Controlling Growth High Uniformity Indium Selenide (In$_2$Se$_3$) Nanowires via the Rapid Thermal Annealing Process at Low Temperature

Ya-Chu Hsu, Yu-Chen Hung and Chiu-Yen Wang *

Abstract
High uniformity Au-catalyzed indium selenide (In$_2$Se$_3$) nanowires are grown with the rapid thermal annealing (RTA) treatment via the vapor-liquid-solid (VLS) mechanism. The diameters of Au-catalyzed In$_2$Se$_3$ nanowires could be controlled with varied thicknesses of Au films, and the uniformity of nanowires is improved via a fast pre-annealing rate, 100 °C/s. Comparing with the slower heating rate, 0.1 °C/s, the average diameters and distributions (standard deviation, SD) of In$_2$Se$_3$ nanowires with and without the RTA process are 97.14 ± 22.95 nm (23.63%) and 119.06 ± 48.75 nm (40.95%), respectively. The in situ annealing TEM is used to study the effect of heating rate on the formation of Au nanoparticles from the as-deposited Au film. The results demonstrate that the average diameters and distributions of Au nanoparticles with and without the RTA process are 19.84 ± 5.96 nm (30.00%) and about 22.06 ± 9.00 nm (40.80%), respectively. It proves that the diameter size, distribution, and uniformity of Au-catalyzed In$_2$Se$_3$ nanowires are reduced and improved via the RTA pre-treated. The systemic study could help to control the size distribution of other nanomaterials through tuning the annealing rate, temperatures of precursor, and growth substrate to control the size distribution of other nanomaterials.

Keywords: In$_2$Se$_3$, Nanowire, Rapid thermal annealing (RTA), In situ annealing TEM

Background
In the past decade, one-dimensional (1D) nanostructure tubes, wires, rods, and belts have become the focal point of the worldwide research in nanotechnology due to their high performance and surface-to-volume ratios, intrinsically associated with low dimensionality, which may lead to unique applications in the various nanoscale device [1, 2]. In particular, 1D semiconductor nanowires (NWs), exhibiting different properties as compared with their bulk or thin film, have shown great potential applications in data storage, computing, and sensing devices [2–4].

Indium selenide (In$_2$Se$_3$) is a black crystalline and very interesting compound semiconductor of the A$^{III}$B$^{VI}$ group with layered structure, which possessed at least five crystal modifications of α (two-layer hexagonal, 2H), β (three-layer rhombohedral, 3R), γ (defect wurtzite in hexagonal, H), δ, and κ [5, 6]. Due to its polymorphism and the related metal-ion defect structure, In$_2$Se$_3$ has attracted substantial attention as a promising semiconductor material for several different applications such as photovoltaic solar cell [7, 8], optoelectronics [9], and ionic battery [10].

The layered structure of In$_2$Se$_3$ is normally consisting of [Se-In-Se-In-Se] sheets stacked with Se atoms along the c-axis [11–15]. The strong intralayer bonding and weak interlayer Van der Waals interaction lead to highly anisotropic structural, electrical, optical, and mechanical properties [16, 17]. Layer-structure In$_2$Se$_3$ nanowires and nanoribbons have been synthesized by using metal nanoparticles as the catalyst via the vapor-liquid-solid (VLS) process [2, 18–20]. The properties of NWs depend not only on their shape anisotropy but also on their crystallographic anisotropy [21]. The vapor-liquid-solid growth mechanism has been demonstrated to control the diameter and growth direction of nanowires [20–24]. Several research results demonstrated that the catalyst is one of an important part to control the morphology of the nanowires. And the crystallographic orientation of a NW

* Correspondence: ChiuYWang@mail.ntust.edu.tw
Department of Material Science and Engineering, National Taiwan University of Science and Technology, Taipei 10607, Taiwan

