

No. 2A-2 LiTaO₃, Lithium tantalate*(M* = 235.887)

1a	Ferroelectricity in LiTaO ₃ was discovered by Matthias and Remeika in 1949.		49Mat
b	phase	II	I
	state	F	P
	crystal system	trigonal	trigonal
	space group	R3c – C _{3v} ^{6 a)}	R $\bar{3}$ c – D _{3d} ^{6 b)}
	Θ_f [°C]	665(5)	
	$P_s \parallel [001]$ of hexagonal unit cell.		64Raz
	$T_{\text{melt}} = 1650$ °C.		54Meg
	$\rho_X = 7.4564 \cdot 10^3$ kg m ⁻³ .		65Bal
	See Fig. 2A-2-001.		67Abr1
	Phase relation: Fig. 2A-2-002.		
	Raman spectra show an amorphous state above 33.2 GPa: see		94Lin
	Vitreous state: see		77Gla
2a	Crystal growth: Czochralski method.		65Bal,
	Flux-grown crystals: see		65Fed
	Producing shaped single crystals by Stepanov (EFG) method (review paper): see		89YeZ
	Ceramics obtained from powders prepared by sol-gel method: see		86Red
	Epitaxial growth of thin films on sapphire substrates by RF magnetron sputtering: see		88Rav
	Synthesis of whiskers from metal alkoxide derived gels: see		92Sai
	Epitaxial growth of thin films on sapphire substrate by excimer laser ablation method: see		93Dei
	Epitaxial thin film growth by metalorganic chemical vapor deposition (CVD): see		93Shi
	Chemical preparation of spherical particle powder: see		93Xie
	Epitaxial growth on sapphire c-substrate by RF magnetron sputtering: see		90Jea1
	Sol-gel derived thin films: see		91Sai
	Epitaxial nonlinear optical films on GaAs: see		90Jea2
	Proton-exchange by pyrophosphoric acid: see		93Hun
	Proton-exchange in melts of acids and acidic salts: see		92Yam1
	Proton-exchange with controlled acidity (review paper): see		89Gan
			90Gan
3a	$a_{\text{hex}} = 5.15428$ Å, $c_{\text{hex}} = 13.78351$ Å and $a_{\text{rh}} = 5.4740$ Å, $\alpha_{\text{rh}} = 56^\circ 10.5'$ at 25 °C.		67Abr1
	Fig. 2A-2-001.		
b	$Z = 4$ in hexagonal unit cell.		67Abr1
	Crystal structure: Tables 2A-2-001...003; Fig. 2A-2-003, Fig. 2A-2-004.		
4	Thermal expansion: $\alpha_a^{(1)} = 0.161 \cdot 10^{-4}$ °C ⁻¹ , $\alpha_a^{(2)} = 0.070 \cdot 10^{-7}$ °C ⁻² ; $\alpha_c^{(1)} = 0.041 \cdot 10^{-4}$ °C ⁻¹ , $\alpha_c^{(2)} = -0.100 \cdot 10^{-7}$ °C ⁻² , where $\Delta a/a = \alpha_a^{(1)} (T - T_0) + \alpha_a^{(2)} (T - T_0)^2$, $\Delta c/c = \alpha_c^{(1)} (T - T_0) + \alpha_c^{(2)} (T - T_0)^2$, with $T_0 = 25$ °C.		71Smi
	Fig. 2A-2-005.		
5a	Dielectric constant: Table 2A-2-004, Table 2A-2-005; Figs. 2A-2-006...2A-2-028; see also Fig. 2A-1-026, Fig. 2A-1-027 in No. 2A-1.		

	Vitreous specimens prepared by roller quenching: see	77Gla
	Curie-Weiss constant: $C = 1.5 \cdot 10^5$ K.	69Yam
c	Spontaneous polarization: $P_s = 0.50$ Cm ⁻² . Fig. 2A-2-029, Fig. 2A-2-030.	68Wem
	Point defects and singularities of low temperature behavior of P_s : see	91Nov
d	Pyroelectricity $p_3 = -17.6 \cdot 10^{-5}$ CK ⁻¹ m ⁻² (primary effect: $-17.8 \cdot 10^{-5}$ CK ⁻¹ m ⁻² , secondary effect: $+0.200 \cdot 10^{-5}$ CK ⁻¹ m ⁻²) at RT; see also Fig. 2A-2-031, Fig. 2A-2-032.	80Bha 75Bee
	Pyroelectric properties of various LiTaO ₃ related ceramics: see	90InH
	Data for calculating figure of merits: see	75Bee
	Point defects and singularities of low temperature behavior of p : see	91Nov
6a	Heat capacity: $C_p = 100$ J K ⁻¹ mol ⁻¹ at 25 °C. Fig. 2A-2-033.	75Bee
7a	Piezoelectricity: Fig. 2A-2-034; see also Table 2A-2-004 and Table 2A-2-006. Pressure dependence of piezoelectric polarization: see	76Gra
8a	Elastic compliance and stiffness: see Table 2A-2-004 in 5a, and Table-2A-006 in 7a; Figs. 2A-2-035...2A-2-040. Acoustic properties: see Table 2A-1-012 in No. 2A-1. Temperature dependence of plate resonators: see Surface acoustic wave: Table 2A-2-007. Temperature dependence: see Effect of film loading: see Surface acoustic wave: The X-cut 112° rotated Y propagation mode has a low bulk spurious response level and a low temperature coefficient of propagation delay time as well as a high coupling coefficient. Table 2A-2-008; Fig. 2A-2-041, Fig. 2A-2-042. Temperature coefficient adjustment of propagation delay time by SiO ₂ coating, see Internal friction: Fig. 2A-2-043. Quantitative evaluation of elastic properties by acoustic microscopy: see	76Bur 72Sch 75Par 77Hir 81Ina 91Kus
9a	Refractive index: Table 2A-2-009; Fig. 2A-2-044, Fig. 2A-2-045. Birefringence: Fig. 2A-2-046. Light transmission: Fig. 2A-2-047. Reflection: Fig. 2A-2-048; see also Fig. 2A-2-024 in 5a. Absorption: Fig. 2A-2-045; see also Fig. 2A-2-025 in 5a, Fig. 2A-1-087 in No. 2A-1. Anomalous refractive index change and recovery of electrooptic coefficient g_{33} in proton-exchanged optical waveguides after annealing: see	92Yuh
b	Electrooptic effect: Table 2A-2-010. Quadratic electrooptic constant: $M_{13}^T = 0.013(3)$ m ⁴ C ⁻² , $M_{33}^T = 0.061(1)$ m ⁴ C ⁻² ($\lambda = 644$ nm). Electrooptical properties in the temperature range 15...950 K: see	68Iwa 92Sha
c	Piezooptic effect: Table 2A-2-011, Table 2A-2-012.	
d	Faraday effect: see Fig. 2A-1-091 in No. 2A-1.	
e	Nonlinear optical property: $d_{22} / d_{36}^{KDP} = 4.4$, $d_{31} / d_{36}^{KDP} = d_{15} / d_{36}^{KDP} = -2.7$, $d_{33} / d_{36}^{KDP} = -41$ at RT for $\lambda = 1.058$ μm. Nonlinear millimeter wave susceptibility:	69Bec, 70Mil

	$d_{33} = 18.62 \cdot 10^{-9} \text{ mV}^{-1}$ for $\lambda = 857 \text{ }\mu\text{m}$ ($f = 35 \text{ GHz}$).	83Ahn
	Nonlinear optical properties of proton-exchanged waveguide: see	94Ahl1
	Blue light generation in a waveguide by quasi-phase-matched second harmonic generation: see	91Miz1
	Characteristics of periodically domain-inverted (by proton-exchange) waveguides for second harmonic generation: see	91Yam
10a	Raman scattering: Table 2A-2-013; Figs. 2A-2-049...2A-2-058; see also	84Men, 87Yan, 87Vor
	New low frequency Raman modes from proton-exchanged crystals: see	95WuX
	Raman scattering in Nd-doped crystal: see	94WuX1
	Raman spectra showed that LiTaO ₃ becomes amorphous above 33.2 GPa: see	94Lin
	Raman scattering under high pressure: see	86Jay
	Hyper Raman scattering: Fig. 2A-2-059, Fig. 2A-2-060.	
	Stimulated Raman scattering by femtosecond laser pulses: Fig. 2A-2-061, Fig. 2A-2-062.	
	Observation of THz phonon-polariton beats by femtosecond laser pulses: see	92Bak
	Brillouin scattering: Table 2A-2-014.	
	Rayleigh scattering: Fig. 2A-2-063, see also	86Zha
11	Resistivity: $\rho = 3.6 \cdot 10^8 \text{ }\Omega\text{m}$ at 25 °C, 1 kHz.	75Bee
	Conductivity: Fig. 2A-2-064, Fig. 2A-2-065; see also Fig. 2A-2-007 in 5a.	
	Bulk and surface resistivity: see	94Len
	Humidity dependence of surface resistance: see	92Mae
	Luminescence from Cr ³⁺ ion: see	69Gla
	Exoelectron emission: see	80Ros
	Seebeck effect: Fig. 2A-2-066.	
	Photovoltaic effect: Fig. 2A-2-067.	
13a	NMR: $e^2qQ/h = 77.6(5) \text{ kHz}$ for ⁷ Li, $e^2qQ/h = 60.2(3) \text{ kHz}$ for ⁸ Li. Fig. 2A-2-073.	68Pet 77Dub
	NMR of Li: see also	85Sen
b	ESR: Table 2A-2-015, Table 2A-2-016. ESR of Ti ³⁺ : $g_{\parallel} = 1.948(3)$, $g_{\perp} = 1.827(3)$. ENDOR of Fe ³⁺ : $g = 1.995(2)$ (nearly isotropic). $3B_2^0 = 9900(100) \text{ MHz}$. ESR spin Hamiltonian parameters of Co ²⁺ : $g_{\parallel} = 2.38$, $g_{\perp} = 5.02$, $A_{\parallel} \approx 0$, $A_{\perp} = 133 \cdot 10^{-2} \text{ m}^{-1}$.	94Thi 89Sot 87Don
c	Mössbauer effect (⁵⁷ Fe): $e^2qQ(7/2) = 46.4(2) \text{ mm s}^{-1} = 9.59(4) \cdot 10^{-7} \text{ eV}$, $\delta = -17.95(10) \text{ mm s}^{-1}$. Figs. 2A-2-069...2A-2-072. Mössbauer effect of ⁵⁷ Co: see	78Wor 90Leu, 90And
14	Small ionic displacements under ac electric fields applied along the <i>c</i> -axis have been observed by X-ray diffraction. EXAFS of Co and Fe doped crystals: see	82Fuj 91Cat1
15a	Domain structure: see Observation of 180° domain walls by high resolution electron microscopy: see Poling method with an aid of electron beams: see	66Lev 82Lin 86Hay

Domain thickness: Fig. 2A-2-073.	
Domain inversion using proton-exchange followed by heat treatment: see	94Miz1
Periodic domain inversion by electron beam scanning: see	94Gup
Fabrication of periodically domain-inverted structure by proton-exchange and heat treatment: see	91Miz2
Characteristics of periodically domain-inverted (by proton-exchange) waveguides for second harmonic generation: see	91Yam
Fabrication of periodically domain-inverted structure by proton-exchange with pyrophosphoric acid: see	92Miz1
Quasi-phase-matched second-harmonic generation using a periodically domain-inverted waveguide: see	92Miz2
Fabrication of periodically inverted domain structures using proton-exchange: see	92Mak

16 Investigation of the effect of internal heat transfer on growth behavior in Czochralski crystal growth process: see	94Oka
Direct bonding of single crystals: see	94Tom
Fabrication of periodically inverted domain structures using proton-exchange: see	92Mak
Vapor diffused Zn-doped optical waveguides: see	88Yoo
Fabrication of alternating LiNbO ₃ /LiTaO ₃ layers: see	87Kan
Surface layer: surface breakdown by laser pulses: see	72Zve
Electron spectroscopic studies of surfaces: see	81Rit
Etchant: mixture of hydrofluoric acid and nitric acid at its boiling point: see	66Lev
Optical waveguide: many fabrication techniques have been reported;	
by out-diffusion, see	73Kam
by in-diffusion, see	74Ham,
	74Nod
by sputtering , see	73Gia
a review paper, see	75Tie
Formation of ion exchanged LiTaO ₃ :Cu waveguides: see	91Bob
Blue light generation by frequency doubling of a laser diode in a periodically domain-inverted waveguide: see	92Yam2
Quasi-phase-matched waveguides fabricated by domain inversion with proton-exchange and heat treatment technique: see	92Yam3
Quasi-phase-matched second-harmonic generation in a waveguide: see	94Miz2
Blue light generation in a waveguide by quasi-phase-matched second harmonic generation: see	91Miz1
Quasi-phase-matched second-harmonic generation using a periodically domain-inverted waveguide: see	92Miz2
Optical damage: change of refractive indices occurs in the region illuminated by laser beams: see	66Ash
Optical waveguide fabrication by proton-exchange: see	83Spi
Photoinduced light scattering: see	94WuX2
Photorefractive effects in proton-exchanged optical waveguides: see	92McW
X-ray topographic study: see	82Yas1
¹⁸¹ Hf → ¹⁸¹ Ta perturbed-angular correlation spectroscopy: Fig. 2A-2-074.	

Table 2A-2-001. LiTaO₃. Fractional atomic coordinates and anisotropic temperature parameters at various temperatures refined by the method of least squares fitting in space group R3c [73Abr]. b_{ij} is defined by Eq. (b) in Introduction.

Parameter	297 K	760 K	820 K	885 K	940 K
$z(\text{Li})$	0.2790(17)	0.2812(26)	0.2818(19)	0.2765(15)	0.2761(14)
$b_{11}(\text{Li})$	0.0106(61)	0.0349(101)	0.0205(56)	0.0354(68)	0.0578(82)
$b_{33}(\text{Li})$	0.0043(10)	0.0071(21)	0.0093(18)	0.0047(12)	0.0077(18)
$b_{11}(\text{Ta})$	0.0055(17)	0.0062(17)	0.0085(14)	0.0118(16)	0.0195(13)
$b_{33}(\text{Ta})$	0.0003(1)	0.0007(2)	0.0009(1)	0.0011(1)	0.0017(1)
$x(\text{O})$	0.0501(5)	−0.0543(5)	0.0547(4)	0.0557(5)	0.0540(5)
$y(\text{O})$	0.3436(3)	0.3396(5)	0.3394(4)	0.3381(5)	0.3345(11)
$z(\text{O})$	0.0687(3)	0.0756(3)	0.0786(2)	0.0835(3)	0.0817(5)
$b_{11}(\text{O})$	0.0056(12)	0.0126(12)	0.0162(12)	0.0206(19)	0.0159(15)
$b_{22}(\text{O})$	0.0038(14)	0.0099(11)	0.0109(10)	0.0118(11)	0.0149(8)
$b_{33}(\text{O})$	0.0011(2)	0.0014(2)	0.0020(2)	0.0021(1)	0.0027(1)
$b_{12}(\text{O})$	0.0019(12)	0.0054(10)	0.0069(9)	0.0105(15)	0.0052(15)
$b_{13}(\text{O})$	−0.0004(4)	−0.0010(4)	−0.0008(3)	−0.0021(4)	−0.0018(6)
$b_{23}(\text{O})$	−0.0007(2)	−0.0021(2)	−0.0022(2)	−0.0031(1)	−0.0032(1)
R	0.0335	0.0526	0.0431	0.0486	0.0320

Table 2A-2-002. LiTaO₃. Fractional atomic coordinates and anisotropic temperature parameters at 885 K and 940 K in centrosymmetric space group $R\bar{3}c$ (a) and $R\bar{3}$ (b) [73Abr]. b_{ij} is defined by Eq. (b) in Introduction.

Parameter	885 K (a)	940 K (a)	940 K (b)
$z(\text{Li})$	0.2833(30)	0.2772(15)	0.2501(46)
$b_{11}(\text{Li})$	0.0820(209)	0.0535(84)	0.0551(105)
$b_{33}(\text{Li})$	0.0063(14)	0.0075(17)	0.0270(42)
$b_{11}(\text{Ta})$	0.0151(23)	0.0198(14)	0.0200(21)
$b_{33}(\text{Ta})$	0.0009(2)	0.0017(1)	0.0016(2)
$x(\text{O})$	0.0531(5)	0.0531(2)	0.0534(4)
$y(\text{O})$	1/3	1/3	0.3339(5)
$z(\text{O})$	1/12	1/12	0.0834(2)
$b_{11}(\text{O})$	0.0200(22)	0.0271(12)	0.0182(10)
$b_{22}(\text{O})$	0.0100(16)	0.0158(9)	0.0152(10)
$b_{33}(\text{O})$	0.0021(2)	0.0027(1)	0.0028(1)
$b_{12}(\text{O})$	$b_{22}/2$	$b_{22}/2$	0.0075(7)
$b_{13}(\text{O})$	$b_{23}/2$	$b_{23}/2$	−0.0016(3)
$b_{23}(\text{O})$	−0.0030(2)	−0.0033(2)	−0.0032(2)
R	0.0544	0.0339	0.0349

Table 2A-2-003. LiTaO₃, Interatomic distances and angles at 24 °C [67Abr1, 67Abr2].

Distance ^{a)} [Å]		Distance ^{a)} [Å]		Distance ^{b)} [Å]		Distance ^{b)} [Å]	
Ta–Ta	3.759(0)	O–O	2.734(25)	Ta–Ta	3.759(0)	O–O	2.724(3)
Li–Li	3.759(0)		2.792(6)	Li–Li	3.759(0)		2.791(1)
Ta–O	1.891(10)		2.812(7)	Ta–O	1.908(3)		2.826(1)
	2.070(10)		2.823(22)		2.073(3)		2.870(3)
			3.064(9)				3.039(2)
Ta–Li	3.003(40)		3.425(18)	Ta–Li	3.046(23)		3.378(4)
	3.058(9)				3.068(6)		
	3.374(19)	Li–O	2.076(16)		3.354(11)	Li–O	2.041(7)
	3.888(40)		2.293(32)		3.845(23)		2.312(16)
Angle ^{a)} [°]		Angle ^{a)} [°]		Angle ^{b)} [°]		Angle ^{b)} [°]	
O–Ta–O	82.7(5)	O–Li–O	73.2(12)	O–Ta–O	82.2(1)	O–Li–O	72.2(6)
	89.5(2)		79.2(5)		88.9(1)		79.5(2)
	90.4(2)		88.9(6)		90.4(1)		88.3(3)
	96.6(4)		111.1(10)		97.5(2)		111.9(5)
Average	89.8	Average	88.1	Average	89.8	Average	87.9

^{a)} X-ray diffraction [67Abr1]. ^{b)} Neutron diffraction [67Abr2].

