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Capillary flow of amorphous metal for high performance electrode

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Metallic glass (MG) assists electrical contact of screen-printed silver electrodes and leads to comparable electrode performance to that of electroplated electrodes. For high electrode performance, MG needs to be infiltrated into nanometer-scale cavities between Ag particles and reacts with them. Here, we show that the MG in the supercooled state can fill the gap between Ag particles within a remarkably short time due to capillary effect. The flow behavior of the MG is revealed by computational fluid dynamics and density functional theory simulation. Also, we suggest the formation mechanism of the Ag electrodes, and demonstrate the criteria of MG for higher electrode performance. Consequently, when $\text{Al}_{85}\text{Ni}_5\text{Y}_8\text{Co}_2$ MG is added in the Ag electrodes, cell efficiency is enhanced up to 20.30% which is the highest efficiency reported so far for screen-printed interdigitated back contact solar cells. These results show the possibility for the replacement of electroplating process to screen-printing process.

Due to the strong demands for high performance electrodes in modern electronic devices, there have been many efforts to form electrodes using various methods such as screen printing, electroplating and deposition processes¹⁻⁴. Among these processes, screen printing process has the great advantage in commercializing the products due to its much higher cost-effectiveness⁵. However, the electrodes formed by the printing process exhibit lower electrical conductance than those by electroplating and deposition processes. To enhance the electrode performance of the printed electrodes, a special glass frit is to be included in the Ag paste, and should satisfy the following conditions for highly conductive contact formation: 1) the glass frit should be able to fill the cavity between the Ag particles at the temperature as low as possible; 2) it should react with Ag particles and Si substrate as fast as possible; and 3) it should have electrical resistance as low as possible. So far, oxide glass (OG) frit has been used to assist the Ag sintering and the contact formation between the electrode and the Si substrate⁶. Nevertheless, there is a critical limitation for the OG frit to form a high performance electrode, since the OG has a very poor electrical conductance. To overcome the limitation of the electrodes formed by the printing process, we have replaced the OG with metallic glass (MG)⁷⁻⁹. Since MGs consist of mostly metallic elements, their conductance is approximately 10^{14} times higher than that of the OGs^{10,11}. Moreover, MGs have some desirable characteristics such as the presence of a wide super-cooled liquid (SCL) region¹² and the outstanding thermoplastic formability which can satisfy the above-mentioned conditions critically required for enhanced solar cell efficiency¹³⁻¹⁶. In particular, it has been shown that even nanometer-sized channels can be filled with SCL in a controllable way when the viscosity is low enough and the capillary effect is present¹⁶. Here, we show that the application of Al-based MG instead of the OG in forming the printed electrodes can significantly enhance the solar cell efficiency comparable to that obtained by electroplating or deposition process. Since the details of the flow behaviour of the MG in the Ag printing electrodes have not been disclosed yet, we performed CFD (Computational Fluid Dynamics) and DFT (Density Functional Theory) simulation. Furthermore, we have



proposed the electrode contact formation mechanism, and demonstrated the criteria of the MG to enhance the performance of the silver printed electrodes.

Results

Flow behavior of metallic glass. Generally, the electrode paste consists of conductive metal particle, glass frit, additives and organic vehicles¹⁷. We selected Ag as an element for the conductive particle and Al-based MG particle as an alternative to the OG frit particle due to its superior electrical conductance. Among the glass forming Al-based alloys, the $\text{Al}_{85}\text{Ni}_5\text{Y}_8\text{Co}_2$ alloy was selected since the SCL region (ΔT_x) during heating is relatively large ($\Delta T_x = T_x - T_g$: 24 K, glass transition temperature T_g : 535 K, crystallization onset temperature T_x : 559 K at a heating rate of 40 K/min, Fig. 1a)⁸. The electrode paste was prepared by mixing Ag particles, $\text{Al}_{85}\text{Ni}_5\text{Y}_8\text{Co}_2$ MG particles and organic vehicles. The electrode pattern was prepared by screen printing the paste on single crystalline Si wafers and then fired by rapid heating from room temperature up to 873 K with a heating rate of 75 K/s.

Our approach for achieving high cell efficiency is based on the unique characteristics of the metallic glass: the metallic glass in the SCL state can infiltrate into a nanometer-scale cavity when a proper wetting condition is provided, i.e. wetting angle (θ) is $< 90^\circ$, and viscosity of the SCL is low, i.e. the SCL region is wide¹⁶. Kumar et al. reported a modified Hagen-Poiseuille's equation to determine the pressure required for MG to flow into nanometer-scale mould¹⁶. A prerequisite to obtain a high cell efficiency is that the MG should spontaneously fill the cavity between the Ag particles during the early stage of the firing process. Since there is no external pressure during the firing process, the condition for the SCL to fill the channel between the Ag particles can be derived by modifying the equation suggested by Kumar et al. (Supplementary Information I):

$$t_{\text{fill}} = \frac{3\eta d}{\gamma \cos \theta} \leq \frac{\Delta T_x}{R_{\text{heat}}} = t_{\text{SCL}} \quad (1)$$

