

Supporting Information

A molten salt-based nitridation approach for synthesizing nanostructured InN electrode materials

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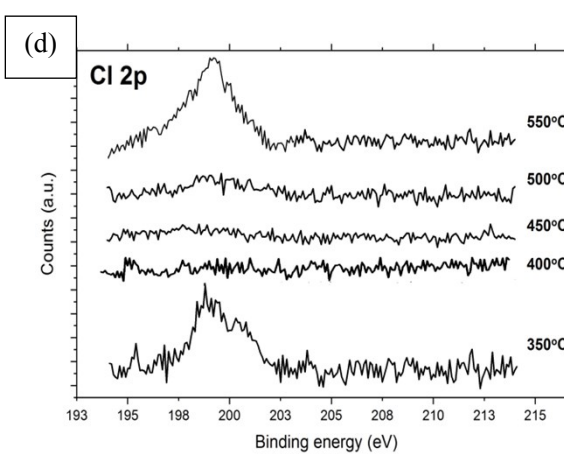
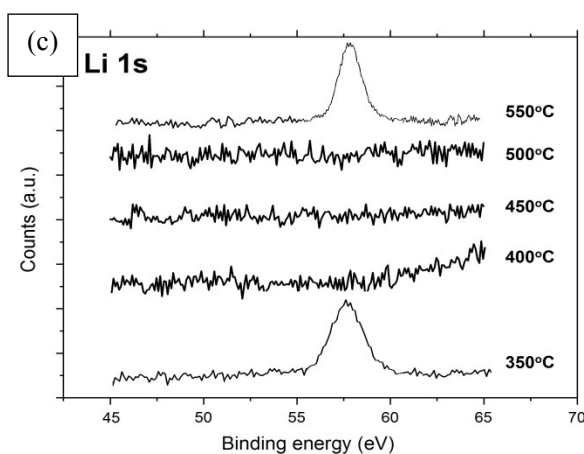
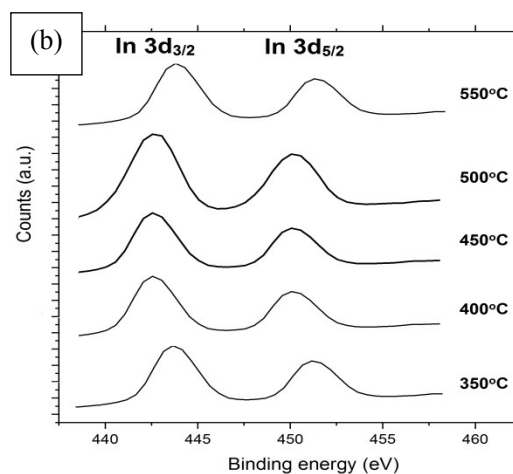
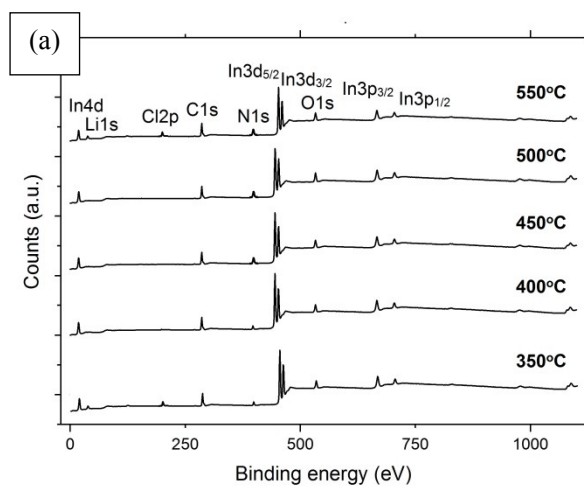
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Contents

