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A. Ahmine, M. Velazquez, V. Nagirnyi, I. Romet, T. Duffar. Conventional Czochralski growth of large Li2MoO4 single crystals. Journal of Crystal Growth, 2023, 578, pp.126420. 10.1016/j.jcrysgro.2021.126420. hal-03417415

HAL Id: hal-03417415

https://hal.science/hal-03417415

Submitted on 5 Nov 2021

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Conventional Czochralski growth of large Li₂MoO₄ single

crystals

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Abstract

Lithium molybdate single crystals up to one kilogram have been grown by a conventional Czochralski process. The growth configuration (geometry, coil, crucible, insulation casing) was optimized by numerical simulation of heat and mass transfer. Dislocations are shown to belong to the basal glide system. Their density ($\leq 10^4~\rm cm^{-2}$) and luminescence properties indicate crystal quality similar to that of previously reported crystals grown by Czochralski technique. Numerical simulation of thermo-elastic stresses suggest that the crack observed in one of these crystals is rather due to a mechanical accident than to internal stresses during growth.

Keywords

B1-Lithium molybdate; A2-Czochralski method; A1-Dislocations, luminescence; A1-Computer simulation, stresses.

1 Introduction

The growth of bulk Li₂MoO₄ crystals for the large-scale astroparticle physics experiments that are foreseen by the end of this decade is proceeding apace. The general context of this R&D effort is related to the quest of the ultimate components of matter, like the neutralino, which is considered as a viable candidate to be the corpuscular component of the dark matter halo of our galaxy [1, 2]. In the underground dark matter direct detection sites, it would also be useful to characterize the ultimate background noise constituted by fast neutrons which mimics the nuclear recoils possibly induced by Weakly Interacting Massive Particle (WIMP)-nucleus collisions. And more broadly speaking, it would be interesting to develop transportable neutron spectrometers based on heat-scintillation cryogenic bolometers [3]. Another important R&D effort of these last years is focused on the search for neutrinoless double beta decays (0ν2β), predicted but never observed, which would prove that neutrinos are their own antiparticles, and so, a new kind of matter, and would also help understanding the origin of their mass [4, 5, 6, 7]. In order to detect these extremely rare events, double readout detector modules are built with a core made of a bulk crystal. One or several atomic constituents of these crystals turn out to be an isotope that exhibits the highest capture cross section or emission properties. All the kinetic energy of the neutralino, or of the fast neutron, is collected and recovered as heat and light. In the case of 0v2\beta decays, the crystal itself is the source of the decay to be detected exactly in the same way. The crystal is typically several hundred grams heavy, and as the huge mass of the overall detector is to be cooled down to cryogenic temperatures in 20-25 days, it is necessary to develop crystals that have low specific heat and high thermal conductivity, that is, thousands of crystals of high quality. Of course, the scintillation light must be emitted in the spectral range where cheap and 100%-efficient light detectors exist, typically between 400 nm and 1.7 µm. Besides, because of the cost of rare isotopes like ¹⁰⁰Mo or ⁶Li, it is necessary to develop purification processes of the initial powders used for the synthesis and recycling processes of the remaining growth loads [8]. The purification, recycling and growth processes must be fast enough to comply with the constrained time schedules of these large-scale experiments.

Li₂MoO₄ crystals have been grown by the vertical Bridgman technique in a three-zone furnace [9, 10], however crystals appeared to be colored, polluted by inclusions and cracked. Barinova et al. [11], proposed a technique of growth from aqueous solution, with strong acoustic mixing in order to destroy the oxide clusters occurring in the liquid [12]; while this technique shows a great potential, only millimetric crystals have been grown so far.

Czochralski technique was used for the growth of the very first single crystals in order to measure their physical properties [13]. This process has been developed, from crystals 34 g in mass [14] to more than 700 g [15]. This research effort led to the Low Temperature Gradient Czochralski (LTG-Cz) method, which is now the reference process for the growth of Li₂MoO₄ scintillating crystals [15-17]. Rationale for the development of this low gradient technology was the potential vaporization or reduction of the melt constituents in case of too high temperature and crack risks in case of large temperature gradients in the solid [18]. In this technique, the crystal grows inside a totally closed Pt chamber, which prevents volatilization [16] and allows a mass yield of 80% [8]. This chamber is placed in a three zone furnace and authors claim that crystals are grown under a gradient as low as 1 K.cm⁻¹ [15-18]. However, numerical simulations of heat transfer suggest that it is in fact much larger [19, 20]. The best bolometric results were obtained with crystals produced in this way [6].