© The Author(s). 2017 Open Access This article is distributed under the terms of the Creative Commons Attribution 4.0 International License (http://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made.
is thermodynamically determined at the liquid-solid (LS) interface within the eutectic liquid droplet of a given size and geometry during the initial nucleation [25, 26]. Also, previous studies have shown that the synthesis of highly uniform semiconductor NWs can be achieved through using the well-defined nanoclusters as catalysts in a VLS growth process [25]. Controlling the growth temperature of the Au-In alloy droplet catalyst can determine the segregation concentrations of In and Se atoms in the Au-In alloy droplet which then affects the diameter of the nanowires. However, Au-catalyzed In$_2$Se$_3$ nanowires are usually grown at relatively high temperature, > 650 °C. According to the Au-In phase diagram, the eutectic temperature is about 530 °C, and the In and/or Se will be precipitated from the Au-In liquid alloy, then reacted with Se to grow the In$_2$Se$_3$ NWs [27]. In this work, rapid thermal annealing (RTA) is used to make the Au film transfer to uniform Au nanoparticles. Furthermore, lower precursor and growth temperature are chosen to reduce the diameter of nanowires and prevent the VS-grown In$_2$Se$_3$ byproducts. Interestingly, the thinner In$_2$Se$_3$ NWs can obtain the diameter by controlling the growth temperature as low as 550 °C. The in situ annealing transmission electron microscopy (TEM) is used to study the effect of heating rate on the Au nanoparticle formation from the as-deposited Au film.

**Experimental**

The In$_2$Se$_3$ NWs were synthesized in a quartz-tube furnace system with a two-temperature zone. Traditionally, the In$_2$Se$_3$ powder (99.9%, CERAC) was used as a precursor then placed upstream in the middle of the tube at 800 °C (heating rate is 0.01 °C/s). The SiO$_2$/Si(100) substrate is coated with a 2.0 nm thick gold film which was placed downstream. The SiO$_2$/Si(100) coated with a 2.0-nm-thick gold film is annealed by RTA, at 550 °C (heating rate is 100 °C/s), then the substrate is loaded into the growth furnace tube to grow In$_2$Se$_3$ nanowires with a flow rate of argon gas at 25 sccm and pressure of 1 Torr. The temperatures of the In$_2$Se$_3$ precursor powder at the upstream and the Au-coated substrate at the downstream (growth zone) were ramped up to 800 °C (1.2 °C/s) and 550 °C (0.1 °C/s), respectively, and kept for 30 min. The morphologies and microstructure of the In$_2$Se$_3$ NWs were characterized by scanning electron microscope (SEM, JEOL JSM-6500F) and transmission electron microscopy (TEM, FEI Tecnai™ G2 F20 Field Emission Gun) operating at 200 kV. The chemical composition confirmed by energy dispersive X-Ray spectrometer (EDS) is equipped in TEM. The phase of the In$_2$Se$_3$ NWs is confirmed with an X-ray diffractometer (XRD, D8 DISCOVER SSS Multi-Function High Power). In situ annealing TEM was used to study the effect of heating on Au nanoparticle formation. To prepare in situ heating TEM samples, a 2.0 nm Au film is deposited on a square opening of SiO$_2$/Si$_3$N$_4$ thin film. The thicknesses of the SiO$_2$ and Si$_3$N$_4$ film are 30 and 60 nm, respectively. The 2.0 nm Au film is deposited on the SiO$_2$ side, then loaded into TEM to heat with a heating holder (Gatan 652 double tilt heating holder) in the TEM.

![Fig. 1](image_url) **Fig. 1** a Illustration of a two-zone quartz-tube furnace. The In$_2$Se$_3$ powder was used as a precursor and placed upstream in the middle of the tube at 800 °C, and SiO$_2$/Si(100) coated with a 2.0 nm thick Au film was placed downstream and the argon gas as the carrier gas. b and c are the SEM images of In$_2$Se$_3$ nanowires which were grown on the substrate with and without the RTA process, respectively. d A typical XRD spectrum of the Au-catalyzed α-In$_2$Se$_3$ NWs. The lattice constants are $a = 4.025$ Å and $c = 19.235$ Å (JCPDS card, No. 34–1279).
Results and Discussion