Table 2A-2-004. LiTaO₃. Elastic, piezoelectric, dielectric constants and their temperature coefficients. For definition of $a^{(1)}$, $a^{(2)}$, see original papers [71Smi, 67War1, 69Yam].

	Absolute quantities			Normalized temperature coefficients [71Smi]	
	[71Smi]	[67War1]	[69Yam]	$a^{(1)}$	$a^{(2)}$
Elastic stiffness	[$\cdot 10^{11}$ Nm ⁻²]	[$\cdot 10^{11}$ Nm ⁻²]	[$\cdot 10^{11}$ Nm ⁻²]	[$\cdot 10^{-4}$ K ⁻¹]	[$\cdot 10^{-7}$ K ⁻²]
c_{11}^E	2.298	2.33	2.28	-1.03	0.77
c_{12}^E	0.440	0.47	0.31	-3.41	-1.18
c_{13}^E	0.812	0.80	0.74	-0.50	6.00
c_{14}^E	-0.104	-0.11	-0.12	6.67	16.7
c_{33}^E	2.798	2.75	2.71	-0.96	-3.21
c_{44}^E	0.968	0.94	0.96	-0.43	1.67
c_{66}^E	0.929	0.93	0.98	-0.47	1.24
Elastic compliances	[$\cdot 10^{-12}$ m ² N ⁻¹]	[$\cdot 10^{-12}$ m ² N ⁻¹]	[$\cdot 10^{-12}$ m ² N ⁻¹]		
s_{11}^E	4.930	4.87	4.86	1.11	0.966
s_{12}^E	-0.519	-0.58	-0.29	-3.83	-16.9
s_{13}^E	-1.280	-1.25	-1.24	2.14	12.8
s_{14}^E	0.588	0.64	0.63	7.74	15.0
s_{33}^E	4.317	4.36	4.36	1.24	6.72
s_{44}^E	10.46	10.8	10.5	0.60	-1.18
s_{66}^E	10.90	10.9	10.3	0.64	-0.73
Piezoelectric stress constants	[Cm ⁻²]	[Cm ⁻²]	[Cm ⁻²]		
e_{15}	2.72	2.6	2.7	-1.32	-7.17
e_{22}	1.67	1.6	2.0	-0.60	-6.28
e_{31}	-0.38	0.0	-0.1	0.87	51.8
e_{33}	1.09	1.9	2.0	1.54	1.41
Piezoelectric strain constants	[$\cdot 10^{-11}$ CN ⁻¹]	[$\cdot 10^{-11}$ CN ⁻¹]	[$\cdot 10^{-11}$ CN ⁻¹]		
d_{15}	2.64	2.6	2.6	-1.31	-9.64
d_{22}	0.75	0.7	0.85	-1.32	-9.79
d_{31}	-0.30	-0.2	-0.30	+3.27	43.1
d_{33}	0.57	0.8	0.92	+2.74	118.0
Dielectric constants					
κ_{11}^S	42.6	41	41	3.29	4.28
κ_{33}^S	42.8	43	42	11.6	78.0
κ_{11}^T	53.6	51	53	2.11	0.100
κ_{33}^T	43.4	45	44	11.47	80.8

Table 2A-2-005. LiTaO₃. Values of dielectric constant at 293 K. Parameter: f . ^{a)}[88Tom], ^{b)}[75Tea], ^{c)}[70Bar], ^{d)}[67Kam].

f	Capacitance measurements ^{a)}					^{b)}
	10 kHz	100 kHz	1 MHz	10 MHz	13 MHz	1 GHz
κ_{11}	53.5	53.5		41.7	40.6	40.3
κ_{33}	42.4	42.2	42.0	39.2	38.8	41.4
	Infrared reflectivity ^{c)}			Raman scattering ^{d)}		
κ_{11}^S	41.5				41	
κ_{33}^S	37.6				43	

Table 2A-2-006. LiTaO₃. $c_{\lambda\mu}$, $e_{i\lambda}$. $c_{\lambda\mu}$: elastic stiffness; $e_{i\lambda}$: piezoelectric stress constant.

Ref.	94Tak	91Mur	67War1
c_{11} [$\cdot 10^{11}$ Nm ⁻²]	2.332	2.178	2.33
c_{13}	0.837	0.643	0.80
c_{14}	-0.106	-0.114	-0.11
c_{33}	2.781	2.494	2.75
c_{44}	0.948	0.963	0.94
c_{66}	0.935	0.950	0.93
e_{15} [Cm ⁻²]	2.56	(2.6)	2.6
e_{22}	1.74	(1.6)	1.6
e_{33}	1.80	(1.9)	1.9
e_{31}	-0.21	(0.0)	0.0

Table 2A-2-007. LiTaO₃. Summary of design data for microwave surface acoustic wave devices [73Sza]. prop: propagation, Vac attn: (surface wave) attenuation in vacuum, Sw: surface wave, bw: bandwidth.

Orientation	Z cut Y prop	Y cut Z prop	22° ro- tated cut X prop	Orientation	Z cut Y prop	Y cut Z prop	22° ro- tated cut X prop
Surface wave velocity v_{∞} [m s ⁻¹]	3329	3230	3302	3 dB Air prop loss time delay at 1 GHz, A [μs]	3.0	2.6	
Estimate of electro- magnetic to acoustic coupling $\Delta v/v_{\infty}$	0.0059	0.0033	0.0027	Slope of electro- mechanical power flow curve $\partial\phi/\partial\theta$ ^{b)}	-1.241	-0.211	+0.764
Measured value of $\Delta v/v_{\infty}$ ^{a)}	0.0045	0.0036		Slope of electro- mechanical power flow curve $\partial\phi/\partial\mu$ ^{c)}	+0.556	-0.229	0
Power flow angle ϕ [°] (electromechanical)	0	0	0	3 dB Beam steering loss time delay B [μs]	4.4	15.6	13.3
Temperature coefficient of delay $\frac{1}{\tau} \frac{\partial\tau}{\partial T}$ [$\cdot 10^{-6}\text{K}^{-1}$]	69.0	35.0	68.0	3 dB Diffraction loss time delay at 1 GHz, C [μs]	11.3	3.6	1.6
Sw attn in air at 1 GHz [dB/μs]	1.0	1.14		Material figure of merit F_M $F_M = ABC \left(\frac{\Delta v}{v_{\infty}} \right)^2$	0.005	0.002	
Air loading at 1 GHz [dB/μs]	0.23	0.20		Time-bw product Figure of merit F_{TB} $F_{TB} = \frac{4 \cdot 10^3 ABC}{v_{\infty}} \cdot \left(\frac{\Delta v}{v_{\infty}} \right)^2$	0.006	0.002	
Vac attn at 1 GHz [dB/μs]	0.77	0.94					

^{a)} [72Sch2]. ^{b)} θ : direction of propagation. ^{c)} μ : direction of plate normal.

Table 2A-2-008. LiTaO₃. Surface acoustic wave properties of *X*-cut 112° rotated *Y* propagation mode [82Yas2]. TCD: temperature coefficient of propagation delay time. IDT: interdigital transducer.

v_s	k_s^2	Coupling per unit length of IDT	TCD
3295 ms ⁻¹	8·10 ⁻³	6.5 pF/cm pair	-18·10 ⁻⁶ K ⁻¹

Table 2A-2-009. LiTaO₃, Refractive indices [65Bon].

λ [μm]	n_o	n_e	λ [μm]	n_o	n_e
0.45	2.2420	2.2468	2.00	2.1066	2.1115
0.50	2.2160	2.2205	2.20	2.1009	2.1053
0.60	2.1834	2.1878	2.40	2.0951	2.0993
0.70	2.1652	2.1696	2.60	2.0891	2.0936
0.80	2.1538	2.1578	2.80	2.0825	2.0871
0.90	2.1454	2.1493	3.00	2.0755	2.0299
1.00	2.1391	2.1432	3.20	2.0680	2.0727
1.20	2.1305	2.1341	3.40	2.0601	2.0649
1.40	2.1236	2.1273	3.60	2.0513	2.0561
1.60	2.1174	2.1213	3.80	2.0424	2.0473
1.80	2.1120	2.1170	4.00	2.0335	2.0377

Table 2A-2-010. LiTaO₃. Electrooptic constants.

	r_{13}	r_{33}	r_{51}	r_{22}	r_c	λ	Ref.
	[$\cdot 10^{-12}$ m V ⁻¹]					[μ m]	
r^T					22	0.633	66Len
	8.4	30.5				0.633	72Onu
r^S	7	30.3	20	≈ 1		0.633	66Len
	4.5	27	15	≈ 0.3		3.39	66Tur

Table 2A-2-011. LiTaO₃. Piezooptic constants for $\lambda = 633$ nm.

	[76Ava]	[67Dix]		[76Ava]	[67Dix]
p_{11}	−0.081(3)	0.0804	p_{33}	−0.044(4)	0.150
p_{12}	0.081(3)	0.0804	p_{41}	−0.085(6)	0.024
p_{13}	0.093(2)	0.094	p_{44}	0.028(2)	0.022
p_{14}	−0.026(2)	0.031	p_{66}	−0.081(4)	
p_{31}	0.089(4)	0.086			

Table 2A-2-012. LiTaO₃. Pressure derivatives of refractive indices [81Cor].

λ [nm]	dn_o/dp [$\cdot 10^{-2}$ GPa ⁻¹]	dn_e/dp [$\cdot 10^{-2}$ GPa ⁻¹]
509.0	0.088	0.209
546.1	0.099	0.221
598.3	0.109	0.227

Table 2A-2-013. LiTaO₃. Assignment of Raman active modes (wave number shifts in cm⁻¹) in the ferroelectric phase (290 K). The superscripts \perp , \parallel and $/$ indicate directions of phonon propagation perpendicular, parallel, and at 45° relative to the unique axis.

A₁ Modes:

[67Kam] TO	LO	[76Pen1] TO	LO	[87Yan] TO	LO	[88Rap] TO	LO
201	245	81	82	206	252	203 ^{\perp}	250 ^{\parallel}
253	347	186	200	253	354	252 ^{\perp}	287 ^{\parallel}
356	399	202	246	356	402	356 ^{\perp}	438 ^{\parallel}
600	864	250	354	600	865	597 ^{\perp}	864 ^{\parallel}
		355	401				
		458	465				
		594	660				
		661	750				
		751	866				

E Modes:

[67Kam] TO	LO	[76Pen1] TO	LO	[87Yan] TO	LO	[88Rap] TO	LO
74	80	69	70	142	142	142 ^{\perp/\parallel}	142 ^{\perp}
140	163	141	200	142	190	(142 ^{\perp?)}	194 ^{\perp}
206	248	249	275	206	206	207 ^{\perp/\parallel}	278 ^{\perp}
251	278	316	345	252	280	(314 ^{\perp?)}	314 ^{\perp}
316	318	345	379	315	315	314 ^{\perp/\parallel}	344 ^{\perp}
383	452	382	459	315	380	355 ^{\perp/\parallel}	378 ^{\perp}
462	474	462	470	382	463	382 ^{\perp/\parallel}	445 ^{\perp}
596	648	591	659	463	463	463 ^{\perp/\parallel}	463 ^{\perp}
662	870	660	866	592	866	592 ^{\perp/\parallel}	865 ^{\perp}

Table 2A-2-014. LiTaO₃. Brillouin scattering [71Kha]. Geometry [for example $z(xz)y$] from left to right: propagation direction of the incident light, polarization of the incident light, polarization of the scattered light and propagation direction of the scattered light. Direction: phonon propagation direction, $\Delta\nu$: phonon frequency, *: intense lines, v_{exp} : experimental phonon velocity, P: phonon polarization, ql: quasi-longitudinal, qt: quasi-transverse, t: transverse.

Geometry	Direction	$\Delta\nu/c$ [$\cdot 10^2 \text{ m}^{-1}$]	v_{exp} [$\cdot 10^3 \text{ m s}^{-1}$]	P
$x(zx)\bar{y}$	$k_x = -\frac{\sqrt{2}}{2}; k_y = -\frac{\sqrt{2}}{2}$	$\begin{cases} 1.15 \\ 0.72 \end{cases}$	$\begin{cases} 6.89 \\ 4.32 \end{cases}$	$\begin{cases} \text{ql} \\ \text{qt} \end{cases}$
$x(zz)\bar{y}$	$k_z=0$	1.10	6.71	ql
$y(zy)x$	$k_x = \frac{\sqrt{2}}{2}; k_y = -\frac{\sqrt{2}}{2}$	0.66*	3.96	qt
$y(zz)x$	$k_z=0$	1.10	6.71	ql
$z(xz)y$	$k_x=0; k_y = \frac{\sqrt{2}}{2}$	0.58*	3.48	t
$z(xx)y$	$k_z = -\frac{\sqrt{2}}{2}$	1.21	7.11	ql
$x(xz)\bar{y}$	$k_x=0$	0.59*	3.54	t
$x(xx)\bar{y}$	$k_y = -\frac{\sqrt{2}}{2}$	1.23	7.23	ql
$x(yz)\bar{y}$		0.59	3.54	qt
$x(yx)\bar{y}$	$k_z = -\frac{\sqrt{2}}{2}$	0.61	3.58	qt
$z(yz)x$	$k_x = -\frac{\sqrt{2}}{2}; k_y=0$	0.62*	3.72	qt
$z(yy)x$	$k_z = -\frac{\sqrt{2}}{2}$	$\begin{cases} 0.64 \\ 1.22 \end{cases}$	$\begin{cases} 3.76 \\ 7.17 \end{cases}$	$\begin{cases} \text{qt} \\ \text{ql} \end{cases}$

Table 2A-2-015. LiTaO₃, ESR data.

Paramagnetic center	Cr ³⁺	Mn ²⁺
H	(4)	(8)
ν [GHz]		X and K band
T		RT
g -factor	$g_{\parallel} = 1.971$ $g_{\perp} = 3.863$	$g_{\parallel} = g_{\perp} = 1.995(5)$
FS [$\cdot 10^{-2} \text{ m}^{-1}$]	$D = 4300(500)$	$b_{20} = 1694(14)$ $b_{40} = -12(16)$
HFS [$\cdot 10^{-2} \text{ m}^{-1}$]		$A_{\parallel} = -78.2(5)$ $A_{\perp} = -76.4(5)$
Ref.	68Bur	68Dan

Table 2A-2-016. LiTaO₃. Spin Hamiltonian parameters for Cu²⁺ center [85Kha]. $T = 77$ K.

Principal values of g -tensor	Principal values of A -tensor [$\cdot 10^{-2} \text{ m}^{-1}$]
$g_z = 2.396(4)$	$A_z = 70(1)$
$g_y = 2.099(5)$	$A_y = 50(2)$
$g_x = 2.050(4)$	$A_x = 56(1)$
$g'_{\parallel} = 2.19(1)$	
$g'_{\perp} = 2.16(1)$	

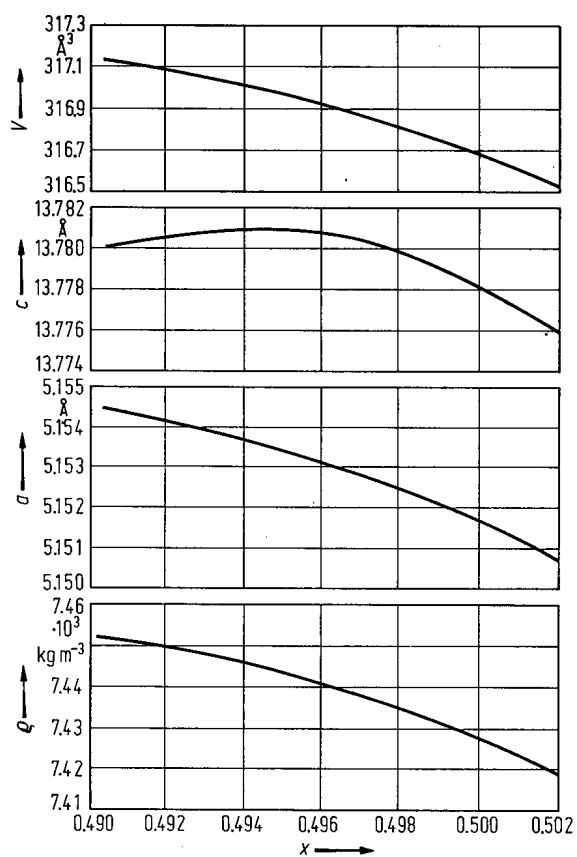


Fig. 2A-2-001. LiTaO₃. Lattice parameters and density of crystals grown from x Li₂O·(1- x)Ta₂O₅ melt vs. x [70Bar1].
 V : unit cell volume, ρ : density.

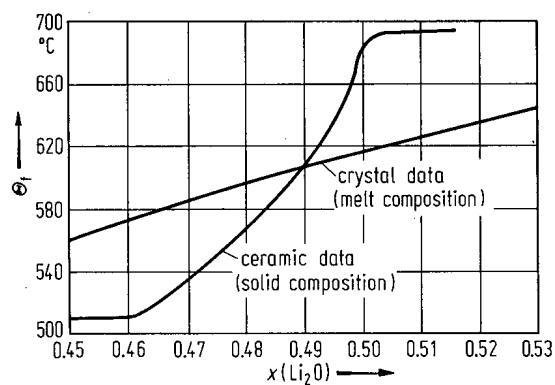


Fig. 2A-2-002. LiTaO₃. Θ_f vs. x . Crystal data: Θ_f of crystals grown from $x \text{ Li}_2\text{O} \cdot (1-x) \text{ Ta}_2\text{O}_5$ melt [67Bal]. Ceramic data: Θ_f of ceramics sintered from $x \text{ Li}_2\text{O} \cdot (1-x) \text{ Ta}_2\text{O}_5$ system [70Bar1]. See also [68Yam1, 71Miy].

