

Review

# Recent Advances in Poly(vinylidene fluoride) and Its Copolymers for Lithium-Ion Battery Separators

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**Abstract:** The separator membrane is an essential component of lithium-ion batteries, separating the anode and cathode, and controlling the number and mobility of the lithium ions. Among the polymer matrices most commonly investigated for battery separators are poly(vinylidene fluoride) (PVDF) and its copolymers poly(vinylidene fluoride-co-trifluoroethylene) (PVDF-TrFE), poly(vinylidene fluoride-co-hexafluoropropylene) (PVDF-HFP), and poly(vinylidene fluoride-co-chlorotrifluoroethylene) (PVDF-CTFE), due to their excellent properties such as high polarity and the possibility of controlling the porosity of the materials through binary and ternary polymer/solvent systems, among others. This review presents the recent advances on battery separators based on PVDF and its copolymers for lithium-ion batteries. It is divided into the following sections: single polymer and co-polymers, surface modification, composites, and polymer blends. Further, a critical comparison between those membranes and other separator membranes is presented, as well as the future trends on this area.

**Keywords:** PVDF; copolymers; battery separator; lithium-ion batteries

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## 1. Introduction

In the field of mobile applications, the efficient storage of energy is one of the most critical issues, since there is a fundamental need to maximize the amount of energy stored. This issue can be accomplished by increasing the gravimetric and volumetric energy density of the batteries [1].

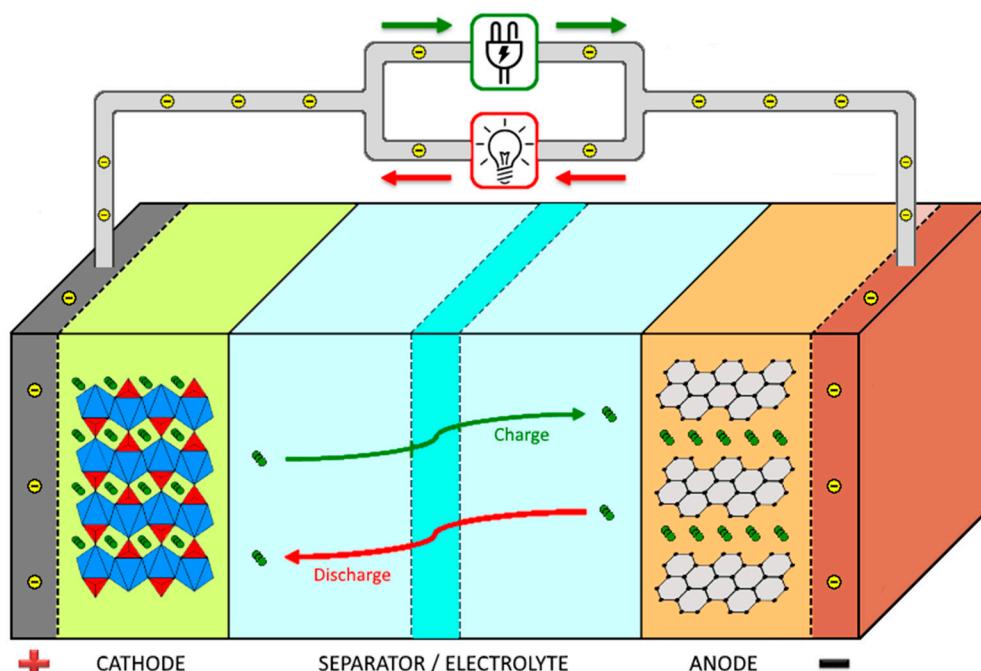
The electrochemical lithium ion battery is used to provide power to a large variety of mobile appliances, such as smartphones, tablets, and laptops, as well as an increasing number of sensors and actuators, which will have a fundamental role in the shaping of the Internet of Things and Industry 4.0 concepts, the main trend for current technological evolution [2]. Lithium ion batteries can also power electric and hybrid vehicles, and take part in the management of renewable energy production, being essential in a more sustainable energy paradigm. As some renewable resources, such as solar and wind, are intermittent over time, storing energy for their use during periods of lack of resources is a critical issue for lithium ion batteries [3,4].

Lithium ion batteries are very suitable for the aforementioned applications due to their advantages with respect to other battery types, as they are lighter, cheaper, have a higher energy density ( $250 \text{ Wh}\cdot\text{kg}^{-1}$ ,  $650 \text{ Wh}\cdot\text{L}^{-1}$ ), lower charge lost, no memory effect, a prolonged service-life, and a higher number of charge/discharge cycles [5].

Furthermore, the global market of lithium ion batteries is currently growing, and it is expected that in 2022, the market value will reach \$46.21 billion, with an annual growth rate of 10.8% [6].

The first commercial lithium ion battery, which was by Sony, entered the market in 1991, with the fundamental contribution of John Goodenough in the development of  $\text{LiCoO}_2$  as the active material for the cathode [7].

The main components of a battery are the anode, the cathode, and the separator, which are represented in Figure 1, together with the working principle of a lithium ion battery.

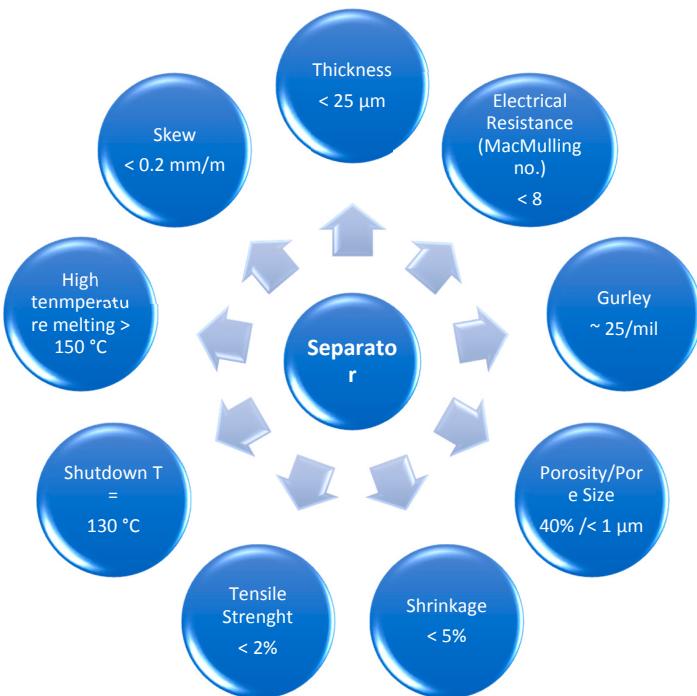


**Figure 1.** Schematic representation of a lithium ion battery and its working operation.

During the discharge process of the battery, the cathode acts as an oxidizing element, receiving electrons from the external electric circuit and being reduced. The anode is the reducing element, releasing electrons to the external electrical circuit, being oxidized during the electrochemical reaction [8].

## 2. Battery Separator: Function, Characteristics, and Types

Separators play a key role in the operation of electrochemical devices. The main purpose of the separator membranes is to separate the cathode from the anode, avoiding the occurrence of short circuits, and controlling the mobility of lithium ions between electrodes. The performance of a separator in a lithium ion battery is determined by some requirements such as porosity, chemical and thermal stability, electrical insulator, wettability, dimensional stability, and resistance to degradation by chemical reagents and electrolytes (Figure 2) [9]. Figure 2 shows the ideal values for the main requirements of a separator membrane.



**Figure 2.** Ideal values for the main requirements of a separator membrane.

There are different types of separators, but the most widely used consist of a polymer matrix embedded by the electrolyte solution, i.e., a liquid electrolyte where salts are dissolved in solvents, water, or organic molecules. The main types of separators are shown in Table 1 [10].

**Table 1.** Types and characteristics of different separators adapted from [10].

Separator	Characteristics	Typical Materials
Microporous	Operates at low temperatures (<100 °C); pore size = 50–100 Å	Nonwoven fibers (cotton, nylon, polyester, glass), polymers (PP, PE, PVC, PTFE), rubber, asbestos, wood
Nonwoven	Resistance to degradation by electrolytes; thickness > 25 µm; pore size = 1–100 µm	Polyolefins (PE, PP, PA, PTFE; PVDF; PVC)
Ion exchange membrane	High chemical resistance; impervious to electrolytes; pore size < 20 Å	PE, PP, Teflon-based films
Supported liquid membrane	Solid matrix with a liquid phase; insolubility in electrolyte; high chemical stability	PP, PSU, PTFE, CA
Polymer electrolyte	Simultaneously separator and electrolyte; high chemical and mechanical integrity	Polyethers, PEO, PPO, lithium salts
Solid ion conductor	simultaneously separator and electrolyte	-

The most commonly used materials as matrix for lithium ion battery separators are polymers, or polymer composites. Some of the most commonly used polymers are poly(propylene) (PP), poly(ethylene) (PE), poly(vinylidene fluoride) (PVDF) and its copolymers, poly(ethylene oxide) (PEO), and poly(acrylonitrile) (PAN) [11]. Some separators are developed by blending two different polymers to improve the characteristics of the membrane. In some cases, nanoparticles are added to the matrix as fillers to increase its mechanical stability or ionic conductivity. In composites separators, the most widely used fillers are oxide ceramics ( $ZrO_2$  [12,13],  $Al_2O_3$  [14,15],  $SiO_2$  [16,17]), carbonaceous fillers (graphene [18], carbon black [19], carbon nanofiber [20]), and ionic liquids [21], among others.

The solvents must possess some requirements to ensure proper battery operation. The properties of a good solvent are high dielectric constant, low viscosity, high chemical stability, and in liquid form over a wide temperature range. For this application, solvents of ethylene carbonate (EC), propylene carbonate (PC), dimethyl carbonate (DMC), diethyl carbonate (DEC), and ethyl methyl carbonate (EMC) are the most commonly used [11].

### 3. Poly(vinylidene fluoride) and Its Copolymers

Considering the different polymer matrices used for battery separators, PVDF and its copolymers (poly(vinylidene fluoride-co-trifluoroethylene), poly(vinylidene fluoride-co-trifluoroethylene) (PVDF-TrFE), poly(vinylidene fluoride-co-hexafluoropropylene), poly(vinylidene fluoride-co-hexafluoropropylene) (PVDF-HFP), and poly(vinylidene fluoride-cochlorotrifluoroethylene) (PVDF-CTFE)) show exceptional properties and characteristics for the development of battery separators, highlighting high polarity, excellent thermal and mechanical properties, wettability by organic solvents, being chemically inert and stable in the cathodic environment, and possessing tailorabile porosity through binary and ternary solvent/non-solvent systems [22,23]. The main properties of these polymers are presented in Table 2 [11].

PVDF and its copolymers are partially fluorinated semi-crystalline polymers where the amorphous phase is located between the crystalline lamellae arranged in spherulites. It can crystallize in different crystalline phase, depending on the temperature and processing conditions [24,25]. In relation to the crystalline phases of PVDF and its copolymers, the most important phases are the  $\beta$ -phase, since it presents ferroelectric, piezoelectric, and pyroelectric properties, and the  $\alpha$ -phase, which is the most stable thermodynamically, when material is obtained directly from the melt [24]. As illustrated in Table 2, PVDF and its polymers are characterized by excellent mechanical properties, good thermal stability up to 100 °C, and a high dielectric constant, which is essential for assisting the ionization of lithium salts.