- Table S1: The melting points of LiCl-KCl molten salt systems.
- Fig. S1: XPS spectra of InN samples: (a) whole region, (b) In 3d, (c) Li 1s, (d) Cl 2p and (e) K 2p.
- Fig. S2: SEM images of InN crystals prepared by the reaction of InCl₃, LiNH₂ and LiCl-KCl at (a) 350°C, (b) 400°C, (c) 450°C, (d) 500°C and (e) 550°C
- Fig. S3: Brunauer-Emmett-Teller (BET) isotherms and Barret-Joyner-Halenda (BJH) pore-size distribution (inset) of InN synthesized at 400°C, 450°C and 500°C.
- Fig. S4: XRD patterns of InN synthesized using InCl₃ and LiNH₂ without LiCl-KCl. The reactions were carried out at 400°C, 450°C, 500°C and 550°C.
- Fig. S5: SEM images of InN synthesized using InCl₃ and LiNH₂ without LiCl-KCl at (a) 400°C, (b) 450°C, (c) 500°C and (d) 550°C.
- Fig. S6: (a) Absorption spectra and (b) band gap determination of InN thin films.
- Fig. S7: Current potential curves of the InN electrodes under Xe lamp illumination with 64R cut filter.
- Fig. S8: Mott-Schottky plots of InN electrodes for 400°C, 450°C and 500°C samples. Donor densities (N_D) are also presented in the figure.

Table S1. The melting points of LiCl-KCl molten salt systems.

| Composition of molten salts (molar ratio) | | Melting point |
|-------------------------------------------|------|---------------|
| LiCl | KCl | |
| 0 | 100% | 770°C |
| 20% | 80% | 707°C |
| 40% | 60% | 589°C |
| 60% | 40% | 353°C |
| 80% | 20% | 450°C |
| 100% | 0 | 605°C |



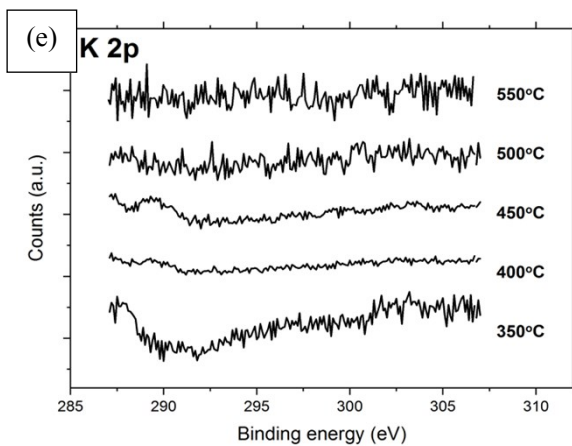


Fig. S1 XPS spectra of InN samples: (a) whole region, (b) In 3d, (c) Li 1s, (d) Cl 2p and (e) K 2p.