Here, t_{fill} is the time required for the SCL to infiltrate into the channel between the Ag particles, η is the viscosity of the SCL, d is the diameter of the Ag particle, γ is the surface tension of the MG, θ is the contact angle between the SCL and the Ag particle, ΔT_x is the SCL region of the MG, R_{heat} is the heating rate during the firing process, and t_{SCL} is the time available for the MG to stay in the SCL region during the firing process. t_{fill} should be smaller than t_{SCL} for the SCL to completely flow into the channel between the particles.

The flow behavior of the $\text{Al}_{85}\text{Ni}_5\text{Y}_8\text{Co}_2$ MG in the Ag electrode paste was simulated using a Computational Fluid Dynamics (CFD) program of Fluent 13.0 (ANSYS Inc.). First, the heating rate dependence of the viscosity was evaluated by Thermo Mechanical Analysis (TMA) (Supplementary Information II). The viscosity data measured at a heating rate of 40 K/min (Fig. 1b) shows that the viscosity

of the MG is $\sim 2 \times 10^{10}$ Pa·s below the T_g , while it is reduced down to $\sim 2 \times 10^8$ Pa·s in the SCL region. Comparison of the viscosity data measured at heating rates of 5, 10, 20, and 40 K/min (Fig. 1c) indicates that the lowest viscosity in the SCL region decreases as the heating rate becomes higher. The lowest viscosity in the SCL region with a heating rate of 75 K/s (same as that used for the firing process) is evaluated to be 1.7×10^5 Pa·s by extrapolation of the measured data in a logarithmic scale (Fig. 1c), which is slightly higher than the previously reported viscosity of 3×10^4 Pa·s obtained at a heating rate of 75 K/s for La-based MG¹⁸. Differential scanning calorimetry (DSC) measurements were also performed at various ramping speed of 5, 10, 20, and 40 K/min to investigate the heating rate dependence of the SCL region. Since the ramping speed during the firing process is 75 K/s, the data were extrapolated using Kissinger's equation¹⁹, and it is derived that T_g and T_x are 564 and 604 K, respectively, at the heating rate of 75 K/s (Supplementary Information III).

The calculation system for the flow behavior consists of 4 layers of close packed MG and Ag particles. Each of the three layers from the bottom contains one MG particle and 32 Ag particles, while the fourth layer consisted of only Ag particles. (Fig. 2a, Supplementary Information IV for the details on the simulation method). The surface tension of the $\text{Al}_{85}\text{Ni}_5\text{Y}_8\text{Co}_2$ MG is calculated to be 1045.7 mN/m using first-principles molecular dynamics (MD) calculation based on the density functional theory (DFT) (Supplementary Information V for the details on the simulation method). Figures 2b–2d show a collection of a series of snap-shots with time at three different cross sections of the simulation model. Here, the viscosity of the MG is assumed to be 10^5 Pa·s and the average MG/Ag and MG/Si interfacial energies are calculated to be 181.8 mN/m, and 961.6 mN/m, respectively using ab initio MD method (Supplementary Information V). The contact angles of the MG on the Ag particle and the Si substrate are 20.1° and 76.6° , when the surface tensions of Ag and Si are 1164.1 mN/m and 1204.1 mN/m at 573 K, respectively^{20,21}. Since the contact angles are below 90° , the MG in the SCL state has a reasonably high driving force to spread on the surface of the Ag particles and the Si substrate by the capillary effect. The time available for the MG to stay in the SCL state is approximately 0.5 sec when the SCL region of the MG is 40 K and the heating rate for the firing process is 75 K/s. It can be seen from Fig. 2c and 2d that the MG in the SCL state begins to flow after 0.1 sec, and almost fill the channel between Ag particles after 0.5 sec, indicating that the SCL fills the channel between the particles within a remarkably short time. Thus, the MG particle in the electrode paste can fill the channel between the Ag particles when it is in the SCL state during the firing process. Figure 2b indicates that the MG particle in the first layer which is in contact with the Si substrate retains the almost spherical shape since the contact angle of the MG on the Si surface is higher than that of the MG on the Ag surface. However, the MG particles in the second and third layer flow into the channel due to the lower contact angle of the MG on the Ag surface.