**Table 2.** Main properties of PVDF and its copolymers [26–28].

Polymer	Melting Temp./°C	Degree of Crystallinity/%	Young Modulus/MPa	Dielectric Constant
PVDF	~170	40–60	1500–3000	6–12
PVDF-TrFE	~120	20–30	1600–2200	18
PVDF-HFP	130–140	15–35	500–1000	11
PVDF-CTFE	~165	15–25	155–200	13

PVDF copolymers have drawn increasing attention for battery separators, as the addition of other monomers to the VDF blocks increases the fluorine content and decreases the degree of crystallinity (Table 2), which is particularly relevant once the uptake of the electrode solution occurs in the amorphous region through a swelling process for accommodating the electrolyte and, as a result, increases the ionic conductivity [29]. The recent literature on PVDF and its battery separator copolymers is structured into four sections dedicated to single polymers, surface modification, composites, and polymer blends, respectively.

The main achievement for PVDF and co-polymers as battery separators was thoroughly reviewed in [11]. Since then, important contributions have been achieved, which are the subject of the present review.

#### 3.1. Single Polymer and Co-Polymers

As already mentioned, one of the main characteristics of PVDF and its co-polymers is their high dielectric permittivity, providing a large affinity with polar electrolytes when compared to other polymers [11]. The main characteristics of the developed PVDF and the copolymer membranes are shown in Table 3.

**Table 3.** Separator membranes based on PVDF and co-polymers, indicating also the main properties, and the main goal/achievement of the investigation.

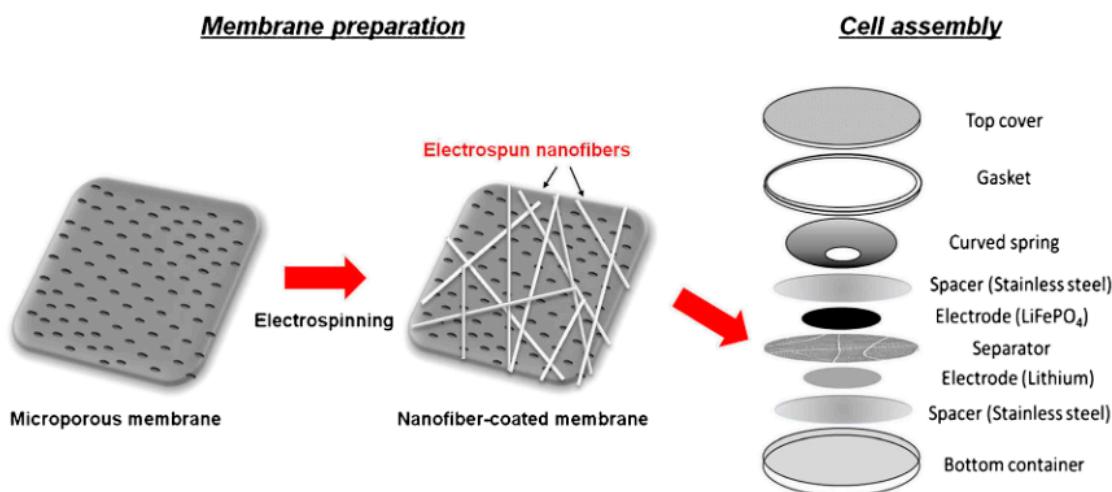
Materials	Electrolyte Solution	Porosity and Uptake (%)	Conductivity ( $\text{S}\cdot\text{cm}^{-1}$ ) and Capacity ( $\text{mAh}\cdot\text{g}^{-1}$ )	Main Goal/Achievement	Reference
PVDF	1 M $(\text{C}_2\text{H}_5)_3\text{CH}_3\text{NBF}_4 + \text{AN}$	-/-	-/-	Study of multistep electrospinning technique on the fabrication of PVDF composite membranes; High specific power.	[30]
PVDF	1 M $\text{LiPF}_6$ in EC:DEC (1:1, <i>w/w</i> )	-/816	$6.83 \times 10^{-4}/101.1$ (0.5C)	Performance comparison with a PVDF-PDA separator; Enhanced cycling performance.	[31]
PVDF	1 M $\text{LiPF}_6$ in EC:DEC (1:1, <i>v/v</i> )	7/-	-/-	Analysis of the migration mechanism of cation and anions through the separator; The separator allows the control of structural stability and ion mobility.	[32]
PVDF	1 M $\text{LiPF}_6$ in EC/DMC/EMCC (1:1:1, <i>w/w/w</i> )	-/-	-/95 (0.2C)	Production of a PVDF membrane; Good capacity retention.	[33]
PVDF	1 M $\text{TEABF}_4$ in AN/PC and 1 M $\text{LiPF}_6$ in EC/DEC	80/-	$1.8 \times 10^{-2}$ (25 °C)/-	Manufacturing of a PVDF separator; Favorable mechanical properties.	[34]
PVDF	1 M $\text{LiBF}_4$ in EC/DMC (50:50 wt. %)	-/-	$4.17 \times 10^{-3}$ (20 °C)/-	Comparison of PVDF membrane performance with Nafigate separators.	[35]
PVDF	1 M $\text{LiPF}_6$ in EC/DMC/DEC (1:1:1)	78.9/427	$1.72 \times 10^{-3}/164.3$ (C/5)	Synthesis of dual asymmetric structure separators; Improved electrolyte uptake and ionic conductivity.	[36]
PVDF	1 M $\text{LiPF}_6$ in EC/DMC/DEC (1:1:1)	-/-	-/447.36 (0.3C)	Production of a solid state SCPC with a PVDF separator; High storage capacity and stability.	[37]
PVDF	-	-/-	-/-	Assembly of a PVDF separator for air-cathode as application in microbial fuel cells; Improved electricity generation.	[38]
PVDF	PVA/ $\text{H}_2\text{SO}_4$	-/-	-/-	Production of a PVDF separator for piezoelectric supercapacitors; High mechanical strength and elevated capacitance.	[39]
PVDF	1 M $\text{NaClO}_4$ in EC/DEC (1:1)	81/34	$7.38 \times 10^{-4}$ (29 °C)/153	Production of an electroactive electrospun PVDF separator for sodium ion batteries.	[40]

Table 3. Cont.

Materials	Electrolyte Solution	Porosity and Uptake (%)	Conductivity ( $\text{S}\cdot\text{cm}^{-1}$ ) and Capacity ( $\text{mAh}\cdot\text{g}^{-1}$ )	Main Goal/Achievement	Reference
PVDF	1 M LiPF <sub>6</sub> in EC/DEC (1:1)	70/66	$1.5 \times 10^{-3}$ /102 (2C)		
PVDF-TrFE	1 M LiPF <sub>6</sub> in EC/DEC (1:1)	72/84	$1.1 \times 10^{-3}$ /118 (2C)	Study of the effect of different PVDF copolymers as lithium ion battery separators.	[41]
PVDF-HFP	1 M LiPF <sub>6</sub> in EC/DEC (1:1)	56/79	$1.3 \times 10^{-1}$ /107 (2C)	Demonstration of the relevance of $\beta$ -phase content.	
PVDF-CTFE	1 M LiPF <sub>6</sub> in EC/DEC (1:1)	59/80	$1.5 \times 10^{-3}$ /85 (2C)		
PVDF	[C <sub>2</sub> mim][NTf <sub>2</sub> ]	20/98	$2.3 \times 10^{-4}$ (25 °C)/74.6 (C/5)	Preparation of PVDF separators using a green solvent and ionic liquid as the electrolyte.	[9]
PVDF-HFP	LiTFSI	48/248	$5.2 \times 10^{-5}$ (20 °C)/-	Application of disiloxane-based electrolytes on PVDF-HFP for the production of gel electrolyte separators; Good thermal and mechanical stability.	[42]
PVDF-HFP	LiNfO/BMImNfO	-/-	$2.61 \times 10^{-2}$ /(100 °C) 138.1 (C/4)	Production of ionic liquid gel polymer electrolytes; High ionic conductivity.	[43]
PVDF-HFP	1 M LiPF <sub>6</sub> in EC/DMC (1:2)	70/247	$3.2 \times 10^{-3}$ (25 °C)/-	Evaluation of the performance of PVDF-HFP, as a single polymer membrane. Understanding of the method of avoiding the formation of beads in the nanofibers of PVDF-HFP; Good electrolyte uptake.	[44]
PVDF-HFP	1 M LiPF <sub>6</sub> in EC:DMC (1:1)	78/86.2	$1.03 \times 10^{-3}$ /145 (0.2C)	Development of a PVDF-HFP gel polymer electrolyte membrane with honeycomb type porous structure; Excellent electrochemical performance.	[45]
PVDF-HFP	1 M LiPF <sub>6</sub> in EC/DEC/EMC (1:1:1)	-/-	-/-	Production of separators with controlled pore structure; Improved rates and cycling performances.	[46]
PVDF-CTFE	1 M LiPF <sub>6</sub> in EC:DMC:EMC (1:1:1, v:v)	74/-	$7.51 \times 10^{-4}$ (25 °C)/147 (0.2C)	Preparation of a nanofiber-coated composite separator by electrospinning; High discharge capacity and good cycling stability.	[47]

Table 3 shows that the electrospinning technique is widely used to produce functional membranes. Thus, electrospun separators have been developed for PVDF-PDA [31], PVDF-HFP [44], and PVDF-CTFE [47].

For the PVDF-CTFE membrane, the cell assembly considered for the battery performance tests is represented in Figure 3.



**Figure 3.** Manufacturing of a testing cell based on PVDF-CTFE separators [47], with copyright permission from Springer Nature.

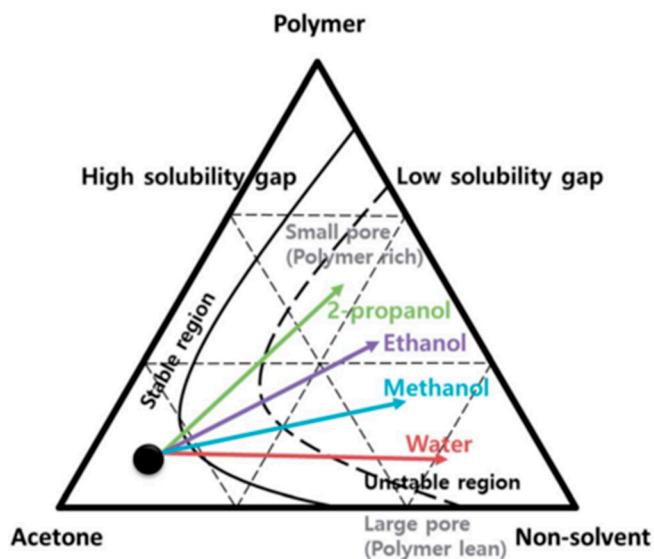
For PVDF-HFP electrospun membranes, it has been demonstrated that a single layer membrane shows good porosity and uptake value, but that the mechanical stability is negatively affected, with the viscosity of the solution playing an important role [44]. Also, a novel gel electrolyte was developed based on PVDF-HFP by the addition of disiloxane into the electrolyte solution [42], leading to a thermally stable separator that is not flammable, thus contributing to safer lithium ion batteries [45]. It this sense, ionic liquids have also been used in electrolyte solutions, improving both safety and the ionic conductivity of the membranes [43].

A multistep electrospinning technique for the production of PVDF membranes for electrical double-layer capacitors has been proposed, allowing for the manufacture of thinner and more densely packed separators [30].

Further, membranes have been developed based on PVDF for air-cathode in microbial fuel cells [38] and piezo-supercapacitors [39]. Dual asymmetric PVDF separators were produced by a thermally-induced phase separation method, in which the large and interconnected pores in the bulk structure ensures an improved electrolyte uptake and ionic conductivity, while the small pores in the surfaces prevent the loss of electrolyte and the growth of lithium dendrites. It is indicated that those separators ensure safer batteries with high discharge capacity and longer cycle life [36].