Figure 1a is the schematic illustration of the quartz-tube furnace system that was used to grow the In$_2$Se$_3$ NWs. Typically, the growth window of Au-catalyzed In$_2$Se$_3$ NWs is 650–750 °C and the precursor In$_2$Se$_3$ is heated at 900–950 °C to provide the source of In and Se via a VLS mechanism [19]. However, the Au-In phase diagram shows that the eutectic temperature of Au-In could be as low as 450–550 °C, depending on the composition of the AuIn$_x$ alloy [28, 29]. It is expected that the diameters of NWs could be controlled by the Au thickness, growth temperature, and the ambient of the furnace. In this work, the temperatures of the growth temperature and In$_2$Se$_3$ precursor powder are set as 550 and 800 °C, respectively. Figure 1b, c are the SEM images of the In$_2$Se$_3$ NWs, grown on the 2.0 nm Au film deposited on a 200-nm SiO$_2$/silicon wafer, with and without rapid thermal annealing (RTA) process, respectively. The bright nanoparticle on the top of the NW can be observed from the inset in Fig. 1b, c, which indicates that the In$_2$Se$_3$ NWs are grown through AuIn$_x$ nanoparticles via the VLS mechanism. The average diameters of In$_2$Se$_3$ NWs (50 nanowires) with and without the RTA process are 97.14 ± 22.95 nm (23.63%) and 119.06 ± 48.75 nm (40.95%), respectively. The average and distribution of the In$_2$Se$_3$ NW diameters with and without the RTA process are conspicuously different. It clearly exhibits that the RTA process could improve the uniformity and may reduce the diameter of In$_2$Se$_3$ NWs [30–32]. Figure 1d is the XRD result of In$_2$Se$_3$ NWs, and all the peaks can be indexed to the hexagonal crystal structure of α-In$_2$Se$_3$ NWs, in which the lattice constants are $a = 4.025$ Å and $c = 19.235$ Å (JCPDS card, no# 34–1279).

Generally, the Au film-coated substrate is loaded into the furnace, the heating rate usually is 1–2 °C/s, then reacts with the precursor to form a low melting point AuIn$_x$ alloy, and the In is segregated as the eutectic alloy is supersaturated to react with Se and grow the In$_2$Se$_3$ NW. The slower heating rate results to poor Au nanoparticle uniformity. Not only the thickness and heating rate of Au film on the substrate but the growth rate of In$_2$Se$_3$ NWs is controlled by the VLS mechanism.

![Fig. 2](image_url) SEM images of In$_2$Se$_3$ nanowires which were grown at a 550 °C, b 600 °C, and c 650 °C, respectively; the scale bars of the inset images (a–c) are 100 nm. d In$_2$Se$_3$ nanowires are grown with the precursor and growth temperature at 850 and 600 °C, respectively.

Table 1 Comparison with other works in terms: growth temperature, precursor temperature, and nanowire diameters

| Growth temperature (°C) | Diameter (nm) | RTA | Precursor | Precursor temperature (°C) | Reference |
|-------------------------|--------------|-----|-----------|---------------------------|-----------|
| 650–700                 | 40–80        | x   | In$_2$Se$_3$ powder | 900–950       | [2]       |
| 690                     | 80–200       | x   | In$_2$Se$_3$ powder | 940           | [5]       |
| 690–740                 | 150          |     | In$_2$Se$_3$ powder | 920           | [26]      |
| 690–740                 | 50–200       |     | In$_2$Se$_3$ powder | 920           | [33]      |
| 690–740                 | 50–200       | x   | In$_2$Se$_3$ powder | 930–950       | [34]      |
| 550                     | 70–150       |     | In$_2$Se$_3$ powder | 800           | in this work |
| 550                     | 80–170       | x   | In$_2$Se$_3$ powder | 800           | in this work |
temperature is also an important factor to control the morphology of nanowires. Figure 2a–c are the SEM images of the In$_2$Se$_3$ NWs after being RTA-treated then grown at 550, 600, and 650 °C, respectively. The corresponding inset images in Fig. 2a–c showed that the In$_2$Se$_3$ NW diameters were 80–100, 100–200, and 300–500 nm, respectively. The results display that the diameter of In$_2$Se$_3$ NWs could be tuned by controlling the growth temperature. Since the growth temperature was raised, the In solubility in the Au catalyst would be increased; that means the In atoms need more amounts to reach the supersaturated concentration. In the same time, the thicker In$_2$Se$_3$ NWs will be grown through the bigger AuIn$_x$ droplets. Figure 2d shows the In$_2$Se$_3$ nanowires grown with the precursor temperature at 850 °C (1.3 °C/s). Both of the Au-catalyzed vapor-liquid-solid growth and self-catalyzed vapor-solid (VS) growth In$_2$Se$_3$ nanomaterials, including nanowires, nanoplates, and film, will be obtained simultaneously. The higher precursor temperature will lead to the higher precursor vapor, and the excessive precursor will lead to the In$_2$Se$_3$ product, which tends to self-nucleate and grow. Compared to other studies, the growth temperature, 550 °C, could be much lower than the general reported, 650–750 °C. Furthermore, the precursor temperature could be reduced to 800 °C to prevent self-catalyzed growth. Table 1 lists the comparison of growing In$_2$Se$_3$ nanowire parameters, including growth substrate temperature (heating rate), growth substrate annealing treatment, precursor temperature, and diameter of nanowires. Due to the lower growth temperature, the byproduct is inhibited such that uniform In$_2$Se$_3$ NWs could be obtained at a relatively low temperature. It clearly displays that the In$_2$Se$_3$ NWs could be grown at the lowest growth temperature and precursor temperature in this work. Furthermore, the results of the RTA process showed better diameter uniformity for the In$_2$Se$_3$ NWs than the conventional system, since the diameter of gold particles were confined.

