

10th Joint Conference on Chemistry

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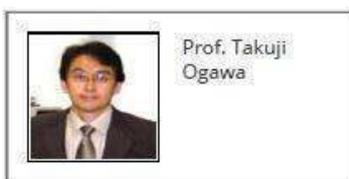
Editorial Materials (preface and photograph)

PREFACE

The 10th Joint Conference on Chemistry is an international conference organized by 4 chemistry departments of 4 universities in central java, Indonesia. The universities are Sebelas Maret University, Diponegoro University, Semarang State University and Soedirman University. The venue was at Solo, Indonesia, at September 8-9, 2015. The total conference participants are 133 including the invited speakers. The conference emphasized the multidisciplinary chemical issue and impact of today's sustainable chemistry which covering the following topics:

- Material innovation for sustainable goals
- Development of renewable and sustainable energy based on chemistry
- New drug design, experimental and theoretical methods
- Green synthesis and characterization of material (from molecule to functionalized materials)
- Catalysis as core technology in industry
- Natural product isolation and optimization

The conference speakers are:



*Single molecular electronic devices and their integration,
Physical Organic Chemistry,
Osaka University
Osaka University, Japan.
Research*



*Catalysis- A powerful means for sustainable developments,
Institute of Inorganic Chemistry
Leipzig University, Germany.
Research*



*Plasma science and technology,
Shizuoka University
Shizuoka University, Japan.
Research*





Prof. Santiago
Gómez Ruiz

In search of novel sustainable systems based on nanostructured materials for different catalytic applications.,
Rey Juan Carlos University,
Móstoles (Madrid), Spain
Research



Dr. Henning
Störz

Current State and Future Prospects Including Monomer Preparation from Biomass
Institute of Agriculture
Technology,
Johann Heinrich Von Thuenen
Institute.
Braunschweig Germany.



Dr. Leny Yuliaty

Photocatalytic hydrogen production,
Ibnu Sina Institute
University Teknologi Malaysia.
Research



Prof. Effendy,
PhD

Molecular Structure Analysis by Single X-ray Diffraction Method,



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Catalytic and sustainable Processes in PERTAMINA,



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Synthesis of $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ by low-temperature solid-state reaction and its microstructure

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Synthesis of $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ by low-temperature solid-state reaction and its microstructure

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Abstract. This study aims to synthesis the Ni-doped $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ ($x = 0; 0.02; 0.04; 0.06; 0.08; 0.1$) by low temperature solid-state reaction. The microstructure of the product was evaluated based on the mole ratio of Ni/Mn of the precursors. The structural analysis of $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ was analyzed by x-ray diffraction equipped with the Direct Method of win PLOT package program and Diamond. It was found that doping with Ni could change the size, crystallinity and microstructure of $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$. The $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ solids have a cubic structure with a space group of $Fd\bar{3}m$. The increase in the doping content does not affect the structure. The particle size of the products is about 150-500 nm. The crystallinity of the solids tends to increase with the increase in Ni content. However, the increase of Ni content in the product causes the lattice parameters of the unit cell to decrease.

1. Introduction

Lithium-ion batteries are widely used for energy storage, portable devices, and electric vehicles. Although many new cathode materials are developed, LiMn_2O_4 is still well recognized as a promising cathode material for these batteries [1]. The spinel-structured LiMn_2O_4 offers attractive alternative material over commercially available LiCoO_2 . LiMn_2O_4 is cheap, non-toxic, and safe [2]. However, it has severe capacity fading in the application [3]. The charge-discharge cycling capability of LiMn_2O_4 at high temperatures can be improved by the substitution of Mn by other transition metals to give the corresponding $\text{LiM}_x\text{Mn}_{2-x}\text{O}_4$ where $M = \text{Co}, \text{Mg}, \text{Al}, \text{Cr}, \text{Ni}, \text{Fe}, \text{Ti}$ and Zn [4]. $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ has a discharge capacity of 140 mAhg^{-1} and open circuit potential of 4.7 V. This energy density is tunable by doping [5].