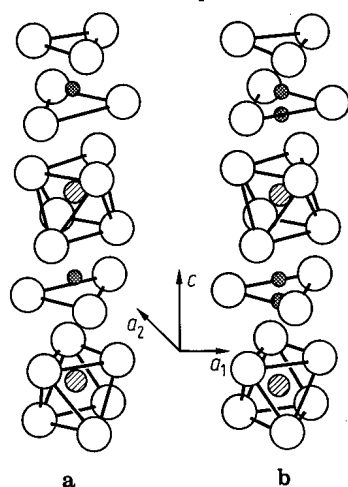


Fig. 2A-2-003. LiTaO₃. **(a)** Polar arrangement of oxygen atom (large circle) octahedra in LiTaO₃ stacked along the trigonal axis. The octahedra contain, in sequence, a Ta atom (hatched circles), no metal atom, and Li atom (cross-hatched circles). **(b)** Arrangement of atoms above the ferroelectric transition temperature within one stack of octahedra. The Ta atom is at an octahedron center and the Li atom is distributed within the next two octahedra [73Abr].

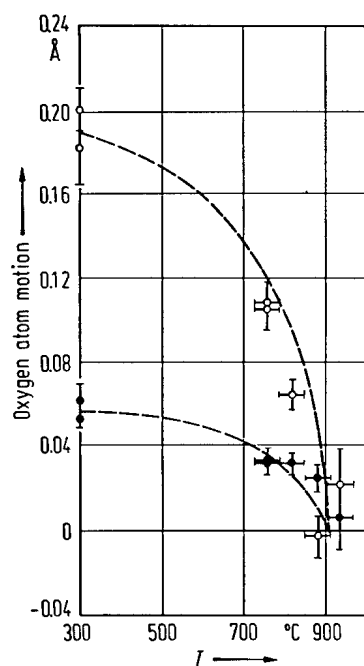


Fig. 2A-2-004. LiTaO₃. Variation of oxygen shifts vs. T [73Abr]. Full circles: $y(O) - 1/3$, open circles: $1/12 - z(O)$. The dashed lines are the variation of P_s with T , normalized to the diffraction data at 298 °C and 907 °C.

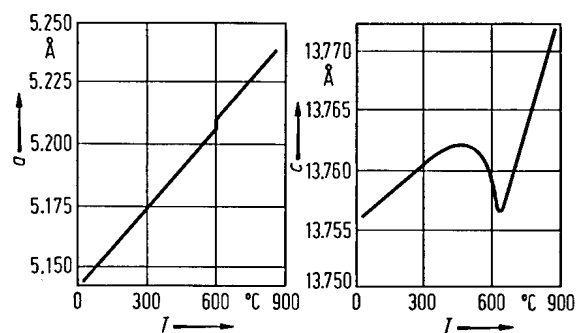


Fig. 2A-2-005. LiTaO₃. a , c vs. T [67Iwa]. Thermal expansion was measured by the dilatometric method.

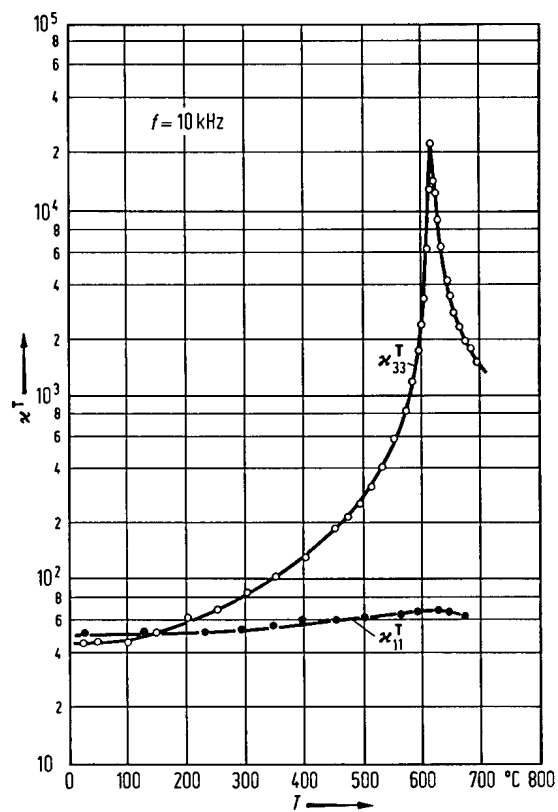


Fig. 2A-2-006. LiTaO₃. κ_{33}^T , κ_{11}^T vs. T [68Yam2]. $f = 10$ kHz.

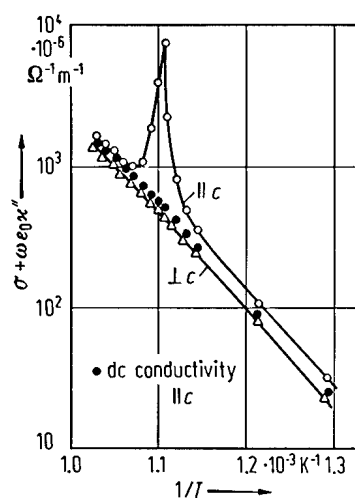


Fig. 2A-2-007. LiTaO₃. $\sigma + \omega\epsilon_0\kappa''$ vs. T^{-1} [69Yam]. Open circles and triangles: $f = 10$ kHz. Full circles: $f = 0$ Hz.

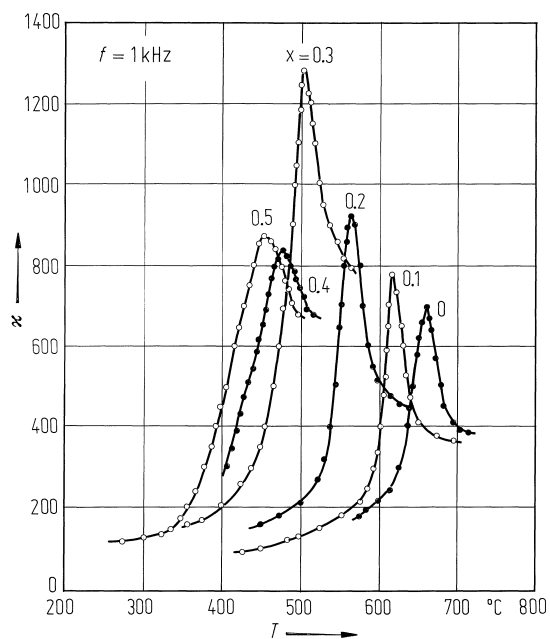


Fig. 2A-2-008. Li_{1-x}Zn_{x/2}TaO₃ (ceramics). κ vs. T [83Tor].
Parameter: x . $f = 1$ kHz.

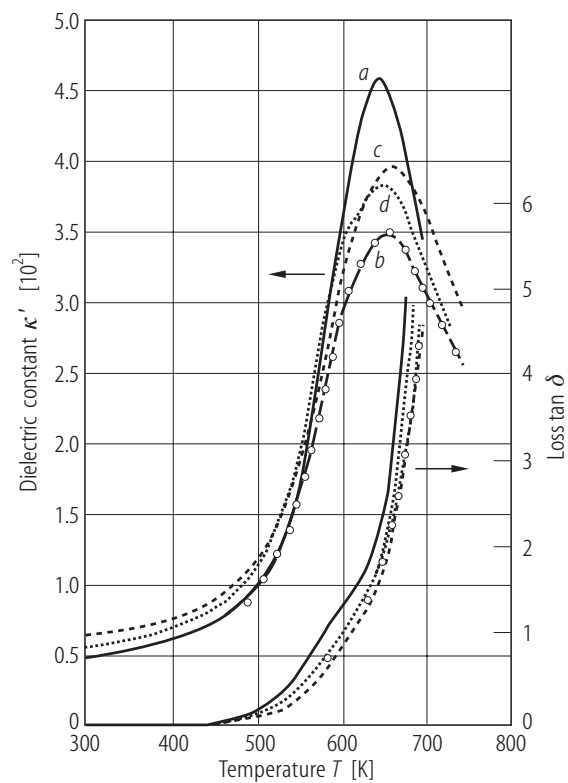


Fig. 2A-2-009. LiTaO₃ (ceramics). κ' , $\tan \delta$ vs. T [90InH].
Sintering time: *a*: 4 h, *b*: 2 h, *c*: 0.5 h, *d*: 0.25 h. $f = 1$ kHz.

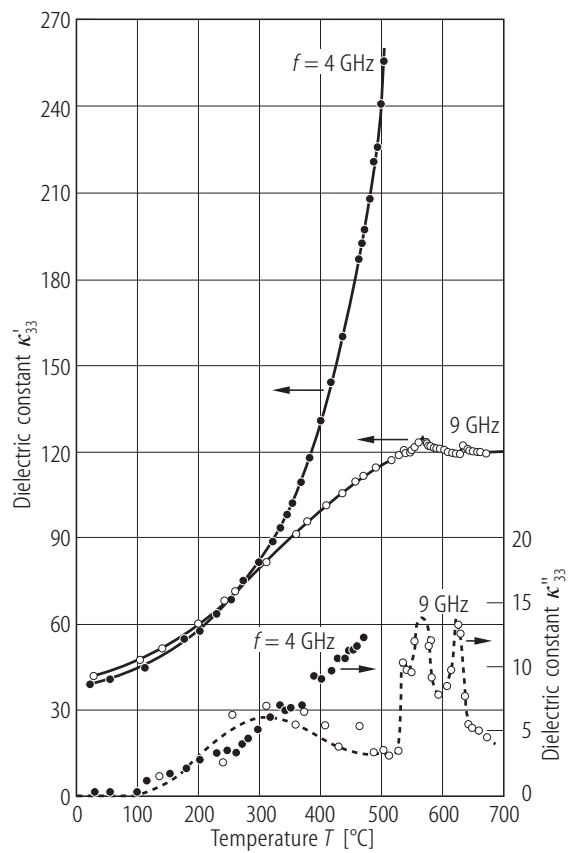


Fig. 2A-2-010. LiTaO₃. κ'_{33} , κ''_{33} vs. T [70Saw]. Parameter: f .

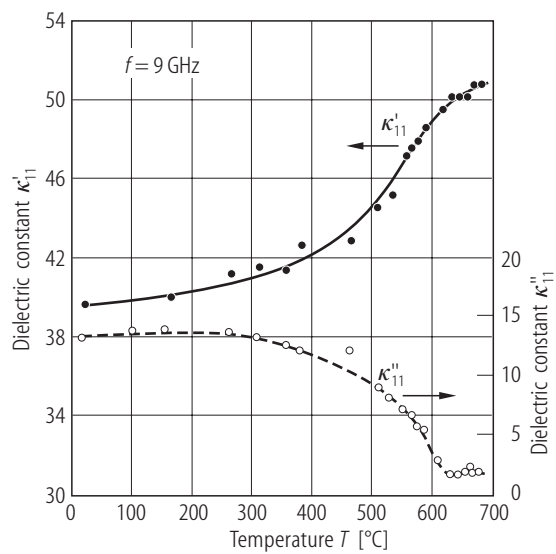


Fig. 2A-2-011. LiTaO₃. κ'_{11} , κ''_{11} vs. T [70Saw]. $f = 9$ GHz.

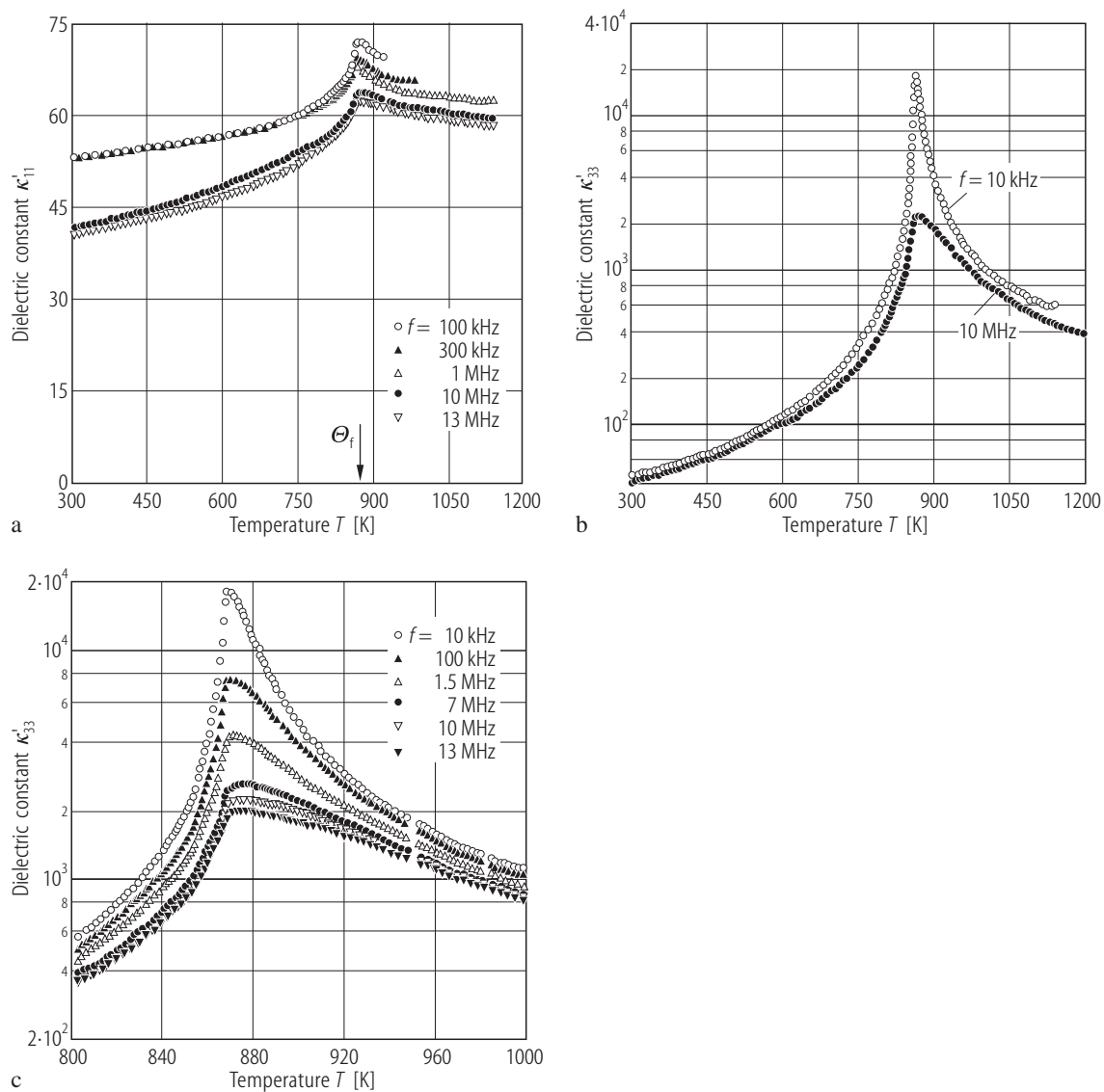


Fig. 2A-2-012. LiTaO₃. κ'_{ii} vs. T [88Tom]. Parameter: f .

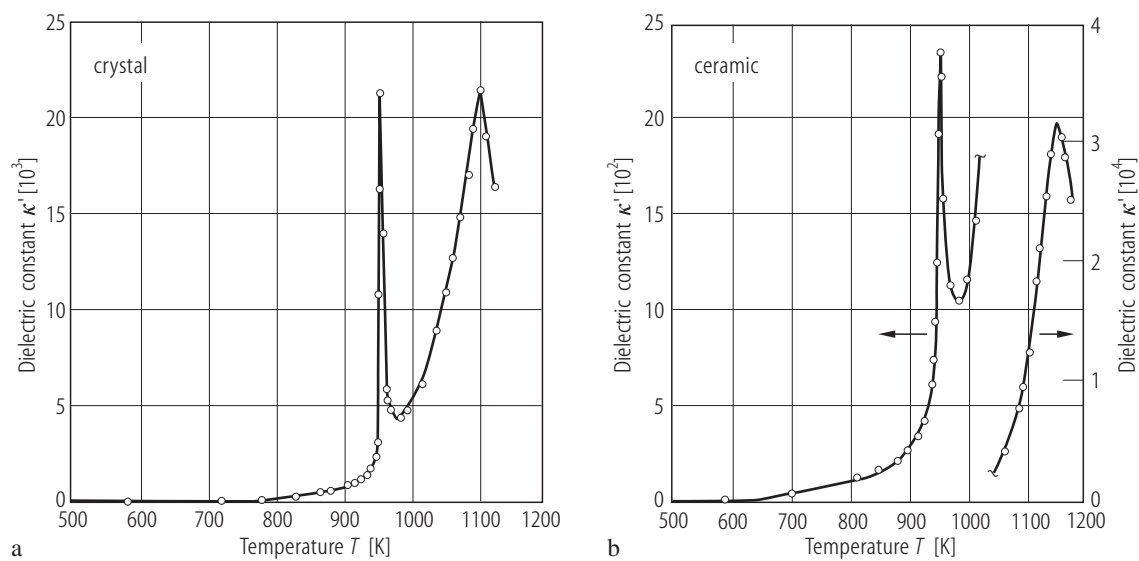


Fig. 2A-2-013. LiTaO₃. κ' vs. T [90YeZ]. $f = 1$ kHz. (a) single crystal. (b) ceramics.