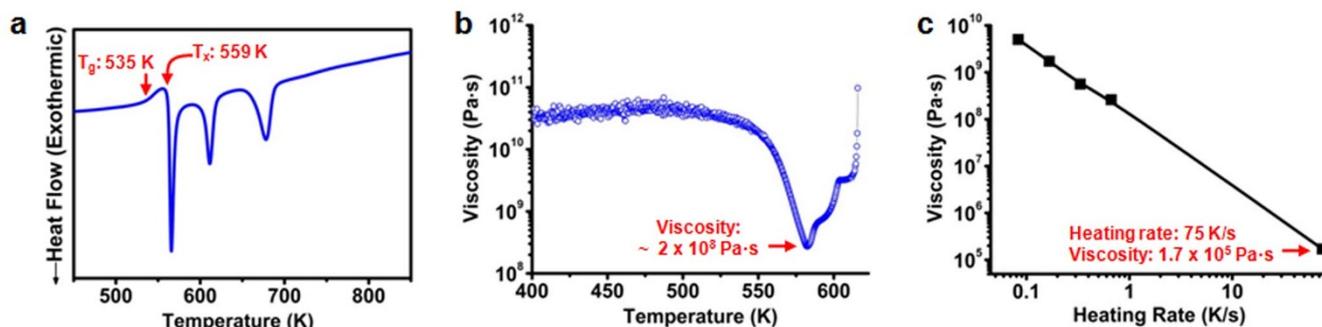


Figure 1 | Viscosity of $\text{Al}_{85}\text{Ni}_5\text{Y}_8\text{Co}_2$ MG (Metallic Glass). (a) Differential Scanning Calorimetry (DSC) curve and (b) temperature dependence of the viscosity of the $\text{Al}_{85}\text{Ni}_5\text{Y}_8\text{Co}_2$ MG at a heating rate of 40 K/min. (c) Effect of heating rate on the viscosity of the $\text{Al}_{85}\text{Ni}_5\text{Y}_8\text{Co}_2$ MG. The viscosity at a heating rate of 75 K/s is evaluated to be 1.7×10^5 Pa·s by extrapolation of the measured data in a logarithmic scale.

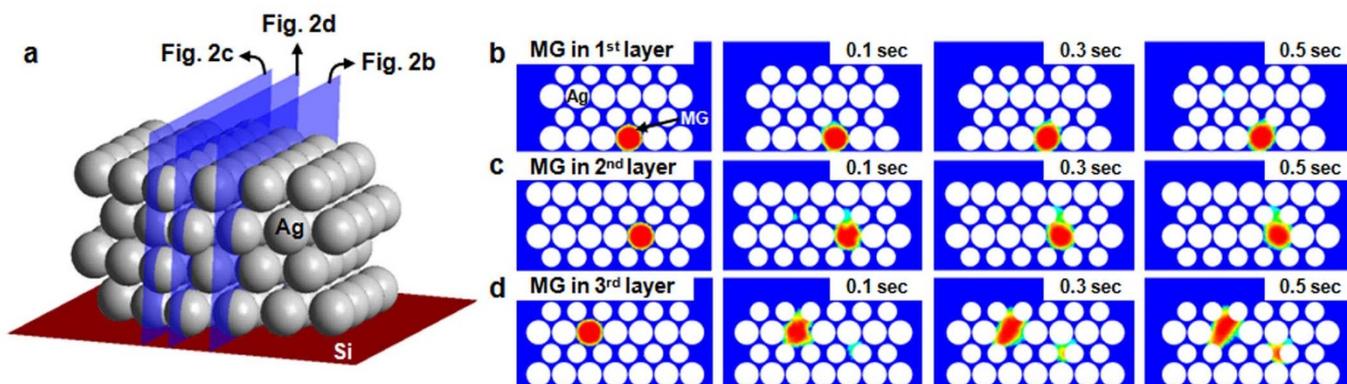


Figure 2 | Simulation of MG (Metallic Glass) flow behavior using CFD (Computational Fluid Dynamics) method. (a) Ag electrode model consists of MG and Ag particles on the Si substrate. Each of the three layers from the bottom contains one MG particle and 32 Ag particles, while the fourth layer has only Ag particles. Grey balls, dark red plate, and blue plates represent Ag particles, Si substrate, and cross section planes, respectively. MG volume fraction contour plots are shown at cross sections of the Ag electrode model where the MG particles are located in the (b) first, (c) second, and (d) third layer from the bottom. White balls represent Ag particles. MG volume fraction is displayed with the color ranging from blue (minimum value) to red (maximum value) on the contour plots. The MG particles begin to flow after 0.1 sec, and almost fill the gap between the Ag particles after 0.5 sec. Spontaneous flow of the MG occurs by the capillary effect due to the low contact angle ($\sim 20^\circ$) between MG-Ag and the low viscosity ($\sim 10^5$ Pa·s) of the MG in its supercooled liquid region.