It is documented well that electrochemical performance of the battery cathode is affected by the powder properties such as structure, particle size, grain morphology, specific surface area and crystallinity [6]. In addition to its large surface area, fast transport of mass and charge, there are many properties to explore. The large surface area and small particle size can improve solubility in the electrolyte. The alteration in chemical properties may make the electrode materials sensitive to the presence of impurities [7].

The electrochemical property of LiMn_2O_4 -based spinel highly depends on its synthetic routes, such as the Pechini process, sol-gel, emulsion method, the citric method [8], etc. However, most of these



methods involve many treatment processes or expensive reagent, which is time-consuming and high cost for commercial applications. We report here on the synthesis of series of $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ using a low-temperature solid-state reaction at various Ni/Mn mole ratios. The physical properties and microstructures of the synthesized materials were investigated.

2. Experimental Section

2.1. Synthesis of MnO_2 and $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$.

An analytical grade of $\text{Mn}(\text{CH}_3\text{COO})_2$ and $\text{Na}_2\text{S}_2\text{O}_8$ (Aldrich) were used. All other chemicals were used without further purification. In a typical synthesis, $\text{Mn}(\text{CH}_3\text{COO})_2$ and $\text{Na}_2\text{S}_2\text{O}_8$ with a molar ratio of 1:1 were dissolved at room temperature in 80 mL deionized distilled water. The mixture was stirred to form a clear homogeneous solution. The solution was transferred to the flask and heated at 120 °C for 12 h. The obtained powder was subsequently dried at 300 °C for 1 h in the oven.

Synthesis of $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ is the following; LiOH of 0.00143 moles, 0.0000286 moles of $\text{Ni}(\text{CH}_3\text{COO})_2$ and 0.0028314 moles were dispersed into high purity ethanol to form a thick slurry and stirred. The product was separated and dried at room temperature. The above process was repeated three times. The $\text{LiNi}_{0.02}\text{Mn}_{1.98}\text{O}_4$ powder was ignited at 750 °C for 10 hours. The same procedure was followed for synthesizing the materials with different mole ratio.

2.2. Characterization of the $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ microstructure.

The obtained MnO_2 and $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ powders were analyzed using X-ray powder diffractometer (XRD). The XRD patterns were obtained on Rigaku Miniflex 600-Benchtop XRD instrument using Cu K α radiation ($\lambda = 1.5406 \text{ \AA}$) at ambient temperature. The XRD instrument was set to operate at 40 kV and 15 mA. The XRD data was obtained with a 2θ interval ranging from 20° to 90°. The Rietveld analysis was conducted with the Fullprof software package [9] to refine the X-ray diffraction data. The refined parameters are a unit cell, scale factor and full width at half-maximum (FWHM). The SEM images were obtained using JEOL JSM-6510LASEM. The effect of the nickel content on the structure of LiMn_2O_4 was studied using energy dispersive X-ray spectroscopy (EDX). The EDX analysis was also used to analyze the presence of Mn, O and Ni elements in the prepared materials. The Brunauer-Emmett-Teller (BET) data was collected based on adsorption data in relative pressure (P/Po) of about 0.30 [10]. The BET calculation was done in ASAP 2020 V4.01 software from Micrometrics.

3. Results and discussion

3.1. XRD Patterns

Figure 1 shows XRD patterns of produced $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$. High peak intensity suggests that the $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ has good crystallinity. Figure 2 shows the variation of the lattice constant of the materials with different compositions. The Rietveld refinement data obtained using Full Prof indicates that lattice parameter decreases with the increase in doping content. The substitution of Mn^{3+} by Ni^{2+} in the octahedral sites leads to the decrease in the lattice parameters. Note that ionic radius of Ni^{2+} is 0.560 Å, smaller than that of Mn^{3+} , which is 0.645 Å. The difference in the ionic radius between Ni^{2+} and Mn^{3+} is small, therefore only a few of change the lattice parameter.

