Flow-Microreactor-System Controlled Polymerization
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Abstract
Flow microreactors are expected to make a revolutionary change in chemical synthesis involving various fields of polymer synthesis. In fact, extensive studies on cationic polymerization, anionic polymerization, radical polymerization, coordination polymerization, polycondensation, and ring-opening polymerization using flow microreactor systems have opened new possibilities in polymer chemistry and polymer industry. This article provides, in a concise form, a current overall picture of polymerization using flow microreactors.

1 Introduction
Microtechnology is no longer the field of electronics, but is now moving into many different areas of science and engineering, including mechanics, optics, and fluids, because it provides better efficiencies while also answering the demands of society for conserving resources and energy. Following remarkable advances in microfabrication technology, microtechnology has been used in the field of chemistry since the 1990s. Micro devices that are used for chemical reactions are called microreactors; reactors with micrometer-size channels for conducting chemical reactions.

Microreactors are normally set up as flow-type reactors (flow microreactors) with a continuous flow of a solution through the reaction chamber. Nowadays, chemical synthesis in flow microreactors has received significant research interests from both academia and industry [1-24]. Recent investigations revealed significant features of flow microreactor systems involving fast mixing stemming from short diffusion paths and fast heat transfer by virtue of high surface-to-volume ratios, which are advantageous to increase the selectivity of chemical reactions [25-28]. Short residence time in a microchannel is beneficial for controlling highly reactive intermediates [29-45]. By taking advantages of such features of flow microreactor systems, various chemical reactions for organic synthesis have been developed so far [46-56].

Polymerization reactions that convert small molecules into macro molecules by repeating chemical reactions are a fascinating field in the applications of flow microreactors. Extensive studies on cationic polymerization, anionic polymerization, radical polymerization, coordination polymerization, polycondensation, and ring-opening polymerization using flow microreactor systems have been carried out so far. Reviews on various polymerization methods in microreactors have been reported by Hessel et al., Wilms et al., Mcquade et al. and Serra et al [57-61]. The main objective of this article is to provide, in a
concise form, a current overall picture of polymerization using flow microreactors, especially from a viewpoint of industrial applications. Because of space limitations, polymer particle synthesis will not be discussed here. Serra and Chang has reported an excellent review on synthesis of polymer particles with an improved control of their sizes, size distributions, morphologies and compositions using microreactors [62].

1.1 Characteristic Features of Flow Microreactors

Flow microreactors can influence the very essence of chemical reactions, because of the following characteristic features derived from their small size and flow nature.

(1) Fast mixing [63, 64]: Many chemical reactions involve combining two substances, and for this reason mixing to achieve homogeneity in solution is very important, especially for fast reactions. Mixing occurs due to molecular diffusion. According to the theory of molecular diffusion, the time needed for molecular diffusion is proportional to the square of the length of the diffusion path. Therefore, the marked shortening of the diffusion path in a microreactor results in a mixing speed that is unobtainable in a macroreactor. The working principle of a typical micromixer (i.e., a multilamination-type micromixer [65]) is shown in Figure 1. The solutions to be mixed are distributed into several segments of flows of small width by using the microstructure. The mixing takes place through the interfaces of the segments of flows by virtue of the short diffusion path.

![Figure 1 Working principles of a multilamination-type micromixer.](image)

(2) Temperature control: Heat is transferred between the interior and exterior of a reactor via the reactor surface according to the theory of heat transfer. Therefore, area per unit volume of the reactor is a crucial factor for heat transfer. Generally, volume is equal to the length cubed, while surface area is equal to length squared. When the length is shortened, surface-to-volume ratio increases. Thus, a feature of microspaces compared to macrospaces is that they have larger surface-to-volume ratios (Figure 2). Because microreactors have a greater surface area per unit volume than macroreactors, heat transfer occurs rapidly in a flow microreactor, enabling fast cooling/heating and, hence, precise temperature control.
Another characteristic feature of microreactors derived from their much greater surface-to-volume ratios is that they make phase-boundary reactions such as gas/liquid, liquid/liquid, or solid/liquid reactions more efficient. This feature of flow microreactors is also advantageous for photochemical [66-75] and electrochemical [76-86] reactions, which have received significant attention from a viewpoint of environmentally benign synthesis.

(3) Residence time control: The length of time that the solution remains inside the reactor is called the residence time. In flow reactors, the residence time increases with the length of the channel and decreases with the flow speed. In flow microreactors the residence time can be greatly reduced by shortening the length of the microchannels. This feature of flow microreactors is extremely useful in controlling reactions involving unstable short-lived reactive intermediates. Unstable reactive species can be transferred to another location to be used in the next reaction before they decompose (Figure 3). By taking advantage of this feature, chemical transformations that are very difficult or impossible in macroreactors can be achieved in microreactors [87-90].

Figure 2 Numerical aspects of decreasing size.

Figure 3 Principle of generation and reaction of unstable short-lived reactive intermediates based on residence time control in a flow microreactor.

1.2 Cationic Polymerization
1.2.1 Basic Principles of Cationic Polymerization of Vinyl Monomers
Cationic polymerization [91, 92] is one of the most fundamental methods for synthesizing polymers. Although there are several types of cationic
polymerization, the most important one is cationic polymerization of vinyl monomers having a cation stabilizing group (Y) (Scheme 1).