In situ annealing TEM is used to study the effect of heating rate on Au nanoparticle formation and nanowire growth. Figure 3a is the TEM image of the as-deposited 2-nm Au film on the SiO$_2$/Si$_3$N$_4$ window, annealing with 0.1 °C/s and 100 °C/s to 550 °C and holding for 30 min.

Fig. 3  a  2.0 nm  Au film at room temperature.  b  The gold film by RTA at 550 °C in 100 °C/s.  c  The gold film was ramped up to the 550 °C in 0.1 °C/s.  d  TEM image of an as-synthesized individual α-In$_2$Se$_3$ nanowire, with an Au nanoparticle tip. SAED pattern of the α-In$_2$Se$_3$ nanowires (inset).  e  The corresponding HRTEM image of d shows the growth direction of the nanowire is along the [001].  f  and g  are the EDS spectra of the selected α-In$_2$Se$_3$ nanowire taken from the body part and the tip part, respectively.
The uniformity of the \( \text{In}_2\text{Se}_3 \) nanowires could be analyzed by the VLS mechanism. Further, the lower precursor and growth temperatures 800 and 550 °C, respectively, are provided to grow the Au-catalyzed self-catalyzed \( \text{In}_2\text{Se}_3 \) nanoplates. Lower temperature will lead to lower precursor concentration and low energy, and then the nucleation of self-catalyzed \( \text{In}_2\text{Se}_3 \) nanowires could be inhibited the \( \text{In}_2\text{Se}_3 \) nanoplate byproduct.

**Table 2** The average particle size and standard deviation (SD) of Au nanoparticles annealed with RTA (100 °C/s) and without RTA (0.1 °C/s) through the in situ annealing TEM

| Annealing condition | Average particle size (nm) | SD (nm) | SD (%) |
|---------------------|---------------------------|---------|--------|
| 550° with RTA (100 °C/s) | 19.84 | 5.96 | 30.00 |
| 550° without RTA (0.1 °C/s) | 22.06 | 9.00 | 40.80 |

Figure 3b, c are the results of Au nanoparticle formation with the heating rate at 100 °C/s and 0.1 °C/s, respectively. According to the in situ annealing TEM result, Au nanoparticle average size and distribution are analyzed and listed in Table 2. Briefly, the smaller average size and better uniformity of Au nanoparticles could be achieved through the faster heating rate. Figure 3d is the TEM image of a representative \( \text{In}_2\text{Se}_3 \) nanowire after being RTA-treated then grown at 550 °C. The result shows that the typical diameter of the nanowire is about 100 nm, and the inset is the corresponding select area electron diffraction (SAED) pattern. Figure 3e shows the high-resolution transmission electron microscopy (HRTEM) image of the corresponding \( \text{In}_2\text{Se}_3 \) NW that was taken from the [010] zone axis which has lattice spacing of 0.35 and 0.48 nm and can be indexed to the \( d \)-spacing of the (100) and (004) planes, which demonstrates that the \( \text{In}_2\text{Se}_3 \) NW is growing along the [001] direction. The EDS analyses are taken from the top and stem; the results are shown in Fig. 3f, g. The Cu and C signals are contributed from the carbon-coated copper TEM grid. Figure 3f which is taken from the stem is composed of In and Se only, and the atomic ratio of In/Se is approximately 2/3. Figure 3g is the EDS result of the top nanoparticle compositions, including In and Au. The additional Au signal proved that the \( \text{In}_2\text{Se}_3 \) nanowires are grown via the vapor-liquid-solid (VLS) mechanism. According to the TEM analyses, SEAD, and HRTEM, the VLS-grown nanowires can be identified as \( \alpha \)-phase of \( \text{In}_2\text{Se}_3 \).

