 \leq x \leq 1$; $0 \leq y \leq 1$; $0 \leq z \leq 1$; $-1 \leq q \leq 1$. The method includes contacting hydrazine, a source of Cu, and a source of at least one of S and Se forming solution A; contacting hydrazine, a source of Sn, a source of at least one of S and Se, and a source of Zn forming dispersion B; mixing solution A and dispersion B under conditions sufficient to form a dispersion which includes Zn-containing solid particles; applying the dispersion onto a substrate to form a thin layer of the dispersion on the substrate; and annealing at a temperature, pressure, and length of time sufficient to form the kesterite film. An annealing composition and a photovoltaic device including the kesterite film formed by the above method are also provided.



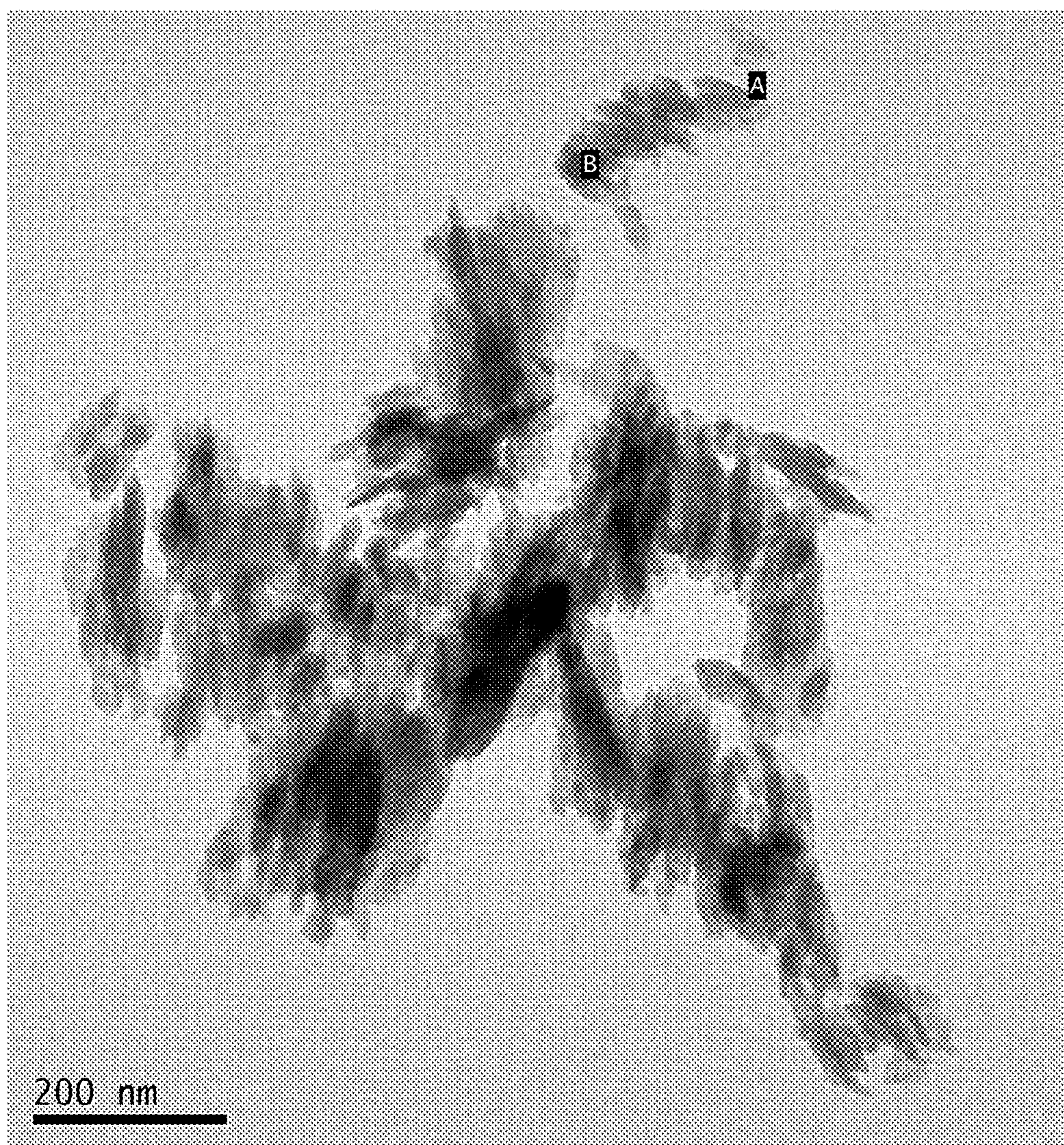


Figure 1

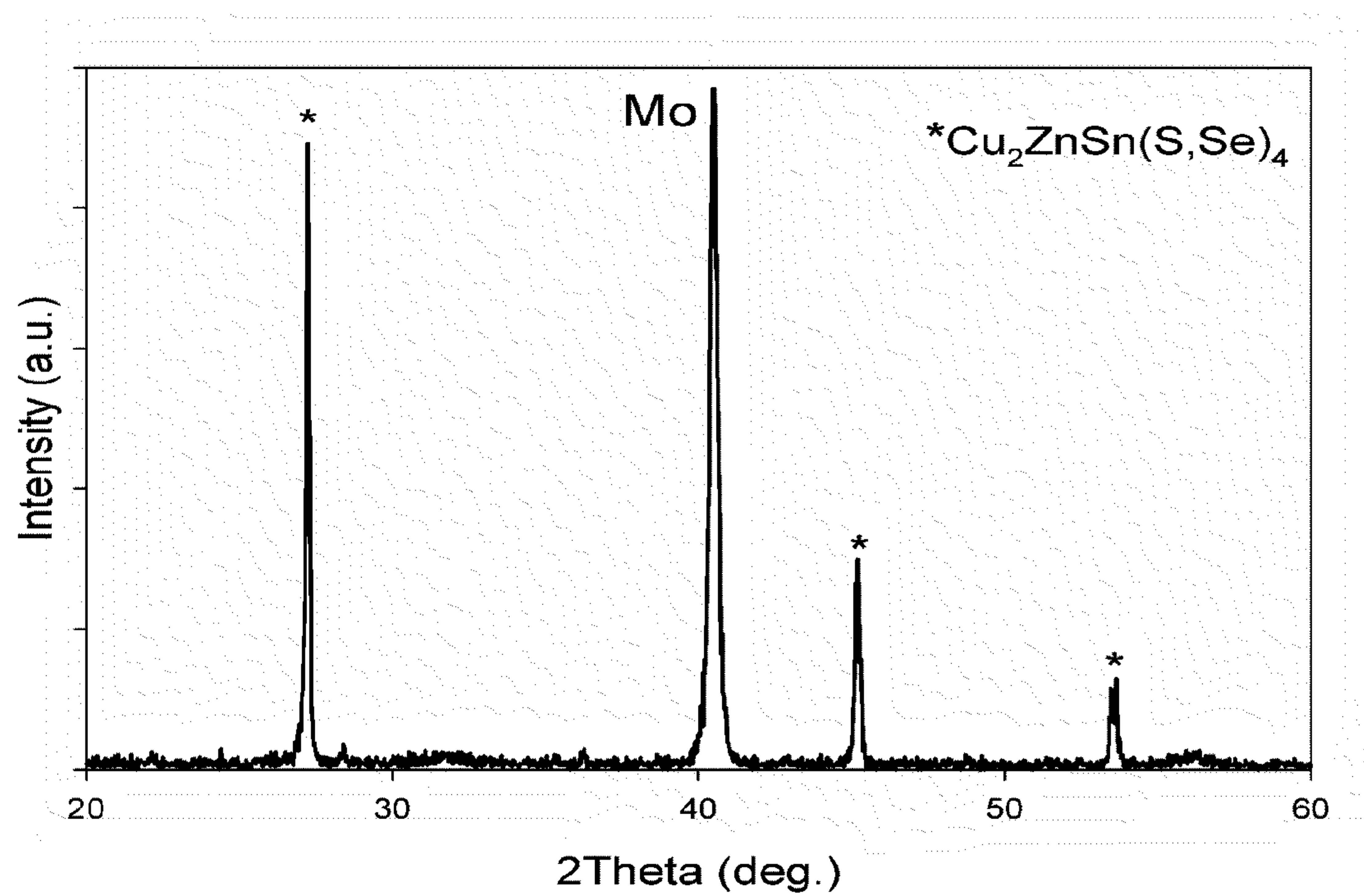


Figure 2

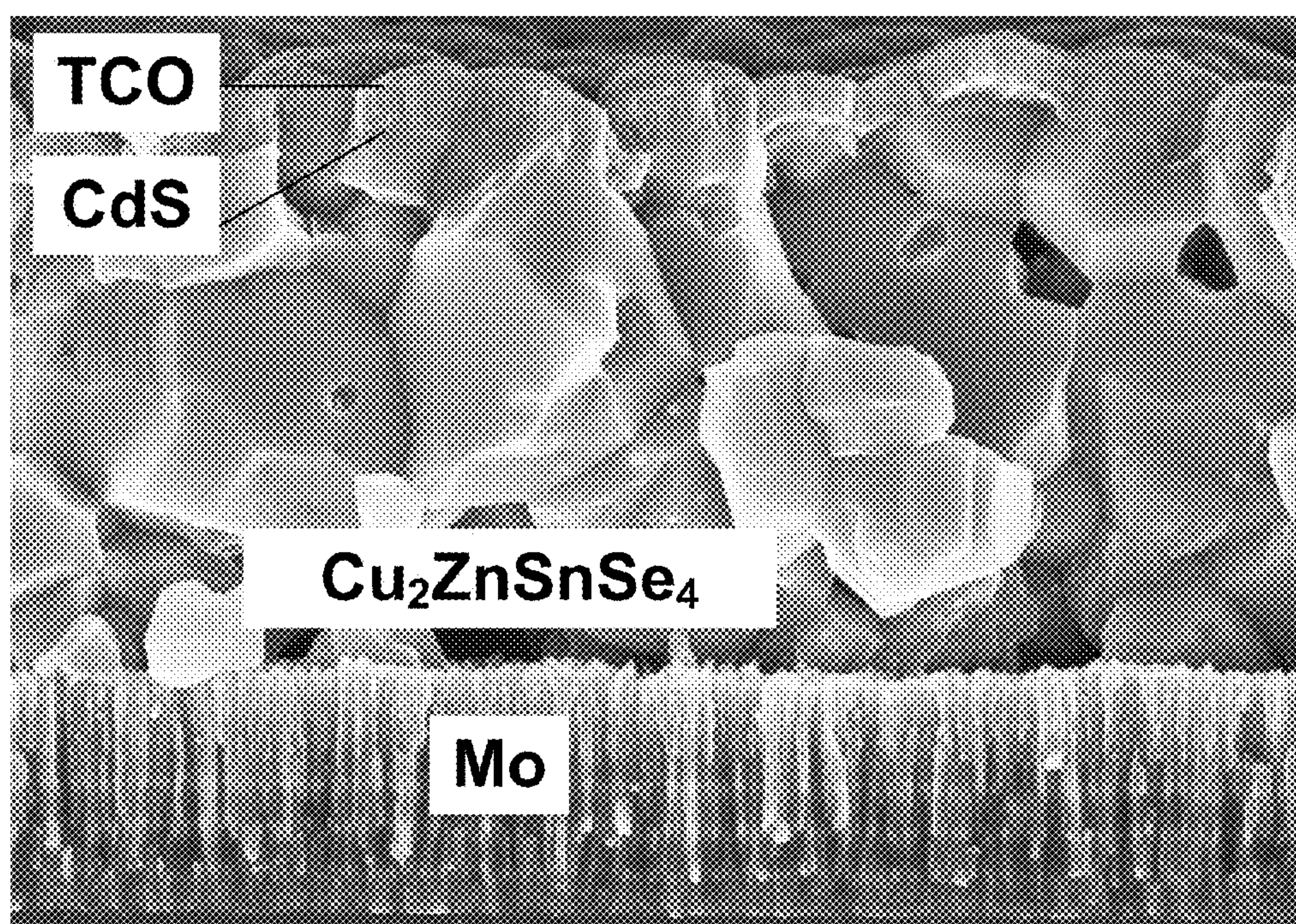


Figure 3

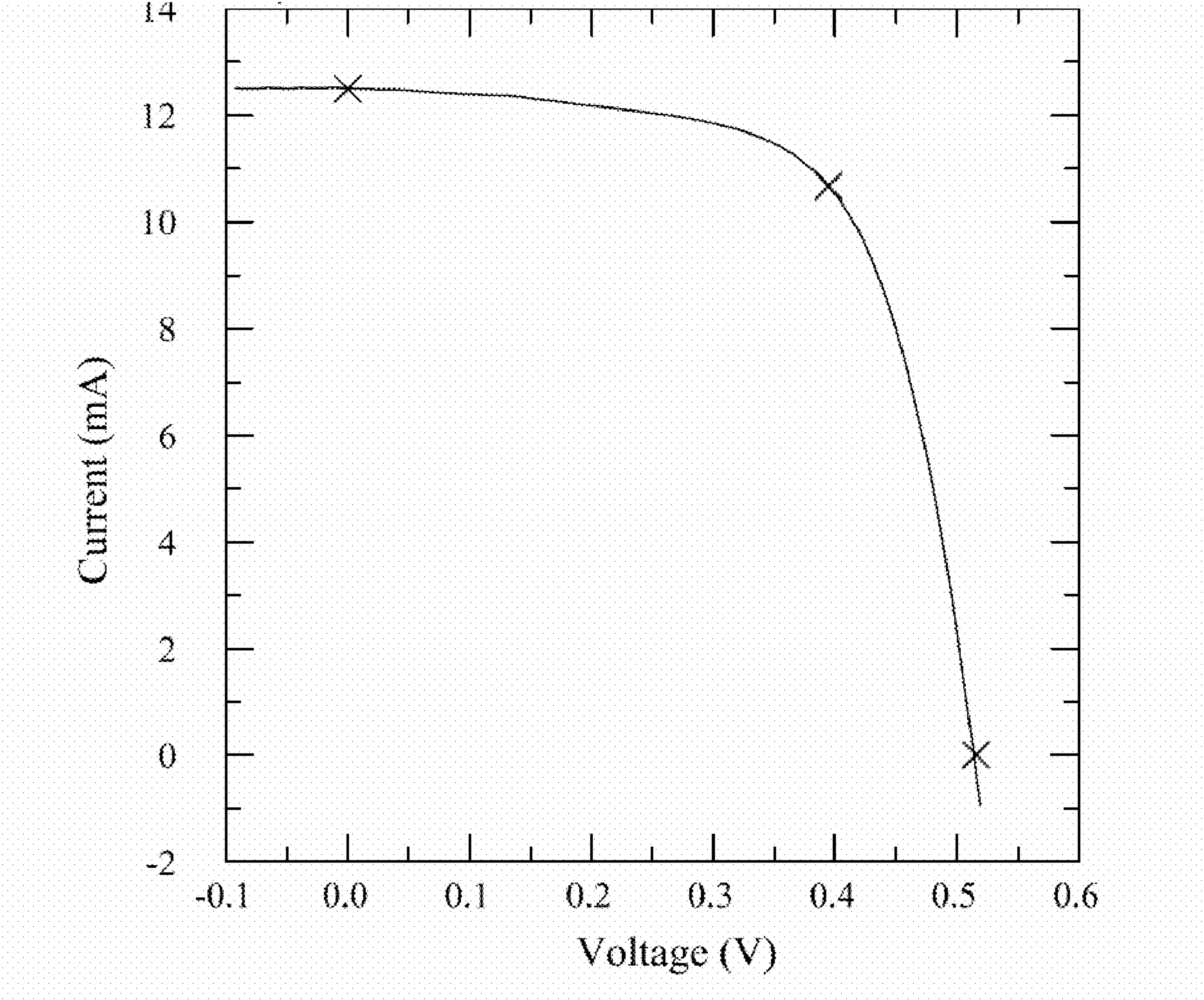


Figure 4

METHOD OF FORMING SEMICONDUCTOR FILM AND PHOTOVOLTAIC DEVICE INCLUDING THE FILM

BACKGROUND OF THE INVENTION

[0001] 1. Field of the Invention

[0002] The present invention relates to a liquid-based method for deposition of inorganic films having Cu, Zn, Sn, and at least one of S and Se, and more particularly to a method of deposition of kesterite-type Cu—Zn—Sn—(Se,S) materials and improved photovoltaic devices based on these films.