A further step towards the development of more environmentally friendly PVDF separator membranes was proposed by using DMPU as a solvent for PVDF, and IL [C<sub>2</sub>mim][NTf<sub>2</sub>] as an electrolyte. The use of the IL increased the ionic conductivity and discharge capacity of the membrane when compared with separators using conventional electrolytes [9].

Porous PVDF-HFP membranes were prepared with non-solvents using the phase inversion technique. When selecting different types of non-solvents such as water, methanol, ethanol, and propanol, and their contents in acetone, it was possible to control the size of the pores (Figure 4) [46].



**Figure 4.** Phase diagram of the ternary mixture—PVDF-HFP, acetone, and non-solvent—in order to control PVDF-HFP membrane morphology [46], with copyright permission from the Royal Society of Chemistry.

Finally, a correlation between the  $\beta$ -phase content of the separators, and the rate capability and cyclability of the batteries was demonstrated for different PVDF co-polymers, showing that the PVDF-TrFE membrane has the best battery performance for the highest  $\beta$ -phase content (100%) [41].

Thus, it is observed that for single (co)polymer membranes, the main focus is to tailor the morphology to obtain good uptake without mechanical deterioration, and to improve the interaction between the electrolyte solution and the separator membrane.

### 3.2. Surface Modification of the Separator Membranes

Typically, surface modification of the membranes is carried out to improve specific properties such as wettability, and thermal and mechanical stability. PVDF membranes have been prepared after different surface modifications, but also have been used to modify the properties of other polymer membranes, as presented in Table 4.

**Table 4.** Surface modifications on PVDF and co-polymers, indicating also the main properties, goal and achievement.

Materials	Electrolyte Solution	Porosity and Uptake (%)	Conductivity ( $\text{S}\cdot\text{cm}^{-1}$ ) and Capacity ( $\text{mAh}\cdot\text{g}^{-1}$ )	Main Goal/Achievement	Ref
PVDF (plasma-treated)	1 M LiPF <sub>6</sub> in EC/DMC (1:1)	-/1200	-/-	Study of the effect of plasma treatment in PVDF separators; Improved electrolyte uptake and mechanical properties.	[48]
PE/PVDF	1 M LiPF <sub>6</sub> in EC:EMC:DEC (1:1:1, <i>w:w:w</i> )	-/-	$0.89 \times 10^{-3}$ (25 °C)/-	Investigation into the pore formation process in a coating layer for separators; Enhanced ionic conductivity.	[49]
PE/PVDF	1.10 M LiPF <sub>6</sub> in EC/PC/EP (3:1:6, <i>v:v:v</i> )	-/-	-/1436 (0.2C)	Study of the electrochemical performance of PE/PVDF separators; Enhanced cycling performance.	[50]
PVDF/PP	1 M LiPF <sub>6</sub> in EC/DMC (1:1)	58/140	$5.9 \times 10^{-4}$ /145 (0.5C)	Coating of PVDF particles in the surface of a PP membrane; Increased electrolyte uptake.	[51]
PET/PVDF	1 M LiPF <sub>6</sub> in EC/DEC/DMC (1:1:1, <i>w/w/w</i> )	-/-	$8.36 \times 10^{-3}$ /-	Investigation of the performance of a hot-pressed PET/PVDF separator; Excellent mechanical behavior.	[52]
PVDF/HEC	1 M LiPF <sub>6</sub> in EC/DMC/EMC (1:1:1)	-/135.4	$8.8 \times 10^{-4}$ (25 °C)/140	Preparation of a PVDF/HEC/PVDF membrane with a sandwich structure; High electrolyte uptake and ionic conductivity.	[53]
PVDF/PMMA	1 M LiTFSI in DME/DOL (1:1)	-/294	$1.95 \times 10^{-3}$ (25 °C)/1711.8	Preparation of a sandwiched GPE based on PVDF and PMMA for lithium-sulfur batteries; High discharge capacity and cycle stability.	[54]
PDA/PVDF	1 M LiPF <sub>6</sub> in EC:DEC (1:1, wt:wt)	-/1160	$9.62 \times 10^{-4}$ /104.5 (0.5C)	Prove that the PDA coating can be promising for manufacturing electrospun nanofiber separators; Better cycling performance and elevated power capability.	[31]
PE/(PVDF/Al <sub>2</sub> O <sub>3</sub> )	1 M LiPF <sub>6</sub> in EC/DEC (1:1)	60.3/125, 314	$1.14\text{--}1.23 \times 10^{-3}$ /-	Development of a multilayer coating for separators; Improvement of thermal stability and electrolyte wetting.	[55]
PI/PVDF/PI	1 M LiPF <sub>6</sub> in EC/DEC/DMC (1:1:1)	83/476	$3.46 \times 10^{-3}$ /114.8 (0.5C)	Production of an electrospun sandwich-type separator; Superior porosity, electrolyte uptake, and ionic conductivity.	[56]
PVDF-HFP	1 M NaClO <sub>4</sub> in EC/PC (1:1)	-/-	$3.8 \times 10^{-3}$ /291.1 (0.2C)	Development of a PVDF-HFP-coated GF separator for sodium ion batteries; Good cycling performance.	[57]

**Table 4.** *Cont.*

Materials	Electrolyte Solution	Porosity and Uptake (%)	Conductivity ( $\text{S}\cdot\text{cm}^{-1}$ ) and Capacity ( $\text{mAh}\cdot\text{g}^{-1}$ )	Main Goal/Achievement	Ref
PVDF-HFP	1 M LiPF <sub>6</sub> in DMC/EMC/EC (1:1:1)	53.5/106.9	$8.34 \times 10^{-4}$ /131.33 (5C)	Study of the effect of the drying temperature on the performance of the separator.	[58]
PP/(PVDF-HFP/SiO <sub>2</sub> )	1 M LiPF <sub>6</sub> in DEC/EC (1:1, v/v)	-/-	$7.2 \times 10^{-4}$ /-	Analysis on the effect of a PVDF-HFP/SiO <sub>2</sub> coating layer for PP separators; Better electrolyte uptake and ionic conductivity.	[59]
PMMA/PVDF-HFP	1 M LiPF <sub>6</sub> in EC:DMC (1:1)	-/342	$1.31 \times 10^{-3}$ /143 (0.2C)	Investigation and analysis of a produced PMMA/PVDF-HFP electrolyte membrane; Exceptional thermal and electrochemical stability.	[60]
PVDF-HFP/PDA	LiPF <sub>6</sub> in EC/DEC/DMC (1:1:1)	72.8/254	$1.40 \times 10^{-3}$ (20 °C)/-	Production of a PVDF-HFP/PDA separator by a dip-coating method.	[12]
PVDF-HFP/PET	1 M LiClO <sub>4</sub> in DMSO	-/282	$6.39 \times 10^{-3}$ (25 °C)/158 (0.1C)	Combination of PVDF-HFP with a SiO <sub>2</sub> nanoparticle-modified PET matrix; Improved thermal stability, electrolyte uptake, and ionic conductivity.	[61]
PP/(AlO <sub>2</sub> /PVDF-HFP)	1 M LiPF <sub>6</sub> in EC/DEC (1:1, v/v)	-/-	$7.95 \times 10^{-4}$ /98.6 (0.2C)	Inspection of the performance of a separator for PP membrane coating; Improved thermal stability.	[62]
γ-Al <sub>2</sub> O <sub>3</sub> /PVDF-HFP/TTT	1 M LiClO <sub>4</sub> in EC/DEC (1:1)	-/157	$1.3 \times 10^{-3}$ /~100 (0.5C)	Dip coating of a PE separator with γ-Al <sub>2</sub> O <sub>3</sub> /PVDF-HFP/TTT; Increased electrolyte uptake and ionic conductivity.	[13]
PP/PE/PP/PVDF-co-CTFE	1 M LiPF <sub>6</sub> in EC/DMC/DEC (1:1:1, v:v:v)	-/-	-/-	Fabrication of PVDF-co-CTFE nanofiber coatings for improving the performance of polyolefin separators; High electrolyte uptake and good wettability.	[63]

The most commonly used surface modification is the use of PVDF and its copolymers for the coating of other polymers such as polyethylene porous separators. Thus, the coating of PE with a  $\text{Al}_2\text{O}_3$  ceramic layer and a PVDF electrospun nanofiber layer leads to enhanced electrolyte uptake, improved capacity discharge, and cycle life [55]. Similarly, a PDA coating on PVDF improves hydrophilicity, enhancing electrolyte uptake and ionic conductivity of the separator [31].

A typical surface modification technique, such as plasma treatment, allows significant improvement of the electrolyte uptake of PVDF electrospun membranes [48].

A hot-pressing technique was proposed to develop PET/PVDF separators, with improved mechanical behavior properties [52].

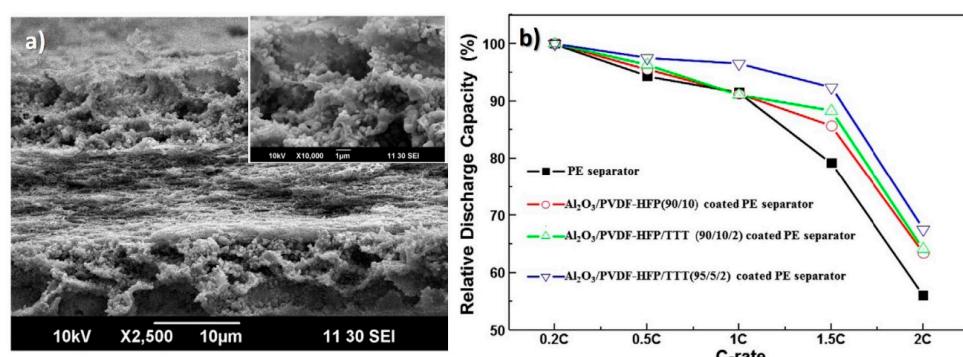
The preparation of a PVDF/PMMA/PVDF separator showed great potential for its use in lithium-sulfur batteries, showing high initial discharge capacity and cycle stability, also reducing cell polarization and suppressing the shuttle effect, which is described as the transport of soluble polysulfides between both electrodes and the associated charge [54].

A composite membrane with a PVDF/HEC/PVDF sandwich structure was developed, leading to higher electrolyte uptake, ionic conductivity, and cycling performance. It is also greener and safer because of the fire-retardant behavior of its components [58].

For PVDF-HFP membranes, several coatings have been applied, such as  $\text{ZrO}_2$  nanoparticles [64], PP polymer [59], PMMA polymer [60], PDA layer [12], and  $\text{SiO}_2$ -modified PET [61], leading mainly to improved electrolyte uptake.

Surface modifications are also achieved by modifying the drying temperature of PVDF-HFP/PET separators prepared by dip-coating, with a drying temperature of 80 °C improving cycle and rate performances with respect to batteries with a conventional PP separator [58].

The dip-coating of a PE separator with  $\gamma\text{-Al}_2\text{O}_3/\text{PVDF-HFP}/\text{TTT}$ , proved to increase electrolyte uptake and ionic conductivity when compared with conventional membranes, as shown in Figure 5 where its microstructure and cycling performance are presented. The discharge performance was also enhanced as well as the thermal resistance [13].



**Figure 5.** (a) Cross-section scanning electron microscopy (SEM) images of the  $\gamma\text{-Al}_2\text{O}_3/\text{PVDF-HFP}/\text{TTT}(95/5/2)$ -coated PE separator and (b) relative discharge capacities as a function of the C-rate [13], with copyright permission from Elsevier.