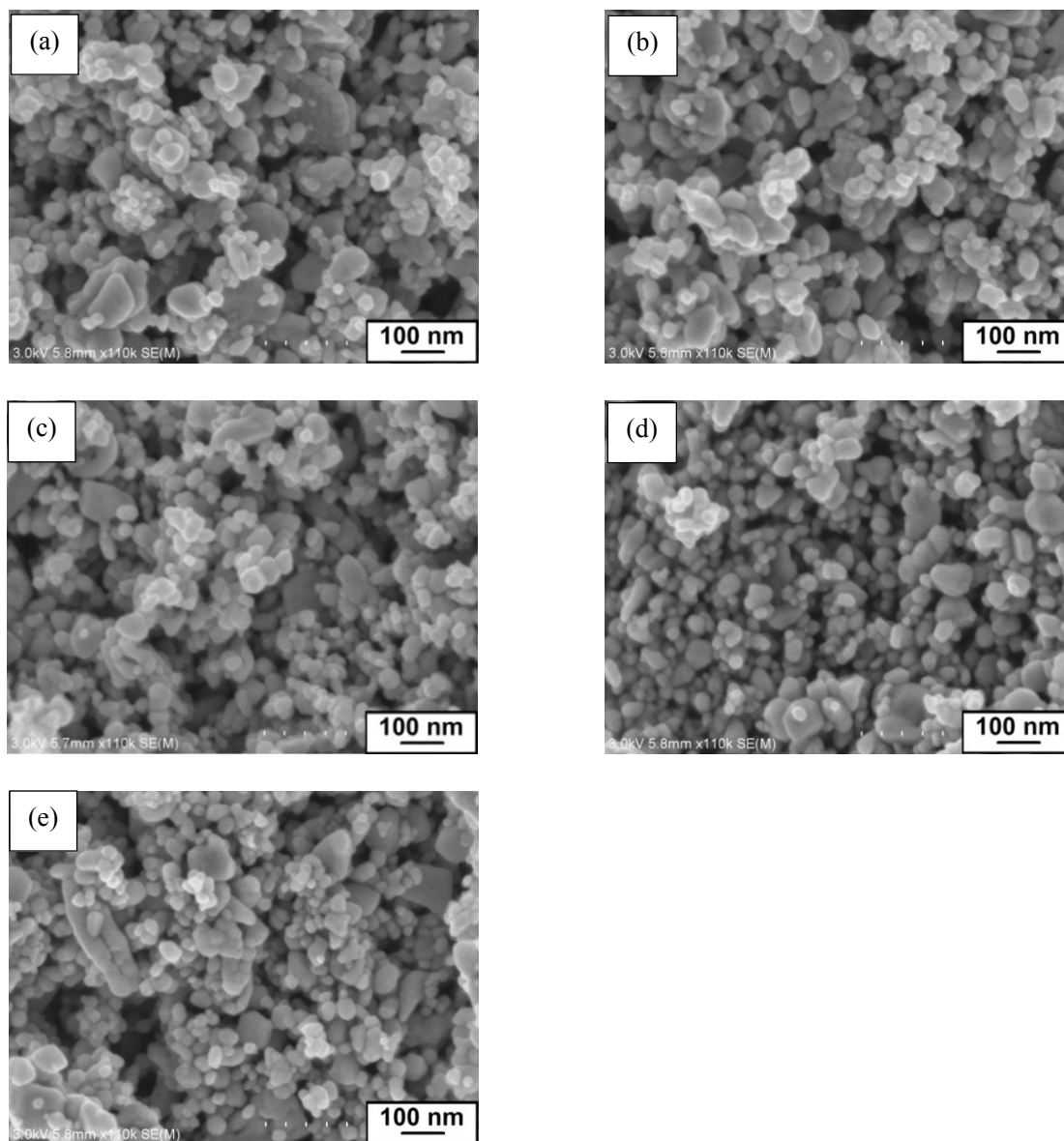


Fig. S2 SEM images of InN crystals prepared by the reaction of InCl_3 , LiNH_2 and LiCl-KCl at (a) 350°C, (b) 400°C, (c) 450°C, (d) 500°C and (e) 550°C.