**Conclusions**

The lower precursor and growth temperatures 800 and 550 °C, respectively, are provided to grow the Au-catalyzed \( \text{In}_2\text{Se}_3 \) nanowires by the VLS mechanism. Furthermore, the uniformity of the \( \text{In}_2\text{Se}_3 \) nanowires could be improved by RTA treatment to reduce the size and distribution of Au nanoparticles. The in situ annealing TEM is used to study the effect of heating rate on Au film transfer to Au nanoparticle. The lower precursor and growth temperatures could reduce the formation of self-catalyzed \( \text{In}_2\text{Se}_3 \) nanoplates. Lower temperature will lead to lower precursor concentration and low energy, and then the nucleation of self-catalyzed \( \text{In}_2\text{Se}_3 \) nanowires could be inhibited the \( \text{In}_2\text{Se}_3 \) nanopllate byproduct.
19. Peng H, Xie C, Schoen DT, Cui Y (2008) Large anisotropy of electrical properties in layer-structured In$_2$Se$_3$ nanowires. Nano Lett 8:1511
20. Lai K, Peng H, Kundhikanjana W, Schoen DT, Xie C, Meister S, Cui Y, Kelly MA, Shen ZX (2009) Nanoscale electronic inhomogeneity in In$_2$Se$_3$ nanoribbons revealed by microwave impedance microscopy. Nano Lett 9:1265
21. Kuykendall T, Pauzauskie PJ, Zhang Y, Goldberger J, Sirbuly D, Denlinger J, Yang P (2004) Crystallographic alignment of high-density gallium nitride nanowire arrays. Nat Mater 3:524
22. Lieber CM (2003) Nanoscale science and technology: building a big future from small things. MRS Bull 28:486
23. Yang P (2005) The chemistry and physics of semiconductor nanowires. MRS Bull 30:85
24. Wacaser BA, Dick KA, Johansson J, Borgström MT, Deppert K, Samuelson L (2009) Preferential interface nucleation: an expansion of the VLS growth mechanism for nanowires. Adv Mater 20:153
25. Wu Y, Cui Y, Huynh L, Barrelet CJ, Bell DC, Lieber CM (2004) Controlled growth and structures of molecular-scale silicon nanowires. Nano Lett 4:433
26. Kang D, Rim T, Baek CK, Meysappon M, Lee JS (2013) Investigation of electromigration in In$_2$Se$_3$ nanowire for phase change memory devices. Appl Phys Lett 103:233504
27. Wu Y, Yang P (2001) Direct observation of vapor-liquid-solid nanowire growth. J Am Chem Soc 123:3165
28. Dick KA, Deppert K, Mårtensson T, Mandl B, Samuelson L, Seifert W (2005) Failure of the vapor-liquid-solid mechanism in Au-assisted MOVPE growth of InAs nanowires. Nano Lett 5:761
29. Bernstein L (1966) Semiconductor joining by the solid-liquid-interdiffusion (SLID) process I. J Electrochem Soc 113:1282
30. Lin YK, Ting HW, Wang CY, Gwo S, Chou LJ, Tsai CJ, Chen LJ (2013) Au nanocrystal array/silicon nanoantennas as wavelength-selective photoswitches. J Mater Chem C 1:3593
31. Ting HW, Lin YK, Wu YJ, Chou LJ, Tsai CJ, Chen LJ (2013) Large area controllable hexagonal close-packed single-crystalline metal nanocrystal arrays with localized surface plasmon resonance response. J Mater Chem C 1:3593
32. Wu HW, Chen LJ, Tsai C (2006) Self-assembled epitaxial silicon nanowires grown along easy-glide directions on Si(001). J Micro & Nano Letters 1:25
33. Kang D, Rim T, Baek CK, Meysappon M, Lee JS (2014) Thermally phase-transformed In$_2$Se$_3$ nanowires for highly sensitive photodetectors. Small 10:3795
34. Li Q, Li Y, Gao J, Hwang SD, Sun XH (2011) High performance single In$_2$Se$_3$ nanowire photodetector. Appl Phys Lett 99:243105