[0003] 2. Description of Related Art

[0004] Large-scale production of photovoltaic devices requires high-throughput technologies and abundant environmentally friendly materials. Thin-film chalcogenide-based solar cells provide a promising pathway to cost parity between photovoltaic and conventional energy sources.

[0005] Currently, only Cu(In,Ga)(S,Se)₂ and CdTe technologies have reached commercial production and offer over 10 percent power conversion efficiency. These technologies generally employ (i) indium and tellurium, which are relatively rare elements in the earth's crust, or (ii) cadmium, which is a highly toxic heavy metal.

[0006] Copper-zinc-tin-chalcogenide kesterites have been investigated as potential alternatives because they are based on abundant elements. However, photovoltaic cells with kesterites, even when produced using high cost vacuum-based methods, have so far achieved at best only <6.7 percent efficiencies, see Katagiri, H. et. al. Development of CZTS-based thin film solar cells; Thin Solid Films 517, 2455-2460 (2009).

[0007] K. Tanaka, M. Oonuki, N. Moritake, H. Uchiki, Solar Energy Mater. Sol. Cells 2009, 93, 583-587 describe a solution-based approach for an indium-free material which produced a photovoltaic device with efficiency of only 1%.

[0008] T. Todorov, M. Kita, J. Carda, P. Escrivano, Thin Solid Films 2009, 517, 2541-2544 describe a deposition approach based on quaternary Cu—Zn—Sn—S precursors formed by reacting metal acetates and chlorides with elemental sulfur in ethylene glycol at 170° C.