Basically, surface modifications are essential for improve the electrolyte wettability of the separators, and are realized in several polymer membranes of single and multiple layers with many polymers (PP, PET, PMMA, etc.) and filler nanoparticles.

### 3.3. Composite Membranes

Polymer composites are used to improve battery performance by incorporating suitable fillers, such as oxides ceramic, zeolites, and carbon nanotubes, among others, with the objective of increasing ionic conductivity, mechanical strength, and thermal stability. The main properties of composite separator membranes based on PVDF and its copolymers are presented in Table 5.

**Table 5.** Polymer composites based on PVDF and co-polymers with main properties, goal, and achievement.

Materials	Fillers	Electrolyte Solution	Porosity and Uptake (%)	Conductivity ( $S \cdot cm^{-1}$ ) and Capacity ( $mAh \cdot g^{-1}$ )	Main Goal/Achievement	Ref.
PVDF	Al <sub>2</sub> O <sub>3</sub>	1 M LiPF <sub>6</sub> in EC/DEC/DMC (1:1:1)	55.8/153.5	$2.23 \times 10^{-3}$ (25 °C)/114.2	Production of a composite PVDF/Al <sub>2</sub> O <sub>3</sub> ; High thermal stability and ionic conductivity, low discharge capacity decay.	[15]
PVDF	Al <sub>2</sub> O <sub>3</sub>	EC/DMC (1:1)	-/230	$1.24 \times 10^{-3}$ /151.97 (C)	Core-shell composite nonwoven separator of PVDF-HFP@Al <sub>2</sub> O <sub>3</sub> ; high heat resistance up to 200 °C without any shrinkage,	[65]
PVDF	Al <sub>2</sub> O <sub>3</sub>	1 M LiPF <sub>6</sub> in EC/DEC (1:1, v:v)	67/230	$1.49 \times 10^{-3}$ /146.3 (0.2C)	Separator-cathode assembly with PVDF/Al <sub>2</sub> O <sub>3</sub> ; Good electrochemical performance.	[66]
PVDF	AlO(OH) nanoparticles	1 M LiPF <sub>6</sub> in EC/DEC (3:7)	-/65	-/-	Ceramic separator based on boehmite nanoparticles; Improved safety and wettability.	[67]
PVDF	BC	1 M LiTFSI in EC/DEC (1:1)	-/-	$4.2 \times 10^{-3}$ (30 °C)/-	Preparation of GPEs based on cross-linkers; High ionic conductivity and thermal stability.	[68]
PVDF	Carbon	1 M LiTFSI and 0.1 M LiNO <sub>3</sub> in DOL/DME (1:1)	-/-	-/827 (0.5C)	PVDF-C separator by phase inversion technique; Superior rate performance and stability.	[69]
PVDF	CNF	1 M LiTFSI in DOL/DME (1:1)	-/119	-/1739.2 (C)	Production of CNF/PVDF separators for Li-S batteries Great battery discharge capacity and cycling stability.	[20]
PVDF	Cellulose acetate/Al(OH) <sub>3</sub>	1 M LiPF <sub>6</sub> in EC/DMC/EMC (1:1:1)	68.6/403.9	$2.85 \times 10^{-3}$ /151.97 (C)	Environmental friendly materials in a separator; High electrolyte uptake, ionic conductivity and cycling performance.	[70]
PVDF	DNA-CTMA	LiAsF <sub>6</sub> in EC/EMC/DMC	-/-	-/-	PVDF/DNA-CTMA membrane as a solid polymer/gel electrolyte separator; Improved thermal and mechanical properties.	[71]
PVDF	LiPVAOB	1 M LiPF <sub>6</sub> in EC/DMC/EMCC (1:1:1, w:w:w)	-/88.5	$2.6 \times 10^{-4}$ /120 (0.2C)	Composite gel polymer electrolyte PVDF/LiPVAOB membrane; Good ionic conductivity.	[33]
PVDF	Nanoclays/PVP	1 M LiPF <sub>6</sub> in EC/DMC (1:1)	87.4/553.3	-/-	Study of the influence of solvents in the separator High porosity and uptake.	[72]

Table 5. Cont.

Materials	Fillers	Electrolyte Solution	Porosity and Uptake (%)	Conductivity ( $S \cdot cm^{-1}$ ) and Capacity ( $mAh \cdot g^{-1}$ )	Main Goal/Achievement	Ref.
PVDF	NCC	1 M LiPF <sub>6</sub> in EC/DMC (1:1)	-/-	$3.73 \times 10^{-3}$ (25 °C)/-	Preparation of NCC-PVDF separators by phase inversion; Improved wettability and mechanical properties.	[73]
PVDF	MA groups	1 M LiPF <sub>6</sub> in EC/DMC/EMC (1:1:1)	67.4/-	$1.48 \times 10^{-3}$ /136 (0.2C)	Study of the addition of MA groups to the PVDF structure; High ionic conductivity.	[74]
PVDF	MMT	1 M LiPF <sub>6</sub> in EC/EMC/DEC (1:1:1)	84.08/333	$4.20 \times 10^{-3}$ (25 °C)/144	Effect of different contents of MMT filler in PVDF separators; High ionic conductivity and porosity.	[75]
PVDF	MOF-808	-	-/-	$1.56 \times 10^{-4}$ (65 °C)/-	Production of a MOF/polymer membrane; Good mechanical properties and durability.	[76]
PVDF	Octaphenyl-POSS	1 M LiPF <sub>6</sub> in EC/DMC/EMC (1:1:1)	66.1/912	$4.2 \times 10^{-3}$ /145.8 (0.5C)	Electrospun membrane with octaphenyl-POSS particles; Increased uptake and porosity, high ionic conductivity.	[77]
PVDF	Polyether (PEGDA+PEGMEA)	1 M LiPF <sub>6</sub> in EC/DMC/EMC (1:1:1)	-/230	$\sim 1.4 \times 10^{-3}$ (25 °C)/93 (0.5C)	Preparation of GPEs with PVDF and polyethers.	[78]
PVDF	PMIA	1 M LiPF <sub>6</sub> in EC/DMC/EMC (1:1:1, <i>w:w:w</i> )	-/-	$8.1 \times 10^{-4}$ /135.29 (0.2C)	Composite sandwich type separator, by electrospinning; High capacity retention and good rate performance.	[79]
PVDF	P-PAEK	1 M LiPF <sub>6</sub> in EC/DMC (1:1)	71.7/123.7	/141.6 (C/2)	Development of a P-PAEK/PVDF separator High wettability and electrolyte uptake.	[80]
PVDF	PFSA	1 M LiPF <sub>6</sub> in EC/DMC/EMC (1:1:1)	-/-	$1.53 \times 10^{-3}$ /137.9 (C)	PVDF/PFSA blend membrane; High stability and discharge capacity.	[81]
PVDF	rGO	1 M LiTFSI + 0.1 M LiNO <sub>3</sub> in DME/DOL (1:1)	71/380	/646	Double-layer PVDF/rGO membrane by electrospinning; High safety and cycling stability.	[82]
PVDF	SiO <sub>2</sub>	1 M LiPF <sub>6</sub> in EC/DMC/EMC (1:1:1)	54.1/279.5	-/175.7	Synthesis of a composite separator with SiO <sub>2</sub> ; High wettability, uptake and thermal/mechanical stability.	[17]
PVDF	SiO <sub>2</sub>	1 M LiPF <sub>6</sub> in EC/EMC (1:1 in volume)	70/370	$2.6 \times 10^{-3}$ /132 (C)	Addition of SiO <sub>2</sub> nanoparticles on PVDF membranes; Improvement of wettability and ionic conductivity.	[83]
PVDF	SiO <sub>2</sub>	1 M LiPF <sub>6</sub> in EC/DEC (1:1, <i>v:v</i> )	85/646	$7.47 \times 10^{-3}$ /159 (0.2C)	Electrospun PVDF/SiO <sub>2</sub> composite separator; Excellent thermal stability and high ionic conductivity.	[84]
PVDF	SnO <sub>2</sub>	1 M LiPF <sub>6</sub> in EC/DMC (1:1 <i>w:w</i> )	-/-	-/-	Use of SnO <sub>2</sub> nanoparticles in a PVDF electrospun separator; Good cycling performance.	[85]

Table 5. Cont.

Materials	Fillers	Electrolyte Solution	Porosity and Uptake (%)	Conductivity ( $S \cdot cm^{-1}$ ) and Capacity ( $mAh \cdot g^{-1}$ )	Main Goal/Achievement	Ref.
PVDF	ZnO	1 M LiPF <sub>6</sub> in EC/EMC (1:2)	-/-	-/-	Piezo-separator for integration on a self-charging power cell; Enhanced electrochemical performance.	[86]
PVDF	ZnO	1 M LiPF <sub>6</sub> in EC/DEC (1:1)	-/-	-/-	Piezo-separator for self-charging power cells; Stable and efficient performance.	[87]
PVDF	ZrO <sub>2</sub> /PEO	1 M LiTFSI in DOL/DME (1:1)	-/147.3	$3.2 \times 10^{-4}$ (25 °C)/1429 (0.2C)	GPE for lithium-sulfur batteries; High discharge capacity and rate performance.	[88]
PVDF-HFP	Al <sub>2</sub> O <sub>3</sub>	0.5 M NaTf/EMITf	-/-	$6.3-6.8 \times 10^{-3}$ (25 °C)/-	Introduction of Al <sub>2</sub> O <sub>3</sub> in a gel polymer electrolyte; Improved mechanical properties.	[14]
PVDF-HFP	Al <sub>2</sub> O <sub>3</sub>	1 M LiPF <sub>6</sub> in EC/DEC +2% VC	-/372	$1.3 \times 10^{-3}$ /155 (0.5C)	Colloidal Al <sub>2</sub> O <sub>3</sub> composite separator; enhancement of the mechanical strength of the PVDF-HFP separator.	[89]
PVDF-HFP	Al <sub>2</sub> O <sub>3</sub>	1 M LiPF <sub>6</sub> in EC/DMC/EMC ( <i>v:v:v</i> = 1:1:1)	-/420	$4.7 \times 10^{-4}$ /109 (4C)	Production of a low cost membrane, with a simple and easy scalable manufacturing process; High electrolyte uptake and good electrochemical stability and performance.	[90]
PVDF-HFP	Al(OH) <sub>3</sub>	1.15 M LiPF <sub>6</sub> in EC/EMC (3:7, <i>v:v</i> )	84/127	$10^{-3}$ /81 (C/2)	Upgrading the battery safety operation by the addition of metal hydroxides in composite separators; Suitable electrolyte uptake.	[91]
PVDF-HFP	Al <sub>2</sub> O <sub>3</sub> /CMC	1 M LiPF <sub>6</sub> in EC/DEC/PC/EMC (2:3:1:3)	42.7/-	$9.3 \times 10^{-4}$ (25 °C)/-	Composite separator with Al <sub>2</sub> O <sub>3</sub> /CMC; Safer and more stable separators.	[92]
PVDF-HFP	BN	1 M LiPF <sub>6</sub> in EC/DEC (1:1)	-/-	-/150 (0.2C)	3D separator; improved cycling stability with lower voltage polarization	[93]
PVDF-HFP	CA	1 M LiPF <sub>6</sub> in EC/DMC	85/310	$1.89 \times 10^{-3}$ /136 (8C)	Porous and honeycomb-structured membrane; higher lithium-ion transference number and improved rate performance	[94]
PVDF-HFP	Clay	1 M LiPF <sub>6</sub> in EC/DEC/EMC (1:1:1, <i>v:v:v</i> )	-/-	$1.49 \times 10^{-3}$ /-	New technique to incorporate clay sheets in a PVDF-HFP matrix, as separator; Thermal stability and higher ionic conductivity.	[95]
PVDF-HFP	EMImNFO-LiNFO	-	-/-	$3.92 \times 10^{-4}$ /(20 °C) 57 (C)	Introduction of anion-based IL and lithium salt in a GPE; High thermal stability, good electrochemical properties.	[96]
PVDF-HFP	GO	1 M LiPF <sub>6</sub> in EC/DEC/EMC (1:1:1)	-/71	$1.115 \times 10^{-3}$ (25 °C)/-	Addition of GO in separators to increase thermal properties; improved electrochemical and mechanical properties.	[97]

Table 5. Cont.

