

Giant Enhancement of Ferroelectricity in Strained BaTiO₃ Thin Films

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Epitaxial ferroelectric thin films have different properties from the bulk single crystals due to epitaxial and thermal strain and it offers an opportunity to modify the various ferroelectric properties. We deposited fully commensurate BaTiO₃ ($a = 3.992 \text{ \AA}$) ferroelectric thin films on novel substrates including (110) DyScO₃ ($a = 3.944 \text{ \AA}$) and (110) GdScO₃ ($a = 3.97 \text{ \AA}$) by pulsed laser deposition (PLD) and molecular beam epitaxy (MBE) and observed ferroelectric transition using temperature dependent (room temperature to 900 °C) four-circle x-ray diffraction and second harmonic generation measurements. Biaxial compressive strain dramatically enhanced the ferroelectric properties—both the ferroelectric transition temperature (T_c) and remanent polarization (P_r)—of BaTiO₃ thin films. This strain, imposed by commensurate epitaxy, can result in a T_c nearly 500 K higher and a P_r at least 170% higher than BaTiO₃ single crystals. This is the largest increase in T_c ever reported for a ferroelectric. These giant property enhancements are consistent with thermodynamic predictions. This work demonstrates a route to a lead-free ferroelectric with the high T_c and high P_r needed for ferroelectric memories, and also a general means for achieving extraordinary physical properties in thin films through strain engineering.