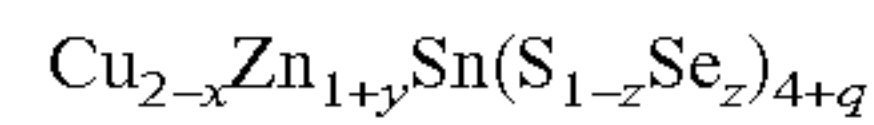
[0009] Guo et. al, J. AM. CHEM. SOC. 2009, 131, 11672-11673 have reported films deposited by a similar approach, subsequently subjected to selenization treatment. They have also reported that devices based on the Cu₂ZnSnS₃Se_{1-y} films yield efficiencies of 0.74%, a level that is lower than the above solution approach for Cu₂ZnSnS₄.

[0010] However, there are no reports of hydrazine-based deposition approaches of depositing homogeneous chalcogenide layers from dispersions of metal chalcogenides in systems that are not strictly soluble in hydrazine. Further, there are no reports to extend the nanoparticle- and microparticle-based approaches to systems without organic binders in a manner that particle-based precursors can readily react with solution component and form large-grained films with good electrical characteristics.

SUMMARY OF THE INVENTION

[0011] The disadvantages associated with the prior art are overcome by embodiments of the present invention by the introduction of hybrid precursor inks including both dissolved and solid components of the targeted material, where the dissolved component acts as a binder, without introducing extraneous elements into the solution that would generally end up as impurities in the final film.