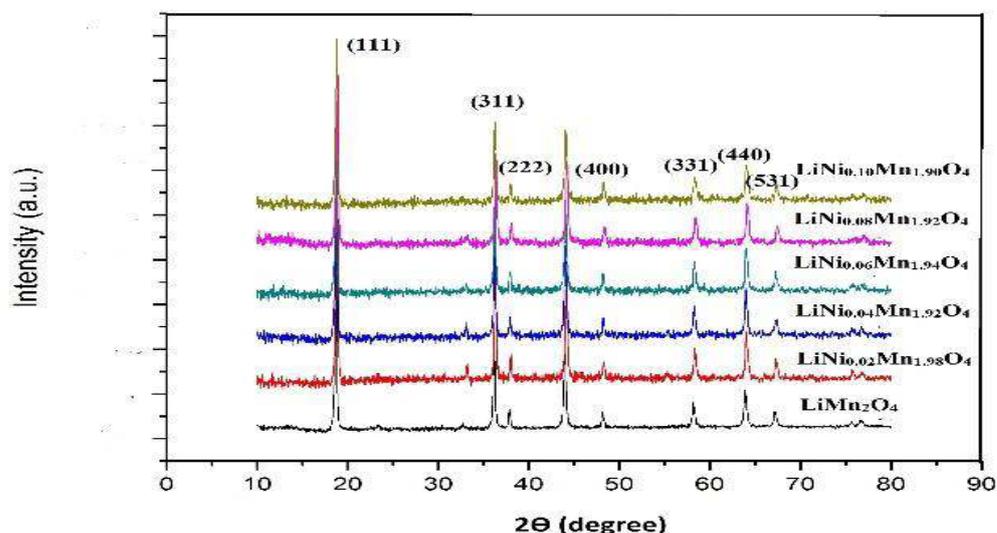


Figure 1. XRD Patterns of $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ Ni/Mn mole ratios

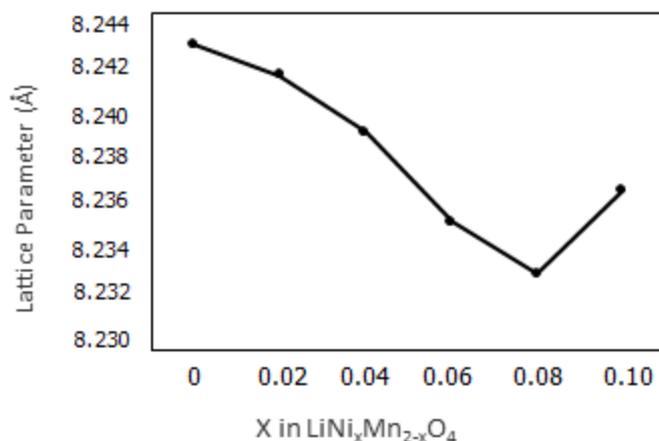


Figure 2. Variation of lattice parameter with x in $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$

Figure 3 shows the XRD patterns of $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$. The products have a cubic phase with space group of $Fd\bar{3}m$ as indicated by results of analysis using winPLOTR package program and Diamond. The results of Rietveld are shown in Fig. 3. The experimental points are given as dot (.) and theoretical data are shown as solid line. The difference between theoretical and experimental data is shown as a bottom line. The vertical lines represent the Bragg's allowed peaks. The X-ray diffraction data show at $31\text{--}33^\circ$, which can be assigned to a Mn_2O_3 phase. The presence of Ni in the structure causes the appearance of NiMnO_3 , which is indicated by the presence of the peaks at $31\text{--}33^\circ$ [11]. It seems that lithium ions occupy the tetrahedral (8a) site. After the modification, the lattice shrinkage is due to the smaller ionic radii of Ni^{2+} , which replaced the Mn^{3+} at 16d sites. On the other hand, Ni^{2+} substituting part Mn^{3+} can enhance the content of Mn^{4+} to keep the charge balance, and Mn^{4+} is with smaller ionic radii than that of Mn^{3+} [12]. Oxygen atoms are arranged in the cubic-closed packing 32e.