Materials	Fillers	Electrolyte Solution	Porosity and Uptake (%)	Conductivity ( $\text{S}\cdot\text{cm}^{-1}$ ) and Capacity ( $\text{mAh}\cdot\text{g}^{-1}$ )	Main Goal/Achievement	Ref.
PVDF-HFP	Graphene	1 M LiPF <sub>6</sub> in EC/DMC/EMC (1:1:1)	88/470	$3.61 \times 10^{-3}$ /149 (C)	PVDF-HFP/graphene GPE by NIPS; Increased porosity, uptake and ionic conductivity.	[18]
PVDF-HFP	HMSS	1 M LiPF <sub>6</sub> in EC/DEC (1:1)	~70/285	$2.57 \times 10^{-3}$ (25 °C)/-	Development of PVDF-HFP with HMSS separators; Increased wettability and porosity.	[98]
PVDF-HFP	Li <sub>1.3</sub> Al <sub>0.3</sub> Ti <sub>1.7</sub> (PO <sub>4</sub> ) <sub>3</sub>	1 M LiTFSI + 0.25 M LiNO <sub>3</sub> in DME/DOL (1:1)	34/143.9	$8.8 \times 10^{-4}$ (25 °C)/1614	Ceramic/polymer membrane for lithium-sulfur cells; High ionic conductivity and discharge capacity.	[99]
PVDF-HFP	LiTSFI/SN	-	-/-	$1.97 \times 10^{-3}$ (20 °C)/-	Production of supercapacitors with GO electrodes and GPE; High ionic conductivity.	[100]
PVDF-HFP	LLTO	1 M LiPF <sub>6</sub> in EC/DMC/EMC (1:1:1)	69.8/497	$13.897 \times 10^{-3}$ (25 °C)/155.56	Incorporation of LLTO in a PVDF-HFP separator; Improved ionic conductivity.	[101]
PVDF-HFP	PI	1 M LiPF <sub>6</sub> in EC/DMC (1:1)	73/350	$1.46 \times 10^{-3}$ /-	Evaluation of a bicomponent electrospinning method to produce the separator, Good physical properties and improved electrochemical stability.	[102]
PVDF-HFP	PET/SiO <sub>2</sub>	1 M LiPF <sub>6</sub> in EC/DEC (1:1)	60/-	$9.3 \times 10^{-4}$ /-	Separator with an organized porous structure, with benefits for cell operation at high C-rates; Excellent cell performance.	[103]
PVDF-HFP	MgAl <sub>2</sub> O <sub>4</sub>	1 M LiPF <sub>6</sub> in EC:DEC (1:1, v:v)	-/-	$2.80 \times 10^{-3}$ /140 (0.1C)	Influence of different quantities of the MgAl <sub>2</sub> O <sub>4</sub> filler in the membrane; Good ionic conductivity.	[104]
PVDF-HFP	MgAl <sub>2</sub> O <sub>4</sub>	1 M LiPF <sub>6</sub> in EC/DEC (1:1, w:w)	60/81	$10^{-3}$ (30 °C)/140 (C/10)	MgAl <sub>2</sub> O <sub>4</sub> as filler of thin and flexible separator; Good thermal stability and stable cycling performance.	[105]
PVDF-HFP	Mg(OH) <sub>2</sub>	1.15 M LiPF <sub>6</sub> in EC/EMC (3:7, v:v)	64/115	$8.08 \times 10^{-4}$ /105 (C/2)	Upgrading the battery safety operation by the addition of metal hydroxides in composite separators; High thermal stability and good capacity retention.	[106]
PVDF-HFP	MMT	1 M LiPF <sub>6</sub> in EC/DEC (1:1, v:v)	40/251	$9.01 \times 10^{-4}$ /105 (0.1C)	Use of montmorillonite as filler; High thermal stability and stable cycling performance.	[107]
PVDF-HFP	NaA	1 M LiPF <sub>6</sub> in EC/DEC (1:1, v:v)	65/194	$2.1 \times 10^{-3}$ /-	Separator with incorporation of NaA zeolite; Excellent thermal stability and wettability.	[108]

Table 5. Cont.

Materials	Fillers	Electrolyte Solution	Porosity and Uptake (%)	Conductivity ( $S \cdot cm^{-1}$ ) and Capacity ( $mAh \cdot g^{-1}$ )	Main Goal/Achievement	Ref.
PVDF-HFP	NaAlO <sub>2</sub>	0.5 M NaTf/EMITf	-/-	$5.5\text{--}6.5 \times 10^{-3}$ (25 °C)/-	Introduction of NaAlO <sub>2</sub> in a gel polymer electrolyte; Improved ionic conductivity.	[14]
PVDF-HFP	m-SBA15	1 M LiPF <sub>6</sub> in EC/DEC (1:1)	-/82.83	$3.23 \times 10^{-3}$ /156 (0.1C)	A PVDF-HFP composite membrane with m-SBA15 as filler; High coulomb efficiency.	[109]
PVDF-HFP	m-SBA15	1 M LiPF <sub>6</sub> in EC/DEC (1:1)	-/85.36	$3.78 \times 10^{-3}$ /198.6 (0.1C)	Effect of the addition of a silica filler on a PVDF-HFP composite matrix separator; High coulomb efficiency.	[110]
PVDF-HFP	OIL	1 M LiPF <sub>6</sub> in EC/DEC (1:1)	-/13	$2 \times 10^{-3}$ (25 °C)/141 (C)	Synthesis of OIL from a phenolic epoxy resin; Non-flammability, good cell performance.	[111]
PVDF-HFP	SiO <sub>2</sub>	1 M LiPF <sub>6</sub> in EC/DMC (1:2)	65.41/217	-/124.5 (C)	Synthesis of dual asymmetric structure separators with SiO <sub>2</sub> particles; High thermal stability and electrolyte uptake.	[16]
PVDF-HFP	SiO <sub>2</sub>	1 M LiPF <sub>6</sub> in DMC/EMC/DC/VC (46.08:22.91:27.22:3.79)	26.7/202	$8.47 \times 10^{-4}$ (25 °C)/154.4	Composite separator with SiO <sub>2</sub> ; Improved thermal stability and cycling performance.	[112]
PVDF-HFP	TiO <sub>2</sub>	1 M LiPF <sub>6</sub> in EC/DMC/EMC (1:1:1, v:v:v)	58/330	$3.45 \times 10^{-3}$ /122 (10C)	Evaluation of the performance of a nanocomposite polymer membrane with addition of TiO <sub>2</sub> ; Excellent electrochemical performance.	[85]
PVDF-HFP	ZrO <sub>2</sub>	1 M LiPF <sub>6</sub> in EC/DEC (1:1)	71/182	$1.48 \times 10^{-3} S \cdot cm^{-1}$ (25 °C)/126.8 mAhg <sup>-1</sup> (0.5C)	Preparation of ZrO <sub>2</sub> /PVDF-HFP by the dip-coating method High wettability, ionic conductivity, and thermal resistance.	[113]
PVDF-HFP	ZrO <sub>2</sub>	1 M LiPF <sub>6</sub> in EC/EMC (1:3)	-/-	$2.06 \times 10^{-3}$ (25 °C)/149.7	Improvement of the electrochemical properties of a electrospun membrane High uptake and ionic conductivity.	[114]
PVDF-HFP	ZrO <sub>2</sub>	1 M LiPF <sub>6</sub> in EC/DEC/DMC (1:1:1)	87.53/351.2	$3.2 \times 10^{-4}$ /646 (0.2C)	Inorganic fibers as substrates to separators; High thermal stability and good mechanical properties.	[115]
PVDF-HFP	ZrO <sub>2</sub>	1 M LiPF <sub>6</sub> in EC/DMC (1:1)	60/160	$10^{-3}$ (25 °C)/75 (C)	Development of thin and flexible ZrO <sub>2</sub> separators High porosity and thermal stability.	[116]

**Table 5.** *Cont.*

Materials	Fillers	Electrolyte Solution	Porosity and Uptake (%)	Conductivity ( $S \cdot cm^{-1}$ ) and Capacity ( $mAh \cdot g^{-1}$ )	Main Goal/Achievement	Ref.
PVDF-HFP	ZrO <sub>2</sub>	1 M LiPF <sub>6</sub> in EC/DMC (1:1)	95.7/481	$2.695 \times 10^{-3}$ (25 °C)/-	Incorporation of ZrO <sub>2</sub> in PVDF-HFP electrospun membranes; High ionic conductivity and cycling stability.	[117]
PVP/PVDF	Black carbon nanoparticles	6 M KOH	-/-	-/-	Production of separators for supercapacitor applications Improved thermal and mechanical properties.	[19]
PP/PVDF-HFP	PMMA	1 M LiPF <sub>6</sub> in EC/DMC (1:1, v:v)	77.9/212	$1.57 \times 10^{-3}$ /138 (0.2C)	Physical and electrochemical performances of a PP/PVDF-HFP/PMMA composite separator; Enhanced thermal stability and electrolyte uptake.	[52]
PP/PVDF-HFP	SiO <sub>2</sub>	1 M LiPF <sub>6</sub> in EC/DEC (1:1, v:v)	-/290	$1.76 \times 10^{-3}$ /150 (0.2C)	PP/PVDF-HFP separator, with the inclusion of SiO <sub>2</sub> nanoparticles; Favorable chemical stability and discharge capacity.	[118]
PI/PVDF-HFP	TiO <sub>2</sub>	1 M LiPF <sub>6</sub> in EC/DEC (1:1, v:v)	-/-	$1.88 \times 10^{-3}$ /161 (0.5C)	Electrospun PI/PVDF-HFP membrane, with addition of TiO <sub>2</sub> nanoparticles; Excellent electrochemical properties.	[119]

Several fillers such as n-butanol [90], SiO<sub>2</sub> [103], ZnO [86] MgAl<sub>2</sub>O<sub>4</sub> [105], and MMT [107] particles were used with PVDF and its copolymer composites in order to improve thermal and mechanical stability as well as the ionic conductivity value.

Mechanical improvement of separators has been achieved by developing sandwich-type composite separators, by a successive electrospinning method and based on PMIA [79].

The addition of DNA-CTMA in a PVDF matrix allows the development of flexible membranes, with interesting mechanical properties, highlighting its favorable stretch properties, allowing foldable separators with elevated elasticity [71].

The addition of cellulose nanoparticles in the separator structure proved to significantly increase the mechanical strength of the membrane. It also improves the wettability and induces the  $\beta$ -phase formation in PVDF. However, the presence of NCC reduces the ionic conductivity of the membrane [73].

The use of SiO<sub>2</sub> nanoparticles in a PVDF electrospun separator can raise the mechanical strength of the membrane, thus leading to a more tough and durable battery [106].

Improved security operation for lithium ion batteries, due to suitable flammability resistance, has been addressed by developing PVDF/LiPVAOB composites membranes [33].

