

Formation of SiO₂ buffer layer for LiNbO₃ thin films growth

Citation for published version (APA): Vakulov, Z. E., Klimin, V. S., Rezvan, A. A., Tominov, R. V., Korzun, K., Kots, I. N., Polyakova, V. V., & Ageev, O. A. (2019). Formation of SiO_buffer layer for LiNbO_thin films growth. *Journal of Physics: Conference Series*, *1410*(1), Article 012042. https://doi.org/10.1088/1742-6596/1410/1/012042

DOI: 10.1088/1742-6596/1410/1/012042

Document status and date:

Published: 20/12/2019

Document Version:

Publisher's PDF, also known as Version of Record (includes final page, issue and volume numbers)

Please check the document version of this publication:

• A submitted manuscript is the version of the article upon submission and before peer-review. There can be important differences between the submitted version and the official published version of record. People interested in the research are advised to contact the author for the final version of the publication, or visit the DOI to the publisher's website.

• The final author version and the galley proof are versions of the publication after peer review.

 The final published version features the final layout of the paper including the volume, issue and page numbers.

Link to publication

General rights

Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

- · Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain
 You may freely distribute the URL identifying the publication in the public portal.

If the publication is distributed under the terms of Article 25fa of the Dutch Copyright Act, indicated by the "Taverne" license above, please follow below link for the End User Agreement:

www.tue.nl/taverne

Take down policy

If you believe that this document breaches copyright please contact us at:

openaccess@tue.nl

providing details and we will investigate your claim.

PAPER • OPEN ACCESS

Formation of SiO_2 buffer layer for LiNbO_3 thin films growth

To cite this article: Z E Vakulov et al 2019 J. Phys.: Conf. Ser. 1410 012042

View the article online for updates and enhancements.

You may also like

- Microscopic analysis of the chemical reaction between Fe(Te, Se) thin films and underlying CaFe
 A Ichinose, F Nabeshima, I Tsukada et al.
- The role of texturing and thickness of oxide buffer layers in the superconducting properties of Fe(Se.Te) Coated Conductors G Sylva, E Bellingeri, C Bernini et al.
- G Sylva, E Dennigen, C Dernin et al.
- <u>Epitaxial growth technique for single-</u> <u>crystalline PbTiO₃ thin film on Si substrate</u> <u>using an HfO₂ buffer layer</u> Hansol Park, Takeshi Kijima and Hitoshi Tabata



This content was downloaded from IP address 131.155.78.78 on 14/02/2022 at 10:05

Formation of SiO₂ buffer layer for LiNbO₃ thin films growth

Z Vakulov¹, V S Klimin^{1,2}, A A Rezvan², R V Tominov¹, K Korzun³, I N Kots¹, V V Polyakova¹, O A Ageev^{1,2}

1410 (2019) 012042

¹Research and Education Center «Nanotechnology», Sothern Federal University, Taganrog 347922, Russia

²Institute of Nanotechnologies, Electronics and Equipment Engineering, Southern Federal University, Taganrog 347922, Russia

³Department of Applied Physics, Eindhoven University of Technology, 5600 MB, The Netherlands

Abstract. This paper shows the results of study of the effect of SiO_2 buffer layer thickness on the morphological parameters of nanocrystalline LiNbO₃ films formed by pulsed laser deposition. It has been established that with increasing in the thickness of SiO_2 buffer layer from 10 nm to 50 nm, the roughness of LiNbO₃ films decreases from 5.1 nm to 4.4 nm. The minimum value of the grain diameter (118 nm) corresponds to the thickness of the buffer layer equal to 50 nm. The results obtained can be used in the design and manufacture of integrated acousto-optic and piezoelectric devices, as well as sensitive elements of sensors using various effects of surface acoustic waves.

1. Introduction

Modern electronics faced with necessity of creation new principles of power elements formation, since the capabilities of electrochemical power sources cannot fully meet its growing demands [1-4]. Moreover, there are certain difficulties associated with the disposal of waste batteries. In this regard, active research on the modernization of existing structures and technologies of formation, as well as the development of fundamentally new energy sources based on nanomaterials that meet the requirements of efficiency, versatility and environmental safety are being conducted [5-7].

A promising direction in the field of battery development is the development of piezoelectric transducers using mechanical energy of the environment as a source of power supply for wearable electronics (smart watches, mobile phones, pacemakers) [8-13]. Using of such piezoelectric transducers can expand the capabilities of wearable electronics devices, significantly increasing the time of their autonomous operation and reducing the weight and dimensions of the power supply elements.

