High Energy Density “Bezel-less” Lithium-ion Battery Using Solvate Ionic Liquid-based Quasi-solid-state Electrolyte

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ABSTRACT

We propose a novel “bezel-less” lithium-ion battery (LIB) structure that enhances the volumetric energy density of LIB. This structure reduces the extra-space volume for seal and accumulation of current collectors, eliminating the bezel part by 16% compared to the conventional aluminum laminated LIBs. The bezel-less LIB using a Ni-rich positive electrode and a graphite negative electrode recorded an energy of 132 Wh and high energy density of 600 Wh L⁻¹ (256 Wh kg⁻¹). The safety of the proposed bezel-less LIB by a combined use of the quasi-solid-state electrolyte layer was successfully demonstrated in overcharge and collapse tests.

Keywords : High Capacity, High Energy Density, Quasi-solid-state Electrolyte, Safety

1. Introduction

Lithium ion batteries (LIBs) are currently used in a wide range of applications, offering a higher energy density than other electrochemical energy storage devices. For large-scale applications including electric vehicles (EVs), distributed power supply for highly efficient use of renewable energy resources and so forth, further increase of energy density is urgently needed. To meet these requirements, efforts are made in R&D of electrode materials to pair high-capacity positive electrodes (Ni-rich type, Li-rich type, sulfur-based conversion positive electrodes, etc.) and negative electrodes (silicon, Li-metal, etc.). One of the challenges for practical applications is enhancing the cycle life of batteries, which is compromised by the formation of high resistive layers, destabilization of the crystal structures, dissolution into electrolyte liquid, collapse of the microstructure accompanied by repeated volume expansion and shrinkage, dendrite growth, etc.

Another possible approach to enhance the energy density of LIBs for large-scale applications such as EVs is by improving its safety. Conventional organic liquid electrolyte solvents used for LIBs have high vapor pressure and are inflammable, so LIBs can catch fire and explode if unexpected accidents occur. Hence, current rechargeable systems, namely the battery packs, require a safety system and a cooling space, which do not contribute to energy storage but reduce the energy density of the LIB packs. If the constituent materials of LIBs can be enhanced, such safety systems and cooling spaces will become unnecessary and could be removed, thus improving the battery pack efficiency (the ratio of the LIB volume to the overall LIB pack volume) and the energy density of the LIB packs.

On the basis of this LIB pack design principle, we developed a quasi-solid-state electrolyte (QSE) comprised of a solvate ionic liquid and silica particles. In our preceding study, an equimolar mixture of solvate ionic liquid and silica particles was used for the negative electrode active material, which was mixed with styrene-butadiene rubber (SBR) and carboxymethyl cellulose (CMC, 2200, Dicel FineChem, Ltd.) in water. The weight was inhibited. A 10-kWh battery pack built with the developed 100 Wh-class laminated LIB allowed driving a compact EV.

Apart from the enhancing LIB pack energy density with a highly safe QSE, there is also a possibility of improving the battery design to increase the energy density of the LIB itself. In this communication, we propose a bezel-less LIB that eliminates the extra space needed for casing, as shown in Figs. 1(a) and (b). The idea of the bezel-less structure comes from the smartphone displays with extremely narrow casing, which gives users an impression that the product does not have the bezel part. By switching from the conventional aluminum laminate to the bezel-less structure as shown in Figs. 1(c) and (d), we reduced the LIB cell volume by 16% (see the detail of the calculation in Supporting Information). By combining a Ni-rich high capacity positive electrode, a graphite negative electrode and the QSE, the proposed structure has successfully demonstrated high energy density of 600 Wh L⁻¹ (defined by the LIB volume) and high capacity of 132 Wh. High safety of the bezel-less LIB was demonstrated by overcharge and collapse tests, which are parts of the International Electrotechnical Commission (IEC) standards, IEC62660-3.