From the DFT simulation and the experimental results mentioned above, the parameters required for equation (1) can be obtained: viscosity of the SCL: 1.7×10^5 Pa·s, surface tension of the MG: 1045.7 mN/m, contact angle of the MG on the Ag surface: 20.1° , diameter of Ag particle: 1 μm , SCL region: 40 K and heating rate: 75 K/s. Equation (1) indicates that the time required for the SCL flow into the channel between the particles, t_{fill} , is 0.52 sec, while the time available for the MG to stay in the SCL state, t_{SCL} , is 0.53 sec. Therefore, it is concluded that the MG in the SCL state can fill the channel between the particles by the capillary effect during the firing process. Equation (1) further indicates that lower viscosity of the SCL, larger ΔT_x of the MG, and smaller size of the Ag particle is preferred for more efficient filling of the cavity by the capillary effect. Higher heating rate can lower the viscosity, but the time available for the MG to remain in the SCL state becomes shorter, suggesting the presence of optimum heating rate to maximize the infiltration of the SCL.

Electrode formation and contact morphology. Due to the lower T_g and T_x of the $\text{Al}_{85}\text{Ni}_{15}\text{Y}_8\text{Co}_2$ MG⁸ than for other glass forming systems such as Cu- and Zr-based MGs^{22,23}, the Al-MG is suitable for low firing-temperature process. Lower firing temperature is particularly important for the back contact solar cell fabrication in minimizing the thermal degradation of the carrier lifetime in the cell²⁴. The firing peak temperature attempted in the present study (873 K) is about 200 K lower than the typical peak temperature for forming conventional solar cell electrodes using the OG frit.

The electrode formed by the firing process exhibited a continuous network structure implying that the MG frit plays an effective role in sintering of Ag particles (Fig. 3a). Several unique microstructural features can be noticed from the contact achieved with the MG electrode paste. First, the original spherical surface of Ag particles changed into a flat surface, especially at the contact region with the Si emitter in spite of the lower firing peak temperature (Fig. 3b). Second, an ultra-thin layer with a thickness of ~ 5 nm is present between the Ag electrode and the Si emitter (Fig. 3c). The layer is enriched in Al and O as shown in the EDS line profile (Fig. 3d). Such contact morphology is strikingly different from that formed using the OG frit in conventional Si solar cells. When the conventional OG frit is used, most of the Ag electrode and the Si emitter are separated by a much thicker highly resistive glass layer with varying thickness, which is responsible for the high contact resistance and therefore the

low cell efficiency²⁵. Third, no Ag crystallites are observed at the interface (Fig. 3c). This is in contrast to the findings when the conventional OG frit is used: there Ag crystallites precipitate from the supersaturated OG frit upon cooling or regrow by an eutectic reaction^{5,6}. It was generally accepted that a uniform distribution of a large number of Ag crystallites in the interface region has been recognized as a prerequisite for high current flow efficiency in conventional Si solar cells²⁶.

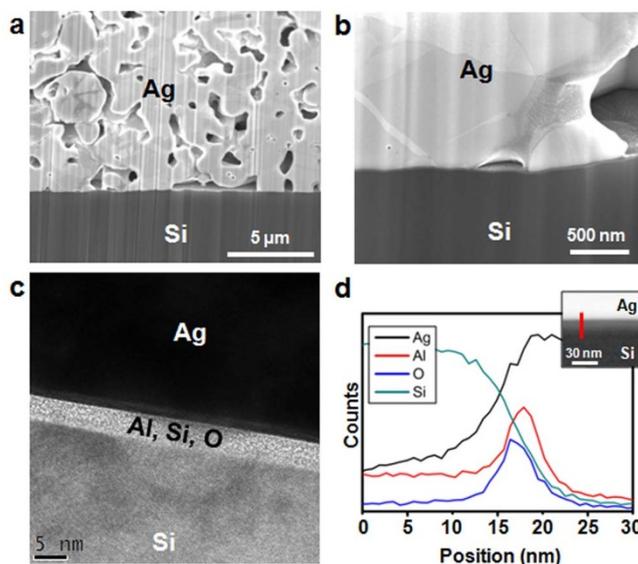


Figure 3 | Cross section of Ag electrode. SEM images obtained from the cross section of the contact between the Ag electrode and the Si wafer showing: (a) Sintered Ag electrode on the Si wafer; (b) Flat surface of the Ag particle in contact with the Si wafer surface. (c) Bright field TEM image obtained from the cross section of the contact between the Ag electrode and the Si wafer; (d) EDS line profile along the red line marked in the inset (dark field TEM image) showing enrichment of Al and O in the interface layer. The ultra-thin interface layer (~ 5 nm) between the Ag electrode and the Si wafer enables efficient carrier transfer through the interface by direct tunneling, thus finally results in the high cell efficiency of the solar cell.