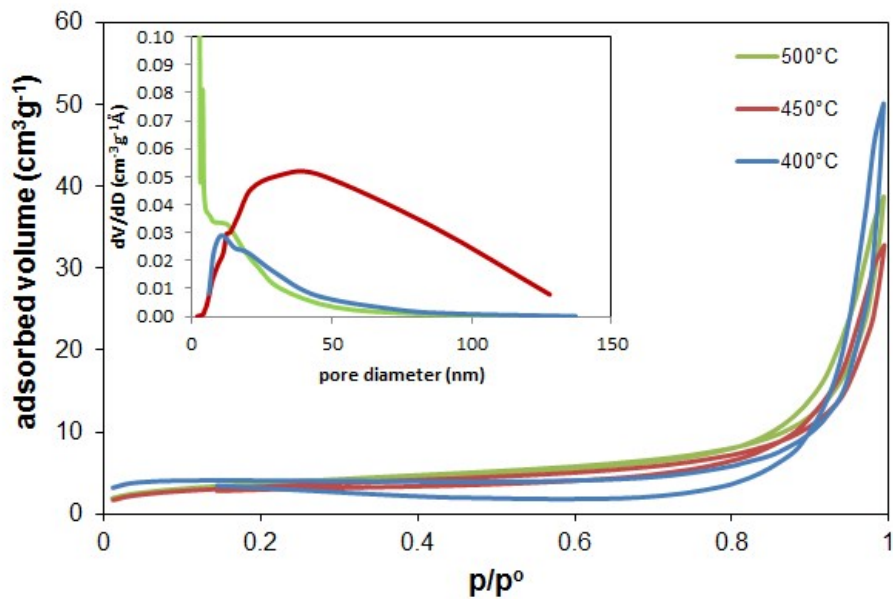


Fig. S3 Brunauer-Emmett-Teller (BET) isotherms and Barret-Joyner-Halenda (BJH) pore-size distribution (inset) of InN synthesized at 400°C, 450°C and 500°C.

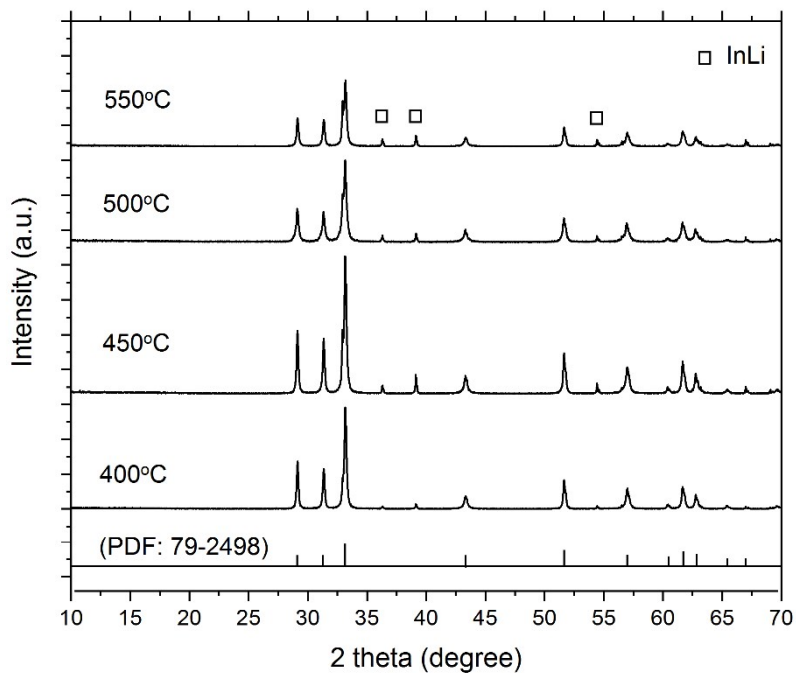


Fig. S4 XRD patterns of InN synthesized using InCl_3 and LiNH_2 without LiCl-KCl . The reactions were carried out at 400°C, 450°C, 500°C and 550°C.