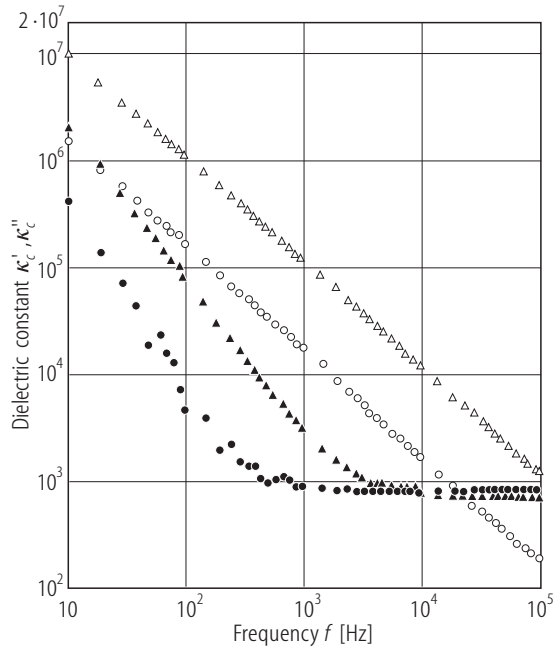


Fig. 2A-2-014. LiTaO₃. κ'_c , κ''_c vs. f [93Tou]. Triangles: proton-exchanged specimen with exchanged layer of about 3 μm thick. Circles: unprocessed specimen. Full triangles, full circles: κ'_c . Open triangles, open circles: κ''_c .

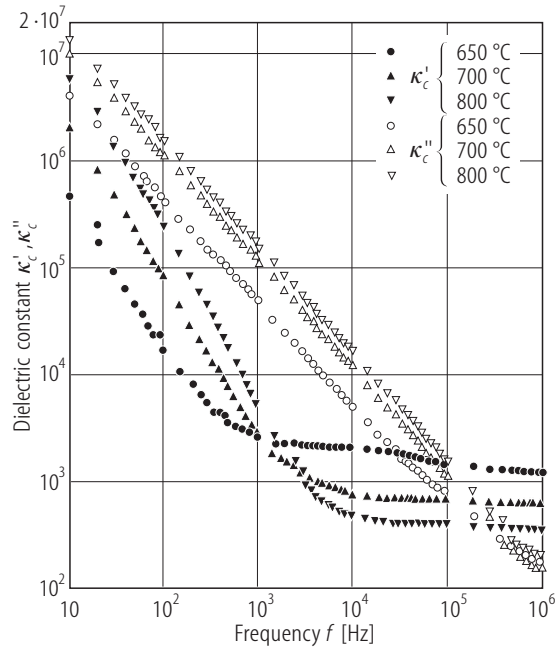


Fig. 2A-2-015. LiTaO₃. κ'_c , κ''_c vs. T of proton-exchanged specimens [93Tou]. Parameter: T . Proton-exchanged layer was estimated to be about 3 μm thick.

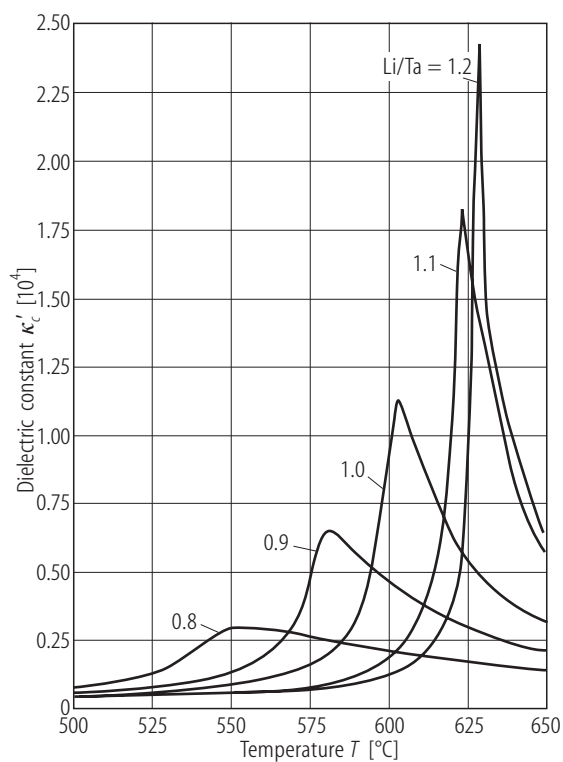


Fig. 2A-2-016. LiTaO₃. κ'_c vs. T in various compositions [71Fuj]. $f = 1$ MHz. Parameter: Li / Ta ratio.

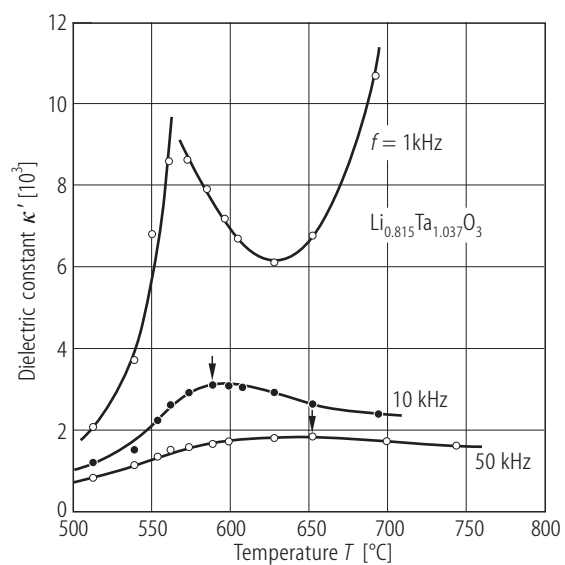


Fig. 2A-2-017. LiTaO₃ (nonstoichiometric ceramic). κ' vs. T [87Hua]. Parameter: f . Arrows indicate maximum κ' points.

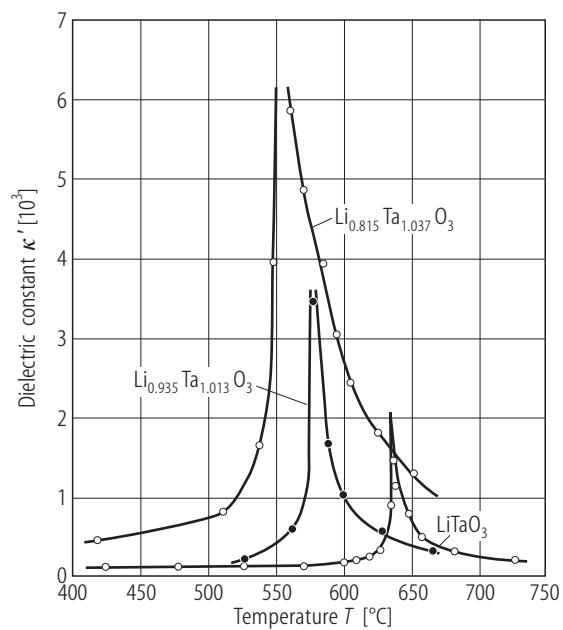


Fig. 2A-2-018. LiTaO₃ (ceramics). κ' vs. T [87Hua].
Parameter: Li/Ta stoichiometry.

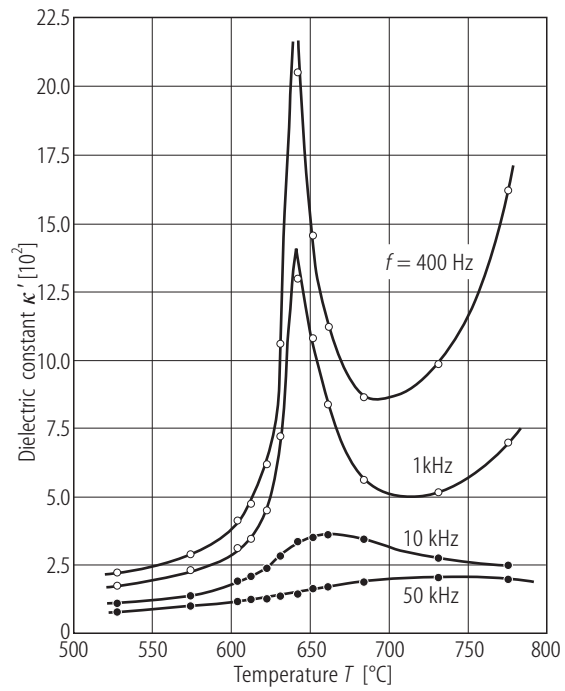


Fig. 2A-2-019. LiTaO₃ (stoichiometric ceramic). κ' vs. T [87Hua]. Parameter: f .

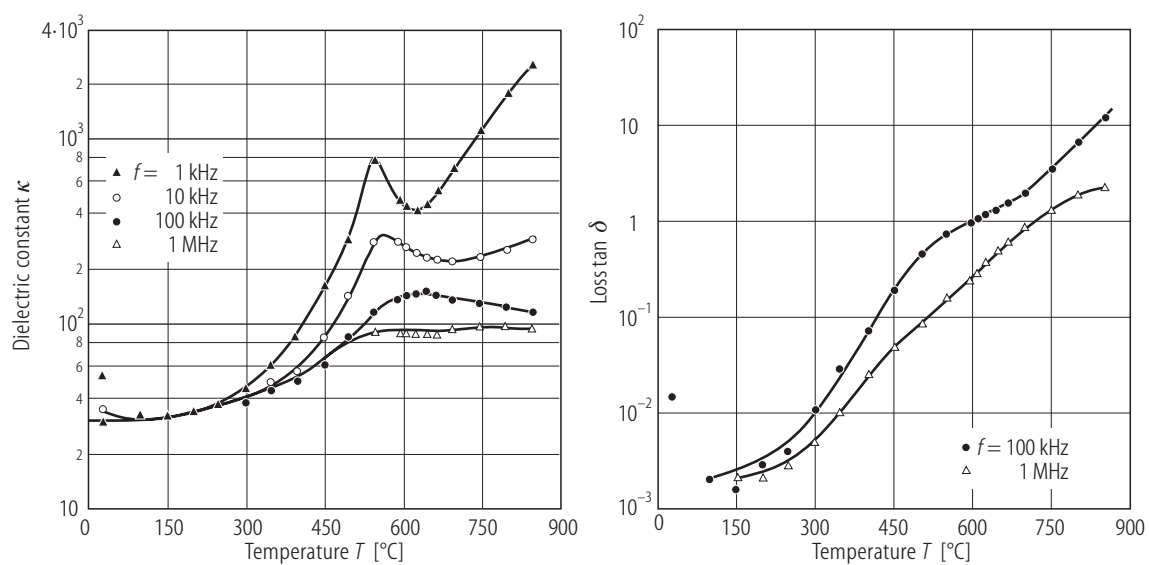


Fig. 2A-2-020. LiTaO₃ (ceramic). κ , $\tan \delta$ vs. T [91Chi]. Parameter: f .

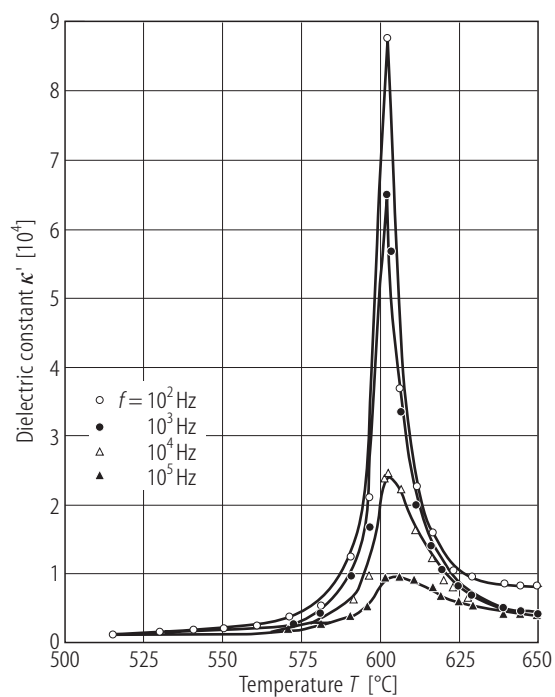


Fig. 2A-2-021. LiTaO₃. κ' vs. T [94Rav]. Parameter: f .

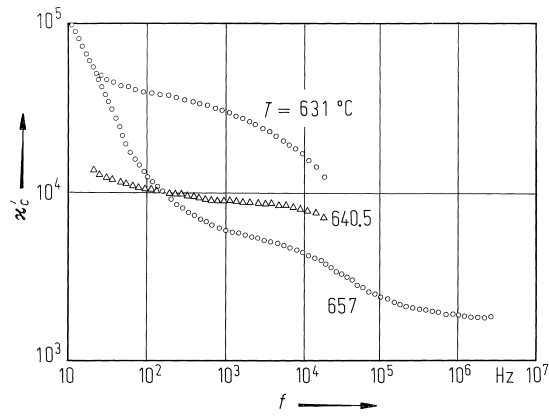


Fig. 2A-2-022. LiTaO₃. κ'_c vs. f [87Pri]. Parameter: T .

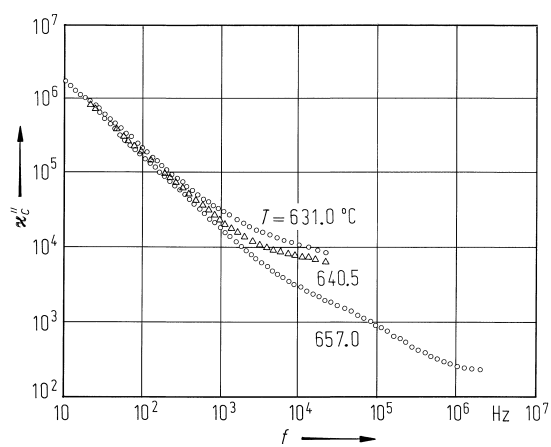


Fig. 2A-2-023. LiTaO₃. κ_c'' vs. f [87Pri]. Parameter: T .

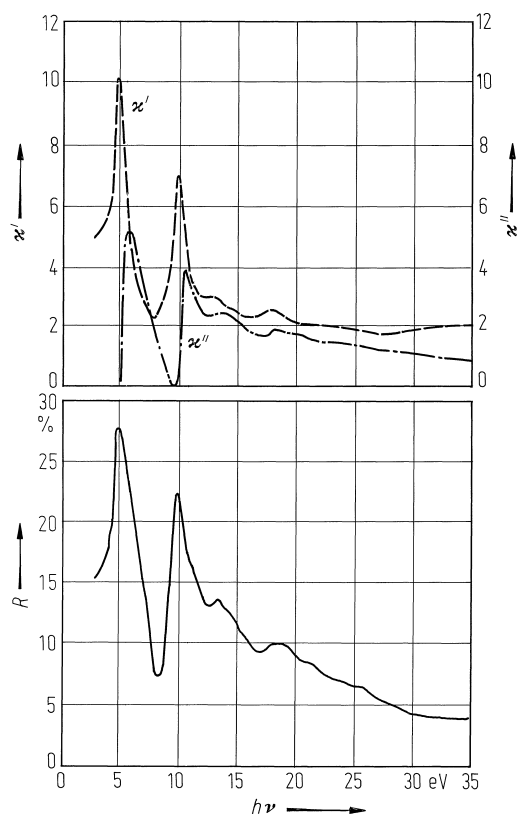


Fig. 2A-2-024. LiTaO₃. R , κ' , κ'' vs. $h\nu$ at 300 K [83Mam].
 $h\nu$: photon energy.

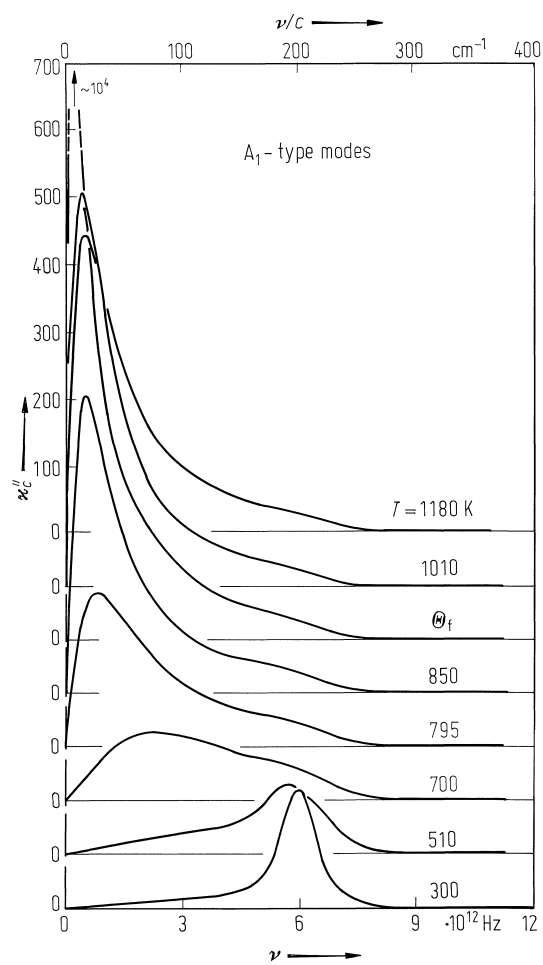


Fig. 2A-2-025. LiTaO₃. κ''_c vs. ν [80Ser]. Parameter: T .

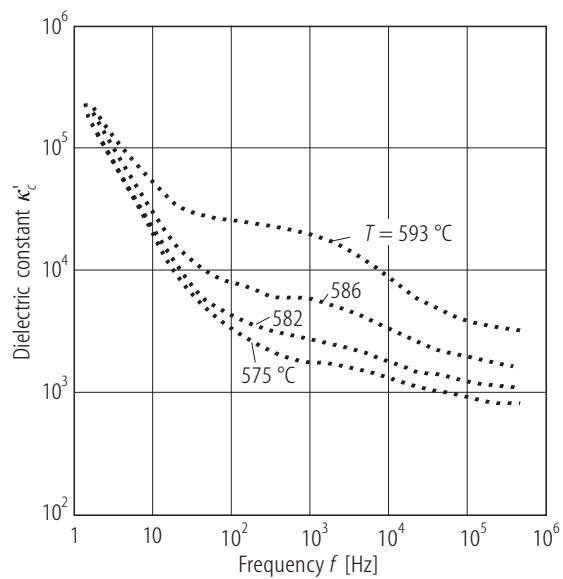


Fig. 2A-2-026. LiTaO₃, κ'_c vs. f [89Sin]. Parameter: T .