2. Experimental

Commercially available LiNi0.8Co0.1Mn0.1O2 (GELON LIB GROUP, denoted by NCM hereafter) with a 20-µm grain diameter was used for the active material of the positive electrode. A positive electrode slurry was prepared by mixing NCM with acetylene black as a conductive additive (HS-100, Denka Company Ltd.), polyvinylidene difluoride (PVdF, KF polymer #9708, Kureha Corp.) as a binder, and N-methylpyrrolidone (NMP) at the desired ratio. The weight ratio of NCM, conductive additive, and a binder was 96:2:2. The resultant slurry was painted on both surfaces of a 15-µm thick aluminum foil current collector, dried, pressed, and then used for the composite positive electrode. The weight of NCM, conductive additive and a binder was 96:2:2. The resultant slurry was painted on both surfaces of a 15-µm thick aluminum foil current collector, dried, pressed, and then used for the composite positive electrode. The weight of the bi-side positive electrode layers was 53 mg cm⁻², and the thickness and the density were 88 µm and 3.0 g cm⁻³, respectively. Natural graphite was used for the negative electrode active material, which was mixed with styrene-butadiene rubber (SBR) and carboxymethyl cellulose (CMC, 2200, Dicel FineChem, Ltd.) in water. The weight
ratio of graphite, SBR and CMC was 98:1:1. The resultant slurry was coated on both surfaces of an 8-µm thick copper current collector, dried, pressed, and then used for the negative electrode. The weight of the bi-side negative electrode coating was 28 mg cm$^{-2}$, and the thickness and the density were 88 µm and 1.6 g cm$^{-3}$, respectively.

To reduce the thickness of the QSE and the electrolyte resistance of the assembled cells, the QSE slurries comprised of silica, poly(vinylidene fluoride hexafluoropropylene) (PVdF-HFP), KF polymer #8500, Kureha Corp.) as a binder and NMP were coated on the positive electrode with a 10-µm thickness. Cross-sectional image was observed for the bi-layer QSE and positive electrode. The cross-section was prepared by using an ion-milling equipment (Hitachi High-Technologies Corp.) with an acceleration voltage of 6 kV for 2 h. Then, field-emission scanning electron microscopy (FE-SEM, S-4300, Hitachi, Ltd.) observation was carried out for the processed cross-section with an acceleration voltage of 5 kV. Owing to fracture of cut surfaces and bending of the electrodes with the QSE layers during stacking at edges, short circuits sometimes occurred. Hence, to avoid the occurrence of the short circuits, a 16-µm thick three layer separator, comprised of polyethylene, polypropylene and polyethylene, (UPZ520, Ube Maxell Co., Ltd.) was placed between the QSE layers of the positive and the negative electrodes, as in a structure schematically illustrated in Fig. 1.

The bi-layer positive electrode with the QSE and the negative electrode were cut into 146 mm × 76 mm and 150 mm × 80 mm, respectively. Then, 38 negative electrodes and 37 positive electrodes were stacked together with the folded tabs and placed in a joint bag, which was prepared using an aluminum laminate in a manner described in Fig. 1(e) with a battery storage space of 160 mm, as schematically illustrated in Fig. 1(a). Note that our bezel-less LIB has a height of 16.5 mm, which is beyond the limit of the press drawing of the conventional aluminum laminates. This is one of the energy density enhancement factors of our bezel-less LIB.

