

## Direct Bonding of LiTaO<sub>3</sub>/Si at Room Temperature Using Self-sputtered Bonding Method

自己スパッタ接合法を用いた常温による LiTaO<sub>3</sub>/Si 直接接合

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### 1. Introduction

LiTaO<sub>3</sub> and LiNbO<sub>3</sub> have been widely used for surface-acoustic-wave (SAW) devices in mobile communication systems, as these materials have high piezoelectric properties. However, as the materials have large temperature coefficient of frequency (TCF), various temperature compensation techniques are currently being studied to reduce the TCF. It has been reported that the TCF characteristics was improved by directly bonding the LiTaO<sub>3</sub> or LiNbO<sub>3</sub> wafer bonded to a support substrate with a low coefficient of thermal expansion (CTE), and the temperature compensated (TC) SAW devices have been already put to practical use by the directly bonded LiTaO<sub>3</sub>/sapphire wafers [1]. Si and glass have been also investigated as the support substrate. The conventional bonding method requires heating process to achieve strong bonds. Since the thermal expansion mismatch between oxide materials such as LiTaO<sub>3</sub> and LiNbO<sub>3</sub> and semiconductors is large, various problems such as cracking, bowing and so on occur during the heating process. Since surface activated bonding (SAB) is a bonding method carried out at room or low temperature [2, 3], the method is suitable for bonding between the dissimilar materials with large thermal expansion mismatch. However, it is difficult to directly bond oxide materials to Si by SAB method [4]. In this paper, we propose the novel bonding process for directly bonding oxide materials to Si at room temperature. We have developed the suitable apparatus for our proposed process and examined the direct bonding of LiTaO<sub>3</sub>/Si wafers. We have evaluated the surface energy and the tensile strength and investigated the bonding interface by transmission electron microscopy (TEM). Moreover, the elemental composition across the bonding interface has been analyzed by energy-dispersive X-ray spectroscopy (EDS).

### 2. Self-sputtered Bonding (SSB) Method

In our proposed bonding process, as shown in Fig. 1, only the lower wafer's surface is etched by Ar plasma, and the thin film of the lower wafer's material is deposited on the upper wafer's surface

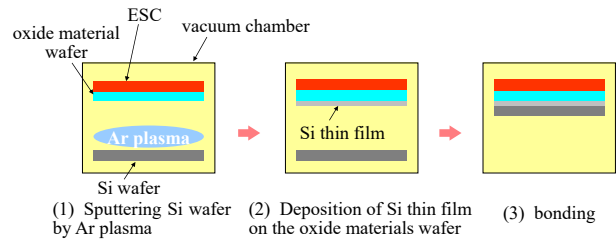


Fig. 1 Bonding process by SSB method.

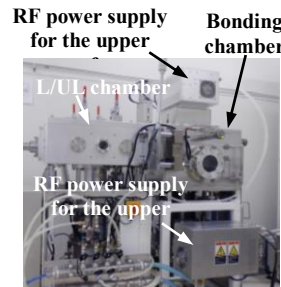


Fig. 2 Bonding apparatus.

Table I. Specification of the apparatus.

Max. wafer diameter	8 inch
Background vacuum pressure	$< 1 \times 10^{-5}$ Pa
RF power for the upper wafer	1000 W
RF power for the lower wafer	500 W
Max. press load	500 kgf

during the etching process. The surface of the lower wafer is activated during Ar plasma etching, and the surface of the film deposited on the upper wafer remains activated due to high vacuum process. The upper and lower wafers are then immediately brought into contact. Therefore, we call this bonding process “Self-sputtered Bonding (SSB)” method. Figure 2 shows the bonding apparatus which we developed for SSB method. The apparatus is comprised of bonding process chamber and load/unload rock chamber. The specification is shown in Table I. Since the apparatus has two radio frequency (RF) power supplies for the upper and lower wafer, each wafer's surface can be individually sputtered, also. The lower wafer can be approached to the upper wafer to contact in the bonding chamber after plasma process.

### 3. Bonding Experiments

4-in 42° Y-X LiTaO<sub>3</sub> wafer (thickness of 350 μm, Yamajyu ceramics) and 4-in Si blanket wafer (thickness of 525 μm) were prepared for this experiments. The LiTaO<sub>3</sub> wafer was held by the electrostatic chuck (ESC) to turn the surface down. The Si wafer was placed on the lower side in the same vacuum chamber. Only Si wafer was etched by Ar plasma produced RF power supply for lower wafer. RF power and irradiation time was 300 W

and 2.5 min, respectively. The background vacuum pressure was lower than approximately  $2 \times 10^{-5}$  Pa before the plasma process. The LiTaO<sub>3</sub> and Si wafers were then immediately brought into contact with a bonding press force of 80 kgf.