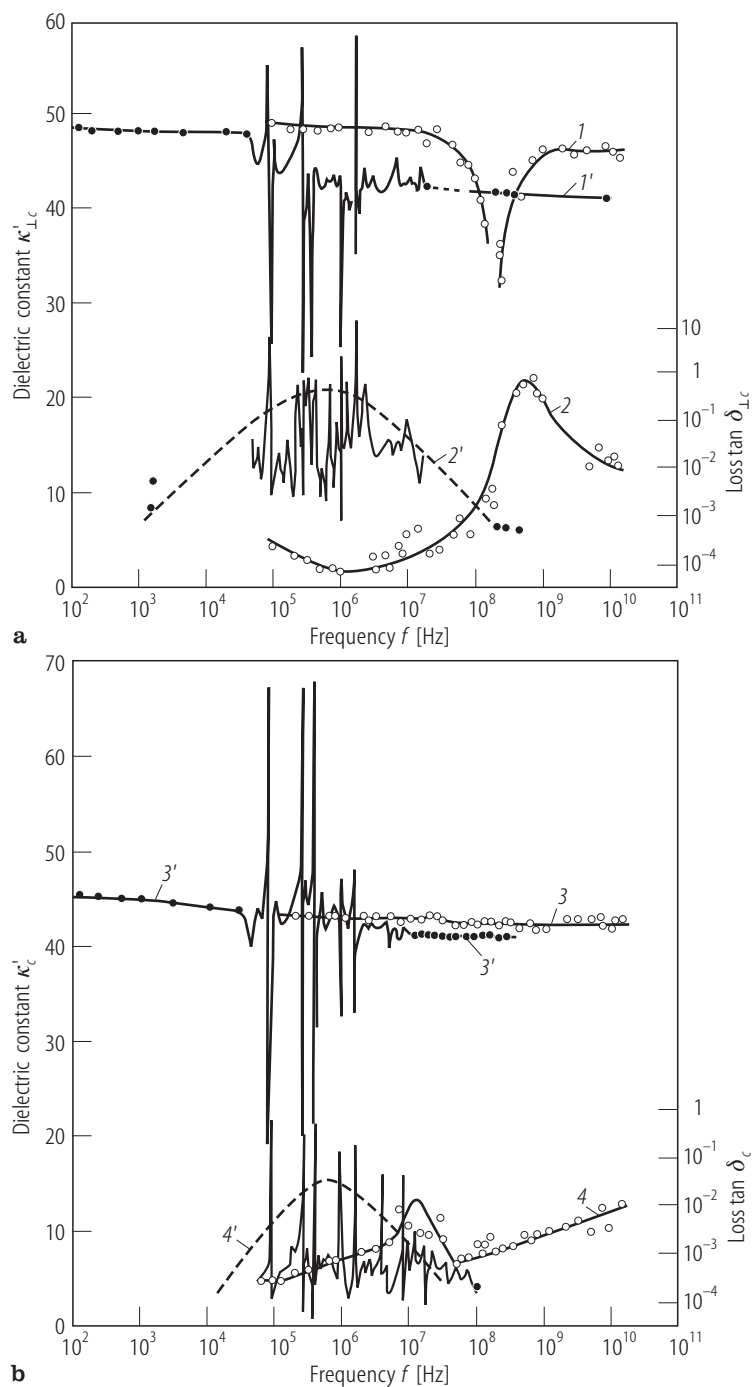


Fig. 2A-2-027. LiTaO₃, κ'_c , $\kappa'_{\perp c}$, $\tan \delta_c$, $\tan \delta_{\perp c}$ vs. f [85Bor]. 1...4: polydomain crystal. 1'...4': single domain crystal.

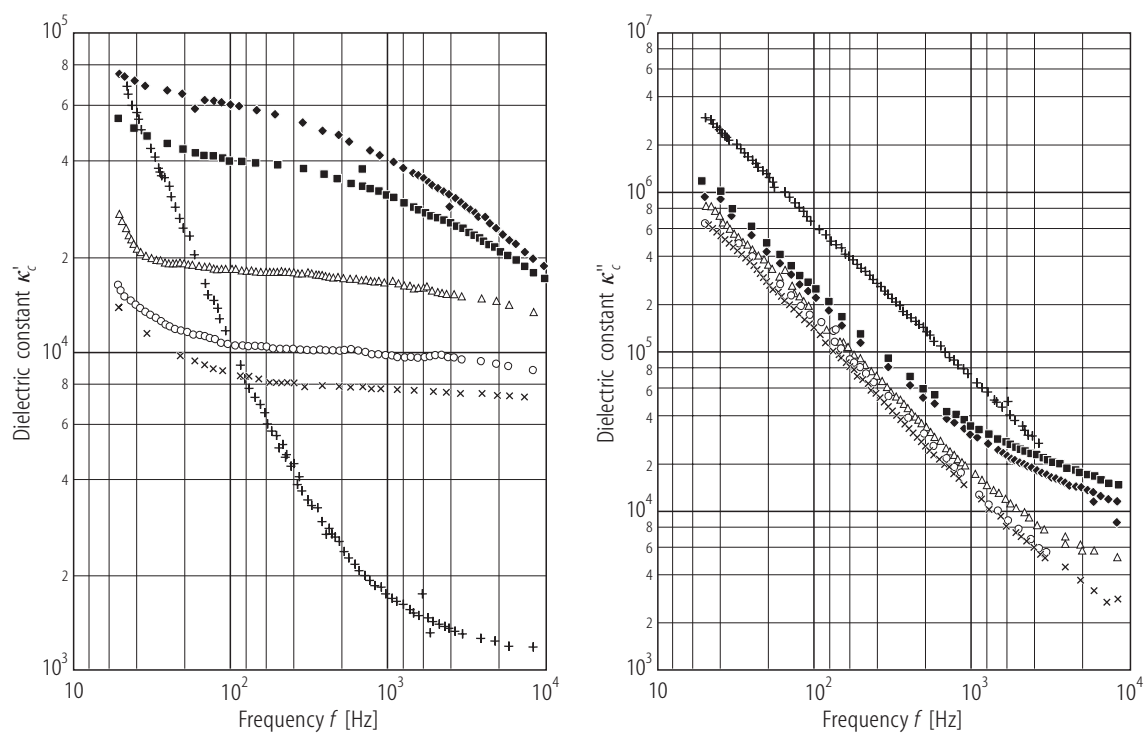


Fig. 2A-2-028. LiTaO₃. κ'_c , κ''_c vs. f [86Pri]. Parameter: T . Plus: 755.5 °C, cross: 643 °C, circle: 643.5 °C, triangle: 635.5 °C, square: 631 °C, diamond: 627 °C.

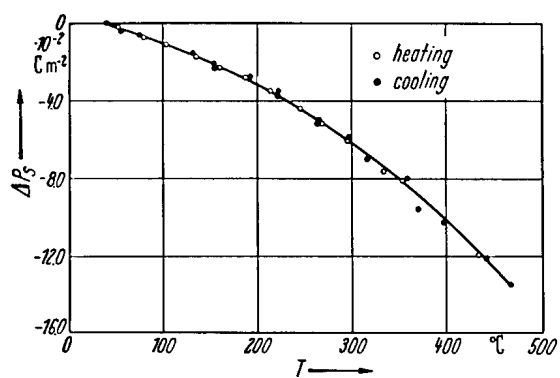


Fig. 2A-2-029. LiTaO₃, ΔP_s vs. T [67Iwa]. $\Delta P_s = P_s(0^\circ \text{C}) - P_s(T)$.

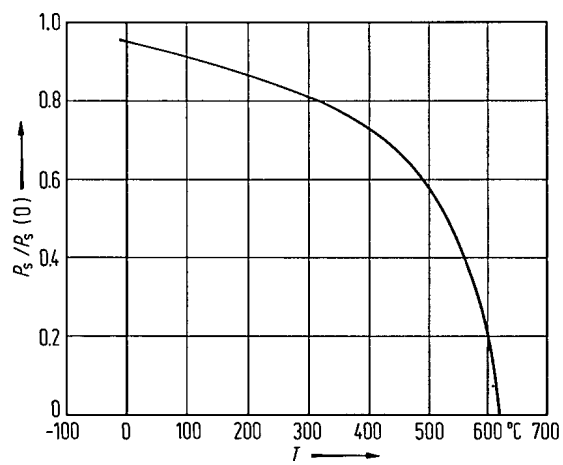


Fig. 2A-2-030. LiTaO₃. $P_s/P_s(0)$ vs. T [68Gla]. $P_s(0)$: spontaneous polarization at 0 K.

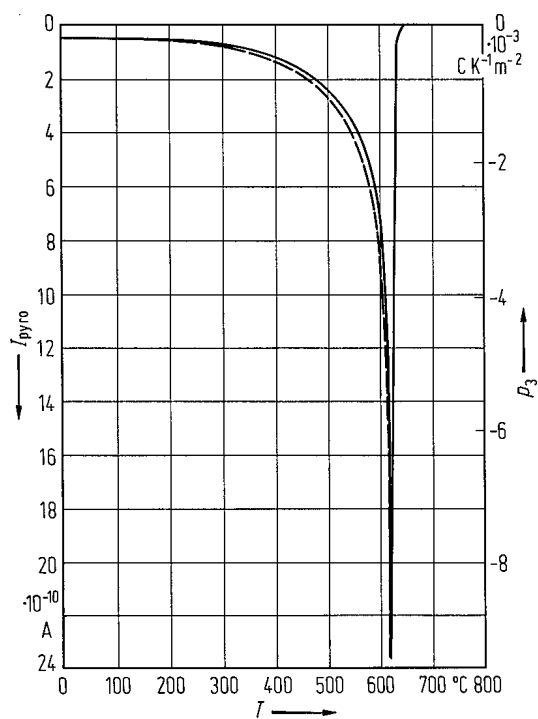


Fig. 2A-2-031. LiTaO₃. I_{pyro} , p_3 vs. T [68Gla]. I_{pyro} : pyroelectric current, p_3 : pyroelectric coefficient along the c -axis. Dashed curve: corrected for temperature variation of c_p and W , where W is the power absorbed.

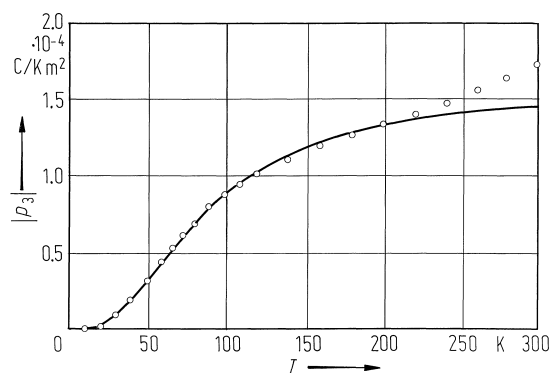


Fig. 2A-2-032. LiTaO₃. $|p_3|$ vs. T [77Lin]. Circles: measured values, full curve: sum of two Einstein functions for modes at 81 and 218 cm⁻¹.

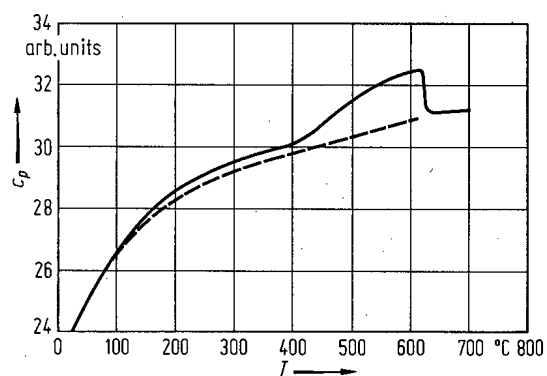


Fig. 2A-2-033. LiTaO₃, c_p vs. T [68Gla]. Dashed curve: background specific heat.

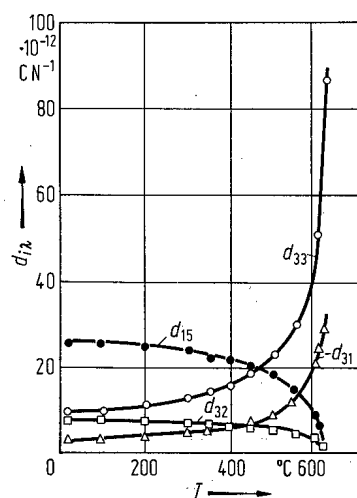


Fig. 2A-2-034. LiTaO₃. d_{ik} vs. T [69Yam]. d_{ik} : piezoelectric constant. d_{32} in the figure should be read as d_{22} .

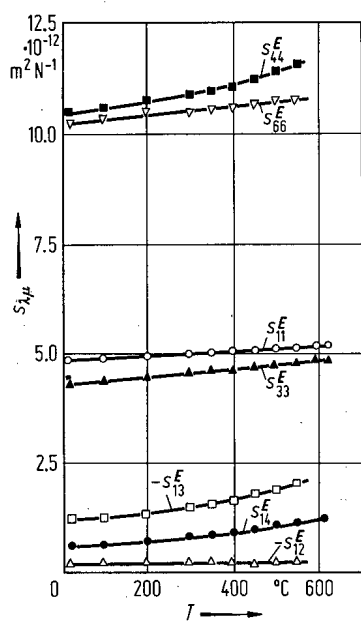


Fig. 2A-2-035. LiTaO₃. $s_{\lambda\mu}$ vs. T [69Yam]. $s_{\lambda\mu}$: elastic compliance.

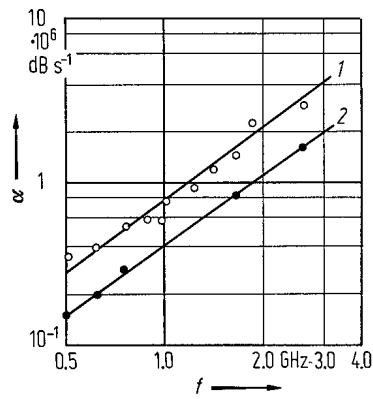


Fig. 2A-2-036. LiTaO₃. α vs. f [69Val]. α : attenuation coefficient of ultrasonic wave along the a -axis. Curve 1: transverse wave, 2: longitudinal wave.

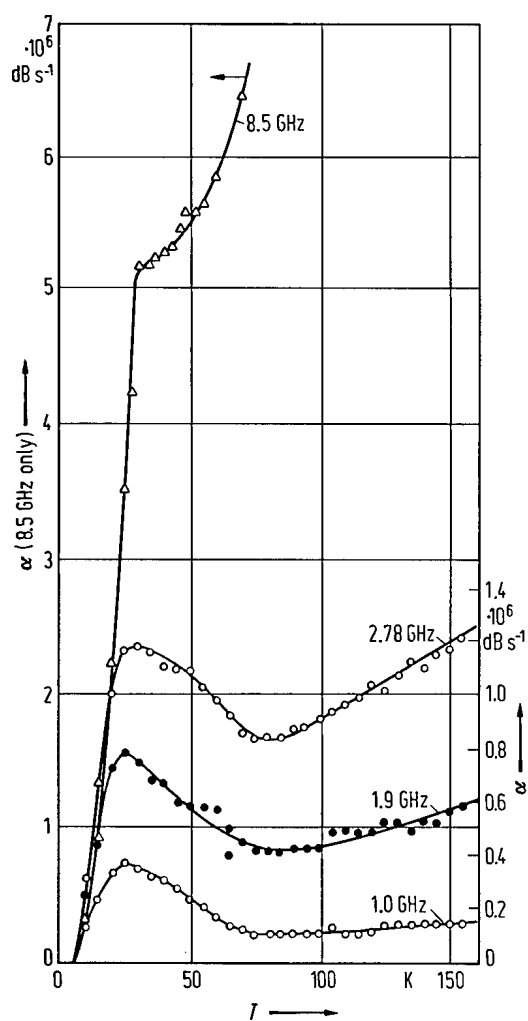


Fig. 2A-2-037. LiTaO₃. α vs. T [70Wor]. α : longitudinal wave attenuation in crystal grown from a melt containing 5 mol % excess of Ta₂O₅. Parameter: f .

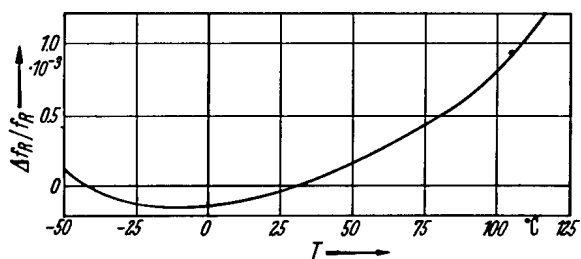


Fig. 2A-2-038. LiTaO₃. $\Delta f_R / f_R$ vs. T [67War2]. f_R : resonant frequency of an x -cut plate. Δf_R : frequency variation compared to f_R at 30 °C.

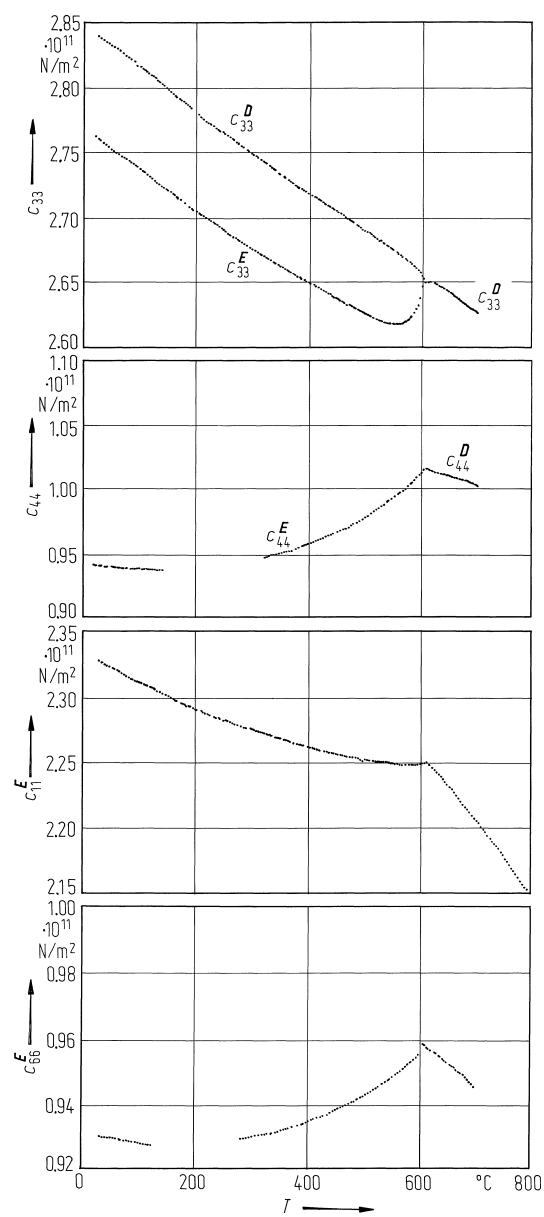


Fig. 2A-2-039. LiTaO₃. $c_{\lambda\mu}$ vs. T [82Tom]. $c_{\lambda\mu}$: elastic stiffness. The crystals were grown from approximately congruent melts. Single domain samples. $f = 20$ MHz.