However, this LGT-Cz technique has some drawbacks, first of all the low growth rate (≤ 0.7 mm.h⁻¹) imposed by the weak temperature gradient [15]. The Pt chamber and crucible are complex in shape and then costly. Because of this chamber, observation of the growing crystal is not possible and growth control remains complex, also due to the three zone furnace [15].

Therefore, a more classical Czochralski process, provided that too high temperatures are avoided, could be a less expensive, more productive technique for the growth of high quality Li₂MoO₄ crystals. In preceding papers [21, 22], numerical simulation was used for optimizing the hot zone of a Czochralski furnace in order to prevent melt overheating or undercooling, avoid large stresses in the crystal and provide a slightly convex solid-liquid interface (seen from the melt). Based on this approach, the growth of several crystals is described. Characterization is performed in terms of dislocation density and luminescence properties. Special attention is paid to the analysis of the thermal stresses in the growing crystal.

2 Growth process and crystals

Li₂MoO₄ is synthetized from stoichiometric mixtures of MoO₃ (Alfa Aesar 5N5) and Li₂CO₃ (Fox-Chemicals, 5N) ($\frac{m_{MoO_3}}{m_{Ll_2}o_3} \approx 1.948$). Powders are first mixed for 12 hours, then heated at 673 K in an alumina crucible for another 12 hours. This mixing-heating process is repeated, with heating at 773 K. Then the obtained powder is mixed again and heated at 873 K for 24 hours, in the Pt crucible (99.99% purity), which will be used for the growth. The obtained partially sintered powder is molten at 993 K for 3 hours in order to reduce its volume and provide room for other powder charges. The entire process is repeated until the Pt crucible (80 mm diameter and 100 mm in height, for a crystal diameter of 50 mm) is full. This heating process, with several steps, is preferred in order to avoid vaporization of MoO₃ at high temperature. Table 1 gives the qualitative chemical evolution of the powder all along the process, measured from X-ray diffraction (XRD) diagrams.

After heating at 673 K	After heating at 773 K	After heating at 873 K (before melting)
Li ₂ MoO ₄ (Large amount)	Li ₂ MoO ₄ (Large amount)	Li ₂ MoO ₄

Li ₄ Mo ₅ O ₁₇ (Small amount)	Li ₄ Mo ₅ O ₁₇ (Small amount)	
MoO ₃ (Large amount)	MoO ₃ (Very few)	
Li ₂ CO ₃ (Large amount)		

Table 1 XRD qualitative analysis of the chemical composition of the $(MoO_3 + Li_2CO_3)$ powder mixture along the heating steps.

The crucible is then placed in the center of an open air vessel, with the various heating and screening elements as defined through the numerical simulation optimization [21]. Figure 1-left shows this heating configuration. It is based on the induction heating of a susceptor element (Kanthal®), which provides much more flexibility than direct coupling to the Pt crucible. The alumina pulling shaft and Pt seed holder are introduced through the top opening, with special care for concentricity of the whole system. A viewing port, also used for pyrometric temperature measurements, exists on the top plate (not shown). The external jacket is continuously cooled with water at a controlled temperature of 28°C. All growth parameters are controlled by automation software (Cyberstar®) through continuous measurement of the crystal weight and adjustment of the induction coil power. Therefore, the sequence of shoulder-cylinder-tail growth phases is programmed in terms of pulling rate, rotation rate and diameter increase-decrease versus time, from the very beginning of the pulling.

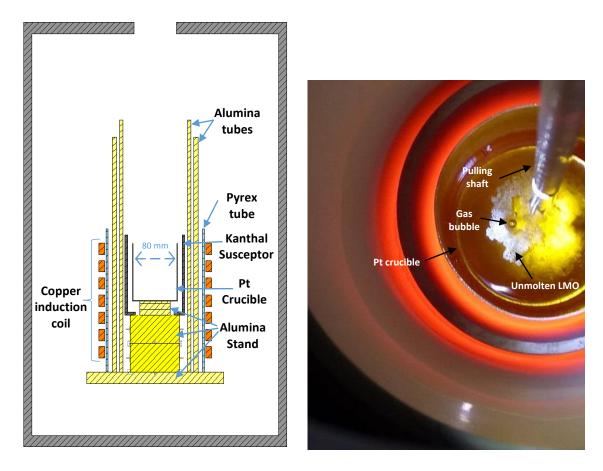


Figure1 Left: Heating and screening elements inside the open air set-up. **Right:** view from the top of the equipment, during the melting phase.