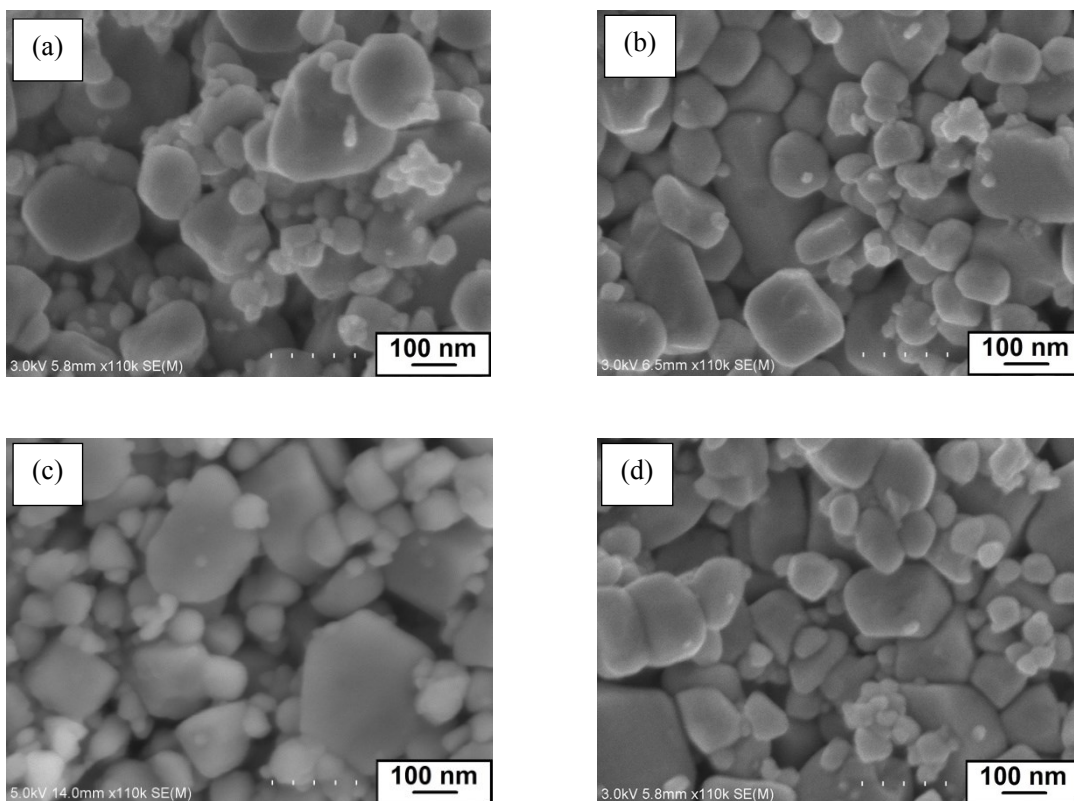


Fig. S5 SEM images of InN synthesized using InCl_3 and LiNH_2 without LiCl-KCl at (a) 400°C, (b) 450°C, (c) 500°C and (d) 550°C.

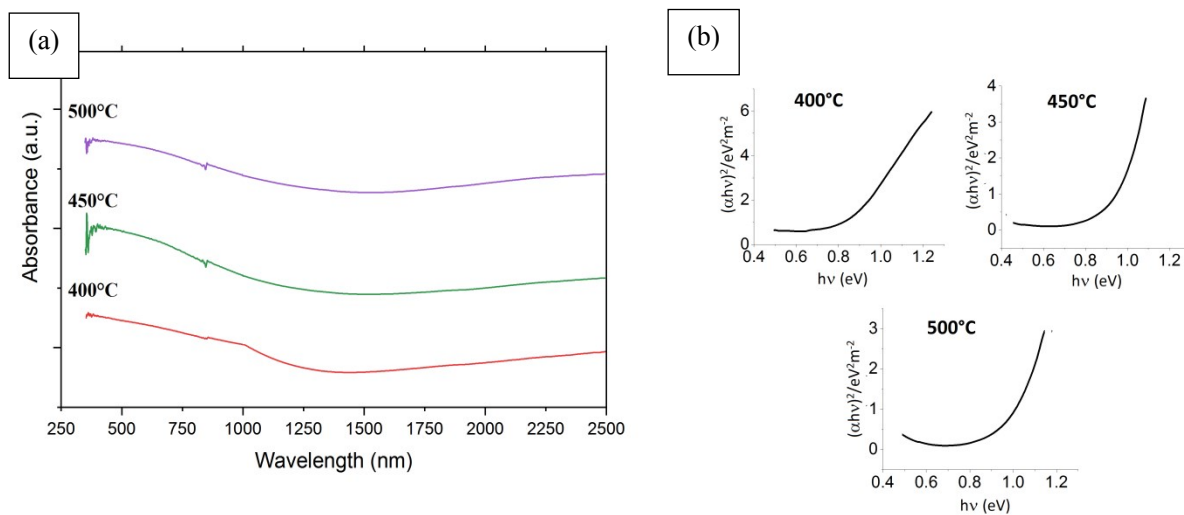


Fig. S6 (a) Absorption spectra and (b) band gap determination of InN thin films.