The direct application of a ceramic suspension of PVDF/Al<sub>2</sub>O<sub>3</sub> in the electrode, resulting in a separator-cathode assembly, enhances the adhesion between these structures, and improves electrochemical cell performance [66].

PVP/PVDF membranes incorporated with carbon black nanoparticles were produced for supercapacitor applications. The separators showed improvements in mechanical properties and dielectric constant values [19].

GPEs based on boron-containing cross-linker proved to have high thermal resistance, maintaining their dimensional stability up to 150 °C, due to their stable PVDF matrix. Also, ionic conductivity and electrochemical stability were improved when compared to commercial separators [68].

Studies on the influence of solvents in nanoclay/PVDF separators showed that using DMAc as a solvent improves the porosity and electrolyte uptake of the membrane when compared with most used solvents such as NMP or DMF. Furthermore, the addition of PVP to the separator structure contributed to increase the pore size and to reduce the degree of crystallinity [72].

The addition of a metal-organic framework to a polymer structure proved to increase the conductivity of the produced membrane without needing an electrolyte. The membrane also showed high durability and good mechanical properties [76].

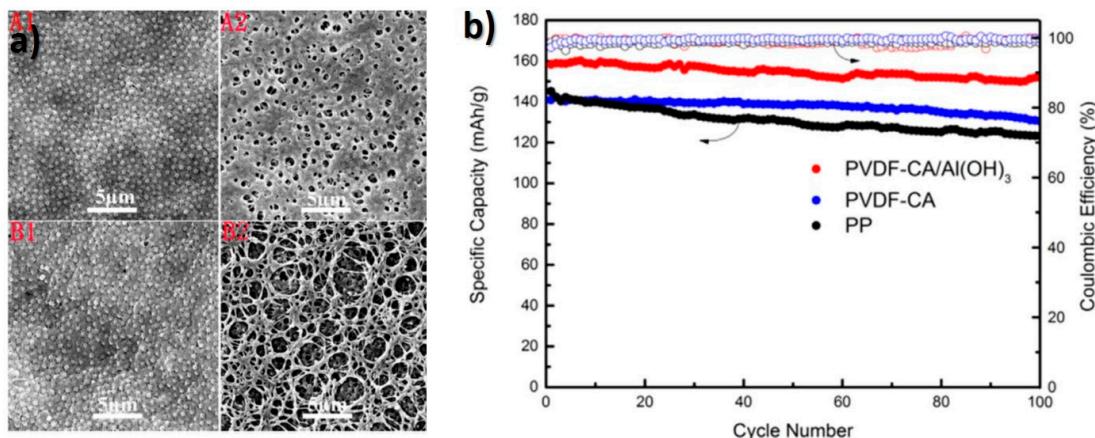
The dipping of PVDF nanofiber membranes into Al<sub>2</sub>O<sub>3</sub> proved to improve the thermal stability of the produced separator and its ionic conductivity. It also shows a low discharge capacity decay, even at high discharge rates [111].

A double-layer separator was prepared with PVDF and reduced graphene oxide, for lithium-sulfur batteries. It is shown that the two layers combined their properties to enhance the thermal stability of the membrane and the cycling performance of the cells [82].

The use of inorganic fibers as substrate for separators lead to improved thermal and mechanical stability when compared to commercial membranes. It was also proven that it enhanced the electrochemical performance of lithium ion cells [115].

CNF/PVDF composite membranes showed greater performance when applied in Li-S batteries, with enhanced cycling stability. The produced batteries retained a capacity of 768.6 mAhg<sup>-1</sup> after 200 cycles at a 0.5 C rate [20]. The development of PVDF-C separators by the phase-inversion method for Li-S batteries also leads to outstanding electrochemical performance results, associated with the presence of the conductive carbon network in the polymer matrix [69].

In the search for more environmental friendly materials, a separator with PVDF, cellulose acetate and Al(HO)<sub>3</sub> particles was developed by non-solvent induced phase separation (NIPS), the microstructure being presented in Figure 6a. This membrane exhibited high porosity, electrolyte uptake, and ionic conductivity, as well as good cycling capacity, even at high C-rates, as demonstrated in Figure 6b [70].



**Figure 6.** (a) SEM images of separators microstructure and (b) cycle performance of cells assembled [70], with copyright permission from Elsevier.

PVDF was also used in the study of the potential of zeolitic imidazolate framework-4 in separators. The prepared membranes showed high thermal stability, porosity, ionic conductivity, and cycling performance when compared with conventional separators [120].

The incorporation of Meldrum's acid groups in the PVDF structure proved to increase the ionic conductivity of the membrane, as well as the cycling performance, in particular at high C-rates [74].

PVDF/PFSA electrospun nanofibers allow the development of membrane with high mechanical stability and ionic conductivity with high discharge capacity and cycling stability [81].

A GPE membrane was developed by blending PVDF with PEO and ZrO<sub>2</sub>. This membrane showed high electrolyte uptake, and excellent rate performance and discharge capacity for application in lithium-sulfur batteries [88].

Electrospun membranes with Octaphenyl-POSS nanoparticles showed a significant improvement in porosity and electrolyte uptake. For a ratio of 2:100 (*w:w*), the separator proved to have high mechanical stability, ionic conductivity, and thermal stability [77].

A nonaflate anion-based IL and lithium salt was introduced on a GPE, allowing the development of a membrane with high thermal stability and electrochemical properties. When used alongside with a LiCoO<sub>2</sub> cathode, this separator also showed good discharge capacity and capacity of retention [96].

The addition of MgAl<sub>2</sub>O<sub>4</sub> as filler in electrospun fibrous PVDF-HFP separators contributes to improving electrochemical performance, with high discharge capacity and excellent cycle life results [104].

The integration of m-SBA15 as filler in a polymer matrix, on the other hand, is advantageous as it decreases the degree of crystallinity of PVDF-HFP, increasing electrolyte uptake and enhancing the ionic conductivity [109,110].

The enhancement of the electrochemical performance has been extensively addressed by composites membranes with TiO<sub>2</sub> nanoparticles [119], and clay nanosheets [95], the latter improving the interfacial areal connection between the polymer structure and clay, facilitating ion transport.

The NaA zeolite is considered to be a very interesting material for incorporation as filler in lithium ion battery separators. It allows the formation of voids in the composite separator structure, which are filled with electrolyte, substantially increasing the ionic conductivity [108].

The safety operation of lithium ion batteries can be upgraded by the addition of metal hydroxides such as Al(OH)<sub>3</sub> and Mg(OH)<sub>2</sub>, in PVDF-HFP composite separators. These metal hydroxides endow a fire-retardant behavior to the cells, due to their natural thermal stability [91].

Kuo et al. synthesized an oligomeric ionic liquid from a phenolic epoxy resin. By blending this ionic liquid with PVDF-HFP, a high performance, non-flammable gel polymer membrane was

obtained. This membrane exhibits high ionic conductivity, although with a low liquid electrolyte uptake (<50%) [111].

The addition of  $ZrO_2$  filler increases the porosity, ionic conductivity, and thermal resistance of the PVDF membranes. The presence of polar constituents and high connected interstitial voids facilitate electrolyte absorption, increasing the ionic conductivity and the performance of the membranes [113]. When a layer of  $ZrO_2$  was added between two layers of PVDF-HFP, the obtained separator presented even better electrochemical properties [114].

Graphene oxide nanosheets incorporated during the phase inversion of PVDF-HFP improve electrochemical battery performances of the produced separators, as well as thermal stability and the mechanical properties of the membrane [97].

HMSS/PVDF-HFP composite separators with improved porosity were developed; the presence of  $SiO_2$  spheres created a well-developed microporous structure, leading to higher wettability and ionic conductivity [98].

The incorporation of a superfine LLTO in a PVDF-HFP separator enhanced the ionic conductivity of the membrane. It was also been shown that a cell with a this type of separator presents improved discharge capacity and rate performance [101].

Bohemite composite separators were produced, exhibiting cycling performances comparable to the conventional ones. These membranes are also safer because of the limitation to Li dendrite formation, preventing the occurrence of short circuits [67].

A comparative study of  $Al_2O_3$  and  $NaAlO_2$  particles in a gel polymer electrolyte proved that  $NaAlO_2$  membranes present higher ionic conductivity than  $Al_2O_3$ , as well as improved mechanical properties [14].

$ZrO_2$  membranes with PVDF-HFP as a binder were produced by solvent casting methods. These separators present high porosity and thermal stability, but show lower mechanical strength than commercially available membranes [116].

A GPE produced by thermal crosslinking of PEGDA and PEGMEA proved to be compatible with lithium ion batteries, with a high coulombic efficiency of 94% after 100 cycles [78].

Liu et al. produced a GPE with PVDF-HFP and graphene via NIPS. The addition of a small concentration of graphene (0.002 wt. %) proved to significantly improve the properties of the membrane by increasing porosity, electrolyte uptake, ionic conductivity, and cycling performance, when compared to commercial separators [18].

Regardless of the fillers type used, Table 5 shows that most of the work is devoted to increasing ionic conductivity and electrochemical performance compared to the pure matrix. In particular, inert oxide ceramics ( $Al_2O_3$ ,  $TiO_2$ ,  $SiO_2$ ,  $ZrO_2$ ) reduce the degree of crystallinity, and enhance mechanical properties and ionic conductivity value. Carbon materials (CNF, Graphene, rGO) improve safety and interfacial stability between electrodes and separator membranes, and lithium fillers such as  $Li_{1.3}Al_{0.3}Ti_{1.7}(PO_4)_3$ , LiTSFI, and LLTO increase ionic conductivity value of the separators.

In addition, there are other fillers types such as zeolites and clays that are being intensely used for the development of separators, allowing the improvement of electrochemical behavior.

### 3.4. Polymer Blend Separator Membranes

Finally, another type of separator membrane are polymer blends where two different polymers with complementary properties are used; for example one showing excellent mechanical properties and the other with a hydrophilic character. The main properties of polymer blends based on PVDF and its copolymer are presented in Table 6.