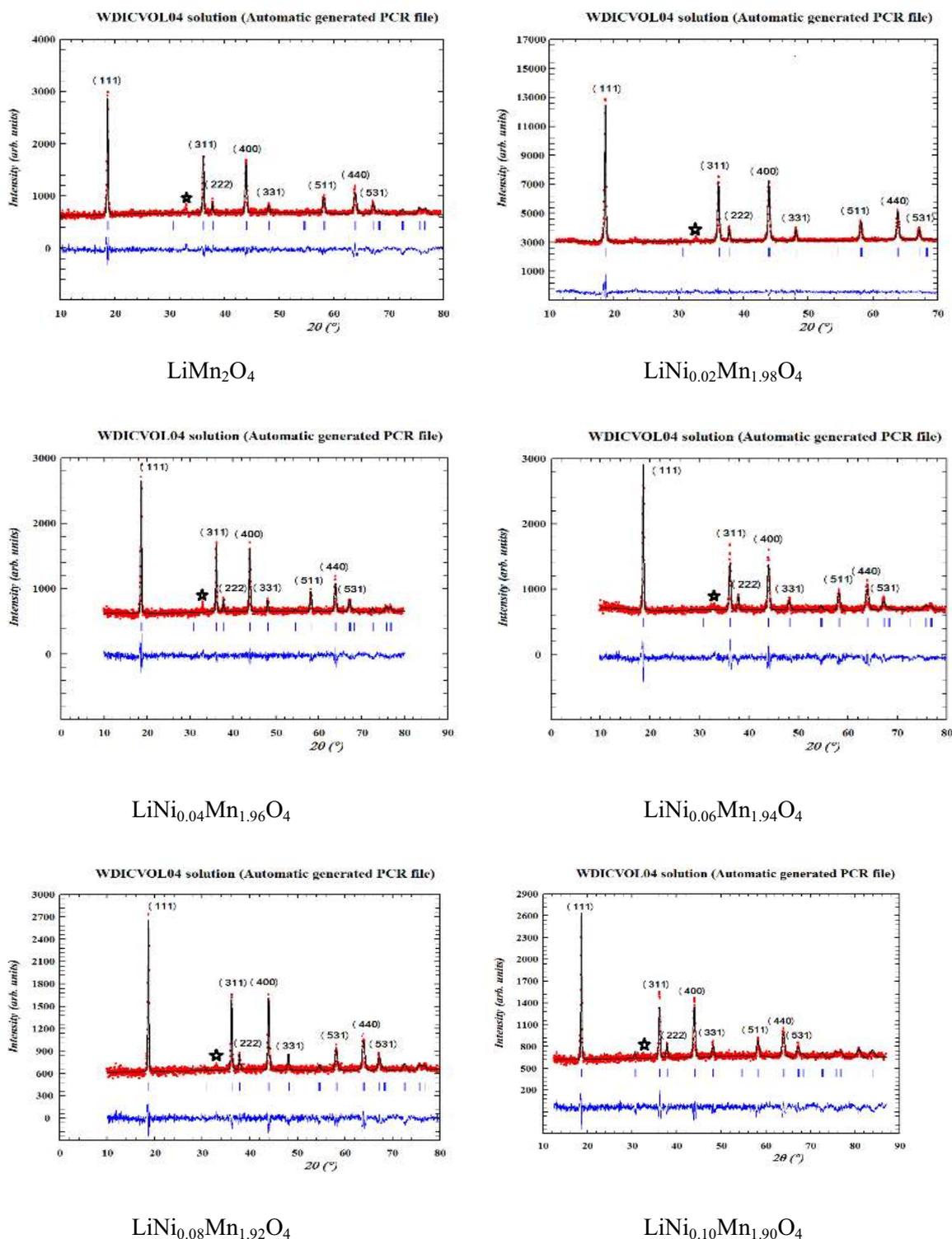


Figure 3 XRD refinement data of $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ with various Ni/Mn mole ratios