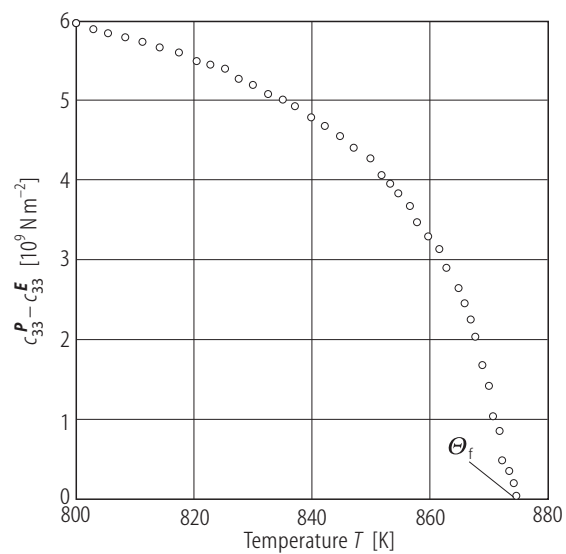


Fig. 2A-2-040. LiTaO₃. $c_{33}^P - c_{33}^E$ vs. T [88Tom].

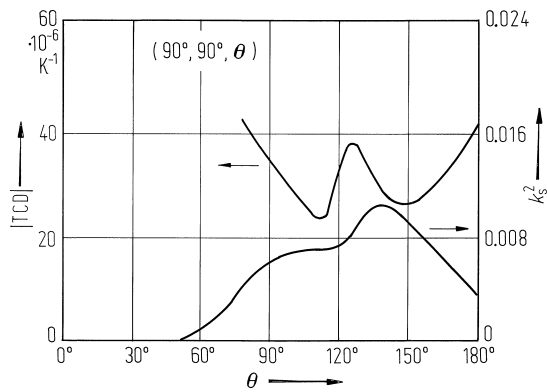


Fig. 2A-2-041. LiTaO₃. TCD, k_s^2 vs. θ [85Hir]. TCD: temperature coefficient of surface acoustic wave delay of X-cut θ rotated Y propagation mode, k_s^2 : surface acoustic coupling constant.

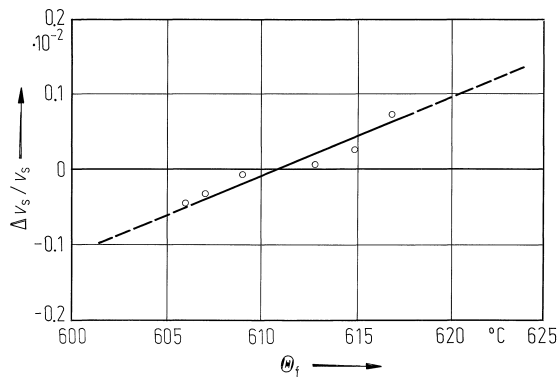


Fig. 2A-2-042. LiTaO₃. $\Delta v_s / v_s$ vs. Θ_f [84Yam]. $\Delta v_s / v_s$: normalized surface acoustic wave velocity change of X-cut 112° rotated Y propagation mode.

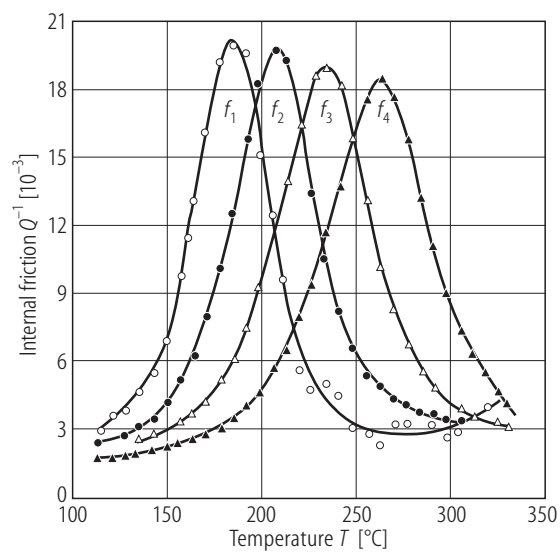


Fig. 2A-2-043. LiTaO₃. Q^{-1} vs. T [90Zha]. Q^{-1} : internal friction. Parameter: f . $f_1 = 0.0473$ Hz, $f_2 = 0.224$ Hz, $f_3 = 1.057$ Hz, $f_4 = 5.00$ Hz.

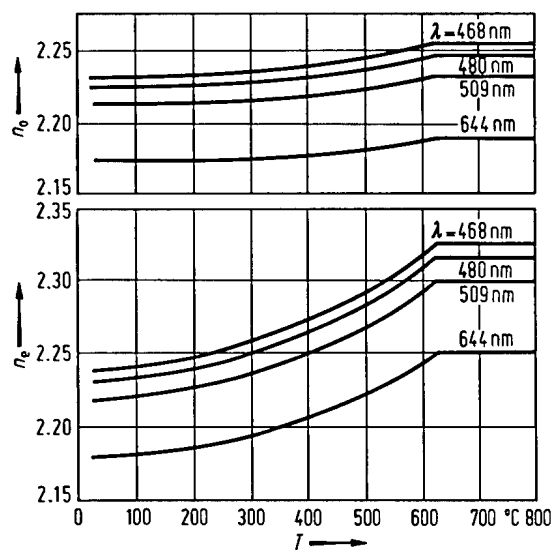


Fig. 2A-2-044. LiTaO₃. n_o , n_e vs. T [68Iwa]. Parameter: λ .

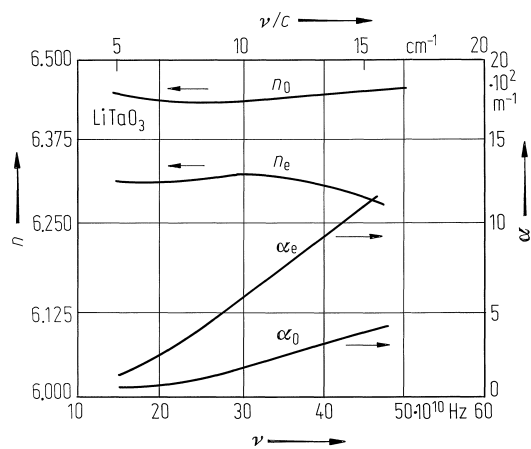


Fig. 2A-2-045. LiTaO₃. n , α vs. ν in the near millimeter wave region [83Bro]. α : power absorption coefficient.

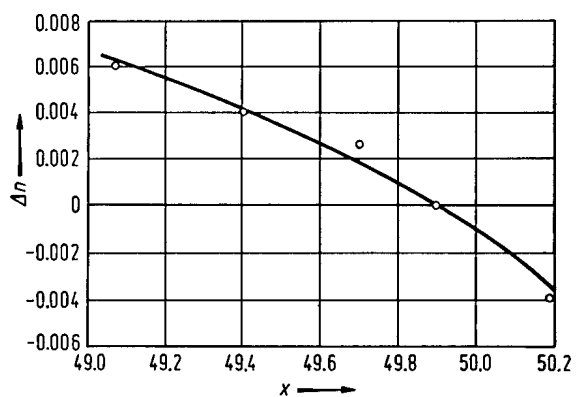


Fig. 2A-2-046. LiTaO_3 . Δn vs. x [70Bar1]. x : mol % Li_2O in the solid sample determined from Θ_f (see Fig. 2A-2-002). Na-D light was used.

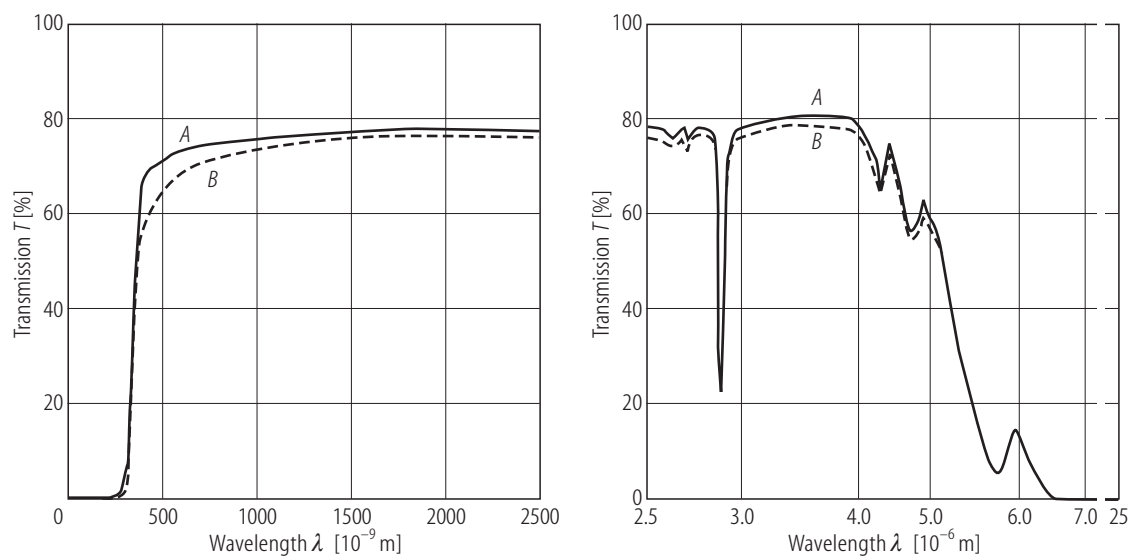


Fig. 2A-2-047. LiTaO₃. T vs. λ [93Wan]. T : light transmission. Sample: x -cut LiTaO₃ single crystal ($d = 3$ mm). A: poled sample; B: annealed sample. Please note changes in horizontal scale.

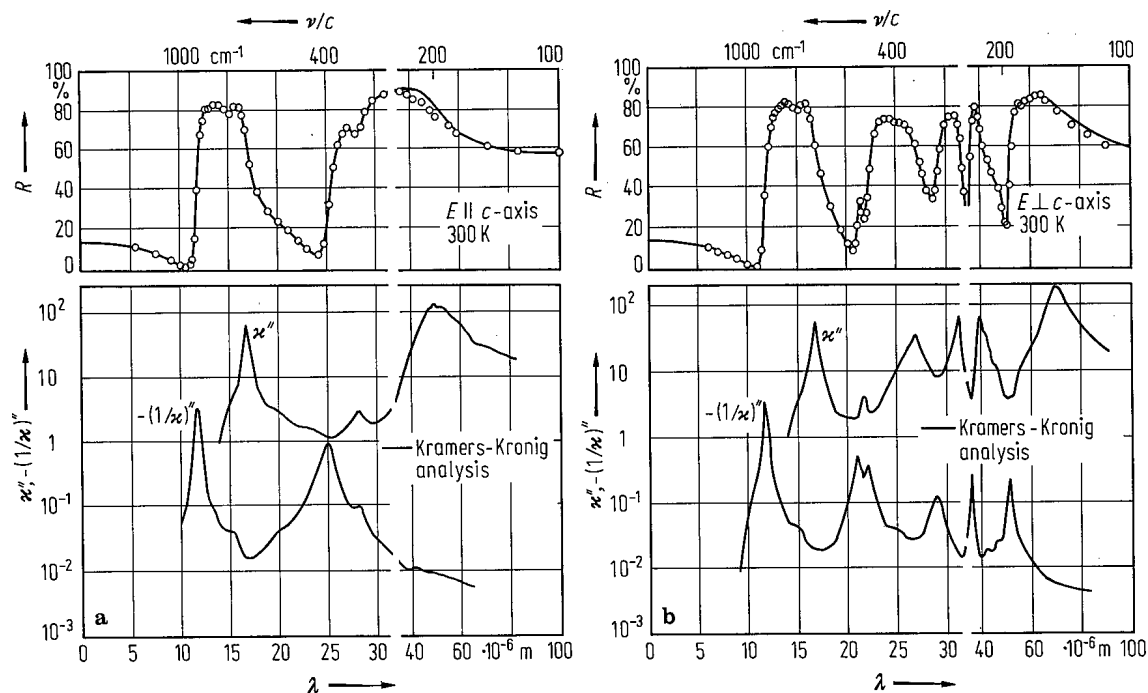


Fig. 2A-2-048. LiTaO₃. R , κ'' , $-(1/\kappa)''$ vs. λ at 300 K [70Bar2]. R : reflectivity, κ'' : imaginary part of κ , $-(1/\kappa)''$: imaginary part of $-1/\kappa$; (a) $E \parallel c$, (b) $E \perp c$. The solid curves through the experimental points are the best oscillator fit.

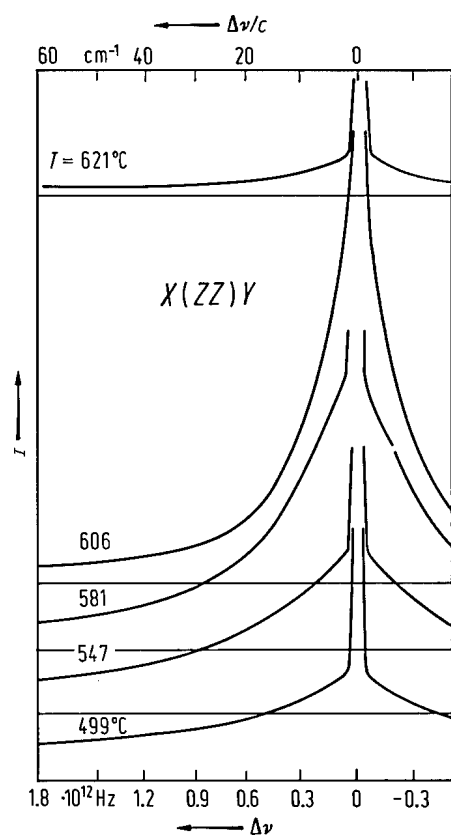


Fig. 2A-2-049. LiTaO_3 . I vs. $\Delta\nu$ [76Pen2]. I : Raman scattering intensity for the polarization geometry of $X(ZZ)Y$, $\Delta\nu$: frequency shift. For other temperatures, see original paper.

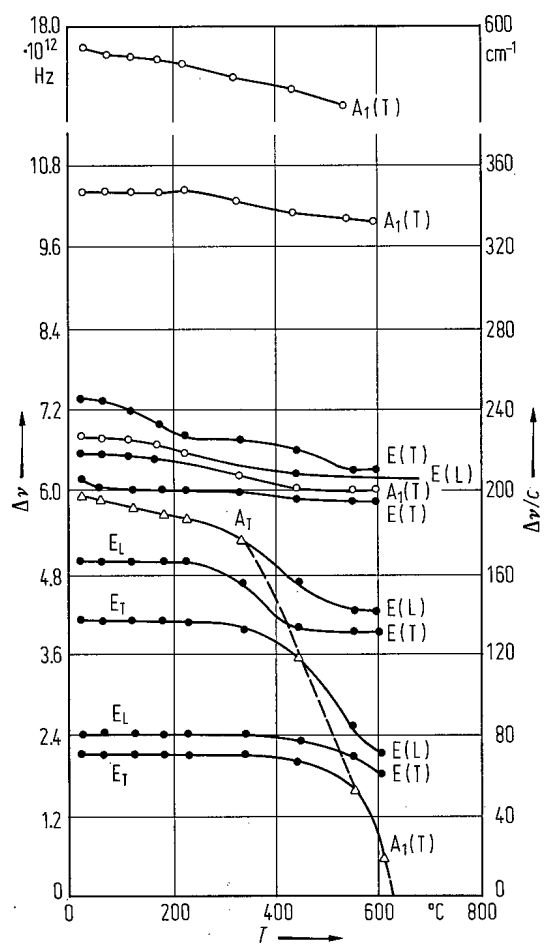


Fig. 2A-2-050. LiTaO₃. $\Delta\nu$ vs. T [68Joh]. $\Delta\nu$: Raman frequency shift. Triangles indicate the soft mode peak. Broken lines are mere extrapolation of the data. See also [76Pen1].

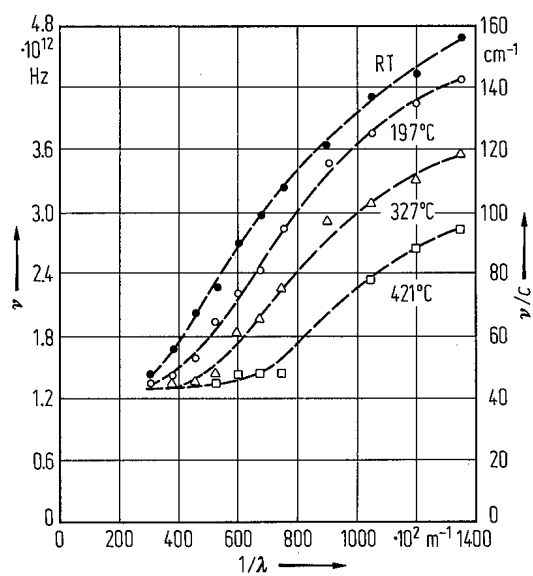


Fig. 2A-2-051. LiTaO₃. ν vs. $1/\lambda$ [76Pen3]. ν : polariton frequency of the lowest A₁ phonon mode. Parameter: T .