**Scheme 1** Cationic polymerization of vinyl monomers.

The initiation usually involves the addition of a cationic species ($A^+$) to a vinyl monomer to produce a carbocationic intermediate associated with a counter anion ($X^-$), which is derived from the initiator. In general, proton acids or carbocations generated from their precursors by acid promoted ionization reactions [93-95], are used as initiators. The carbocationic intermediate thus obtained adds to another molecule of the monomer to give the next carbocationic intermediate, which adds to another monomer (propagation step). The carbocationic intermediates are usually highly reactive and unstable. They undergo a number of side reactions such as chain transfer and termination. Chain transfer to monomer is the most problematic from a view point of molecular weight control and molecular weight distribution control. Namely, the β-proton of the carbocationic intermediate is inherently acidic because of the positive charge on the carbon. On the other hand, monomers used in cationic polymerization are inherently nucleophilic or basic. Therefore, the proton abstraction from the carbocationic intermediate by the monomer is inevitable and is very difficult to suppress (Scheme 2).

**Scheme 2** Chain transfer as a side reaction in cationic polymerization of vinyl monomers.

**1.2.2 Controlled/living Cationic Polymerization of Vinyl Monomers Based on Cation Stabilization**

One of the most important breakthroughs in cationic polymerization is the discovery of living cationic polymerization. The inherent and serious drawback of cationic vinyl polymerization is instability of the carbocationic intermediates, which causes the chain transfer leading to the formation of
polymers of broad molecular weight distribution. Higashimura and Sawamoto et al. proposed and verified experimentally that living cationic polymerization can be attained by stabilizing the carbocationic intermediate by nucleophilic interaction with a suitably nucleophilic counter anion or an externally added Lewis base (B) (Scheme 3) [96-98].

Scheme 3 Stabilization of the carbocationic intermediate by nucleophilic interaction with a suitably nucleophilic counteranion or an externally added Lewis base.

In both methods the positive charge of the carbocationic center is reduced and thereby the acidity of the \( \beta \)-proton is reduced to suppress the chain transfer. As a result, good molecular weight control and molecular weight distribution control are attained. On the basis of the principles, a number of initiating systems have been developed for living cationic polymerization [99].

1.2.3 Controlled/living Cationic Polymerization of Vinyl Ethers Based on Cation Stabilization Using Flow Microreactor Systems [100]

Living cationic polymerization of vinyl ethers initiated by SnCl\(_4\)/RCl catalytic system can be carried out in a continuous microreactor system, which consists of a multilamination micromixer \( \text{M} \) (channel width = 40 \( \mu \text{m} \), IMM) and a micro tube reactor \( \text{R} \) (Figure 4). A solution of a monomer and an initiator’s precursor is mixed with a solution of SnCl\(_4\) using the micromixer at -78 °C and the resulting mixture was allowed to react in the microtube reactor at the same temperature. For example, isobutyl vinyl ether (IBVE) can be polymerized using functionalized initiators to obtain the end-functionalized polymers of narrow molecular weight distribution (Mw/Mn < 1.2). Block copolymerization of IBVE and \( n \)-butyl vinyl ether (NBVE) can also be successfully achieved using a microreactor system consisting of two micromixers and two microtube reactors to obtain the corresponding copolymer of narrow molecular weight distribution (Mw/Mn < 1.3).
1.2.4 Controlled/living Cationic Polymerization of Vinyl Ethers Without Stabilizing Carbocationic Intermediates Using Flow Microreactor Systems

One of the major drawbacks of the controlled/living cationic polymerization based on stabilization of the carbocationic intermediates is slow propagation. Because the concentration of the active propagating species is very low because of the equilibrium between active species and dormant species, overall polymerization reactions are much slower than those without the equilibrium. Another important drawback of the controlled/living polymerization is the use of additives such as Lewis bases. Such additives should remain in polymer products and are generally rather difficult to be removed from the polymer products.

Recently, it has been demonstrated that good molecular weight control and molecular weight distribution control can be attained by using microreactor systems without stabilizing the carbocationic intermediates. The concept of this new technology (flow-microreactor-system-controlled polymerization) is described in the following section.

1.2.4.1 Concept of Flow-microreactor-system-controlled Polymerization Technology

Molecular weight and molecular weight distribution control in polymerization can be seen as control of competitive consecutive reactions as shown in Scheme 4, where A is an initiator and B is a monomer.

Scheme 4 Polymerization as a competitive consecutive reaction.

In the first step, A (initiator) reacts with B (monomer) to produce the first carbocationic intermediate P1. In the second step, P1 reacts with another monomer to produce the second carbocationic intermediate P2. Further reactions lead to polymer formation. If the initiation step to give P1 is faster than propagation steps, there is a chance to get good molecular weight control and molecular weight distribution control based on monomer/initiator ratios. However, this is the case only when the reactions are slower than the mixing and the reactions proceed in a homogeneous solution. If reactions are faster than mixing a significant amount of P2 (and Pn (n>2)) is formed before A is consumed even if $k_1 >> k_2$. In this case, it is meaningless to define concentration of A and B based on the total volume of the solution, because the solution is not homogeneous, and the product selectivity is not determined by kinetics. This problem is similar to the problem of disguised chemical selectivity for competitive consecutive reactions [101], this problem can be solved by extremely fast micromixing. In fact, the enhancement of product selectivity of competitive consecutive reactions [102] such as Friedel-Crafts
reactions [103, 104], [4+2] cycloaddition reactions [105, 106], lithiation of dibromobiphenyls [107-110], iodination of aromatic compounds [111, 112], and the reaction of a Grignard reagent with B(OMe)\(_3\) [113] by virtue of extremely fast micromixing using flow microreactor systems has been reported. These successful results suggest the possibility of molecular weight and molecular weight distribution control by using flow microreactor systems.

In flow-microreactor-system-controlled polymerization extremely fast mixing of an initiator solution and a monomer solution is achieved by extremely fast mixing leading to selective initiation. Propagation proceeds based on the monomer/initiator ratio under homogeneous conditions, and therefore polymers are obtained with good molecular weight and molecular weight distribution control. Minimizing local deviation of temperature by fast heat transfer through the wall of a microreactor by virtue of high surface-to-volume ratios is also important because polymerization processes are usually highly exothermic.