A solvate ionic liquid, an equimolar complex of tetraethylene glycol dimethyl ether (G4, ≥99%, Merck KGaA) and lithium bis(trifluoromethanesulfonyl)amide (LiTFSA, 99.5%, Kishida Chemical Co., Ltd., Li(G4)TFSA, was used for the liquid phase of the QSE). Then, to enhance lithium-ion transport during battery operation while maintaining the ignition point of the mixed liquid, polypropylene carbonate (PC, 99.5%, Kishida Chemical Co., Ltd.) was mixed together. On the basis of systematic investigations of viscosity, ionic conductivity, self-diffusion coefficient and battery performance as a function of PC concentration, we determined the optimal mixing ratio of Li(G4)TFSA-44.5 wt%PC (with the ignition point of 415 K, as opposed to 293–303 K of the conventional organic liquid electrolyte). For establishing a stable solid electrolyte interphase (SEI), 3 wt% of vinylene carbonate (VC, 99.5%, Kishida Chemical Co., Ltd.) was introduced as a primary additive. Then, for inhibition of corrosion of the aluminum positive electrode current collector by the use of LiTFSA, 2.5 wt% of tetrabutyl ammonium hexafluorophosphate (TBAPF$_6$, >98%, Tokyo Chemical Industry Co., Ltd.) was also introduced as a secondary additive. The details of the liquid composition optimization are described in Ref. 6. The resultant mixed liquid was injected into a bezel-less LIB, evacuated, sealed and left for 16 hours for sufficient impregnation. The battery assembly was carried out in a dry room, where a dew point was kept at −30°C.

To inhibit gas evolution by a formation of stable solid-electrolyte interphase (SEI) at the graphite surface and to enhance columbic efficiency at the initial cycle, the charge was started at 0.005 C for 10 hr. Subsequently, the charge rate (c-rate) was increased to 0.05 C (1 C is equal to 36 A, corresponding to 160 mAh g$^{-1}$ where the current was normalized by the weight of the positive electrode active material) until the voltage reached 4.2 V. The voltage was kept at 4.2 V until the current decreased to 0.005 C. Discharge was carried out at 0.05 C, 0.1 C, 0.2 C, and 0.5 C until the voltage decreased to 2.7 V. The volumetric power density was evaluated by the direct-current resistance (DCR) technique. The state-of-charge (SOC) of
the bezel-less LIB was kept at 50%, and then the LIB was discharged at 0.1 C, 0.2 C, and 0.5 C for 10 seconds. The volumetric power density was estimated from the relationship between the voltage drop and current density (or c-rate).

The cycle life of the bezel-less LIB was also evaluated. The discharge capacity at 0.05 C was recorded with an interval of 20 cycles at 0.2 C. The retention ratio of the discharge capacity was defined as a ratio of the discharge capacity to the initial discharge capacity at 0.05 C.

In this study, the overcharge test and the collapse test were carried out to evaluate the safety of the bezel-less LIB. For the overcharge test, the bezel-less LIB was fully charged at 4.2 V and then, the cell was further charged at 1 C until voltage exceeded 5 V (120% of the SOC) or the SOC exceeded 130% (both are safety criteria determined by IEC62660-3). A thermocouple was placed on the LIB, and then the LIB temperature was monitored during the overcharge test. The fully charged bezel-less LIB was used for the collapse test. The cell was fixed on a holder, and then the LIB was collapsed by a semi-circular ball with a rate of 0.1 mm sec⁻¹. A thermocouple was attached onto the LIB surface, and the LIB temperature was recorded during the collapse test.

3. Results & Discussion

In our LIB configuration, the thickness of the QSE layer is one of the key factors determining the safety of LIB. We reported that a 20-µm thick QSE layers on both the positive and negative electrodes (40 µm in total) can inhibit short circuits via a metallic nail, thus minimizing the generation of the joule heat. In addition, the high ignition point of the liquid phase (415 K) contributes to inhibition of firing triggered by the joule heat generation. The QSE layer should be thin as it does not contribute to the energy storage and is one of the factors of the internal resistance increase. When the QSE layer is too thin, a short circuit can easily happen, compromising the safety of the battery. Considering the above, we decided to form a 10-µm thick QSE layer. By optimizing the QSE and slurry compositions and coating conditions, we succeeded in placing a uniform 10-µm thick QSE layer only on the positive electrode layer, as shown in the cross-sectional SEM image in Fig. 2. The volume ratio of SiO₂, P(VdF-HFP) and void was 45.2:6.8:48, estimated from the QSE slurry composition and the size of the coated QSE layer.