**Table 6.** Polymer blends based on PVDF and co-polymers with main properties, goal, and achievement.

Materials	Blends	Electrolyte Solution	Porosity and Uptake (%)	Conductivity ( $S \cdot cm^{-1}$ ) and Capacity ( $mAh \cdot g^{-1}$ )	Main Goal/Achievement	Ref
PVDF	HDPE	1 M LiPF <sub>6</sub> in EC/DEC/DMC (1:1:1)	58/260	$2.54 \times 10^{-3} S \cdot cm^{-1}$ (25 °C)/156.1 mAhg <sup>-1</sup> (0.1C)	Production of a sponge-like PVDF/HDPE film; High ionic conductivity and cycling performance.	[121]
PVDF	HTPB-g-MPEG	1 M LiPF <sub>6</sub> in EC/DMC/EMC (1:1:1)	56/350	$3.1 \times 10^{-3}$ /116 (C)	Enhancement of the stability of entrapped liquid electrolyte and corresponding ion conductivity.	[122]
PVDF	MC	1 M LiPF <sub>6</sub> in EC/DEM/EMC (1:1:1, w:w:w)	-/138.6	$1.5 \times 10^{-3}$ /110 (C)	PVDF composite separator with cellulose material; Excellent electrochemical performance.	[123]
PVDF	MEP	1 M TEABF <sub>4</sub> in AN/PC and 1 M LiPF <sub>6</sub> in EC/DEC	77/-	$1.3 \times 10^{-2}$ /-	Manufacturing by phase inversion, with MEP as a cross-linking agent; Good mechanical strength.	[34]
PVDF	NCC	1 M LiFAP in EC/DMC (1:1)	-/-	-/-	Separators with applications in hybrid electric vehicles; Favorable performance at high-voltage cells.	[124]
PVDF	NCC	1 M LiPF <sub>6</sub> in EC/DMC (1:1)	-/-	-/108 (1C)	Separators with applications in hybrid electric vehicles; Influence on high-rate cell operation.	[125]
PVDF	PAN	1 M LiPF <sub>6</sub> in EC/DMC/DEC (1:1:1)	77.7/414.5	$2.9 \times 10^{-3}$ (25 °C)/-	Improved thermal and mechanical properties; High cycling stability.	[126]
PVDF	PAN	1 M LiPF <sub>6</sub> in EC/DMC/EMC (1:1:1)	-/320	$1.45 \times 10^{-3}$ /145.71 (0.2C)	Production of an electrospun blended membrane; High thermal and mechanical stability.	[127]
PVDF	PBA	1 M LiPF <sub>6</sub> in EC/DEC/DMC (1:1:1)	-/120	$8.1 \times 10^{-4}$ (25 °C)/95 (0.1C)	Preparation of cross-linked PBA/PVDF GPE; Good cycling stability.	[128]
PVDF	PDMS-g-(PPO-PEO)	1 M LiPF <sub>6</sub> in EC/DMC/EMC (1:1:1, w:w:w)	80.1/512	$4.5 \times 10^{-3}$ /120 (1C)	Porous separator; Good electrochemical stability.	[129]
PVDF	PEGDA	1 M LiPF <sub>6</sub> in EC/DMC (1:1)	-/-	$3.3 \times 10^{-3}$ /117 (0.1C)	Separator produced by thermal polymerization; High capacity retention.	[130]
PVDF	PEO	1 M LiPF <sub>6</sub> in EC/DMC (1:1)	/530	-/-	Production of blended membranes by electrospinning; improved conductivity and uptake.	[131]

Table 6. Cont.

Materials	Blends	Electrolyte Solution	Porosity and Uptake (%)	Conductivity ( $S \cdot cm^{-1}$ ) and Capacity ( $mAh \cdot g^{-1}$ )	Main Goal/Achievement	Ref
PVDF	PEO	1 M LiPF <sub>6</sub> in EC/DMC (1:1)	-/527	-/-	Development of electrospun membranes; High electrolyte uptake, low shutdown temperature.	[132]
PVDF	PET	-	80/270	-/-	Synthesis of a hybrid separator; High wettability and electrolyte uptake.	[133]
PVDF	PI	1 M LiPF <sub>6</sub> in EC/PC/DEC/VC (35.4:17.2:45.1:2.3)	-/-	$1.3 \times 10^{-3}$ /141	Preparation of the separator by electrospinning; Improved thermal stability and mechanical properties.	[134]
PVDF	PMMA/CA	1 M LiPF <sub>6</sub> in EC/DMC (1:1, <i>w:w</i> )	99.1/323	-/-	Elevated porosity and electrolyte uptake.	[135]
PVDF	P(MMA-co-PEGMA)	1 M LiPF <sub>6</sub> in EC/EMC/DMC (1:1:1, <i>w:w:w</i> )	-/372	$3.01 \times 10^{-3}$ / -	Porous separator; Improved capacity retention.	[136]
PVDF	PMMA/SiO <sub>2</sub>	-	80.1/293.2	$1.97 \times 10^{-3}$ / -	Evaluation of the effect of a PMMA and SiO <sub>2</sub> blend on a PVDF electrospun membrane as a separator; High electrolyte uptake and improved ionic conductivity.	[137]
PVDF	PVP	1 M Et <sub>4</sub> N-BF <sub>4</sub> /PC	-/360	$1.8 \times 10^{-3}$ (25 °C) / -	Separators for supercapacitors; High uptake and power density.	[138]
PVDF	TAIC	1 M TEABF <sub>4</sub> in AN/PC and 1 M LiPF <sub>6</sub> in EC/DEC	75/-	$1.4 \times 10^{-2}$ / -	Manufacturing of separator by phase inversion, with TAIC as cross-linking agent. High ionic conductivity.	[34]
PVDF-TrFE	PEO	1 M LiTFSI in PC	44.5/107	$5.4 \times 10^{-4}$ / 124 (C/5)	Research on the physical and chemical properties of a PVDF-TrFE/PEO blend Favorable cycling performance.	[139]
PVDF-HFP	CA	1 M LiPF <sub>6</sub> in EC/DMC/EMC (1:1:1, <i>v:v:v</i> )	66.36/355	$6.16 \times 10^{-3}$ / 138 (0.2C)	Investigation of the use of CA from waste cigarette filters, in PVDF-HFP membranes; Good electrochemical performance and excellent thermal stability.	[140]
PVDF-HFP	HDPE	-	71/300	$2.97 \times 10^{-3}$ (25 °C) / 140.5 (C)	Preparation of the separator by non-solvent-induced phase separation; High ionic conductivity.	[141]
PVDF-HFP	PANI	1 M LiPF <sub>6</sub> in EC/DMC (1:1)	83/270	$1.96 \times 10^{-3}$ / -	High thermal stability, electrolyte uptake, and ionic conductivity	[142]

**Table 6.** *Cont.*

Materials	Blends	Electrolyte Solution	Porosity and Uptake (%)	Conductivity ( $S \cdot cm^{-1}$ ) and Capacity ( $mAh \cdot g^{-1}$ )	Main Goal/Achievement	Ref
PVDF-HFP	PEG/PEGDMA	1 M LiClO <sub>4</sub> in EC/DEC (1:1, <i>v:v</i> )	71/212	$1.70 \times 10^{-3}$ / -	Investigation into a strengthened electrospun nanofiber membrane separator; High porosity and electrolyte uptake.	[143]
PVDF-HFP	PLTB	1 M LiPF <sub>6</sub> in EC/DMC (1:1, <i>v:v</i> )	70/260	$1.78 \times 10^{-3}$ / 138 (0.5C)	Excellent electrochemical performance.	[144]
PVDF-HFP	PSx-PEO3	1 M LiTFSI in EC/DMC (1:1, <i>w:w</i> )	-/520	$4.2 \times 10^{-4}$ (20 °C) / 123 (C)	Production of a safe PVDF-HFP blended membrane, which can be sprayed; Elevated electrolyte uptake.	[145]
PVDF-HFP	PVSK	1 M LiTFSI + 0.25 M LiNO <sub>3</sub> in DME/DOL (1:1)	27/-	-/1220	Improved cycling performance.	[146]
PVDF-HFP	PVC	1 M LiPF <sub>6</sub> in EC/DMC (1:2)	62/230	$1.58 \times 10^{-3}$ / 125 (0.1 C)	Tri-layer polymer membrane; Good mechanical and thermal stability.	[44]
PEI/PVDF	x-PEGDA	1 M LiPF <sub>6</sub> in EC/DMC/EMC (1:1:1)	64.6/235.6	$1.38 \times 10^{-3}$ (25 °C) / 160.3 (0.2C)	Production of x-PEGDA-coated PEI/PVDF membranes; high wettability, porosity, and ionic conductivity.	[147]

PVDF composite separators with methyl cellulose as host of gel polymer electrolyte allows the development of low cost and environmentally friendlier separators with excellent mechanical, thermal, and electrochemical performances [123].

A trilayer porous membrane of PVDF-HFP with PVC as the middle layer was developed. It was shown that a good porosity and uptake value can be achieved, though the mechanical stability was negatively affected [44].

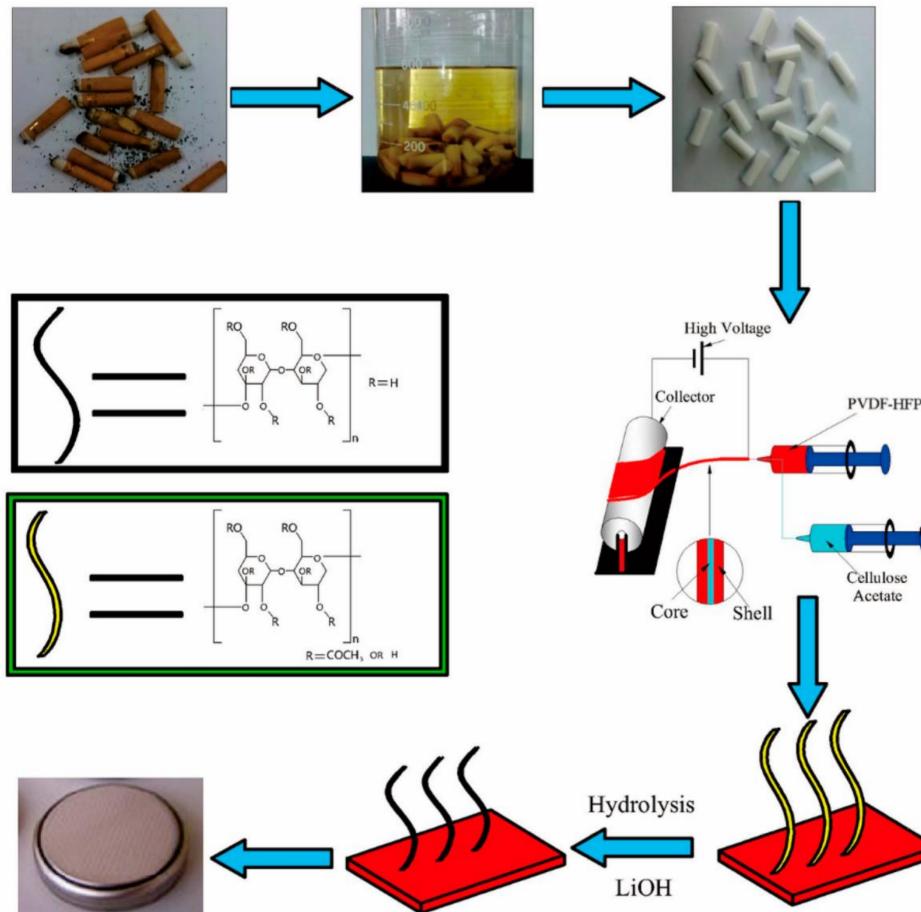
Cells produced with PVDF-NCC separators presented good battery performance at high C-rates, which is very critical for meeting the minimum and maximum power-assist requirements for integration in hybrid electric vehicles [124,125].

A mechanically strengthened electrospun composite PVDF-HFP/PEG/PEGDMA separator was developed. PEG and PEGDMA allow the improvement of the mechanical strength of the composite membrane, which is confirmed by the existence of physical bonded structures [143].

P(MMA-co-PEGMA) and PDMS-g-(PPO-PEO) copolymers within PVDF allow the reduction of the crystallinity of the PVDF matrix, and gently improve the electrolyte uptake, thus leading to an enhanced ionic conductivity [129,136].

PLTB can be successfully used in a PVDF-HFP composite separator. In comparison with a typical PP separator, it is more safe and efficient, due to its thermal and electrochemical stability. This separator is very promising in terms of security operation, because of flame retardant characteristics [144].

An eco-friendly technique to recover cellulose acetate from wasted cigarette filters (Figure 7) was developed, and the material can be integrated in a PVDF/CA membrane for lithium ion batteries, which presents a good performance [140].



**Figure 7.** Preparation of PVDF-HFP/CA nanofiber separators for lithium ion batteries [140], with copyright permission from the American Chemical Society.

PVDF separators were manufactured by a phase inversion technique, with two different cross-linking agents (TAIC and MEP) and with the application of gamma radiation. The produced membranes are characterized by good mechanical behavior and low electrical resistance [34].