1.2.4.2 “Cation Pool” Initiated Polymerization of Vinyl Ethers Using Flow Microreactor Systems

Flow-microreactor-system-controlled cationic polymerization requires extremely reactive initiators, and highly reactive organic cations serve as effective initiators for this method. Usually organic cations such as carbenium ions andonium ions are generated by an acid-promoted reversible process from their precursors. Yoshida et al. have developed the “cation pool” method [114, 115] in which organic cations are generated irreversibly by low temperature electrolysis and are accumulated in relatively high concentration in the absence of nucleophiles. N-Acylium ions [116], alkoxy-carbenium ions [117-122], diarylcarbenium ions [123, 124], and glycosyl triflate [125] have been generated and accumulated by this method. Yoshida et al. reported that N-acyliminium ion pools serve as extremely reactive initiators for cationic polymerization in flow microreactors.

A pool of an N-acyliminium ion is generated by low temperature electrochemical oxidation of its precursor, a \(\alpha\)-silyl-substituted N-acylamine. Cationic polymerization of butyl vinyl ether (NBVE) using the N-acyliminium ion as an initiator in a conventional batch reactor gives the polymer in a quantitative yield after quenching with \(i\)-Pr\(_2\)NH/CH\(_2\)Cl\(_2\), but the molecular weight distribution is broad (M\(_n\) = 5700, M\(_w\)/M\(_n\) = 2.56). The reverse addition sequence (the N-acyliminium ion to the monomer) gives rise to a similar molecular weight distribution (quantitative yield, M\(_n\) = 13100, M\(_w\)/M\(_n\) = 2.25). The simultaneous addition of a monomer solution and a solution of the N-acyliminium ion does not improve the molecular weight distribution control (quantitative yield, M\(_n\) = 24500, M\(_w\)/M\(_n\) = 2.43). The molecular weight, however, strongly depends upon the method of mixing because the rate of the polymerization is so fast. In contrast, the use of a flow microreactor system consisting of two micromixers (M\(_1\) and M\(_2\)) and two microtube reactors (R\(_1\) and R\(_2\)) shown in Figure 5 leads to excellent control of molecular weight and its distribution (M\(_n\) = 6700, M\(_w\)/M\(_n\) = 1.14)
Solutions of \(N\)-acyliminium ion [127] and NBVE are introduced to \(M1\) by the syringe pump technique at -78 \(^{\circ}\)C. Then, the reaction mixture is introduced to a microtube reactor (\(R1\)) (\(\phi = 1.0\) mm, 10 cm), in which the polymerization takes place. In the final stage, \(i\)-Pr\(_2\)NH/CH\(_2\)Cl\(_2\) is introduced at \(M2\) to quench the polymerization. The polymerization takes place quite effectively and is complete within the residence time of 0.05 s to give the polymer with narrow molecular weight distribution (\(Mn = 6700, Mw/Mn = 1.14\)). The molecular weight can be controlled by changing the monomer/initiator ratio. The molecular weight (\(Mn\)) increases linearly with the amount of NBVE, indicating that chain transfer reaction does not play significant roles in this system.

**Figure 5** Flow microreactor system for polymerization of vinyl ether initiated by \(N\)-acyliminium ion (cation pool). \(M1, M2\): micromixer; \(R1, R2\): microtube reactor.

The effect of the flow rate on molecular weight distribution (Table 1, runs 4-6) indicates the importance of mixing, because it is known that mixing efficiency decreases with a decrease in the flow rate in the micromixer [128]. Reaction temperature is also important for controlling molecular weight distribution. \(Mw/Mn\) increases with an increase in the temperature (runs 4, 7, 8, and 9). Anyway, a high level of molecular weight control can be achieved by control of the initiation process by fast micromixing. Precise control of polymerization temperature also seems to be responsible for the remarkable control.

**Table 1** Cationic polymerization of NBVE initiated by \(N\)-acyliminium ion using a flow microreactor system

<table>
<thead>
<tr>
<th>run</th>
<th>monomer (equiv)</th>
<th>flow rate (mL/min)</th>
<th>Temperature ((^{\circ})C)</th>
<th>(Mn)</th>
<th>(Mw/Mn)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>10</td>
<td>5.0</td>
<td>-78</td>
<td>1500</td>
<td>1.40</td>
</tr>
<tr>
<td>2</td>
<td>25</td>
<td>5.0</td>
<td>-78</td>
<td>2900</td>
<td>1.26</td>
</tr>
<tr>
<td>3</td>
<td>35</td>
<td>5.0</td>
<td>-78</td>
<td>4400</td>
<td>1.17</td>
</tr>
<tr>
<td>4</td>
<td>50</td>
<td>5.0</td>
<td>-78</td>
<td>6700</td>
<td>1.14</td>
</tr>
<tr>
<td>5</td>
<td>50</td>
<td>3.0</td>
<td>-78</td>
<td>5600</td>
<td>1.35</td>
</tr>
<tr>
<td>6</td>
<td>50</td>
<td>1.0</td>
<td>-78</td>
<td>6200</td>
<td>2.84</td>
</tr>
<tr>
<td>7</td>
<td>50</td>
<td>5.0</td>
<td>-48</td>
<td>8200</td>
<td>1.30</td>
</tr>
<tr>
<td>8</td>
<td>50</td>
<td>5.0</td>
<td>-27</td>
<td>5500</td>
<td>1.34</td>
</tr>
<tr>
<td>9</td>
<td>50</td>
<td>5.0</td>
<td>0</td>
<td>6500</td>
<td>1.61</td>
</tr>
</tbody>
</table>
The polymer end is living within the residence time of 0.5 s at -78 °C, and can be effectively trapped by allyltrimethylsilane. Moreover, the present “cation pool” initiated polymerization using a microreactor system can be applied to other vinyl ethers such as isobutyl vinyl ether (IBVE) and tert-butyl vinyl ether (TBVE) to obtain the corresponding polymers (Mw/Mn = 1.12 (IBVE), Mw/Mn = 1.50 (TBVE)), though the corresponding polymerization using batch macro reactors results in much poorer molecular weight distribution control (Mw/Mn = 4.31 (IBVE), Mw/Mn = 2.29 (TBVE)).