The creation of devices based on ferroelectric and piezoelectric films at the present stage of technology development is difficult due to the problems associated with the insufficient development of the technology for their production, as well as the need to use additional buffer layers. Most ferroelectric and piezoelectric materials (BaTiO₃, PbTiO₃, LiNbO₃) are multicomponent oxides, which significantly complicates the technological process of their production and their integration with silicon technology of micro- and nanoelectronics [14-18]. Pulsed laser deposition is one of the promising method for the formation of multicomponent oxides films, due to a large number of technological parameters, which allows control the composition of the formed films, as well as their morphological and electrical parameters [19].

The purpose of this work is determination of influence regularities of SiO_2 buffer layer thickness on the morphological parameters of LiNbO₃ nanocrystlline films fabricated by pulsed laser deposition.

2. Materials and Methods

For fabrication of naocrystalline LiNbO₃ films we used nanotechnological cluster complex NANOFAB NTK-9 (NT-MDT, Russia), comprising PLD module Pioneer 180 (Neocera Co., USA). LiNbO₃ target were ablated by excimer KrF laser (λ =248 nm). Energy density on target surface was maintained at 2.0 J/cm². The quantity and frequency of laser pulses were 50 000 and 10 Hz respectively. Target-substrate distance was 115 mm.

LiNbO₃ films were deposited on Si substrates (1 cm x 1 cm) with buffer layer of SiO₂. All substrates were subjected to preliminary purification in inorganic solvents to remove surface contamination. The morphology of the obtained film was researched by Ntegra probe nanolab (NT - MDT, Russia) and scanning electron microscop NOVA Nanolab 600 (FEI Co., Netherlands).

The formation of Si and SiO₂ films was carried out by plasma chemical deposition in a combined discharge plasma. The combined capacitive and inductively coupled plasma discharge allows controlling the granularity and roughness of the resulting dielectric surface. Film growth modes $P_R = 2$ Pa, gas flows $N_{SiH4} = 5 \text{ cm}^3/\text{min}$, $N_{N2O} = 60 \text{ cm}^3/\text{min}$, power of the capacitive plasma source was $W_{RIE} = 10 \text{ V}$, power of the source inductively coupled plasma $W_{ICP} = 400 \text{ V}$, deposition time 54 minutes.

3. Results

Figure 1 shows the results of AFM images of $LiNbO_3$ films obtained on SiO_2 buffer layer with different thickness.



1410 (2019) 012042 doi:10.1088/1742-6596/1410/1/012042



Figure 1 (a, b, c) AFM images and profilograms of LiNbO₃ nanocrystalline films obtained on SiO₂ buffer layer with different thickness: 10 nm (a), 50 nm (b), 100nm (c)

Figure 2 shows the values of surface roughness and grain diameter of nanocrystalline $LiNbO_3$ films obtained on SiO_2 buffer layer with different thickness.



Figure 2 (a, b) Dependences of the morphological parameters of LiNbO₃ nanocrystalline films on the thickness of the SiO₂ buffer layer: grain size (a) and surface roughness (b)

As the thickness of the buffer layer increases from 10 to 100 nm, the roughness of the nanocrystalline $LiNbO_3$ films decreases due to the activation of the process of interatomic interaction of the materials of the formed film. Increasing in the grain size of $LiNbO_3$ films with an increase in the thickness of the buffer layer may be associated with an increase in the grain diameter of the SiO_2 layer and the effect of "inheriting" the morphology of the buffer layer. Figure 3 shows SEM images of $LiNbO_3$ nanocrystalline films obtained on SiO_2 buffer layer with different thickness.



Figure 3 (a, b, c) SEM images of LiNbO₃ nanocrystalline films obtained on SiO₂ buffer layer with different thickness: 10 nm (a), 50 nm (b), and 100 nm (c)

HV WD HFW mode det mag □ 10.00 kV 5.1 mm 2.56 µm SE TLD 100.000 x

4. Discussion and Conclusions

In this work, experimental studies were carried out on the deposition of a buffer layer of SiO_2 by plasma chemical deposition. LiNbO₃ films were obtained by pulsed laser deposition. It has been established that with an increase in the thickness of the buffer layer of SiO_2 from 10 nm to 50 nm, the roughness of LiNbO₃ films decreases from 5.1 nm to 4.4 nm. The effect of the thickness of the buffer layer of SiO_2 on the morphological parameters of LiNbO₃ nanocrystalline films formed by pulsed laser deposition was studied. The obtained films with these roughness indicators can be used in the formation of solar cells, as well as elements of nanophotonic devices.