Electrospun PVDF membranes blended with PMMA/SiO<sub>2</sub> showed good porosity and elevated electrolyte uptake [137]. Blending with PI further enhanced their thermal and mechanical properties, ensuring a better battery performance than commercial PE separators [134].

PVDF/PEO blend membranes show an increase of the ionic conductivity and electrolyte uptake when compared with PVDF membranes. The improved wettability and porosity in x-PEGDA-coated PEI/PVDF membranes has been also reported [147].

PVDF-HFP/HDPE membranes were prepared by non-solvent induced phase separation. This separator presents good cycling performance in lithium ion batteries and a high ionic conductivity [141]. Further studies showed an increased discharge capacity of these membranes, by decreasing the size of the HDPE fillers [121].

PVDF/PAN blend separators were produced by TIPS [126] and electrospinning [127], with improved thermal and mechanical properties. The best PVDF/PAN ratio was 90:10. Despite the lower ionic conductivity when compared with conventional separators, these membranes showed higher cycle and C-rate performance [126].

PVDF/PAN electrospun membrane have excellent dimensional stability even at high temperatures, high electrolyte uptake and ionic conductivity, and superior discharge capacity [127].

The blending of PVDF and PEO in an electrospun membrane proved to increase significantly the electrolyte uptake of the separator, while decreasing the shutdown temperature [132].

Cross-linked PBA/PVDF GPE were prepared by soaking semi-interpenetrating polymer networks with liquid electrolyte. For a PBA/PVDF ratio of 1:0.5, the best results of electrolyte uptake, ionic conductivity, and cycling stability were obtained [128].

A PVDF/PET hybrid separator was produced via a mechanical pressing process. The obtained membrane presented high wettability and electrolyte uptake, while maintaining good thermal stability [133].

The introduction of PANI in a PVDF separator by the breath figure method proved to increase the electrolyte uptake and ionic conductivity of the membrane. The best results were obtained for 30% of PANI, with a uniform pore structure and excellent thermal stability [142].

The use of PVDF-HFP/PVSK membranes in lithium-sulfur batteries has been reported. It has been proved that even small amounts of PVSK (5 wt. %) increase the discharge capacity of the cell and reduce the capacity decay [146].

An increase of the use of natural polymers and biopolymers is observed for the preparation of PVDF and copolymer blends, considering the environmental issues. It is demonstrated in Table 6 that they allow to improve mechanical properties and wettability, and consequently the battery performance. In addition, the use of conductive polymers such as PANI in polymer blends has acquired special attention in recent years, considering that the electrical properties are improved without mechanical deterioration. Typically, the most commonly used PVDF and PVDF-HFP blends are developed with PAN and PEO polymers, allowing the improve thermal and mechanical stability, as well as wettability and ionic conductivity value, respectively.

#### 4. Conclusions and Future Trends

In this review, the latest advances in PVDF-based battery separators for lithium-ion battery applications are presented.

Considering the excellent properties of PVDF and its copolymers as a separation membrane and the importance of the role of the battery separator in battery applications, this review was divided into four different sections—single polymers, surface modification, polymer composites, and blends, where, for each category, the improvement of the main properties of the separators' degree of porosity,

uptake value, mechanical and thermal properties, ionic conductivity, and cycling performance, as well as safety and environmental impacts,- for the different developed materials was presented.

In the single polymer category, PVDF and PVDF-HFP stand out as the most commonly applied polymers produced by various processing techniques, with TIPS and electrospinning methods being the most commonly used to tailor microstructure (degree of porosity and pore size) to improve battery performance.

The number of research papers on surface modifications of the membranes has increased in recent years, as the surface of the polymer membrane strongly affects the uptake process. Surface modification is accomplished by coating hydrophilic polymers or plasma treatment to increase the interaction between the polymer membrane and the electrolytic solution.

Generally, the addition of fillers increases battery performance through the improvement of ionic conductivity in polymer composites, but has not yet demonstrated the best filler for PVDF and its copolymer membranes. The most commonly used fillers are inert oxide ceramics, carbon materials, and lithium fillers. The most improved properties are mechanical properties, interfacial stability, between electrodes and separator membranes and ionic conductivity value, respectively.

In relation to the polymer blends, the appearance of new blends based on natural and conductive polymers within PVDF for battery separators has been observed.

The blends of PVDF and its copolymers widely used are with PAN and PEO polymers, allowing the improvement of mechanical properties and wettability and electric properties, respectively.

The future trends for single polymer separators are to obtain single polymers with a porosity above 50% but a smaller pore size below 500 nm to prevent dendrite growth. Further, it is expected an increase in the use of ionic liquids as electrolytic solution. In relation to surface modifications, the use of poly (ionic liquids) and natural polymers as a surface modification coating of PVDF polymer membranes will be interesting, considering environmental issues.

With respect to polymer composites, future perspectives are related to improving the interaction between polymer matrix and fillers, in order to optimize filler content without decreasing electrical properties or hindering mechanical stability. Also, the use of more than one filler with complementary properties may be the way for improving cycling performance.

The progress with respect to polymer blends is related to the scalability of the fabrication process, increasing the interaction and compatibilization of the two polymers.

In summary, PVDF-based battery separators allow the tailoring of all the properties/characteristics required for a new generation of separator membranes for lithium-ion batteries with high power and excellent cycling performance.

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## List of Symbols and Abbreviations

(C <sub>2</sub> H <sub>5</sub> ) <sub>3</sub> CH <sub>3</sub> NBF <sub>4</sub>	Triethylmethylammonium tetrafluoroborate
[C <sub>2</sub> mim][NTf <sub>2</sub> ]	1-ethyl-3-methylimidazolium bis(trifluoromethylsulfonyl)imide
Al(OH) <sub>3</sub>	Aluminum hydroxide
Al <sub>2</sub> O <sub>3</sub>	Aluminum oxide
AlO(OH)	Bohemite
AN	Acetonitrile
BC	Boron-containing cross-linker
CA	Cellulose acetate
CMC	carboxymethyl cellulose
CNF	Carbon nanofiber
DEC	Diethyl carbonate

DEM	Diethoxymethane
DMAc	Dimethyl acetamide
DMC	Dimethyl carbonate
DME	1,2-dimethoxyethane
DMF	Dimethyl formamide
DMSO	Dimethyl sulfoxide
DNA-CTMA	Deoxyribonucleic acid-cetyltrimethylammonium
DOL	1,3-dioxolane
EC	Ethylene carbonate
EMC	Ethyl methyl carbonate
EMImNfO-LiNfO	1-ethyl-3- methylimidazolium nonafluoro-1-butanesulfonate/lithium nonafluoro-1-butanesulfonate
EMITf	1-ethyl 3-methyl imidazolium trifluoromethane sulfonate
EMITFSI	1-ethyl-3-methyl-imidazolium bis(trifluoromethanesulfonyl)imide
EP	Ethyl propionate
Et <sub>4</sub> N-BF <sub>4</sub>	Tetraethylammonium tetrafluoroboratein
GF	Glass fiber
GO	Graphene oxides
GPE	Gel polymer electrolyte
H <sub>2</sub> SO <sub>4</sub>	Sulfuric acid
HDPE	High density polyethylene
HEC	Hydroxyethyl cellulose
HMSS	Hollow mesoporous silica spheres
HTPB-g-MPEG	Hydroxyl-terminated polybutadiene-grafted-methoxyl polyethylene glycol
KOH	Potassium hydroxide
LiClO <sub>4</sub>	Lithium perchlorate
LiCoO <sub>2</sub>	Lithium cobalt oxide
LiFAP	Lithium Tris(pentafluoroethane)-trifluorophosphate
LiNfO/BMImNfO	Lithium nonafluorobutanesulfonate/1-butyl-3-me-thylimidazolium nonafluorobutanesulfonate
LiNO <sub>3</sub>	Lithium nitrate
LiPF <sub>6</sub>	Lithium hexafluorophosphate
LiPVAOB	Lithium polyvinyl alcohol oxalate borate
Li-S	Lithium-sulfur
LiTFSI	lithium bis(trifluoromethanesulfonyl)imide
LLTO	Li0.33La0.557TiO3
MA	Meldrum's acid
MC	Methyl cellulose
MEP	Ethylene oxide-propylene oxide
Mg(OH) <sub>2</sub>	Magnesium hydroxide
MgAl <sub>2</sub> O <sub>4</sub>	Magnesium aluminate
MMT	Montmorillonite
MOF-808	Zirconium (IV) metal-organic framework
m-SBA 15	Mesoporous silica
NaA	NaA zeolite
NaClO <sub>4</sub>	Sodium perchlorate
NaTf	Sodium trifluoromethane sulfonate
NCC	Nanocrystalline cellulose
NIPS	Non-solvent induced phase separation
NMP	N-methyl-2-pyrrolidone
OIL	Oligomeric ionic liquid (bromide bis(tri-fluoromethane)sulfonimide)
P(MMA-co-PEGMA)	Poly(methyl methacrylate-co-poly(ethylene glycol) methacrylate)
PAN	Polyacrylonitrile
PANI	Polyaniline
PBA	Poly(butyl acrylate)
PC	Propylene carbonate
PDA	Polydopamine
PDMS-g-(PPO-PEO)	Poly(dimethylsiloxane)-graft-poly(propylene oxide)-block-poly(ethylene oxide)
PE	Polyethylene

PEG	Polyethylene glycol
PEGDA	Poly(ethylene glycol)diacrylate
PEGDMA	Polyethylene glycol dimethacrylate
PEGMEA	Poly(ethylene glycol) methyl ether acrylate
PEI	Polyetherimide
PEO	Polyethilene oxide
PET	Polyethylene terephthalate
PFSA	Perflourosulfonic acid
PI	Polyimide
PLTB	Polimeric lithium tartaric acid borate
PMIA	Poly(m-phenylene isophthalamide)
PMMA	Polymethyl methacrylate
POSS	Polyhedral oligomeric silsesquioxane
PP	Polypropylene
P-PAEK	Phenolphthaleine-poly(aryl ether ketone)
PSx-PEO3	Polysiloxane-comb-propyl(triethylene oxide)
PSU	Poly(sulfone)
PTFE	Poly(tetrafluoroethylene)
PVA	Polyvinyl alcohol
PVC	Poly(vinyl chloride)
PVDF	Poly(vinylidene fluoride)
PVDF-co-CTFE	Polyvinylidene fluoride-co-chlorotrifluoroethylene
PVDF-co-HFP	Poly(vinylidene fluoride-co-hexafluoropropylene)
PVDF-HFP	Poly(vinylidene fluoride-co-hexafluoropropene)Poly(vinylidene fluoride-hexafluoropropylene)
PVDF-PE	Polyvinylidene difluoride-coated polyethylene
PVDF-TrFE	Poly(vinylidene fluoride-trifluoroethylene)
PVP	Polyvinylpyrrolidone
PVSK	Polyvinylsulfate potassium salt
rGO	Reduced graphene oxide
SCPC	Self-charging power cell
SiO <sub>2</sub>	Silicon dioxide
SN	Succinonitrile
SnO <sub>2</sub>	Tin oxide
TAIC	Triallyl isocyanurate
TEABF <sub>4</sub>	Tetraethyl ammonium tetrafluoroborate
TiO <sub>2</sub>	Titanium dioxide
TIPS	Thermal-induced phase separation
TTT	1,3,5-triallyl-1,3,5-triazine-2,4,6(1 H,3 H,5 H)-trione
VC	Vinylene carbonate
x-PEGDA	x-polyethylene glycol diacrylate
ZnO	Zinc oxide
ZrO <sub>2</sub>	Zirconium dioxide

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