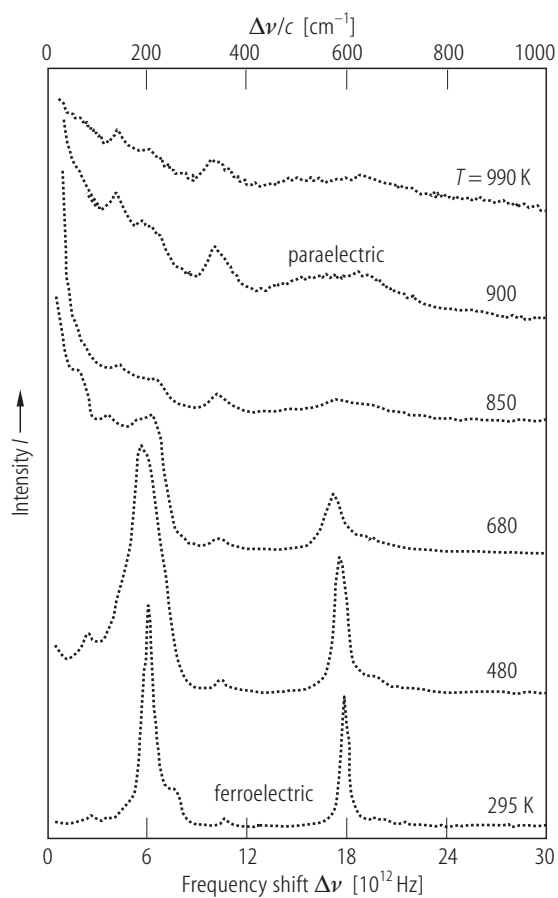


Fig. 2A-2-052. LiTaO₃. *Y(ZZ)X* Raman spectrum of the $A_1(\text{TO})$ (ferroelectric) and A_{1g} (paraelectric) phonons [88Rap]. Parameter: *T*.

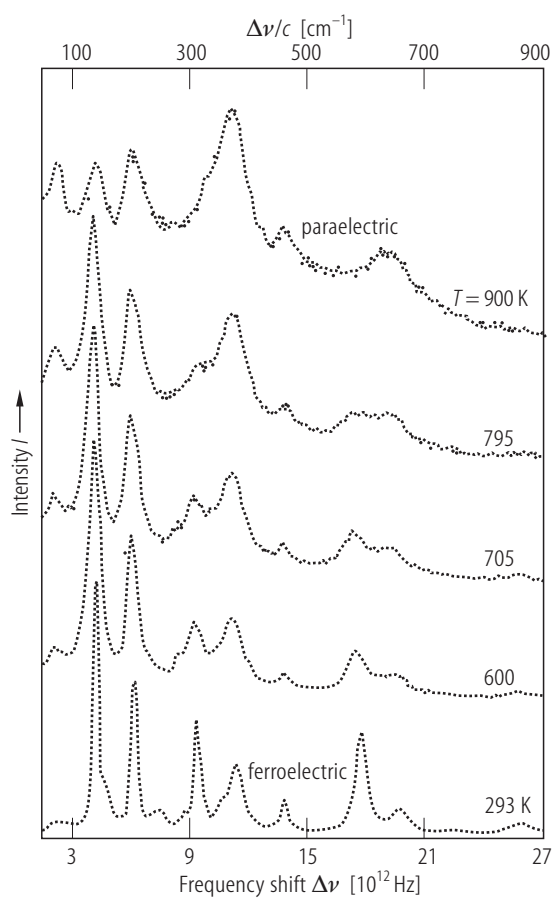


Fig. 2A-2-053. LiTaO₃. Z(XY)Z Raman spectrum of E(TO) phonons and oblique phonons in the ferroelectric and E_g phonons in the paraelectric phase [88Rap]. Parameter: *T*.

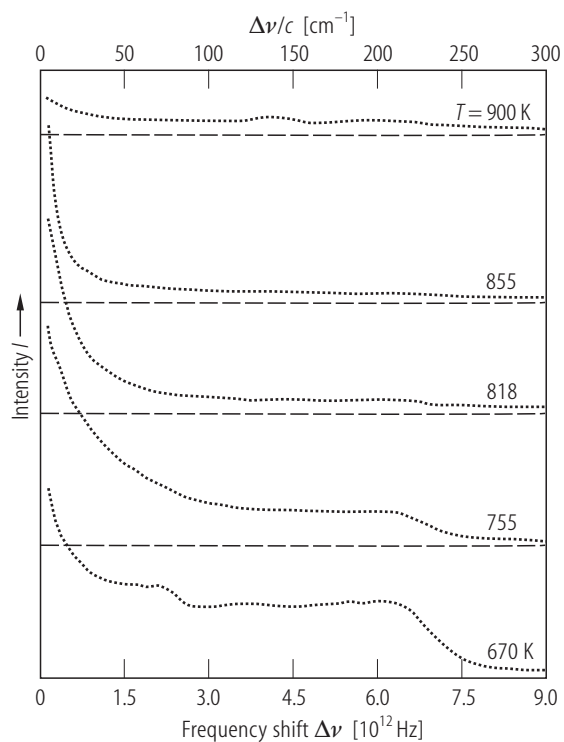


Fig. 2A-2-054. LiTaO₃. Y(ZZ)X Raman spectrum [88Rap].
Parameter: T .

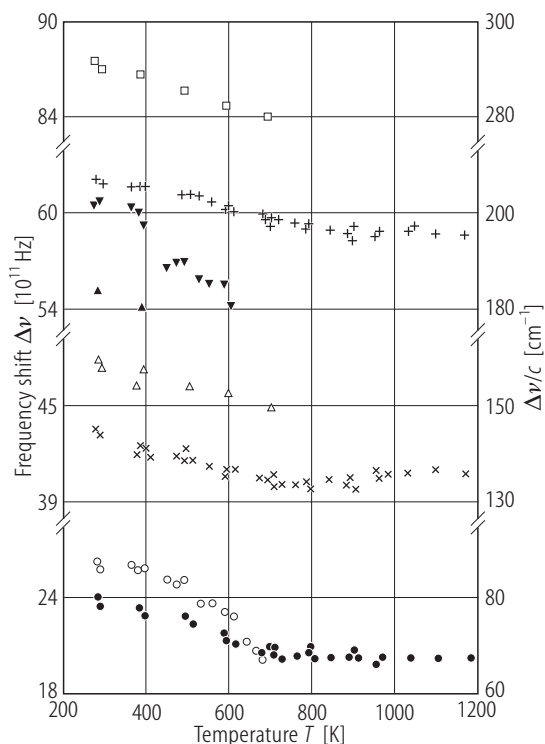


Fig. 2A-2-055. LiTaO₃. $\Delta\nu$ vs. T [88Rap]. $\Delta\nu$: Raman shift. Full and open circles: two-phonon (difference) band; crosses: E mode at 142 cm⁻¹ in ferroelectric phase (E_g in paraelectric phase); open triangles: oblique phonon at 157 cm⁻¹; full triangles: oblique phonon at 183 cm⁻¹; upside down full triangles: A_1 at 203 cm⁻¹; pluses: $E(E_g)$ at 207 cm⁻¹; open squares: oblique phonon at 293 cm⁻¹. All frequencies refer to room temperature.

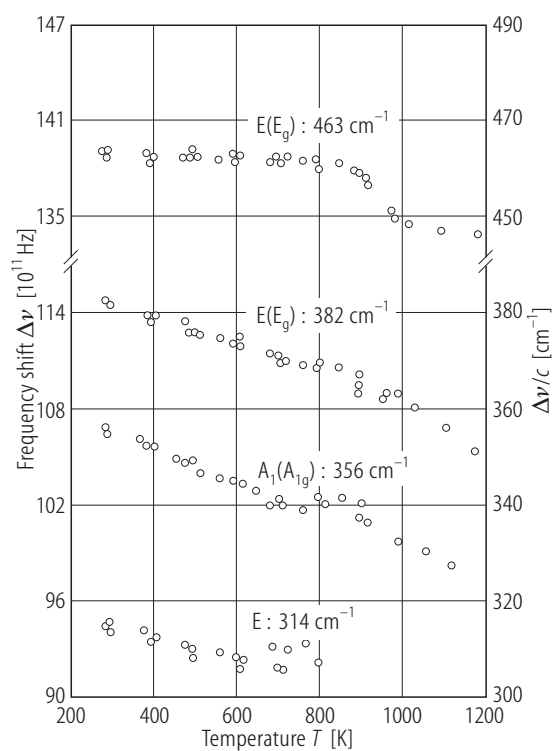


Fig. 2A-2-056. LiTaO₃. $\Delta\nu$ vs. T [88Rap]. $\Delta\nu$: Raman shift.

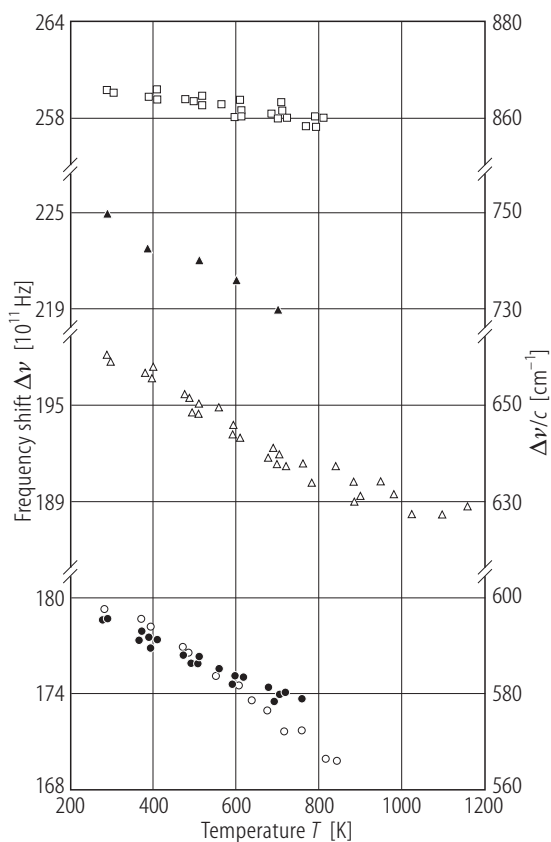


Fig. 2A-2-057. LiTaO₃. $\Delta\nu$ vs. T [88Rap]. $\Delta\nu$: Raman shift. Full circles: E mode at 592 cm^{-1} ; open circles: A₁ at 597 cm^{-1} ; open triangles: two-phonon (summation) band; full triangles: two-phonon (summation) band; open squares: E at 864 cm^{-1} . All frequencies refer to room temperature.

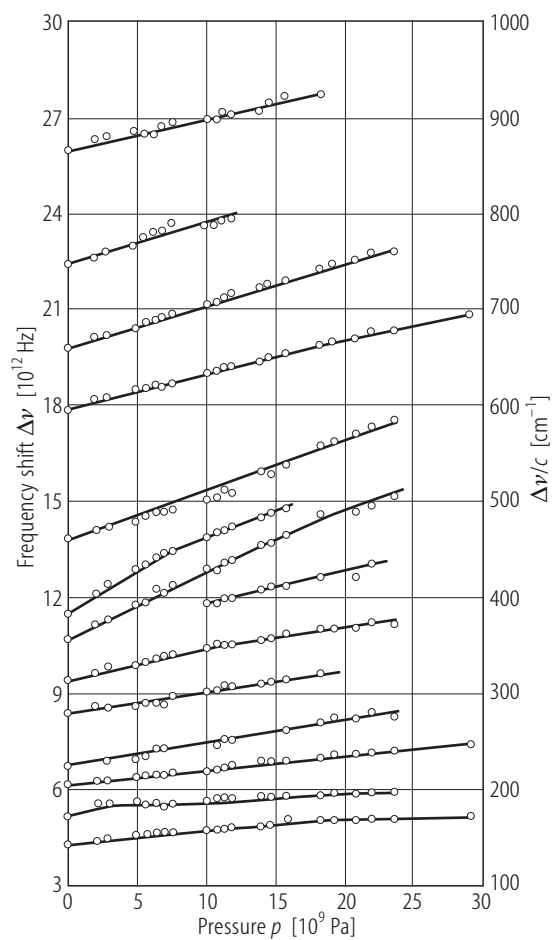


Fig. 2A-2-058. LiTaO₃, $\Delta\nu$ vs. p [94Lin]. $\Delta\nu$: Raman shift.

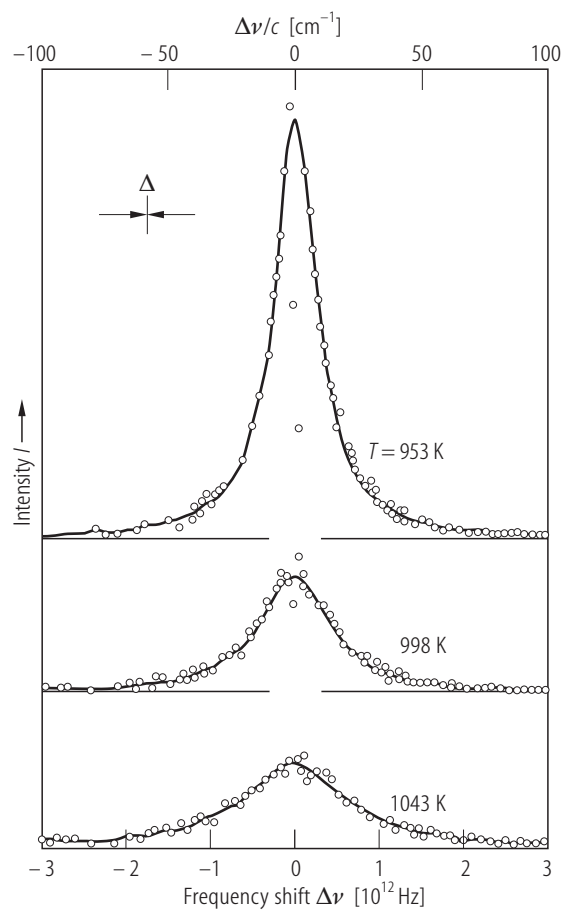


Fig. 2A-2-059. LiTaO₃. Central mode of hyper Raman scattering [94Tez]. I : scattering intensity. Δ : resolution. Parameter: T . $X(ZZ)Y$.

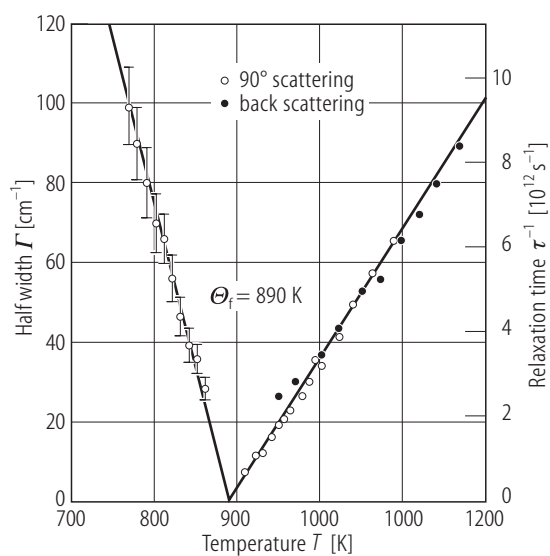


Fig. 2A-2-060. LiTaO₃. Γ vs. T [94Tez]. Γ : FWHM of the central mode of hyper Raman scattering. τ : relaxation time.

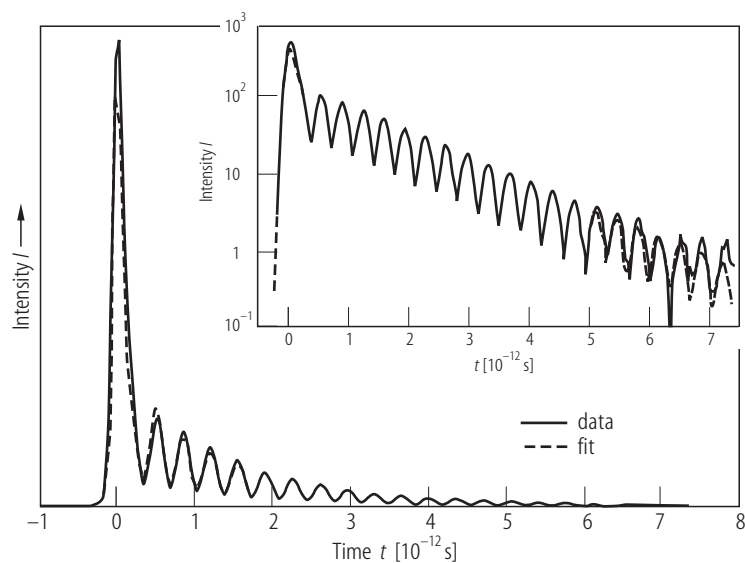


Fig. 2A-2-061. LiTaO₃. Real time observation of A₁ polariton at $q = 1800 \text{ cm}^{-1}$ by a femtosecond pulse laser [95Wie]. I : scattering intensity. Full line: data; dashed line: fit by a single damped oscillator.

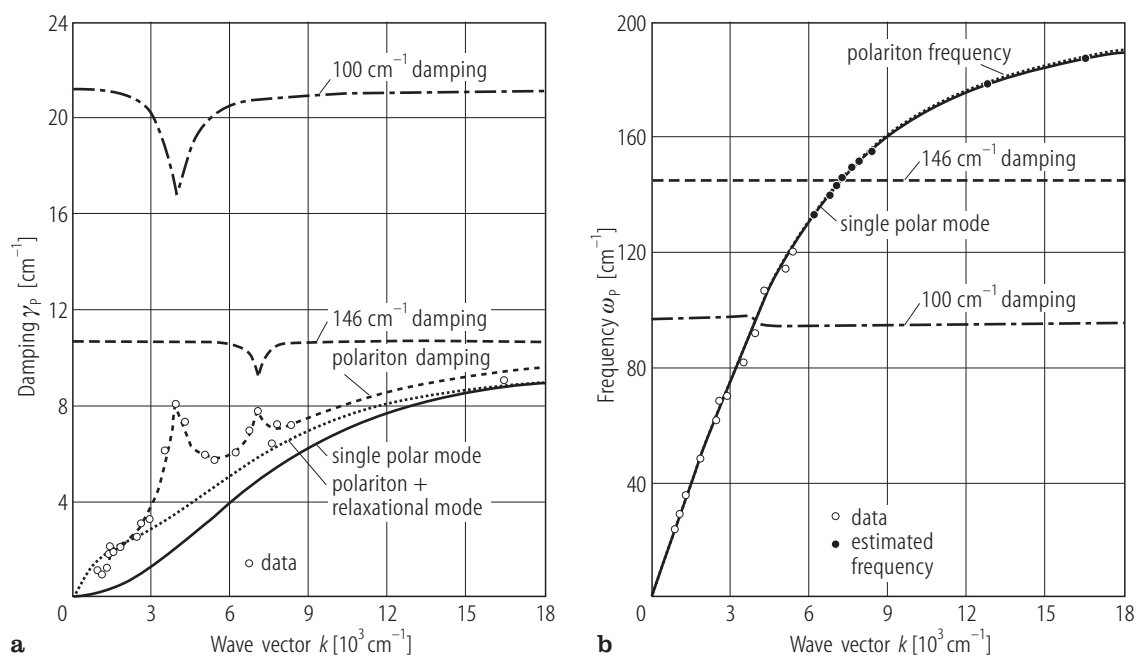


Fig. 2A-2-062. LiTaO₃. γ_p , ω_p vs. k [95Wie]. Femtosecond real time observation of A₁ polariton. γ_p : damping constant; ω_p : frequency. k : wave number vector.