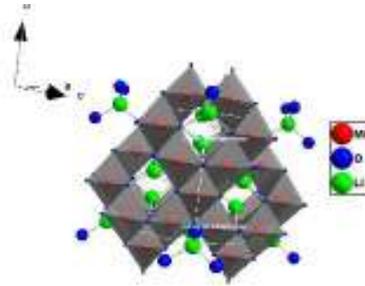


Figure 4. Microstructure of $\text{LiNi}_{0.1}\text{Mn}_{1.9}\text{O}_4$

3.2. SEM Images

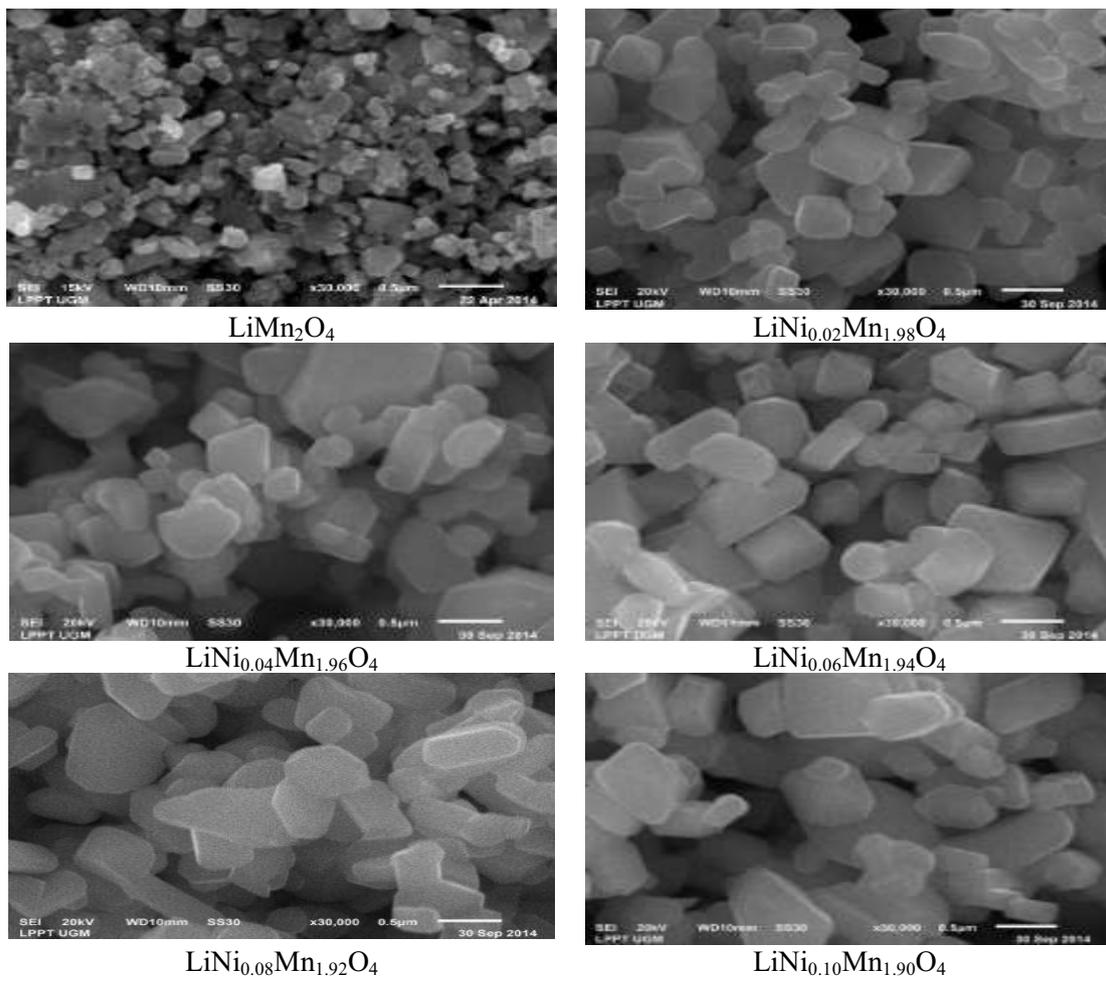


Figure 5. SEM photographs of $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$

The SEM images of the synthesized powders are shown in Figure 5. The average particle size of the nickel-doped samples is smaller than that of undoped LiMn_2O_4 . The doping of $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ with Ni cause the tendency of nucleation process overcomes the tendency of crystal growth. The Ni-doped LiMn_2O_4 powders have irregular particle size. The particle sizes of $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ are in the range of 150 to 500 nm. The increase in the nickel doping leads to increase in LiMn_2O_4 particle size. The morphology is almost similar.