[0012] The present method provides a method of depositing a kesterite film including a compound of the formula:



[0013] wherein $0 \leq x \leq 1$; $0 \leq y \leq 1$; $0 \leq z \leq 1$; $-1 \leq q \leq 1$

[0014] The method includes the steps of contacting hydrazine, a source of Cu, a source of Sn, a source of Zn and a source of at least one of S and Se under conditions sufficient to form a dispersion including Zn-containing solid particles. In a preferred embodiment, the method includes the steps of contacting hydrazine, a source of Cu, and a source of at least one of S and Se forming solution A; contacting hydrazine, a source of Sn, a source of at least one of S and Se, and a source of Zn form dispersion B; mixing solution A and dispersion B under conditions sufficient to form a dispersion including Zn-containing solid particles; applying the dispersion onto a substrate to form a thin layer of the dispersion on the substrate; and annealing at a temperature, pressure, and length of time sufficient to form the kesterite film.

[0015] The present invention further provides a composition, which includes a dispersion of Zn-containing solid particles formed from hydrazine, a source of Cu, a source of Sn, a source of Zn, and a source of at least one of S and Se; which when annealed, forms a compound of the formula: Cu_{2-x}Zn_{1+y}Sn(S_{1-z}Se_z)_{4+q} wherein $0 \leq x \leq 1$; $0 \leq y \leq 1$; $0 \leq z \leq 1$; $-1 \leq q \leq 1$.