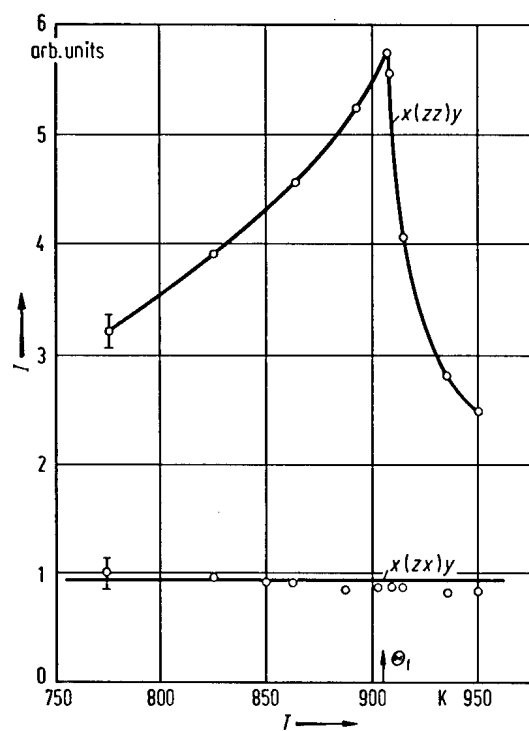


Fig. 2A-2-063. LiTaO₃. I vs. T [68Joh]. I : Rayleigh scattering intensity for the scattering geometry of $x(zz)y$ and $z(zx)y$.

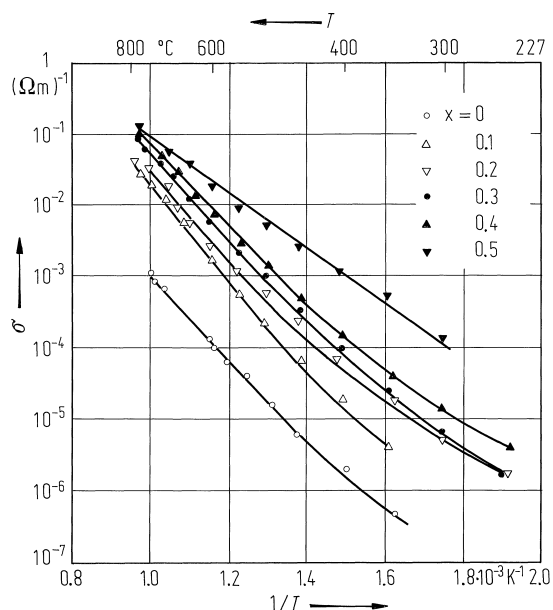


Fig. 2A-2-064. $\text{Li}_{1-x}\text{Ta}_{1-x}\text{W}_x\text{O}_3$ (ceramics). σ vs. $1/T$ [85Kaw]. σ : electrical conductivity. Parameter: composition.

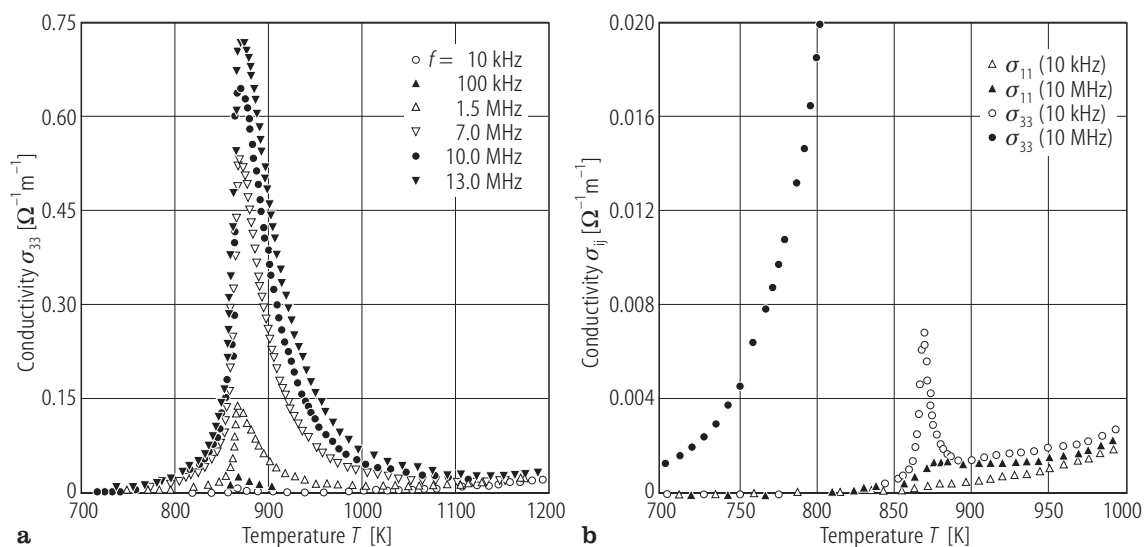


Fig. 2A-2-065. LiTaO₃. σ_{11} , σ_{33} vs. T [88Tom]. σ_{11} , σ_{33} : conductivity along the a -, c -axes, respectively. Parameter: f .

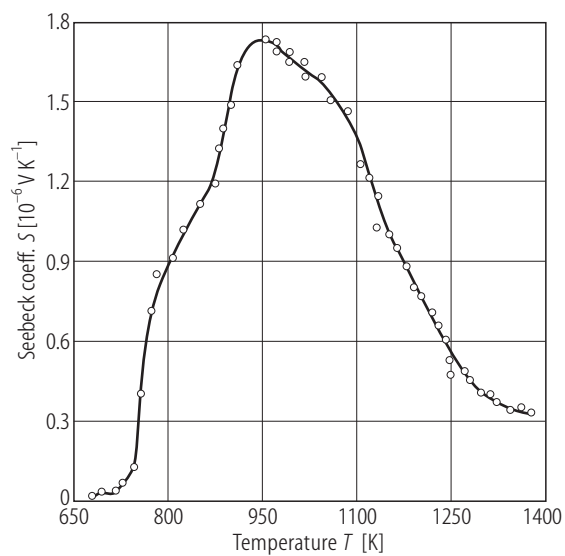


Fig. 2A-2-066. LiTaO₃. S vs. T [88Kha]. S : Seebeck coefficient.

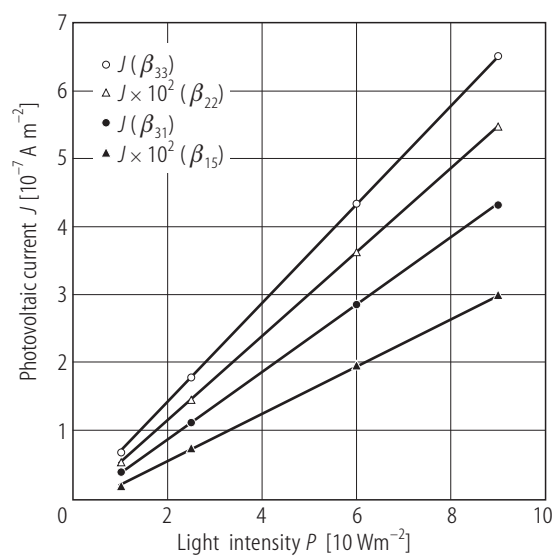


Fig. 2A-2-067. LiTaO₃:Fe. J vs. P [92Kar]. J : photovoltaic current; P : light intensity of photon energy 2.6 eV. For definitions of β_{ik} , see original paper.

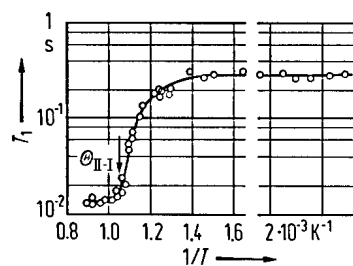


Fig. 2A-2-068. LiTaO₃ (powder). T_1 vs. $1/T$ [74Slo]. T_1 : spin lattice relaxation time of ^7Li . $\nu_L = 9 \text{ MHz}$.

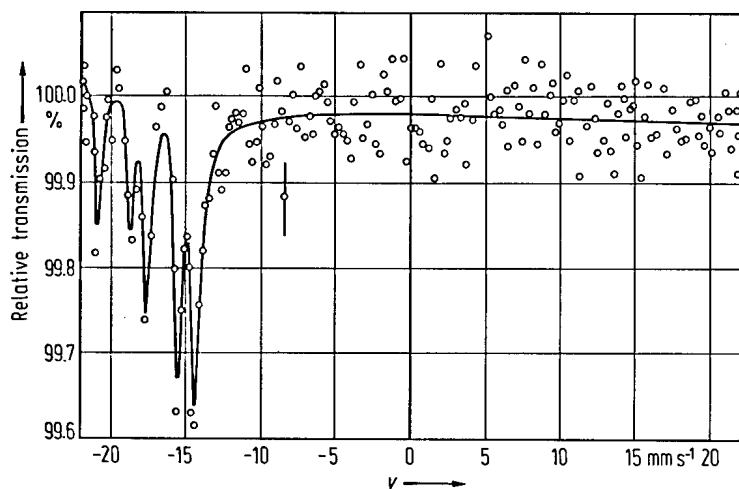


Fig. 2A-2-069. LiTaO₃, Mössbauer spectrum at RT [78Wor]. v : $^{181}\text{W}(\underline{\text{W}})$ source velocity. $^{181}\text{W}(\underline{\text{W}})$ means ^{181}W in W metal.

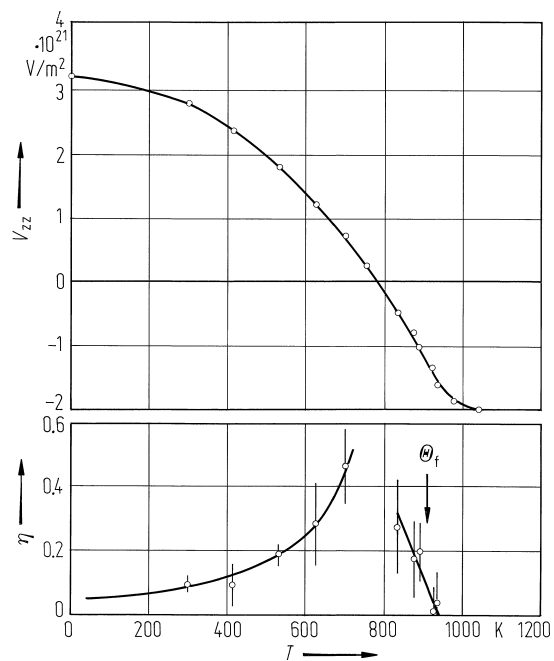


Fig. 2A-2-070. LiTaO₃. V_{zz} , η vs. T [81Loh]. V_{zz} : principal component of the electric field gradient tensor at Ta site; η : asymmetry parameter. ^{181}Ta Mössbauer effect.

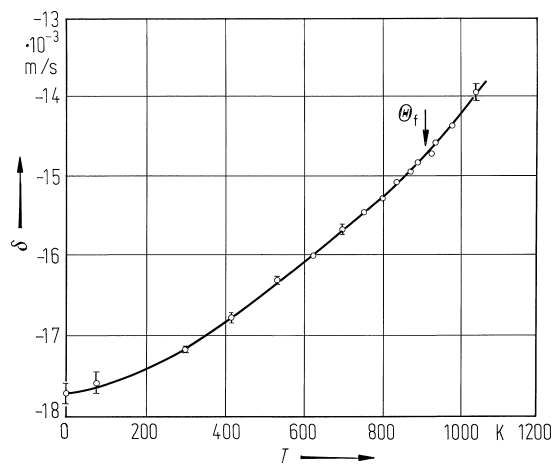


Fig. 2A-2-071. LiTaO₃. δ vs. T [81Loh]. δ : line shift of the ¹⁸¹Ta Mössbauer resonance spectrum relative to a W source.

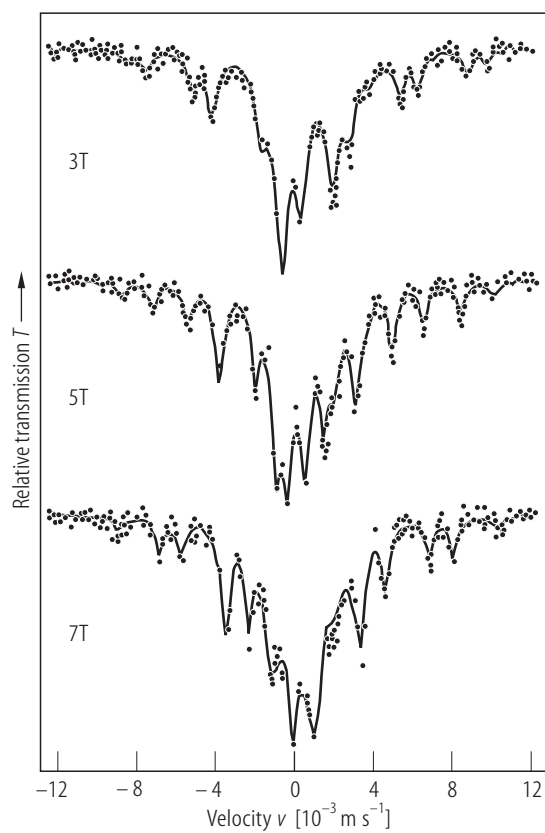


Fig. 2A-2-072. $\text{LiTaO}_3\text{:Fe}$. ^{57}Fe Mössbauer spectra [90Gie]. Single crystal. $H \parallel c$. Parameter: external magnetic field.

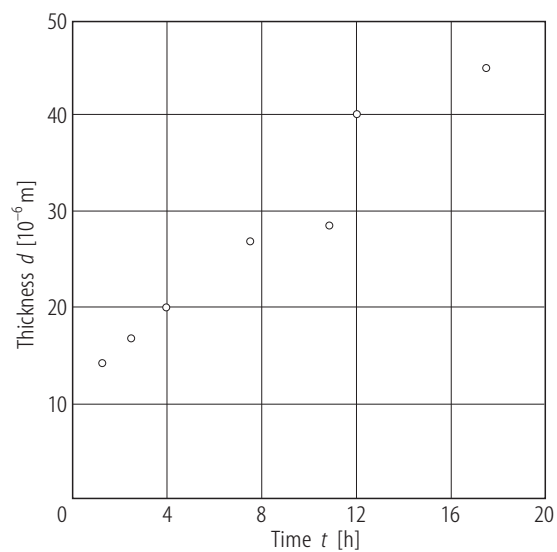


Fig. 2A-2-073. LiTaO₃. d vs. t [94Ahl2]. d : single-domain layer thickness on heat treatment at 1100 °C for z -plate. t : heat treatment time.

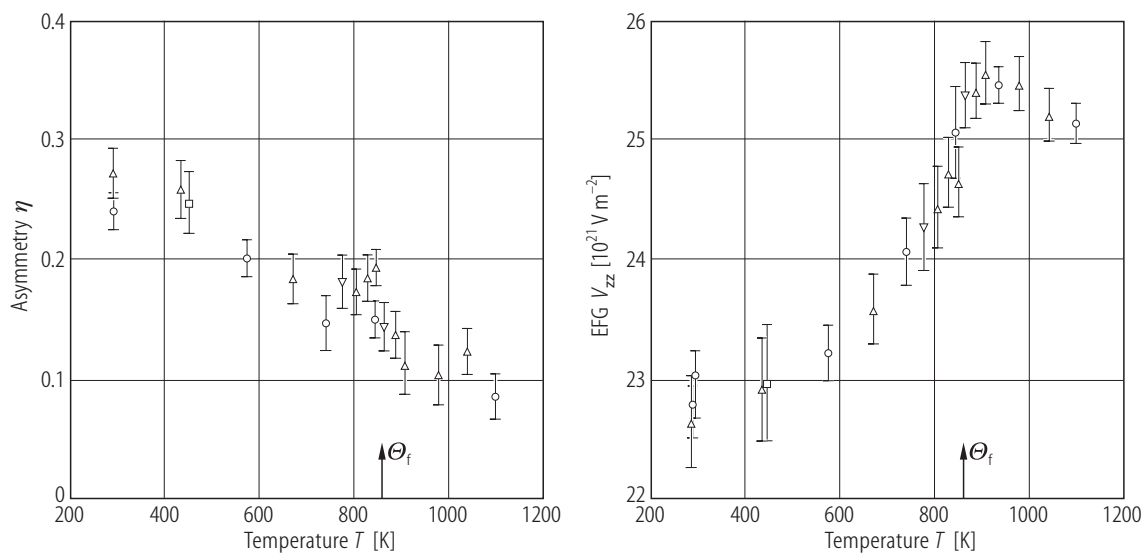


Fig. 2A-2-074. LiTaO₃. V_{zz} , η vs. T [91Cat2]. V_{zz} , η : electric field gradient and asymmetry parameter at Li sites determined by $^{181}\text{Hf} \rightarrow ^{181}\text{Ta}$ perturbed-angular correlation spectroscopy.

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