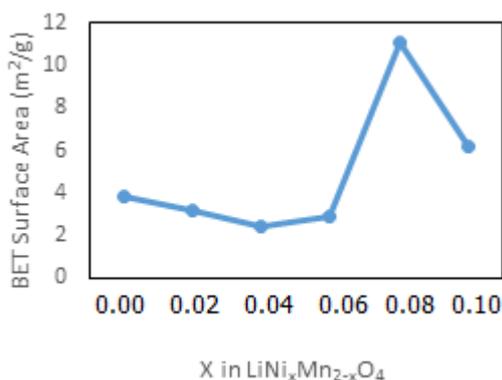


Figure 6. BET surface area of $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$

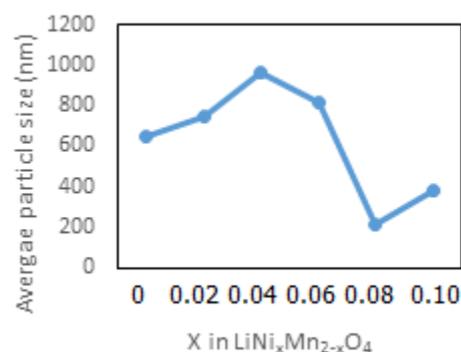


Figure 7. Variation of the average particle size with x in $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$

Figure 6 shows the BET surface area of $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ at various mole ratios. The addition of Ni does not change the BET surface area. However, at a mole ratio of 0.08, there is a noticeable change in the BET surface area from $6 \text{ m}^2/\text{g}$ to $11 \text{ m}^2/\text{g}$. Figure 7 shows the average particle size of the material. The particle size is in the range of 200-900 nm (Table 1). At low doping content, the particle size is relatively constant until it reaches $x = 0.04$. For $x > 0.04$, the particle size decreases along with the increase of doping agent. The particle size is about 300 nm for $x = 0.08$. Good battery performance requires small particle size of cathodes [13]. In general, the particles have a rod shape. The particles are also agglomerated. The agglomerated particles might be beneficial for the production of lithium-ion batteries with high energy capacity [14]. A further test of the energy capacity of $\text{LiNi}_{0.08}\text{Mn}_{1.92}\text{O}_4$ is required. Calculation using Debye-Scherrer equation gives crystallite size of $\text{LiMn}_{2-x}\text{O}_4$ in the range of 30-40 nm. The particle size obtained from the calculation (***) is around 200-900 nm.

Table 1. The crystallite size of the prepared materials as calculated by Debye-Scherrer method, D, and the particle size as calculated from BET surface area data.

x	XRD		BET
	D (nm)*	L (nm)**	$S_{\text{BET}}(\text{m}^2/\text{g})$
0	39.42802	651.2078	3.6454
0.02	41.39063	746.8011	3.1756
0.04	43.1764	965.8867	2.4553
0.06	33.45325	817.3502	2.9015
0.08	36.38719	213.5754	11.104
0.1	36.21143	381.4117	6.2178

*D = $(K\lambda)/B \cdot \cos \Theta$

** L = $6000/(\rho \cdot S_{\text{BET}})$

4. Conclusion

$\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ has been synthesized by the low-temperature solid-state reaction. The $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ powders have a cubic crystal structure with an $Fd3m$ space group. The particle size of the prepared materials is not homogeneous. Doping with Ni leads to changes in size, crystallinity, and microstructure of the products. The average particle size of $\text{LiNi}_x\text{Mn}_{2-x}\text{O}_4$ is about 150-500 nm. The crystallinity of the materials tends to increase with the increase in the Ni doping content. When Ni content in the compound increases, the lattice parameters decrease.

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