## 4. Results and discuss

### 4.1 Bonding results

**Figure 3** shows the boned LiTaO<sub>3</sub>/Si wafers. A void was confirmed, which was caused by the residue on the LiTaO<sub>3</sub> wafer's surface owing to insufficient cleaning before bonding; however, it seems that the SSB method yielded a good bonding. The surface energy of the bonded wafer was evaluated by the crack-opening method [5]. The surface energy of wafers bonded by the normal SAB method was approximately 1 J/m<sup>2</sup>. However, as shown in **Fig. 4a**, a fracture from the LiTaO<sub>3</sub> wafer in the bonded wafers by SSB method occurred when the blade was inserted toward the bonding interface. The bonding strength was also measured by a tensile test. The specimen for the tensile test was prepared by cutting the bonded wafers into  $8 \times 8$ mm<sup>2</sup> sections. The bonding strength was found to be approximately 16 MPa, but the specimen fractured at the glue interface and not at the bonded interface as shown in **Fig. 4b**. This means that the bonding strength of the LiTaO<sub>3</sub>/Si interface is higher than the measured value.

### 4.2 TEM observation and EDS analysis of LiTaO<sub>3</sub>/Si bonding interface

**Figure 5a** shows the cross-sectional TEM image of the LiTaO<sub>3</sub>/Si bonding interface. No microvoids or gaps are observed at the bonding interface, but the intermediate layer is visible. The thickness of this layer was approximately 8 nm. The bonding interface is assumed to be below the center of the intermediate layer as shown in **Fig. 5a**. The layer above the

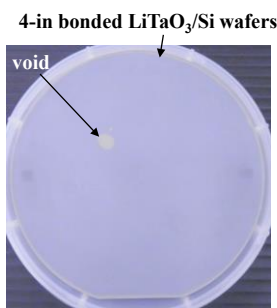


Fig. 3 Bonded LiTaO<sub>3</sub>/Si wafers by SSB method

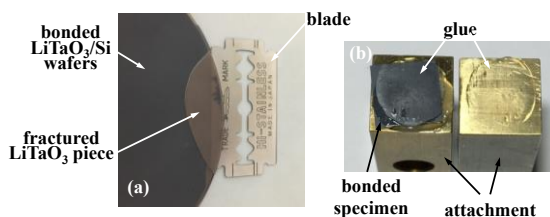


Fig. 4 (a) Fracture image of the bonded LiTaO<sub>3</sub>/Si wafer by opening-crack method. (b) Image of LiTaO<sub>3</sub>/Si bonding specimen fractured by tensile test.

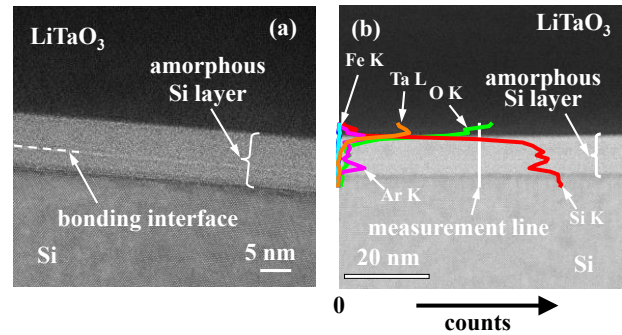


Fig. 5 (a) High resolution TEM cross sectional image of the bonding interface. (b) EDS line analysis along the white line on TEM image.

bonding interface is assumed to be deposited during Ar plasma process, and the lower is assumed to be damaged formed by Ar plasma irradiation [6]. The elemental composition across the LiTaO<sub>3</sub>/Si bonding interface was measured by EDS. **Figure 5b** shows the Si, O, Al, Fe, Ar, and Ta concentrations across the interface. Although O and Ar were slightly detected in the intermediate layer, the layer seems to be amorphous Si layers.

## 5. Conclusion

We have demonstrated a novel bonding method, SSB method, to directly bond oxide material to Si at room temperature, and developed the bonding apparatus for the method. We showed the bonding strength of the bonded LiTaO<sub>3</sub>/Si wafers by the method was higher than that by SAB method. Our proposed method promises to directly bond Si to the oxide materials or the undesirably damaged materials by surface activation at room temperature.

## References

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