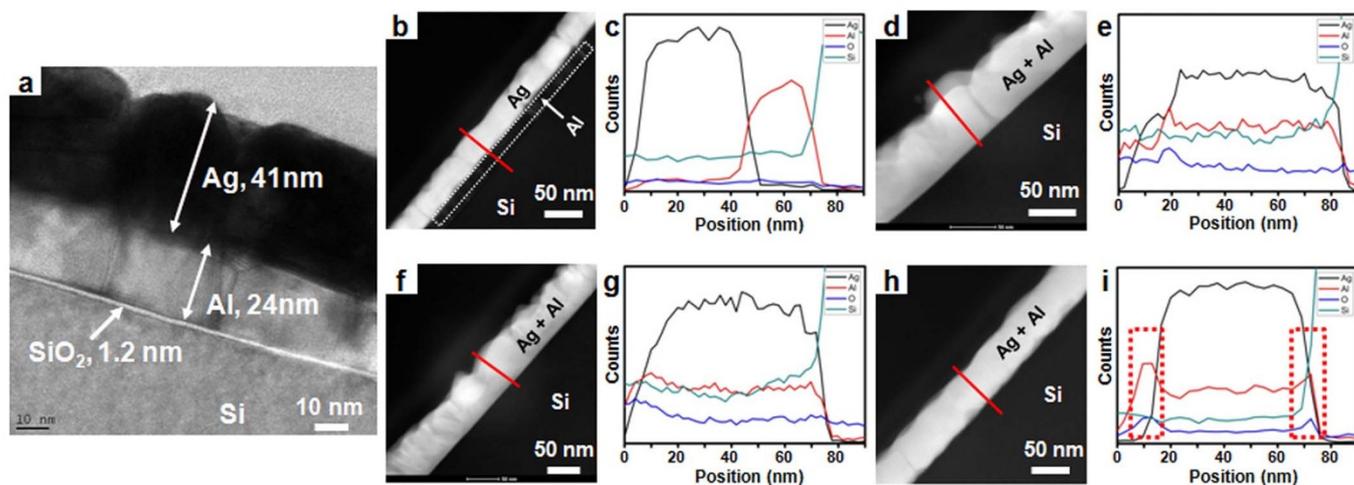


Figure 4 | Mechanism studies in Ag-Al-Si layers. Evaporation of Al and Ag thin layers on the Si substrate. (a) Bright field TEM image showing Ag and Al layers deposited on the Si wafer; (b), (d), (f), (h) Dark field TEM images showing the deposited layer on the Si wafer before heat treatment and after heating up to 573, 673 and 873 K, respectively; (c), (e), (g), (i) EDS line profiles across the deposited layer before heat treatment and after heating up to 573, 673 and 873 K, respectively. (d–e) Interdiffusion of Al and Ag at the interface forms an Al-Ag solid solution layer at 573 K. (h–i) The oxidation of Al occurs by the redox reaction of $4/3\text{Al} + \text{SiO}_2 \rightarrow 2/3\text{Al}_2\text{O}_3 + \text{Si}$ at the interface between the Al-Ag solid solution layer and the Si wafer.

Contact formation mechanism. In order to investigate the contact formation mechanism, a separate experiment was performed by depositing thin Al and Ag layers sequentially on the Si wafer surface using an evaporator (Fig. 4a). The native SiO_2 film with a thickness of ~ 1.2 nm was present on the Si wafer surface. Before heat treatment, the Al and Ag layers were well separated (Figs. 4b, 4c). The deposited samples were heated with a heating rate of 40 K/s up to 573, 673, and 873 K, and held for 1 min before cooling to room temperature. After heating up to 573 K, the composition profiles of Al and Ag across the deposited layer was almost constant (Figs. 4d, 4e), indicating that Al diffuses into the Ag layer due to relatively larger solubility of Al in Ag (13 at% at 573 K). After heating up to 673 K (Figs. 4f, 4g) and 773 K (not shown), same type of composition profile was obtained across the deposited layer. However, after heating up to 873 K (Figs. 4h, 4i), Al and O enriched regions were observed at both ends of the deposited layer (marked in Fig. 4i), indicating that selective oxidation of Al occurs as the temperature becomes higher. At the interface between the deposited layer and the Si wafer, the oxidation of Al occurs by the redox reaction of $4/3\text{Al} + \text{SiO}_2 \rightarrow 2/3\text{Al}_2\text{O}_3 + \text{Si}$, ΔG^0 (J/mol) = $-207010 + 32.42 T$ due to the presence of the native SiO_2 layer. Therefore, the native SiO_2 layer reduces to Si when Al_2O_3 forms²⁷.