The material is slowly melted (Figure 1-Right) and soaked for 24 hours at 1003 K, as measured by the pyrometer on the liquid surface. Seeds oriented [11 $\overline{2}$ 0] and [0001] were used, without significant effect on the growth progress. After careful seeding, growth is usually begun while the measured weight is slightly decreasing, in order to get a seed necking. Then the fully programmed pulling is started. The rotation rate was always 5 rpm and pulling rate 2 mm.h⁻¹. While the extraction process can also be performed by the pulling software, it was usually performed manually (at 30 mm.h⁻¹) in order to avoid a too strong thermal shock to the crystal. The furnace is then slowly cooled down. It should be noted that absolutely no deposit of any kind has been found in the chamber after growth.

Three crystals with a diameter of 50 mm (Figure 2) have been grown under the conditions presented above. Masses scaled from 532 g to 967 g and material yield from 66% to 81.4%.

Two other crystals, with a diameter 40 mm, were grown from 3N LMO powder (Sigma-Aldrich) in a slightly modified configuration. However, they presented a yellowish color and some inclusions (cf. Figure 5).







Figure 2 Li₂MoO₄ crystals, 50 mm in diameter and, respectively, 820 g, 967 g, and 532 g in mass, pulled with the optimized Czochralski configuration.

3 Characterization

X-ray diffraction analyses only show rhombohedral ($R\overline{3}$) Li₂MoO₄ peaks. Several Laue back reflection diagrams, taken on various places on the ingots, have shown that they are single

crystals without twin or grain. As the material is hygroscopic, cutting and polishing are performed under ethanol instead of water.

3-1 Dislocations

Etch pit densities have been measured on polished (roughness $0.15~\mu m$) and acetone-cleaned surfaces randomly cut inside the crystals. Water, chosen with consideration to the crystal hygroscopicity, appeared to be an excellent etchant. Typical etching times, at ambient temperature, range from 20 s to 40 s, depending on the surface orientation. Dislocation densities were automatically measured on the microscopic photographs, using the "particle analysis" ImageJ® routine. This numeric procedure has been validated by comparison with one-by-one pit counting. Figure 3 shows the counting procedure applied to a [11 $\bar{2}$ 0] surface, etched 20 seconds, on which pits appear as small diamonds elongated along the c0 direction. The dislocation density is taken as the mean value of the etch-pit densities of all the areas analyzed on the surface. The results are in the range [0.8 – 1.4] 10^4 cm⁻² for the crystals 50 mm in diameter. The less pure crystals 40 mm in diameter are in the range [1.7 – 2.8] 10^4 cm⁻² with numerous dislocations appearing around precipitates occurring in the material. Energy dispersive shows that these precipitates contain Si, P and V, possibly coming from the 3N raw material and the old, smaller, Pt crucible used for these crystals.

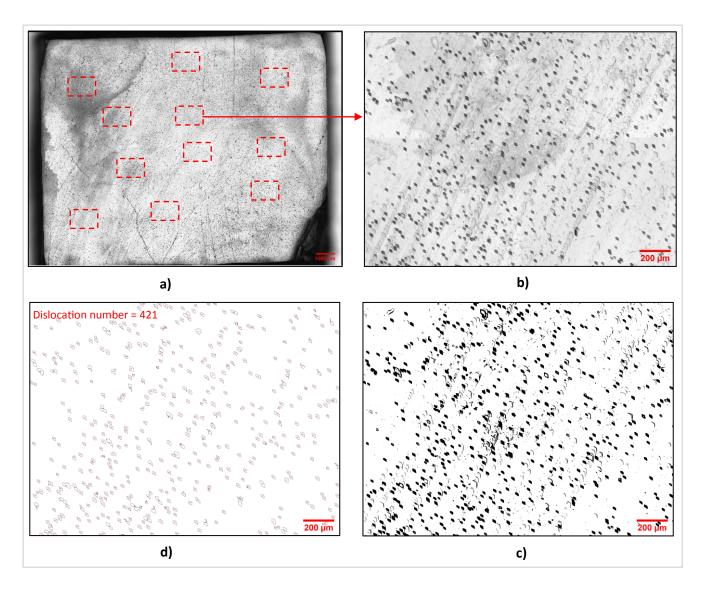


Figure 3 Etch-pit density measurement with ImageJ software. **a**) Full sample $[11\bar{2}0]$ surface; **b**) one of the analyzed areas; **c**) binary (B&W) image; **d**) etch-pit numbering.

Figure 4 shows the etched (0001) and (11 $\bar{2}0$) faces of a cube cut inside a crystal. The (11 $\bar{2}0$) surface, etched for 20 s, shows well defined etch-pits, while the (0001) surface shows only polishing defects and no pits in spite of a 40 s etching time. This result suggests that dislocations in Li₂MoO₄ are oriented along the [11 $\bar{2}0$] direction and contained in the (0001) plane, corresponding to the basal glide system. This is the most common slip system in rhombohedral crystals [23, 24].