[0016] The present invention still further provides a photovoltaic device, including a top electrode having transparent conductive material; an n-type semiconducting layer; a kesterite film on the substrate formed by the above described method; and a substrate having an electrically conductive surface.

[0017] The present invention provides a non-vacuum slurry-based coating method that enables fabrication of Cu₂ZnSn(Se,S)₄ devices with 9.6% percent efficiency, almost an order of magnitude improvement over previous attempts to use high-throughput non-vacuum approaches and sufficient performance enhancement to suggest direct commercial utility for this material system.

[0018] The hybrid deposition process combines advantages of both a recently described hydrazine-based solution approach, enabling the incorporation of select metal chalcogenide anions directly in solution, and particle-based deposition, which allows the incorporation of additional otherwise insoluble components. The described modified slurry method enables the high-throughput low-capital-cost fabrication of thin films of different semiconductor materials, ranging from new chalcogenide-based photovoltaic absorbers to a broader range of materials employed in the optical, electronic and other semiconductor industries.

BRIEF DESCRIPTION OF THE DRAWINGS

[0019] FIG. 1 is a TEM image of solid particles filtered out of the hybrid deposition solution in Example 1.

[0020] FIG. 2 shows an X-ray diffraction pattern of CZTSe film prepared on molybdenum-coated glass substrate according to Example 1.

[0021] FIG. 3 is Scanning electron microscopy image of CZTSe kesterite film of Example 1 in a solar cell prepared according to Example 3.

[0022] FIG. 4 depicts the Current-voltage curve of a CZTSe solar cell, prepared according to Example 3, having $V_{oc}=0.5160$ V, $I_{sc}=12.481$ mA, $J_{sc}=28.612$ mA/cm², Fill Factor=65.43%.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

[0023] The details given in this section are non-limiting and for exemplary purpose only, demonstrating various modes of applying the invention. When components of the invention are defined as containing elements, it must be understood that these elements can be present in either isolated or in compound form, (e.g., a Zn-containing component may contain Zn, ZnS, ZnSe or any other known Zn compound).

[0024] The present invention provides a liquid (ink-based) method for light absorbing semiconductor film formation suitable for ultrahigh throughput production. The ink is based on a hybrid precursor containing at least a dissolved component and at least a solid component (particles) whereas elements from both components are incorporated into the final film composition by a reactive anneal. The dissolved component provides effective binding media surrounding the particles while the particles have stress-relief and crack-deflection effect permitting the deposition of thicker films in a single coating.

[0025] The present invention provides a method for coating a substrate with a Cu—Zn—Sn—(S,Se) based compound film. The method is suitable for ultrahigh-throughput fabrication and overcomes the disadvantages associated with the prior art by:

[0026] Forming a hybrid precursor ink composed of both solid (particle-based) and dissolved precursor component in a suitable liquid media (solvent), where both the solid and dissolved components preferably contain metal atoms and both are aimed to be substantially incorporated into a final nominally single-phase composition;

[0027] Coating and drying the hybrid precursor ink on a substrate to produce a coated substrate. As the solvent(s) evaporate, dissolved components act as binders forming continuous media around the solid particles, which in turn, as non-compressible components have stress-relief and crack-deflecting effect, permitting the formation of a thicker layer in single coating; and

[0028] Heating the coated substrate in a desired atmosphere to induce chemical reaction between the precursor particles and the binding (formerly dissolved) precursor component. The intimate contact between the two phases provides conditions for homogeneous reaction yielding a substantially single-phase film. In specific embodiments, the atmosphere may contain additional reactants, e.g., sulfur or selenium or compounds thereof.

[0029] The obtained film on a substrate can further be used for diverse applications, including improved photovoltaic devices.

[0030] A specific embodiment of the invention provides Cu—Zn—Sn—S(Se) compound films and high-efficiency photovoltaic devices based on these compounds.

[0031] The ink, or functional liquid used to deposit an inorganic, substantially single-phase compound film by a liquid-coating method includes at least one of each 3 following categories:

Solvent, Including a Mixture of Solvents:

[0032] The solvent can be water or non-aqueous liquid, the second being either organic or inorganic liquid. Preferably,

the solvent can be substantially eliminated by evaporation, i.e., it has sufficient vapor pressure below its decomposition temperature and/or upon thermal anneal that it is substantially converted to gas phase and does not leave residual contamination of elements foreign to the targeted composition.

[0033] In the specific embodiment where a non-oxide (e.g., selenide, sulfide, phosphide, nitride, telluride) compound is targeted, particularly when anneals in oxidation atmosphere are undesirable, the solvent and its coordination compounds with the dissolved component preferably do not retain carbon or oxygen, except in the cases when this is strictly desirable.