Based on the results of the simulation and the deposition experiment, we propose that the contact formation during screen printing proceeds via three main steps: formation of SCL wetting layer between Ag particles and crystallization of MG \rightarrow formation of Al-Ag eutectic liquid film \rightarrow sintering of Ag particle and electrode contact formation, as schematically illustrated in Fig. 5. Before the firing process, the Ag and MG particles are randomly distributed in the screen printed paste (Fig. 5a). As the temperature is raised during the firing process, the MG in the paste transforms into the SCL state and fill the gap between Ag particles by the capillary force (Fig. 5b). Since the heating rate is very high during firing, crystallization of the SCL occurs shortly after wetting on the Ag particles. At the same time, interdiffusion of Al and Ag occurs at the interface between the MG wetting layer and the Ag particle (Fig. 5c). As the temperature increases further, binary Al-Ag eutectic reaction occurs at the interface (Al-Ag eutectic temperature, 840 K, Fig. 5d). Since the viscosity of general metals in their liquid state is 1 to 10 mPa·s, it is expected that the Al-Ag eutectic liquid can wet the Ag particle surface, accelerating the sintering of Ag particles (Fig. 5e). Also, the Al-Ag eutectic

liquid layer flattens the original spherical shape of the Ag particles at the contact region with the Si substrate (Fig. 5f). Additional eutectic reactions of binary Al-Si (850 K) and ternary Ag-Al-Si (840 K)²⁸ are possible at the contact region with the Si substrate. The presence of the eutectic liquid at the peak firing temperature leads to the formation of an ultra-thin interface layer between the Ag electrode and the Si wafer (Fig. 3c) during cooling. After cooling the resulting ultra-thin layer is enriched in Al and O as shown in Fig. 3d (Fig. 5g).

During contact formation, there are two important steps: flow of the SCL into the channel between the Ag particle (as can be seen in the simulation result mentioned above); and flow of the eutectic liquid by which the sintering process and contact formation can be completed. Here, the fluidic behavior of the Al-Ag eutectic liquid was predicted using CFD simulation mentioned above. The result shows that the Al-Ag eutectic liquid forms a wetting layer on the Ag particles and the Si wafer surface within a remarkably short time of 0.0003 sec (Fig. 6, Supplementary Information VI for the DFT simulation of the interfacial energies and Supplementary Information VII for the input parameters of the CFD simulation).

Evaluation of cell efficiency. Generally, conventional Ag paste containing the OG frit is sintered by firing up to 1073 K, and the specific resistance of the resulting electrode is 3 – 5 $\mu\Omega\text{cm}^2$ ²⁹. However, in our study, the specific resistance of the electrode is 2.96 $\mu\Omega\text{cm}^2$ in spite of the lower firing temperature of 873 K. Also, Al atoms in the MG frit induce binary and ternary eutectic reactions with Ag and Si atoms, leading to the formation of a ~ 5 nm thick ultra-thin interface layer (Fig. 3c). Such a thin interface layer enables efficient carrier transfer through the interface by direct tunneling, thus significantly decreasing the contact resistance between Ag electrode and Si wafer. The contact resistance of the electrode is measured to be 0.18 and 0.37 $\text{m}\Omega\text{cm}^2$ for p-type emitter and n-type BSF (Back Surface Field), respectively, which is in the same range of 0.1 – 0.3 $\text{m}\Omega\text{cm}^2$ as for electro-plated electrodes. Such an improved contact resistance implies that the cell performance is comparable to that of the cell fabricated using an electroplating method. To evaluate the cell efficiency, interdigitated back contact (IBC) solar cells were fabricated using an electrode paste containing Al-based MG frit. IBC cell design was adopted since it demonstrates the highest efficiency among the Si solar cells developed until now. The higher cell efficiency is mainly attributed to elimination of the