The bezel-less LIB assembled in this study demonstrated repeated operation with high capacity and high coulombic efficiency, as can be seen in the charge-discharge profiles in Fig. 3(a). The bezel-less LIB exhibited an initial charge capacity of 43.7 Ah and a discharge capacity of 36.6 Ah (with coulombic efficiency of 83.7%). At the average voltage of 3.6 V during the discharge, our bezel-less LIB has the energy of 132 Wh. From the bezel-less LIB energy and geometry, a volumetric energy density was calculated to be 600 Wh L⁻¹ (256 Wh kg⁻¹). The bezel-less LIB design can eliminate extra volumes for sealing and accumulation of current collectors, achieving 16% of volume reduction, compared to the conventional aluminum laminates, as schematically shown in Figs. 1(c) and (d) and Fig. S1. This surely contributes to the enhanced energy density. Another advantage of the bezel-less LIB concept is that it does not have thickness restrictions. This is different from a conventional press drawing that have a limitation on thickness to avoid a fracture of aluminum. The spaces for laminates of individual cells in the LIB pack can be reduced, further enhancing energy density of the LIB pack for the large-scale applications such as EVs and stationary uses which require a number of LIBs and laminates.

The bezel-less LIB retained high discharge capacity of 27.0 Ah at 0.5 C, as the discharge profiles shown in Fig. 3(b). As seen in the figure, the discharge capacity monotonically decreased as the c-rate increased. The bezel-less LIB employed a 10-µm QSE, the resistance from the ion transport in thick electrode layers (both positive and negative electrode layers are 88-µm thick) can increase the internal resistance. Hence, further exploration of the liquid phase used for the QSE is required for enhancing the high rate capability in future studies. From the DCR measurement, the output power was quantified. The linear relationship between the voltage drop and current density was confirmed by the DCR measurement, and then the power density was estimated to be 1160 W L⁻¹ at the SOC of 50%. As shown in Fig. 3(c), the 41st discharge capacity retention ratio was as high as 90%, and an expansion of the bezel-less LIB was not obvious after the cycle test.

The safety of the bezel-less LIB was investigated by the overcharge and collapse tests. The bezel-less LIB passed the overcharge test, i.e. inhibition of thermal runaway at voltage of 120% (5.0 V in our LIB) or the SOC of 130%. Figure 4 shows...
As demonstrated in this communication, the novel bezel-less concept enhances the energy density by minimizing the volume of the bezel part of the LIBs. High safety of the bezel-less LIB was successfully demonstrated via the overcharge and collapse test. Development of the QSE sheet that guarantees the insulation at the edge of electrode layers and allows for the LIB assembly without separator would be fruitful for further enhancing an energy density. A combined use of the QSE and the bezel-less concept for the LIB assembly can contribute to enhancing the energy density of the LIB packs used for large-scale applications, including EVs and stationary uses.

4. Conclusions

In this study, we developed a high capacity, high energy density and high power bezel-less LIB with the QSE. The bezel-less LIB demonstrated a capacity of 132 Wh, an energy density of 600 Wh L\(^{-1}\) (256 Wh kg\(^{-1}\)), and an output power of 1160 W L\(^{-1}\). The bezel-less LIB enabled stable and repeated operation. The 41st discharge capacity retention ratio recorded 90%. In addition, the bezel-less LIB passed the overcharge and collapse tests, based on the IEC62660-3 standard. These results suggest that a combined use of the bezel-less concept and the QSE may enhance the energy density of the LIB and the LIB pack for large-scale applications such as EVs and stationary uses.

Supporting Information

The Supporting Information is available on the website at DOI: https://doi.org/10.5796/electrochemistry.19-00076.

Acknowledgment

Part of this study was supported by Innovative Science and Technology Initiative for Security, Acquisition, Technology & Logistics Agency (ATLA), Japan.

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