1.2.4.3 Brønsted Acid Initiated Polymerization of Vinyl Ethers Using Flow Microreactor Systems

Proton addition is one of the most simple and straightforward methods for the initiation of cationic polymerization. Brønsted acids are effective for this purpose. However, if we use a weak Brønsted acid, i.e. a conjugate acid of a strong nucleophilic anion, the addition of a Lewis acid is necessary to establish a reversible activation of a covalent end group for effective propagation. On the other hand, if we use a strong Brønsted acid, i.e. a conjugate acid of an extremely weak nucleophilic anion, the addition of a Lewis base is required to stabilize the carbocationic propagating polymer ends [129]. In the absence of a Lewis base, highly ionic polymer ends are too reactive, and participate in transfer reactions by loss of β-protons leading to a very broad molecular weight distribution.

Trifluoromethanesulfonic acid (TfOH) is an effective initiator for cationic polymerization. For example, TfOH-initiated polymerization of isobutyl vinyl ether (IBVE) in 1,2-dichloroethane using a macro batch reactor [130] is complete within 10 s at -25 °C. The molecular weight distribution is, however, rather broad and Mw/Mn ranges from 2.73 to 4.71, presumably because of chain transfer reactions due to high reactivity of polymer ends. By employing flow microreactor systems consisting of a T-shaped micromixer and a microtube reactor, however, cationic polymerization using a strong Brønsted acid such as TfOH can be accomplished in a highly controlled manner without adding a Lewis base (Figure 6) [131]. The polymerization is complete within the residence time of 0.37-1.5 s at -25 °C (almost quantitative yields). The degree of molecular weight distribution control strongly depends on the inner diameter of the mixer and the flow rate as depicted in Table 2. Mw/Mn decreased with a decrease in the mixer inner diameter, presumably because faster mixing is achieved by a mixer of smaller diameter. Mw/Mn also decreases with an increase in the flow rate, probably because the increase of the flow rate enhanced the mixing efficiency. High level of molecular weight distribution control can be attained even at -25 °C. It is important to note that very low temperatures such as -78 °C, which might be an obstacle to industrial-scale applications, are not required.
Table 2  TfOH initiated polymerization of isobutyl vinyl ether (IBVE) in microreactor system.

<table>
<thead>
<tr>
<th>T-shaped inner diameter (µm)</th>
<th>flow rate (mL/min)</th>
<th>Mn b</th>
<th>Mw/Mn b</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>NBVE</td>
<td>TIOH</td>
<td>2900</td>
</tr>
<tr>
<td>250</td>
<td>2</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>3</td>
<td>3</td>
<td>2400</td>
</tr>
<tr>
<td>4</td>
<td>4</td>
<td>4</td>
<td>1600</td>
</tr>
<tr>
<td>5</td>
<td>5</td>
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<td>1500</td>
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<td>6</td>
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<td>7</td>
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<td>8</td>
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<td>1500</td>
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<td>6</td>
<td>1500</td>
</tr>
<tr>
<td>7</td>
<td>7</td>
<td>7</td>
<td>1400</td>
</tr>
<tr>
<td>800</td>
<td>6</td>
<td>6</td>
<td>2500</td>
</tr>
</tbody>
</table>

One of the advantages of controlled/living polymerization is that the method allows flexible synthesis of structurally defined block copolymers composed of different monomers, offering greater opportunities for synthesis of organic materials with interesting properties.

An example of microreactor systems for block copolymerization is shown in Figure 7. The first monomer IBVE is mixed with TfOH in the first micromixer (M1). Introduction of the second monomer (NBVE or EVE) at the second micromixer M2 results in the formation of the polymer of higher molecular weight with narrow molecular weight distribution [132]. Block copolymerization can be carried out with any combination and with either order of monomer addition as shown in Table 3, demonstrating that the present method serves as a flexible method for the synthesis of block copolymers. Therefore, flow-microreactor-system-controlled polymerization serves as a powerful method for synthesis of structurally well-defined polymers and copolymers in industry.
Figure 7 Flow microreactor system for block copolymerization of vinyl ether initiated by TfOH. M1, M2: micromixer; R1, R2: microtube reactor.

Table 3 Block polymerization of vinyl ether initiated by TfOH using the flow microreactor system.

