

A Thermally Stable Li-ion Battery Electrolyte Using LiTDI

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In recent decades, Li-ion batteries have become ubiquitous, with widespread use not only in portable electronics, but also in commercial airliners, electric vehicles, and even grid-level energy storage systems. Li-ion batteries, however, have been constantly plagued by thermal instabilities with cell performance degrading significantly after short term exposure to moderately elevated temperatures ($\sim 60^\circ\text{C}$). LiPF_6 , the primary lithium salt in state-of-the-art electrolytes, is well known to be the main culprit causing this instability (1-3). In some cases, additives such as LiDFOB have been shown to improve the high temperature performance of Li-ion batteries (4). It is desirable, however, to completely circumvent the persistent challenge of thermal instability by finding a suitable replacement for LiPF_6 that can withstand higher temperatures. Doing so has proven difficult though, as primary salts to replace LiPF_6 must fulfill the many roles necessary for a high performance Li-ion battery electrolyte. These include high conductivity, and stability towards common electrode materials, among others.

A relatively unknown lithium salt, lithium 4,5-dicyano-2-(trifluoromethyl)imidazole (LiTDI) shows promise as a primary lithium salt for use in thermally stable, next-generation electrolytes. Reports have shown that LiTDI-based electrolytes are stable with respect to various cathode materials and have a high conductivity, thus facilitating fast Li^+ cation transport across the cell (5,6). In stark contrast to LiPF_6 , however, TGA and DSC experiments indicate that LiTDI does not degrade or volatilize until a temperature well above 100°C , making it a candidate for Li-ion battery electrolytes with a wider temperature range.

Coin cells containing either LiTDI or LiPF_6 in EC/DEC were cycled at 25°C and 60°C using a $\text{LiNi}_{0.5}\text{Mn}_{0.3}\text{Co}_{0.2}\text{O}_2$ cathode material and Li metal counter electrode (Figure 1). Whereas the state-of-the-art LiPF_6 electrolyte degrades at 60°C —evidenced by the abrupt drop in cell capacity—cells containing the LiTDI electrolyte show an increased capacity at elevated temperature. The relatively stable high capacity of the LiTDI-containing cell at elevated temperature is also noteworthy. LiTDI thus has the potential to be a thermally stable electrolyte salt which may be suitable for replacing LiPF_6 , thus making Li-ion batteries safer and more robust.

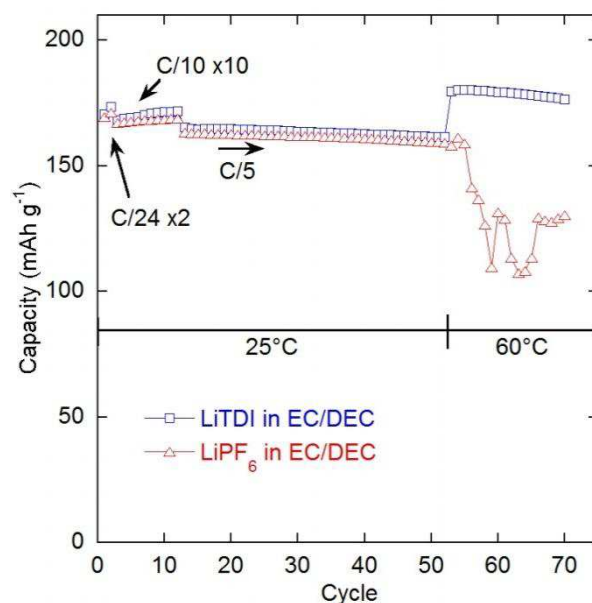


Figure 1. Discharge capacity of $\text{LiNi}_{0.5}\text{Mn}_{0.3}\text{Co}_{0.2}\text{O}_2$ half-cells containing EC/DEC electrolytes with either LiTDI (squares) or LiPF_6 (triangles).

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REFERENCES

1. C. L. Campion, W. Li, and B. L. Lucht, *J. Electrochem. Soc.*, **152**, A2327 (2005).
2. X. Sun, H. S. Lee, X.-Q. Yang, and J. McBreen, *Electrochem. Solid State Lett.*, **5**, A248 (2002).
3. S. E. Sloop, J. K. Pugh, S. Wang, J. B. Kerr, and K. Kinoshita, *Electrochem. Solid State Lett.*, **4**, A42 (2001).
4. M. Xu, L. Zhou, L. Hao, L. Xing, W. Li, and B. L. Lucht, *J. Power Sources*, **196**, 6794 (2011).
5. L. Niedzicki, S. Grugeon, S. Laruelle, P. Judeinstein, M. Bukowska, J. Prejzner, P. Szczecinski, W. Wiczonek, and M. Armand, *J. Power Sources*, **196**, 8696 (2011).
6. L. Niedzicki, G. Z. Zukowska, M. Bukowska, P. Szczecinski, S. Grugeon, S. Laruelle, M. Armand, S. Panero, B. Scrosati, M. Marcinek, and W. Wiczonek, *Electrochim. Acta*, **55**, 1450 (2010).