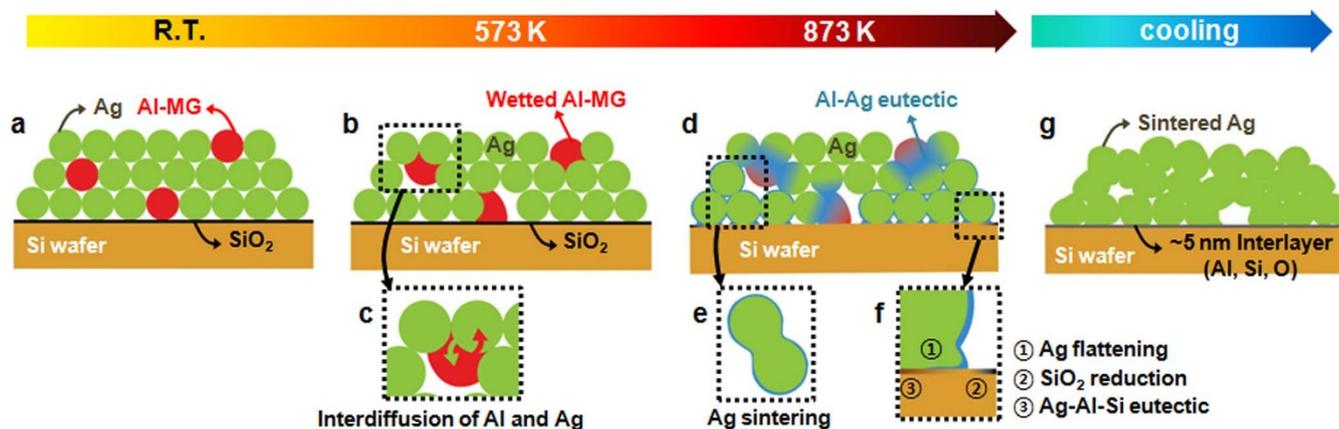


Figure 5 | Electrode contact formation mechanism. (a) Cross section of Al-MG/Ag paste printed on Si wafer. Green, red, yellow, and black represent silver, $\text{Al}_{85}\text{Ni}_5\text{Y}_8\text{Co}_2$ MG (Metallic Glass), Si wafer, and native SiO_2 layer, respectively. (b) Al-MG particle spontaneously flows into the channel between the Ag particles during firing process. Since the contact angle between the MG and the Ag particles is below 90° and the viscosity of the MG is decreases in its SCL (supercooled liquid) region, the MG has a reasonably high driving force to fill the gap between Ag particles by the capillary effect. (c) Interdiffusion of Ag and Al occurs at the interface. (d) At the peak temperature (873 K) of the firing process, binary Al-Ag eutectic reaction occurs at the interface. Blue layer represents the Al-Ag eutectic liquid. (e) The Al-Ag eutectic liquid accelerates the sintering of Ag particles. (f) 1. Al-Ag eutectic liquid layer flattens the originally spherical Ag particle at the contact region with the Si wafer; 2. The native SiO_2 layer on the Si wafer reduces to Si by Al in the Al-Ag eutectic liquid: $4/3\text{Al} + \text{SiO}_2 \rightarrow 2/3\text{Al}_2\text{O}_3 + \text{Si}$; 3. Additional eutectic reactions of binary Al-Si (850 K) and ternary Ag-Al-Si (840 K) occur at the contact region with the Si substrate. (g) The presence of the eutectic liquid at the peak firing temperature leads to the formation of an ultra-thin interface layer between the Ag electrode and the Si wafer during cooling. Grey layer represents the ultra-thin interface layer.

front shading losses by placing all electrodes at the back of the cell. The fabricated cells exhibit an energy conversion efficiency of 20.30% (Table 1), which is the highest record reported for screen-printed IBC solar cells so far^{30,31}. Cell efficiency of 20.30% is acquired using commercialized size (154.8 cm^2) of Si solar cell. Furthermore, we fabricated the solar cell using the same Ag paste for both of p-type emitter and n-type BSF, which would be highly advantageous in commercializing the screen-printed solar cell using Al-based MG frit. Such a high cell efficiency is mainly attributed to the lower

contact resistance which can be achieved by securing a very thin interface layer ($\sim 5 \text{ nm}$, Fig. 3c) beneath the Ag electrode. The thickness of the interface layer becomes much larger, generally over 100 nm, when the conventional OG frit is used³². Comparison of the I-V parameters (Table 1) of the screen printed IBC solar cell fabricated using a MG frit (SPC) and those of the electrodeposition processed solar cell (EPC) shows that the cell performance of the SPC is comparable to that of the EPC. The mechanical stability of the solar cell fabricated using a MG frit is