<table>
<thead>
<tr>
<th>monomer-1</th>
<th>monomer-2</th>
<th>Mn $^b$</th>
<th>Mw/Mn $^b$</th>
</tr>
</thead>
<tbody>
<tr>
<td>IBVE</td>
<td>-</td>
<td>1500</td>
<td>1.18</td>
</tr>
<tr>
<td>IBVE</td>
<td>NBVE</td>
<td>2300</td>
<td>1.43</td>
</tr>
<tr>
<td>IBVE</td>
<td>EVE</td>
<td>2400</td>
<td>1.54</td>
</tr>
<tr>
<td>NBVE</td>
<td>-</td>
<td>1000</td>
<td>1.24</td>
</tr>
<tr>
<td>NBVE</td>
<td>IBVE</td>
<td>1700</td>
<td>1.45</td>
</tr>
<tr>
<td>NBVE</td>
<td>EVE</td>
<td>1900</td>
<td>1.55</td>
</tr>
<tr>
<td>EVE</td>
<td>-</td>
<td>860</td>
<td>1.16</td>
</tr>
<tr>
<td>EVE</td>
<td>IBVE</td>
<td>2100</td>
<td>1.54</td>
</tr>
<tr>
<td>EVE</td>
<td>NBVE</td>
<td>2100</td>
<td>1.41</td>
</tr>
</tbody>
</table>

1.2.5 Cationic Polymerization of Diisopropenylbenzenes Using Flow Microreactor Systems

Polyindanes have received significant research interest because of their remarkable thermal resistance. Extensive work on the synthesis of 1,1,3-trimethyl substituted polyindanes by cationic polymerization of 1,4-diisopropenylbenzenes and their thermal properties has done by Nuyken and coworkers [133, 134]. The polymers of high indane-unit content serve as useful materials having high thermal resistance and low dielectric constants. The use of flow microreactor systems is effective for increasing the indane-unit content. In fact, indane-unit content (99%) was much higher in the flow microreactor system than that (71%) for the batch macro system under similar conditions (70 °C, [1,4-diisopropenylbenzene] = 0.1 M, [TfOH] = 2 M) (Figure 8) [135]. The characteristic features of flow microreactor systems including fast mixing and uniformity of the temperature seem to be responsible.
1.3 Anionic Polymerization

1.3.1 Controlled/living Anionic Polymerization of Vinyl Monomers

Living anionic polymerizations have received significant attentions since Michael Szwarc’s first report in the 1950s [136]. Anionic polymerization [137-141] serves as an important and powerful method for macromolecular engineering permitting the preparation of highly defined block copolymers, star polymers, and further complex architectures, because the anionic polymer ends are living even in the absence of a capping agent [142, 143]. In fact, the anionic growing polymer ends (usually organolithium species) can be utilized for end-functionalization reactions with various electrophiles and block copolymerization with other monomers. Major drawbacks of conventional anionic polymerization in polar solvents in batch macro reactors include the requirement of low temperatures, such as -78 °C [144]. Such a requirement causes severe limitations in the use of this highly useful polymerization in industry. Using nonpolar solvents, the polymerization can be conducted at higher temperatures, but much longer reaction time is needed for completion. It is also problematic that frequently the use of individually manufactured laboratory equipment is necessary.

The kinetic studies on anionic polymerization in a continuous flow mode have also been reported by Szwarc and Schulz et al [145, 146], and Lochmann and Muller et al [147, 148]. However, preparative anionic polymerizations in continuous flow mode have not been studied until recently.

1.3.2 Controlled/living Anionic Polymerization of Styrenes

Anionic polymerization of styrenes is a highly useful technique for the synthesis of polystyrenes with precisely adjustable molecular weights and molecular weight distributions and is applied for the synthesis of structurally well-defined polymers such as end functionalized polymers and block copolymers.

1.3.2.1 Controlled/living Anionic Polymerization of Styrenes in Polar Solvent Using Flow Microreactor Systems [149]

In a conventional anionic polymerization of styrenes in polar solvents in a batch macro reactor, major drawbacks include the requirement of low temperature such as -78 °C. In contrast, Nagaki and Yoshida reported that controlled anionic polymerization of styrene can be conducted under easily accessible conditions such as 0 °C in a polar solvent using a flow microreactor to obtain the polystyrene with narrower molecular weight distribution (Mn = 1200 ~ 20000, Mw/Mn = 1.09 ~ 1.13) (Figure 9) [150]. Moreover, the molecular weight can be easily controlled by changing the flow rates of monomer and initiator solutions. Furthermore, these methods can be applied to styrene derivatives having silyl, methoxy, alkynyl, and alkylthio groups on the benzene ring. Löwe and Frey also reported the anionic polymerization of styrene at 20 °C using a flow
microreactor [151]. Polystyrenes in a broad range of molecular weight with the narrow molecular weight distribution can be obtained within several seconds (Mn = 1700 ~ 70000, Mw/Mn = 1.09 ~ 1.41). It should be noted that strict dryness of the apparatus and high vacuum techniques is needed in the classical batch methods but that such experimental effort can be significantly reduced by using flow microreactors. Residual impurities and moisture can be removed by purging the reactor with solutions of a monomer and an initiator before a solution of a desired polymer product is collected at the outlet of flow microreactors.

Figure 9 A flow microreactor system for anionic polymerization of styrene in THF. M: T-shaped micromixer; R: microtube reactor.

Integration of chemical reactions enhances the power and speed of organic and polymer synthesis, and recently it has been recognized that flow microreactors enable space-integration of reactions [152-157]. On the basis of livingness of the polymer end, structurally well-defined polymers such as end-functionalized polymers and block copolymers can be synthesized using integrated flow microreactor systems consisting of two micromixers and two microtube reactors. For example, functionalization of a living polymer end using chlorosilanes such as chlorotrimethylsilane and chlorodialylvinylsilane is effectively achieved by using integrated flow microreactor systems to obtain polystyrenes bearing the silyl group at the terminal. Block copolymerization can also be achieved using the integrated flow microreactor system at 0 and 24 °C to obtain structurally defined block copolymers composed of two different styrenes in quantitative yields (Figure 10).