[0034] In one specific embodiment, the solvent, for example hydrazine, does not contain carbon or oxygen elements. Other examples of solvents include but are not limited to: alcohols, ethers, glycols, aldehydes, ketones, alkanes, amines, dimethylsulfoxide (DMSO), cyclic compounds, halogenated organic compounds;

Dissolved Components:

[0035] The dissolved components preferably include Cu, and Sn in combination with S and/or Se in total concentration of from about 5 to about 1000 mg/ml. Optionally, the dissolved components include an element, selected from the group: Li, Na, K, Mg, Ca, Sr, Ba, Sb, Bi, and B; and

Solid Component:

[0036] The solid components, referred to herein as “particles,” are defined as solid structures of crystalline or amorphous nature are dispersed in the solvent. The solid components are detectable by any analytical technique known to those skilled in the art, and include techniques, such as, but not limited to, x-ray diffraction; optical (laser) scattering; and optical, electron or atomic force microscopy.

[0037] Preferably the particles are in concentration from about 5 to about 1000 mg/ml.

[0038] The particle size preferably is from about 2 nm to about 2000 nm and ranges therebetween. More preferably, the particle size is from about 10 nm to about 1000 nm and most preferably, the particle size is from about 10 nm to about 500 nm.

[0039] In a preferred embodiment, the dimensions of these particles are represented by the formula:

$$d \geq 2e$$

[0040] wherein d is at least one dimension of the particles; and e is any other dimension of the particles.

[0041] The particles may be of various shapes, e.g., elongated, spherical, rod-like, planar, cubic, tetrapod, flake-like etc., and of various sizes, e.g., 2 nm-100 micron, or otherwise they can be nanoparticles and/or microparticles. However, preferably, the particle size is smaller than the thickness of the film being deposited.

[0042] The particles can be prepared by any standard technique known to the skilled in the art, such as, but not limited to, solution-based, e.g., controlled precipitation, sol-gel, wet atomization, gas-phase reactions, optical, e.g., laser ablation, electrical, e.g., electro explosion, plasma jet, electric arc, or mechanical, e.g., grinding, ablation, milling, and water-jet.

[0043] Optionally, the particles can contain an element selected from the group: Li, Na, K, Mg, Ca, Sr, Ba, Sb, and Bi.

[0044] An advantage of the process is the possibility to deposit highly pure semiconductor layers by printing techniques without the necessity to use organic enhancers, such as, polymeric binders known to produce carbon and/or oxy-

gen contamination upon thermal decomposition. A specific embodiment of the invention is a Cu—Zn—Sn—Se—S based film deposited by use of the hybrid ink and an improved photovoltaic cell base on the film.

[0045] Another advantage of the present invention is to avoid or reduce the necessity of enhancing additives, in particular organic polymers acting as binders, surfactants and/or extenders, as their function can be substantially engineered by adequate introduction of desirable dissolved components that are subsequently incorporated into the final composition.

[0046] Nevertheless, in cases where additive use is desirable or in cases where such additives can be conveniently eliminated, e.g., by thermal anneal in oxidizing atmosphere when oxide materials are targeted, these can be readily used. Therefore, in addition to the above 3 principle components the ink may optionally contain enhancing additives that improve the dispersion of the solid phase and/or the solubility of the liquid phase and/or the rheological properties of the ink.

[0047] Some non-limiting examples of such additives include: binders, viscosity modifiers, pH modifiers, dispersants, wetting agents and/or solubility enhancers, such as, polymers, surface active compounds, complex forming agents, e.g., amines, and acidic and basic substances.

[0048] The deposition of the prepared ink on a substrate can be accomplished by forming a liquid layer of the ink by any standard liquid-coating technique, such as, but not limited to, spin coating, dip coating, doctor blading, curtain coating, slide coating, spraying, slit casting, meniscus coating, screen printing, ink jet printing, pad printing, flexography, and gravure printing.

[0049] The substrate may be made of glass, metal, ceramics, polymers, or a combination thereof including composite materials. In one embodiment the substrate is metal or alloy foil containing as non-limiting examples molybdenum, aluminum, titanium, iron, copper, tungsten, steel or combinations thereof. In another embodiment the metal or alloy foil is coated with an ion diffusion barrier and/or an insulating layer succeeded by a conductive layer. In another embodiment the substrate is polymeric foil with a metallic or other conductive layer, e.g., transparent conductive oxide, carbon) deposited on the top of it. In one preferred embodiment, regardless of the nature of the underlying substrate material or materials, the surface contacting the liquid layer contains molybdenum.

[0050] After a liquid layer of the ink is deposited on the surface of the substrate, the solvent is subjected to evaporation by means of exposure to ambient or controlled atmosphere or vacuum that may be accompanied with a thermal treatment, referred to as preliminary anneal, to fabricate substrate coated with a hybrid precursor including discrete particles and surrounding media. This surrounding media is formed by solidification of the dissolved component.

[0051] The substrate thus coated with a hybrid precursor is then subjected to recrystallization or reactive anneal producing a nominally single-phase material. The nominally single-phase material preferably contains at least 80% of the targeted compound, more preferably at least 90% and even more preferably at least 95% by mass of the targeted compound.

[0052] The reactive anneal can be carried out in inert, e.g., nitrogen, helium or argon, atmosphere or reactive atmosphere inducing oxidation, reduction or otherwise introduction or elimination of a specific element into the final composition. The reactive atmosphere may contain as non-limiting examples S and Se. The thermal anneal is carried at tempera-

tures from about 200° C. to about 800° C., preferably from about 400° C. to about 600° C. Most preferably, the anneal temperature is from about 500 to about 600° C.