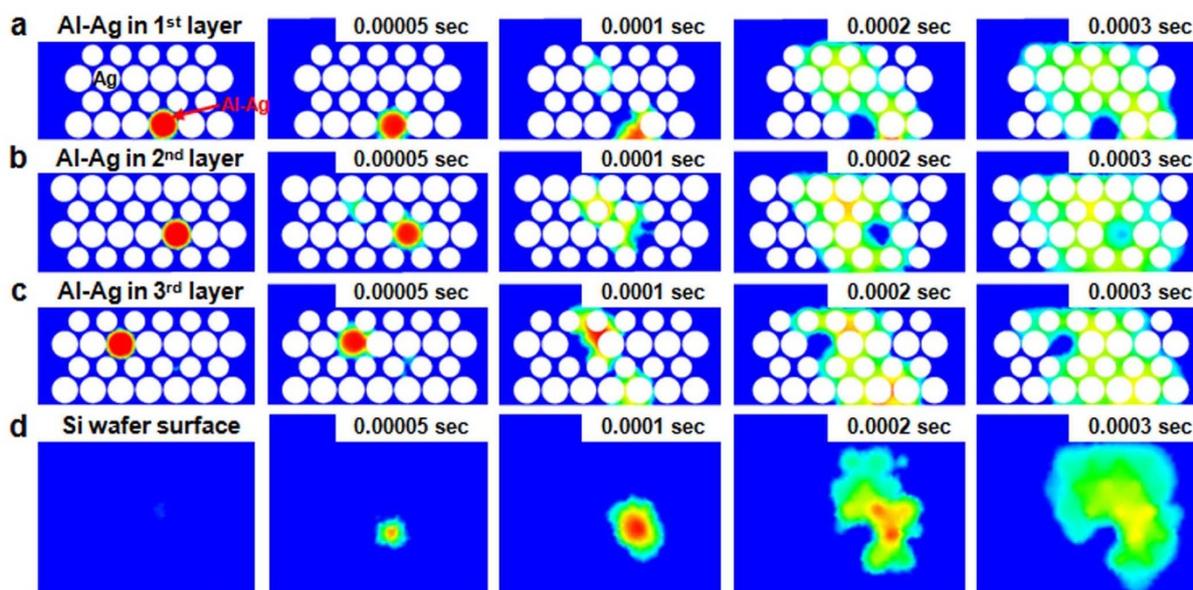


Figure 6 | Simulation of fluidic behavior of Al-Ag eutectic alloy. Al-Ag liquid alloy volume fraction contour plots at cross sections of the Ag electrode model (Same model described in Fig. 2a except that the MG balls are substituted by Al-Ag balls) where the Al-Ag alloys are located in the (a) first, (b) second, and (c) third layer from the bottom. White balls represent Ag particles. Volume fraction of Al-Ag alloy is displayed with the color ranging from blue (minimum value) to red (maximum value) on the contour plots. (d) Al-Ag alloy volume fraction contour plots of Si substrate. Since formation of the eutectic liquids are possible at the peak firing temperature (873 K), the eutectic liquid can form wetting layer on the Ag particles and on the Si substrate in a remarkably short time ($\sim 0.0003 \text{ sec}$).



Table 1 | Comparison of I–V parameters of solar cells fabricated using screen printing method and electroplating method (cell area: 154.8 cm²)

	$J_{sc}^{[a]}$	$V_{oc}^{[b]}$	$R_s^{[c]}$	$R_{sh}^{[d]}$	FF ^[e]	η ^[f]
Electroplating method	40.8	680	0.63	$\geq 6,000$	79.35	22.02
Screen printing method	40.8	672	1.11	$\geq 10,000$	74.26	20.30

[a] J_{sc} (mA/cm²): short-circuit current density, [b] V_{oc} (mV): open-circuit voltage, [c] R_s (Ω cm²): series resistance, [d] R_{sh} (Ω cm²): shunt resistance, [e] FF (%): fill factor, [f] η (%): cell efficiency

not affected by the crystallized metallic glass which is known to be brittle, since eutectic reactions between crystallized product, Ag and Si occurs during the firing process. Therefore, the use of the MG frit in the electrode paste can significantly enhance the competitiveness of the screen printing process which will eventually become superior to that of electroplating or deposition processes.

Discussion

We have achieved a significant improvement in the quality of the contact between the Ag electrode and the Si emitter using a MG frit in the paste for screen printing. The dramatic improvement of the contact quality is attributed to: (i) the spontaneous flowability of the SCL to fill the cavity between the Ag particles; (ii) the efficiency in sintering to form a sound Ag electrode on the emitter surface; and (iii) the existence of eutectic reactions enabling to form an ultra-thin layer and a larger contact area between the Ag electrode and the Si emitter. The solar cell fabricated using such a good contact electrode exhibits an energy conversion efficiency of 20.30%, which is the highest efficiency reported so far for screen printed solar cells.

Methods

The electrode paste was prepared by mixing Ag particles (diameter: $\sim 1 \mu\text{m}$), $\text{Al}_{85}\text{Ni}_5\text{Y}_8\text{Co}_2$ MG particles (diameter: $< 5 \mu\text{m}$, fabricated in Phoenix Scientific Industries Ltd.) and organic vehicles. Butyl carbitol and ethyl cellulose-based chemicals were used for the organic vehicles. The volume fraction of Al-based MG in the electrode paste was $\sim 6 \text{ vol}\%$ excluding the organic vehicles. The pre-mixed paste was 3-roll-milled, and then re-mixed in the paste mixer to remove air bubbles. The electrode pattern was prepared by screen printing the paste on single crystalline Si wafers, followed by drying in an IR-belt furnace at 423 K for 5 min to evaporate the organics in the paste. The dried wafer was fired using a rapid thermal process in an IR-belt furnace from room temperature up to 873 K with a ramping speed of 75 K/s.

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Author contributions

S.Y.K., J.M.P., K.H.P., K.R.L., E.S.L. fabricated the metallic glasses and analyzed their properties; S.J.K., S.S.J. fabricated the Ag/MG paste and measured its electrical properties; S.J.K., S.S.J., S.C.P. fabricated the solar cell and measured the cell efficiency; J.H.L., I.Y.S., J.C.P. carried out the microscopy experiments; S.Y.K., S.M.L., I.T.H., D.H.K., E.S.L. designed the experiments and analyzed the experimental results; S.Y.K., E.A.C. calculated the deformation behavior of metallic glasses and interfacial energies. S.Y.K., S.J.K., W.T.K., J.E., D.H.K., E.S.L. wrote the paper.



Additional information

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