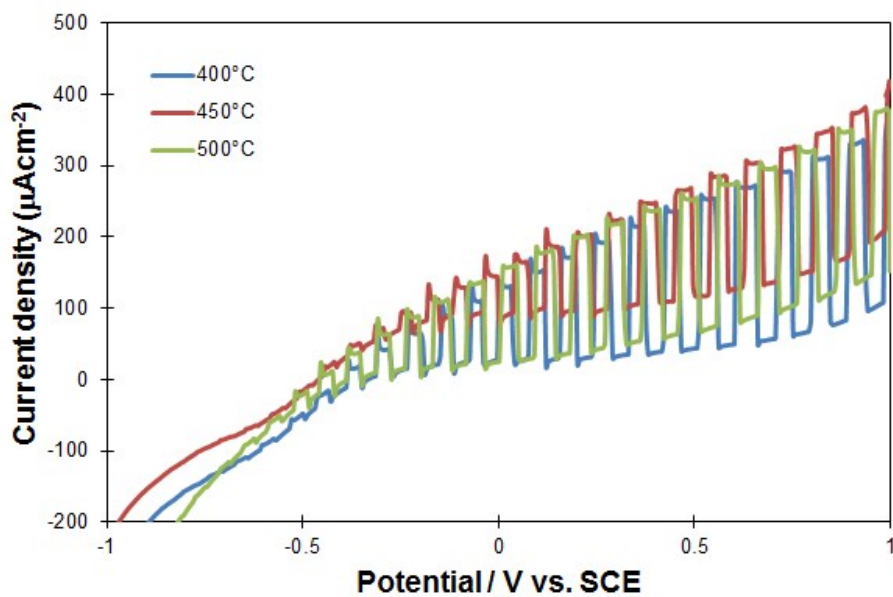


Fig. S7 Current potential curves of the InN electrodes under Xe lamp illumination with 64R cut filter.

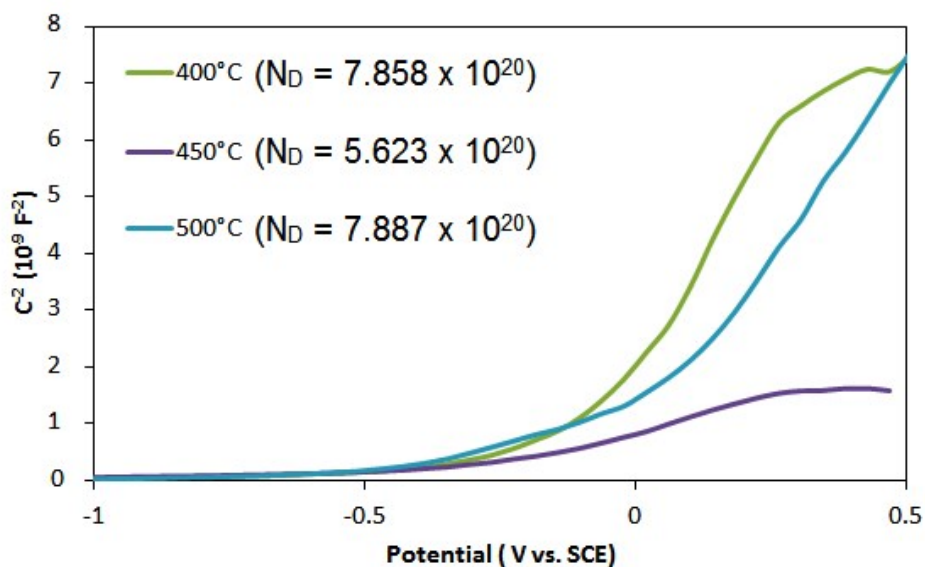


Fig. S8 Mott-Schottky plots of InN electrodes for 400°C, 450°C and 500°C samples. Donor densities (N_D) are also presented in the figure.