[0053] The preliminary and/or reactive anneal can be carried out by any technique known to the skilled in the art, including but not limited to: furnace, hot plate, infrared or visible radiation, e.g., laser, lamp furnace, rapid thermal anneal unit, resistive heating of the substrate, heated gas stream, flame burner, electric arc and plasma jet. The duration of this anneal can vary depending on the process and typically is from about 0.1 sec. to about 72 hr.

[0054] The intimate contact between the two components of the hybrid precursor for most embodiments allows to limit the anneal duration to less than 30 min.

[0055] The obtained film on substrate may then be used for the desired application, such as, optical, electrical, anti-friction, bactericidal, catalytic, photo-catalytic, electromagnetic shielding, wear-resistance, and diffusion barrier.

Example 1

Preparation of $\text{Cu}_2\text{ZnSn}(\text{S},\text{Se})_4$ Film

[0056] All operations were performed in nitrogen-filled glove box. The deposition solution was prepared in two parts in glass vials under magnetic stirring: **A1**, by dissolving Cu_2S , 0.573 g and sulfur, 0.232 g in 3 ml of hydrazine and **B1**, by mixing SnSe , 0.790 g, Se, 1.736 g and Zn, 0.32 g with 7 ml of hydrazine. After 3 days under magnetic stirring, solution **A** had an orange transparent aspect, while **B1** was dark green and opaque. Solutions **A1** and **B1** were mixed (**C1**) before deposition.

[0057] A sample of the mixed solution was filtered through a syringe filter and the filtered particles were observed by Transmission Electron Microscopy (FIG. 1). Particles are elongated with dimensions that can be represented by the formula $d \geq 2e$ wherein d is at least one dimension of the particles and e is any other dimension of the particles. For example, where d can be the length and e can be the width. EDX analysis confirmed presence of Zn and Se in the solid particles and the X-ray powder diffraction pattern matched that for ZnSeN_2H_4 , JCPDS 37-0619 (FIG. 2).

[0058] Films were deposited on soda lime glass substrates coated with 700 nm molybdenum by spin coating at 800 rpm and heated at 540° C. for 2 minutes. The coating and heating cycle was repeated 5 times before a final anneal was carried out for 10 minutes.

Example 2

Preparation of $\text{Cu}_2\text{ZnSn}(\text{S},\text{Se})_4$ Film

[0059] Repeating the procedure of Example 1, atmosphere containing elemental sulfur vapor (0.12 g/l N_2) was used for the final anneal.

Example 3

Photovoltaic Devices Prepared by the Method of the Present Invention

[0060] Solar cells were fabricated from the above-described $\text{Cu}_2\text{ZnSn}(\text{Se},\text{S})_4$ films by deposition of 60 nm CdS buffer layer by chemical bath deposition, 100 nm insulating ZnO and 130 nm ITO (indium-doped zinc oxide) by sputter-

ing (FIG. 2). In addition to the shown structure, Ni/Al metal contacts and 110 nm MgF coatings were deposited by electron-beam evaporation.

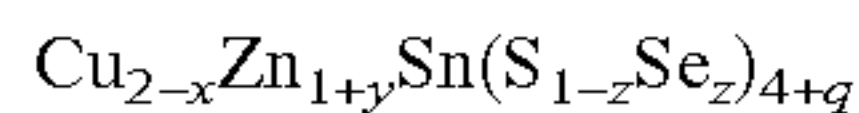
[0061] Photovoltaic performance was measured (NREL CERTIFIED, FIG. 4) under ASTM G173 global spectrum, yielding 9.3% efficiency with films prepared according to Example 1 and 9.66% total area, including metal contacts, i.e., about 5% of the total area, conversion efficiency with films prepared according to example 2, with $V_{oc}=0.5160$ V, $I_{sc}=12.481$ mA, $J_{sc}=28.612$ mA/cm², Fill Factor=65.43% (FIG. 4).

[0062] The potential advantage of this process is that it would be much lower cost than the traditional approaches. This performance is a world record for the category of indium and cadmium-free photovoltaic absorbers deposited by any method, including vacuum-based and makes this material already eligible for commercial consideration.

[0063] The present invention has been described with particular reference to the preferred embodiments. It should be understood that variations and modifications thereof can be devised by those skilled in the art without departing from the spirit and scope of the present invention. Accordingly, the present invention embraces all such alternatives, modifications and variations that fall within the scope of the appended claims.

What is claimed is:

1. A method of depositing a kesterite film comprising a compound of the formula:



wherein $0 \leq x \leq 1$; $0 \leq y \leq 1$; $0 \leq z \leq 1$; $-1 \leq q \leq 1$

said method comprising the steps of:

contacting hydrazine, a source of Cu, a source of Sn, a source of at least one of S and Se, and a source of Zn forming a dispersion comprising Zn-containing solid particles;
applying said dispersion onto a substrate to form a thin layer; and
annealing at a temperature, pressure, and length of time sufficient to form said kesterite film.

2. The method of claim 1, wherein said solution further comprises at least one of a solvent, an amine, a polymer, and an additive.

3. The method of claim 1, wherein x, y, z and q respectively are: $0 \leq x \leq 0.5$; $0 \leq y \leq 0.5$; $0 \leq z \leq 1$; $-0.5 \leq q \leq 0.5$.