Figure 10 An integrated flow microreactor system for anionic block copolymerization of styrenes in THF. M1, M2: micromixer; R1, R2: microtube reactor.
End-functionalization with epoxides is also popular, since epoxides high have reactivity toward nucleophiles by virtue of ring strain. Use of functionalized epoxides enables a further transformation after deprotection. For example, polymerization of styrene followed by end-functionalization with the various glycidyl ethers having acetal structures such as ethoxy ethyl glycidyl ether (EEGE), 1,2-isopropylidene glyceryl glycidyl ether (IGG), and trans-2-phenyl-1,3-dioxane glycidyl ether (PDGE) can be accomplished using a flow microreactor system (Figure 11) [158]. The acetal and ketal protecting groups in the glycidyl ethers are stable toward the highly reactive carbanionic living polymer ends but they can be easily cleaved under acidic conditions to afford multihydroxyl end-functionalized polymers (Figure 12).

**Figure 11** A flow microreactor system for anionic polymerization of styrene in THF initiated by s-BuLi and subsequent functionalization reaction with epoxides. M1, M2: micromixer; R1, R2: microtube reactor.

**Figure 12** Synthesis of multihydroxyl end-functionalized polystyrenes

The syntheses of various branched polymers with complex architectures such as star polymers and dendrimer-like star-branched polymers based on living anionic polymerization have been studied extensively. These branched polymers have attracted much attention from theoretical, synthetic, and practical viewpoints.
because of their unique and interesting properties in solution, melt, and solid states. Block copolymers having different polymer chains on a core are especially interesting. To synthesize such a structure, selective 1:1 reaction of a living polymer chain and a poly-functional core molecule is essential in the first step. In a conventional batch macro reactor, an excess amount of poly-functional core should be used to obtain the monosubstituted compound selectively [159-161]. This requirement is problematic because an excess amount of functional core should remain unchanged in the first step, and therefore, it should be removed before proceeding to the second step. The use of a flow microreactor system serves as a powerful method for solving this problem (disguised chemical selectivity [102-113]). As shown in Figure 13, the end functionalization with one equivalent of dichlorodimethylsilane leads to selective formation of a product having a single polymer chain on silicon (Mn = 1400, Mw/Mn = 1.13), although use of a batch macro reactor leads to lower controllability (Mn = 1300, Mw/Mn = 1.21). Extremely fast 1:1 micromixing of the living polymer chain and dichlorodimethylsilane enables the selective introduction of a single polymer chain into silicon. Therefore, the subsequent reaction with another living polymer chain using an integrated flow microreactor system gives block copolymers having two different polymer chains on a silicon core. The chlorosilane having a single polymer chain can be used for the subsequent reaction with alcohols and Grignard reagents.

**Figure 13** An integrated flow microreactor system for the synthesis of block copolymers having two different polymer chains on a silicon core (M1, M2, M3, M4: T-shaped micromixers; R1, R2, R3, R4: microtube reactors).

### 1.3.2.2 Controlled/living Anionic Polymerization of Styrenes in Nonpolar Solvent Using Flow Microreactor Systems

Anionic polymerization of styrenes can be conducted in nonpolar solvents at room temperature in a macro batch reactor. However, much longer reaction time is needed for completion. It is also important to carry out the polymerization with < 20% by volume styrene because the reactions with > 20% by volume styrene may result in rapid increase in reaction temperature, potentially
causing danger. The use of flow microreactor is effective for solving the problem. In fact, controlled anionic polymerization of styrene initiated by s-BuLi in cyclohexane as a nonpolar solvent can be conducted at 80 °C by using a flow microreactor system to obtain polystyrenes in quantitative yields within 1 ~ 5 min (Figure 14). The controlled polymerization of styrene in cyclohexane under high monomer concentration (25~42%, by volume styrene) at 60 °C can be achieved by using an aluminium-polyimide microfluidic device (Figure 15) [162]. Moreover, the molecular weight distribution of polymers is influence by the channel patterns ((straight, periodically pinched, obtuse zigzag, and acute zigzag channels).

Figure 14 A flow microreactor system for anionic polymerization of styrene in cyclohexane at 80 °C initiated by s-BuLi. M: T-shaped micromixer; R: microtube reactor.

Figure 15 A aluminium-polyimide microfluidic device for anionic polymerization of styrene initiated by s-BuLi in cyclohexane at high concentrations at 60 °C.

1.3.3 Controlled/living Anionic Polymerization of Alkyl Methacrylates Using Flow Microreactor Systems

Synthesis of poly(alkyl methacrylate)s with well-defined structures has received significant research interests as versatile materials such as plastics, adhesives and elastomers containing a number of different reactive functions. The use of living anionic polymerization of alkyl methacrylates is very popular for this purpose. Living anionic polymerization of alkyl methacrylates does not need a capping agent, and therefore is very fast compared to living radical polymerization. However, synthesis of poly(alkyl methacrylate)s via anionic polymerization using a conventional batch macro reactor should be carried out at low temperatures such as -78 °C to obtain polymers of narrow molecular-weight distribution [163, 164]. The requirement of such low temperatures causes several limitations in the use of this highly useful polymerization in industry. However, if the problem is solved, living anionic polymerization of alkyl methacrylate serves as a powerful method for synthesis of poly(alkyl methacrylate)s.
Controlled anionic polymerization of alkyl methacrylates initiated by 1,1-diphenylhexyllithium using a flow microreactor gives the corresponding poly(alkyl methacrylate)s with high level of control of molecular weight under easily accessible temperatures compared with conventional batch macro polymerization (-28 °C (methyl methacrylate (MMA): Mw/Mn = 1.16), 0 °C (butyl methacrylate (BuMA): Mw/Mn = 1.24), 24 °C (tert-butyl methacrylate (t-BuMA): Mw/Mn = 1.12). Precise control of the reaction temperature and fast mixing of a monomer and an initiator seem to be responsible (Figure 16) [165].