Acknowledgments

This work was financially supported by the Russian Foundation for Basic Research (project № 18-29-11019 mk) and Grant of the President of the Russian Federation No. MK-3512.2019.8. The work was done on the equipment of the Research and Education Centre «Nanotechnology», Southern Federal University.

References

[1] Schlachter F 2013 Proc. Natl. Acad. Sci. USA 110 5273

[2] Granqvist, C.G. 2007 Solar Energy Materials and Solar Cells **91** (17) pp. 1529-1598

Jayaraj, M.K., Antony, A., Ramachandran, M. 002 Bulletin of Materials Science, 25 (3) pp. 227-230

[3] Kots I N, Kolomiitsev A S, Lisitsyn S G, Polyakova V V, Klimin V S, Ageev O A 2019 Russian Microelectronics **48(2)** p 72-79

[4] Kon, M., Song, P.K., Mitsui, A., Shigesato, Y. 2002 Japanese Journal of Applied Physics, Part 1: Regular Papers and Short Notes and Review Papers, **41** (10) pp. 6174-6179

[5] Liu, D.-S., Wu, C.-Y., Sheu, C.-S., Tsai, F.-C., Li, C.-H. 2006 Japanese Journal of Applied

Physics, Part 1: Regular Papers and Short Notes and Review Papers, **45 (4 B)**, pp. 3531-3536

[6] Krivchenko, V.A., Lopaev, D.V., Pashchenko, P.V., Pirogov, V.G., Rakhimov, A.T., Suetin, N.V., Trifonov, A.S. 2008 *Technical Physics*, **53** (8), pp. 1065-1069

[7] Kim H S, Kim J H, Kim J 2011 Int. J. Pr. Eng. Man. 12 1129

[8] Klimin V S, Vakulov Z E, Tominov R V, Varzarev Y N, Clemente I E, Miakonkikh A V, Rudenko K V, Ageev O A 2019 Proc. of SPIE **11022** 110221E-2

[9] le Hung, N., Ahn, E., Jung, H., Kim, H., Kim, D. 2010 Journal of the Korean Physical Society, **57** (**61**), pp. 1784-1788.

[10] Dang, W.L., Fu, Y.Q., Luo, J.K., Flewitt, A.J., Milne, W.I. 2007 Superlattices and Microstructures, 42 (1-6), pp. 89-93

[11] Klimin V S, Tominov R V, Avilov V I, Dukhan D D, Rezvan A A, Zamburg E G, Smirnov V A, Ageev O A 2019 Proc. of SPIE **11022** 110220E-1

[12] Youssef, S., Combette, P., Podlecki, J., Al Asmar, R., Foucaran, A. 2009 Crystal Growth and Design, 9 (2), pp. 1088-1094

[13] Minami, T., Miyata, T., Yamamoto, T., Toda, H. 2000 Journal of Vacuum Science and Technology A: Vacuum, Surfaces and Films, **18** (4 II), pp. 1584-1589.

[14] Klimin V S, Rezvan A A, Ageev O A 2019 Proc. of SPIE 11022 1102220-1

[15] Vakulov Z E, Varzarev Y N, Gusev E Y, Skrylev A V, Panich A E, Miakonkikh A V,

Klemente I E, Rudenko K V, Konoplev B G, Ageev O A 2019 Russ. Microelectron. **48** 59

[16] Klimin V S, Kots I N, Polyakova V V, Rezvan A A, Ageev O A 2019 Proc. of SPIE 11022 110221R-4

[17] Chang, Z., Gaoyuan, M., Dostanko, A.P., Golosov, D.A., Zavatskiy, S.M. 2010 J. Appl. Opt., **31**, p. 855.

[18] Golosov, D.A., Zavadski, S.M., Svadkovski, I.V., Melnikov, S.N. 2010 Vakuum. Tekh. Tekhnol., 20, p. 227

[19] Subramanyam, T.K., Srinivasulu Naidu, B., Uthanna, S. 2000 *Crystal Research and Technology* **35** (10) pp. 1193-1202.