4. The method of claim 1, wherein said source of Cu is at least one of Cu₂S and Cu₂Se.

5. The method of claim 1, wherein said source of Sn is at least one of SnSe, SnS, SnSe₂, and SnS₂.

6. The method of claim 1, wherein said source of Zn is at least one of metallic zinc, ZnS, and ZnSe.

7. The method of claim 1, wherein said source of Zn comprises particles having a particle size from about 2 nm to about 2000 nm and ranges therebetween.

8. The method of claim 7, wherein the dimension of said particles are represented by the formula:

$$d \geq 2e$$

wherein d is at least one dimension of said particles; and e is any other dimension of said particles.

9. The method of claim 1, wherein said source of S is selected from the group consisting of: elemental sulfur, CuS, Cu₂S, SnS, SnS₂, ZnS, and a mixture thereof.

10. The method of claim 1, wherein said source of Se is selected from the group consisting of: elemental selenium, SnSe₂, and SnSe.

11. The method of claim 1, wherein said substrate is selected from the group consisting of: metal foil, glass, ceramics, aluminum foil coated with a layer of molybdenum, a polymer, and a combination thereof.

12. The method of claim 1, wherein said substrate is coated with a transparent conductive coating.

13. The method of claim 1, wherein the step of applying is carried out by a method selected from the group consisting of: spin coating, dip coating, doctor blading, curtain coating, slide coating, spraying, slit casting, meniscus coating, screen printing, ink jet printing, pad printing, flexographic printing, and gravure printing.

14. The method of claim 1, wherein the step of annealing is carried out at a temperature from about 200° C. to about 800° C. and ranges therebetween.

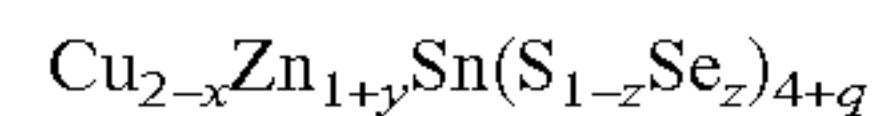
15. The method of claim 1, wherein said anneal is carried out in an atmosphere comprising:

at least one of: N₂, Ar, He, forming gas, and a mixture thereof; and optionally
vapors of at least one of: sulfur, selenium, and a compound thereof.

16. The method of claim 2, wherein said additive contains a metal, or compound thereof, wherein said metal is selected from the group consisting of: Li, Na, K, Mg, Ca, Sr, Ba, Sb, and Bi, in amount from about 0.01 to about 5 mass %.

17. The method of claim 1, wherein the concentration of each of said sources is from about 5 to about 1000 mg/ml.

18. A method of depositing a kesterite film comprising a compound of the formula:



wherein $0 \leq x \leq 1$; $0 \leq y \leq 1$; $0 \leq z \leq 1$; $-1 \leq q \leq 1$

said method comprising the steps of:

contacting hydrazine, a source of Cu, and a source of at least one of S and Se forming solution A;
contacting hydrazine, a source of Sn, a source of at least one of S and Se, and a source of Zn forming dispersion B;

mixing said solution A and said dispersion B under conditions sufficient to form a dispersion comprising Zn-containing solid particles;

applying said dispersion onto a substrate to form a thin layer; and

annealing at a temperature, pressure, and length of time sufficient to form said kesterite film.

19. A composition, comprising:

a dispersion of Zn-containing solid particles formed from hydrazine, a source of Cu, a source of Sn, a source of Zn, and a source of at least one of S and Se; which when annealed, forms a compound of the formula: $Cu_{2-x}Zn_{1+y}Sn(S_{1-z}Se_z)_{4+q}$ wherein $0 \leq x \leq 1$; $0 \leq y \leq 1$; $0 \leq z \leq 1$; $-1 \leq q \leq 1$.

20. The composition of claim 19, wherein said solution further comprises at least one of a solvent, an amine, a polymer, and an additive.

21. The composition of claim 19, wherein x, y, z and q respectively are: $0 \leq x \leq 0.5$; $0 \leq y \leq 0.5$; $0 \leq z \leq 1$; $-0.5 \leq q \leq 0.5$.

22. The composition of claim 19, wherein said source of Zn comprises particles having a particle size from about 2 nm to about 2000 nm and ranges therebetween.

23. The composition of claim **22**, wherein the dimension of said particles are represented by the formula:

$$d \geq 2e$$

wherein d is at least one dimension of said particles; and e is any other dimension.

24. The composition claim **18**, wherein the concentration of each of said sources is from about 5 to about 1000 mg/ml.

25. A photovoltaic device, comprising:

a top electrode having transparent conductive material;
an n-type semiconducting layer;
a kesterite film on said substrate formed by the method in claim **1**; and
a substrate having an electrically conductive surface.

26. A photovoltaic device of claim **25**, wherein:

said substrate is coated with a molybdenum layer and is selected from the group consisting of: glass, plastic, polymer, ceramic, and aluminum foil;

said n-type semiconducting layer has at least one of: ZnS, CdS, InS, oxides thereof, and selenides thereof;

said transparent conductive material is selected from the group consisting of: doped ZnO, Indium-tin oxide, doped tin oxide, and carbon nanotubes.

27. The photovoltaic device of claim **25**, comprising: a plurality of electrically interconnected photovoltaic devices forming a photovoltaic module.

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