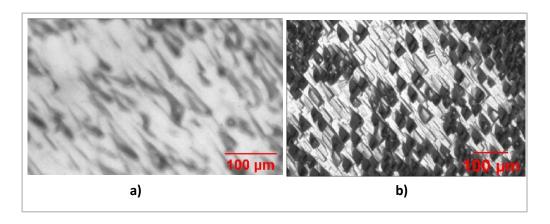


Figure 4 Photos of two etched faces of a Li₂MoO₄ cube. a) [0001] face, b) $[11\bar{2}0]$ face.

3-2 Luminescence properties

Luminescence features of the grown crystals have been studied by the techniques described in [25]. They are similar to those in Li₂MoO₄ crystals grown by the conventional Czochralski and Low-Temperature-Gradient Czochralski methods [26]. The emission spectrum at 4.2 K is represented by a non-elementary band in the region of 1.5-3 eV (Figure 5). The peak position of the emission band varies from 2.1 to 2.2 eV depending on the exciting photon energy (compare curves 1 and 2). The high-energy component of the emission is excited more efficiently in the band near 4.4 eV, while the low-energy part is more intense under excitation deeper in the fundamental absorption region (curves 3 and 4). According to the time-resolved spectroscopy studies, the emission spectrum of Li₂MoO₄ crystals comprises two strongly overlapping bands peaking at 2.08 and 2.25 eV, which are attributed, respectively, to the radiative decay of self-trapped excitons (STEs) and excitons stabilized by crystal structure defects, probably oxygen vacancies [26]. The ratio of their intensities depends on the excitation energy and number of structural defects in the sample. The decay times of the STE and defect-related emissions are substantially different at 4.2 K (130 and 35 µs, respectively), which allows reliable distinguishing of the corresponding emissions in time-resolved spectroscopy experiments. The ratio of the integrated intensity of the STE emission to that of the defect related emission, both calculated from a decay curve recorded at excitation 4.75 eV,

was suggested in [26] as a measure of the relative number of defect-related centers in a crystal. Its value was equal to 4.7 in the best Li_2MoO_4 crystals grown by Low-Temperature-Gradient Czochralski technique and 2.7 in the case of growth by the conventional Czochralski method studied in [26]. The decay curve recorded for a present crystal at 4.2 K is shown in the inset of Figure 5. It can be reliably decomposed into two decay components with decay times τ_1 =130 and τ_2 =35 μ s, whereas the ratio of their intensities S_1/S_2 =2.3 is close to 2.7. This shows clearly that large-size single crystals grown here by an optimized Czochralski technique possess a relative number of lattice defects and structural quality similar to those of the crystals grown by the conventional Czochralski method.

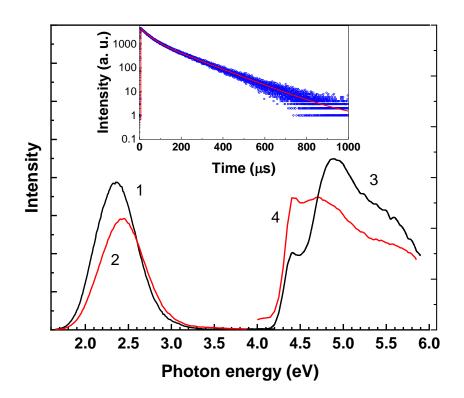


Figure 5 Emission spectra of a Li₂MoO₄ crystal measured at excitation energies 4.8 (1) and 4.6 eV (2) and excitation spectra for 2.7 eV (3) and 1.95 eV (4) emissions. Inset: the decay curve measured at excitation energy 4.75 eV (blue circles) and fitting of the experimental curve with two decay components of 35 and 130 μs (red solid line).

4 Stresses in the growing crystal

Metal molybdates are known as fragile materials where cracks are expected to occur during growth [18]. As depicted in figure 6, one of the 40 mm diameter crystal, grown from 3N commercial molybdate, has shown a transverse crack at the end of growth. The crack plane was characterized as (0001) by Laue pattern analysis, which is the weakest plane in such crystals. The specific orientation is due to seed at 15° from $< 11\overline{2}0 >$.



Figure 6 Li₂MoO₄ crystal, 40 mm in diameter, and 366 g in mass, showing crack during growth process.