![Figure 16 A flow microreactor system for anionic polymerization of alkyl methacrylates initiated by 1,1-diphenylhexyllithium. M1, M2: T-shaped micromixer; R1, R2: microtube reactor.](image)

Livingness of the reactive carbanionic polymer end is important to produce end-functionalized polymers and block copolymers. Livingness of the polymer end in a flow microreactor system can be verified as shown in Figure 17. A solution of an alkyl methacrylate and that of 1,1-diphenylhexyllithium are mixed micromixer M1, and the polymerization is carried out in microtube reactor R1. Then, a solution of the same monomer is introduced at micromixer M2, which is connected to microtube reactor R2 where the sequential polymerization takes place. By changing the length of R1 with a fixed flow rate, the effect of the residence time in R1 can be examined. The Mn increases by the addition of the second monomer solution. However, an increase in the residence time in R1 causes an increase in the Mw/Mn, presumably because of decomposition of the polymer end (Figure 18). By choosing an appropriate residence time in R1 (MMA: 2.95 s, BuMA: 0.825 s), the sequential polymerization can be successfully carried out without significant decomposition of the living polymer end [166]. The polymer chain end is really living within such residence times. Moreover, the subsequent reaction of the living polymer end with a different alkyl methacrylate leads to the formation of a block copolymer having narrow molecular-weight distribution (Table 4).
Figure 17 An integrated flow microreactor system for the sequential anionic polymerization of alkyl methacrylates initiated by 1,1-diphenylhexyllithium. M1, M2: T-shaped micromixer; R1, R2: microtube reactor.

Figure 18 Size exclusion chromatography traces of polymers obtained in the integrated flow microreactor system. Effect of residence time on the molecular weight distribution. (a) methyl methacrylate - methyl methacrylate, (b) butyl methacrylate - butyl methacrylate.

Table 4 Block copolymerization of alkyl methacrylates initiated by 1,1-diphenylhexyllithium using the integrated flow microreactor system.

<table>
<thead>
<tr>
<th>monomer-1</th>
<th>monomer-2</th>
<th>Mn</th>
<th>Mw/Mn</th>
</tr>
</thead>
<tbody>
<tr>
<td>MMA</td>
<td>-</td>
<td>3200</td>
<td>1.17</td>
</tr>
<tr>
<td>MMA</td>
<td>MMA</td>
<td>8600</td>
<td>1.29</td>
</tr>
<tr>
<td>BuMA</td>
<td>-</td>
<td>4700</td>
<td>1.22</td>
</tr>
<tr>
<td>BuMA</td>
<td>BuMA</td>
<td>9900</td>
<td>1.39</td>
</tr>
<tr>
<td>BuMA</td>
<td>tBuMA</td>
<td>9000</td>
<td>1.31</td>
</tr>
<tr>
<td>tBuMA</td>
<td>-</td>
<td>5300</td>
<td>1.13</td>
</tr>
<tr>
<td>tBuMA</td>
<td>tBuMA</td>
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<td>1.13</td>
</tr>
<tr>
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</tr>
<tr>
<td>tBuMA</td>
<td>MMA</td>
<td>8400</td>
<td>1.15</td>
</tr>
</tbody>
</table>
1.3.4 Controlled/living Anionic Block Copolymerization of Styrenes and Alkyl Methacrylates Using Integrated Flow Microreactor Systems

As mentioned above, flow microreactors are effective for accomplishing the controlled anionic polymerization of styrenes and alkyl methacrylates. A high level of molecular weight distribution control can be achieved under easily accessible conditions such as 24 to -28 °C by virtue of the characteristic features of flow microreactors including fast mixing, fast heat transfer, and precise residence time control. Another advantage of flow-microreactor-controlled polymerization is easy modulation of flow microreactors to integrate polymerization reactions. In fact, by using integrated flow microreactor systems, the polystyrene living polymer end, which is produced by butyllithium initiated anionic polymerization of styrene can be effectively trapped with 1,1-diphenylethylene, and the resulting organolithium species can be used as a macro initiator for anionic polymerization of alkyl methacrylates. Therefore, styrene-alkyl methacrylate diblock copolymers can be synthesized with high level of molecular weight distribution control at easily accessible temperatures such as 24 to -28 °C (Figure 19) [167]. Moreover, triblock copolymers can be also synthesized by sequential introduction of styrene and two different alkyl methacrylates in a similar manner (styrene – tert-butyl methacrylate – methyl methacrylate triblock copolymer: Mn = 8800, Mw/Mn = 1.23, styrene – tert-butyl methacrylate – butyl methacrylate triblock copolymer: Mn = 9000, Mw/Mn = 1.35).

![Figure 19](image)

**Figure 19** Integrated flow microreactor system for the anionic block copolymerization of styrene and alkyl methacrylates initiated by s-BuLi. M1, M2, M3: T-shaped micromixer; R1, R2, R3: microtube reactor.

1.4 Radical Polymerization
1.4.1 Free Radical Polymerization Using Flow Microreactor Systems

Free radical polymerization is an important process for the industrial synthesis of macromolecules because free radicals are compatible with a wide variety of functional groups that do not survive in ionic and metal-catalyzed polymerization [168]. Because free-radical polymerization is usually highly
exothermic, the precise temperature control is essential for performing free radical polymerization in a highly controlled manner. Therefore, the polymerization in a conventional batch macro reactor often suffers from a low level of molecular weight distribution control because of inefficient heat removal and the lack of homogeneity of the reaction temperature. Therefore, the controllability of the reaction temperature is a major concern in free radical polymerization from viewpoint of industrial applications. As a matter of fact, heat removal capacity is often a limiting factor in polymerizations in batch macro reactors. Therefore, the advantage of flow microreactors for radical polymerization is obvious, because they enable fast heat transfer.