In order to understand the origin of this crack better, stresses occurring during growth have been calculated by finite element numerical simulation of heat transfer and thermo-elasticity, by the Comsol ® software. The modelling procedure is the same as presented in [21, 22], applied to the specific set-up configuration used for growing this crystal. Several pseudo-stationary calculations (i.e. taking into account pulling rate and latent heat release) have been performed. Figure 7 shows the temperature field at three times during the growth. The

interface deflection is small, < 4 mm. The temperature gradient in the solid reaches a maximum of 50 K.cm⁻¹, and is always situated close to the triple phase line.

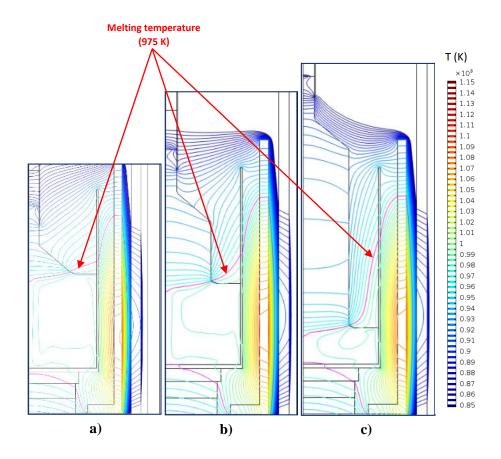


Figure 7 Temperature field in the crystal and surroundings at three times during growth: a) end of shoulder, b) 3cm of cylinder, c) end of growth.

Thermo-elastic stress calculations are based on the computed axi-symmetric temperature field and take into account anisotropic dilatation coefficient [27] and elastic constants [28]. Figure 8 shows the distribution of tensile stresses projected on the crack plane observed on figure 6.

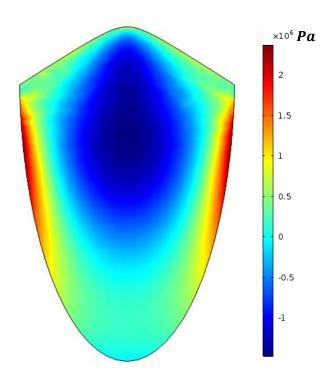


Figure 8 Compressive and tensile stresses during growth, projected on the fracture plane of the crystal shown on figure 6.

It can be seen that the maximal tensile stress is 2 MPa, below the fracture stress measured at high temperature as 3 MPa [28]. This maximum is located on the cylinder side, far from the upper or lower ends of the crack, where it was likely to begin. This result suggests that the crack was rather due to some unexpected vibration or shock occurring during the growth and that its expansion was helped by the precipitates observed in the crystal.

5 Conclusion

After the optimization of the synthesis process, three Li₂MoO₄ crystals, 50 mm in diameter and up to 1 kg in mass, have been successfully grown by a conventional Czochralski process, at a velocity of 2 mm.h⁻¹ and a mass yield larger than 80%. It should be noted that these parameters have not been optimized during this study: in view of the fluency with which the crystals have been pulled, they can certainly be increased. No chemical deposit has been

observed in the setup after growth, so that material evaporation, if any, can be considered as negligible.

This growth ease and reproducibility is certainly due to the combination of a careful optimization of the furnace design, by numerical simulation, and the efficiency of the Cyberstar® control software. By the way, the crystals shown on figure 3 have been produced after only ten experiments, including mechanical and power adjustment. This demonstrates the usefulness of numerical simulation optimization, in terms of time, manpower and money saving.

Dislocation densities are less than 10^4 cm⁻² for the best crystals and the luminescence properties are similar to those of previously reported smaller crystals grown by a conventional Czochralski process.

One smaller crystal, grown from less pure raw material and having inclusions, has shown a crack after growth. Computation of thermo-elastic stresses during this growth show that the crack was likely due to unexpected shock occurring during the growth, rather than to too high thermo-elastic tensile stresses.

All in all it appears that the numerically optimized conventional Czochralski pulling is an efficient process for the mass production of Li₂MoO₄ single crystals. One bolometer has been built from one crystal and is now under evaluation at the Canfranc underground laboratory.

Acknowledgments

This research has been financially supported by the ANR-Clymène project, ref. ANR-16-CE08-0018-02. The support of the Estonian Research Council (project PUT PRG111) and ERDF funding in Estonia granted to the Centre of Excellence TK141 "Advanced materials and high-technology devices for sustainable energetics, sensorics and nanoelectronics (HiTechDevices)" (grant no. 2014-2020.4.01.15-0011) is also gratefully acknowledged.

Growths were performed with a Cyberstar pulling system and control software, provided by the Cristalinnov technical platform (Sainte Hélène du Lac 73800, France). Laue diffraction experiments were performed with the help of H. Roussel, from the Grenoble-INP CMTC characterization platform.

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