Iwasaki and Yoshida reported the free radical polymerization of various monomers using flow microreactor systems. Polymerization of butyl acrylate (BA) gives the polymer of much smaller Mw/Mn than that obtained with batch macro reactor because of a much higher heat-removal efficiency of the flow microreactor (Figure 20) [169]. For the polymerization of benzyl methacrylate ($n$BMA) and methyl methacrylate (MMA), the effect of the flow microreactor on molecular weight distribution control is smaller than the case of BA. For the polymerization of vinyl benzoate (VBz) and styrene, no appreciable effect is observed. The tendency indicates that the flow microreactor is quite effective for highly exothermic polymerization such as that of BA, but it is not so effective for less exothermic polymerization. Similar results on the polymerization of styrene have been reported by Leveson et al [170]. Moreover, a microchemical pilot plant for radical polymerization of MMA has been built by numbering up eight microtube reactors (vide infra) [171].

![Figure 20](image_url)  
**Figure 20** A flow microreactor system for the free radical polymerization initiated by AIBN and relative rate of the polymerization in the flow microreactor. M: T-shaped micromixer; R1, R2: microtube reactor.
Effects of mixing in radical polymerization of methyl methacrylate are interesting [172]. The use of 5 mm static mixer leads to fouling in the reactor. In contrast, the use of IMM micromixer with 36 lamellae of 25 µm thickness results in the reduction of the fouling. The numbering-up approach enables production of 2000 tons per year without the fouling problem [173].

Serra et al. studied the outstanding effect of mixing on conversion, molecular weight and polydispersity in free radical polymerizations of styrene by a numerical simulation using different micromixer geometries [174,175].

Latex production by miniemulsion polymerizations [176-178] in continuous tubular reactors has also been reported by McKenna et al [179].

1.4.2 Living Radical Polymerization Using Flow Microreactor Systems

The major drawback of free radical polymerization is the low controllability of macromolecular structures, and the polymers have broad molecular weight distribution. Living radical polymerization [180-184] has been developed to overcome the problem. Because more than 50% of polymers are produced via free radical processes in industry, living radical polymerization offers a possible solution to synthesizing polymers of narrow molecular weight distribution [185]. Various methods including the atom transfer radical polymerization (ATRP) [186-191], the reversible addition-fragmentation chain transfer radical polymerization (RAFT) [192-194], the nitroxide-mediated radical polymerization (NMP) [195-197], organoiodine-mediated radical polymerization (IRP) [198], cobalt-mediated polymerization [199] and organotellurium-, antimony-, or bismuth-mediated living radical polymerization (TERP) [200-205] have been developed for conducting living radical polymerization.

1.4.2.1 Atom Transfer Radical Polymerization (ATRP) Using Flow Microreactor Systems

Atom transfer radical polymerization (ATRP) [186-191] is one of the most extensively studied living radical polymerization because of its simplicity and broad applicability, and serves as a method for synthesizing previously inaccessible well-defined nanostructured polymeric materials [206]. One of the main drawbacks is low catalytic efficiency and deep color of the final product. In general, the use of 0.1% ~ 1% catalyst results in the formation of the colored polymer containing the residual catalyst. Therefore, after polymerization, additional purification by passing the solution through silica- or alumina-gel is needed to remove the catalyst from product. A possible solution to this problem is the use of supported catalysts.

A continuous column reactor packed with silica-gel supported CuBr-HMTETA catalyst for ATRP of methyl methacrylate (MMA) exhibits high catalyst retention, high catalytic activity, and good stability up to 100 h (Figure 21). Moreover, the product solution is colorless [207]. In addition, the molecular weight of the resulting PMMA can be controlled by simply changing the flow rate of MMA. The polymerization in the reactor is in first order with respect to
monomer concentration, typical for ATRP process. However, the polydispersity index of the resulting polymer is about 1.8, which is larger than that for the polymer prepared using the same supported catalyst in batch (about 1.1). This is presumably because back-mixing in the column and trapping of polymer chains in the silica gel pores take place. The system in which two column reactors are connected in series has been developed and is applied to block copolymerization of MMA with \( n \)-butyl methacrylate (\( n \)BMA) (Figure 22) [208].

**Figure 21** CuBr-HMTETA-silica gel packed column reactor for the atom transfer radical polymerization of MMA initiated by methyl \( \alpha \)-bromoacetate.

**Figure 22** CuBr-HMTETA-silica gel packed column reactor for the atom transfer radical block copolymerization of MMA with butyl methacrylate (\( n \)BMA) initiated by methyl \( \alpha \)-bromoacetate.

Photopolymerized microfluidic device consisting of two inlets, active mixing chamber containing a magnetic stir bar, a single reaction channel (500 × 600 µm), and one outlet has been used for ATRP reaction of 2-hydroxypropyl methacrylate (HPMA) initiated by methyl 2-bromopropionate (Figure 23) [209]. The molecular weight can be controlled by changing the residence time at the different flow rates (Table 5). The kinetics and polymer properties are similar to those for the batch reactions reported in the literature [210]. Furthermore, a block copolymer poly(ethylene oxide-block-2-hydroxypropyl methacrylate) (PEO-b-HPMA) can be prepared by the copolymerization of HPMA with a PEO macroinitiator prepared by reaction of the terminal hydroxy group of the PEO-OH with a two-fold molar excess of 2-bromo-isobutyryl bromide and triethylamine (Figure 24). The stoichiometry of the reactants can be easily changed by varying relative flow rates [211].
Table 5: Microchannel polymerization of HPMA at different pumping rates and initiator concentrations.

<table>
<thead>
<tr>
<th>[MBP]/[HPMA]</th>
<th>Pump rate (µl/h)</th>
<th>Residence time (h)</th>
<th>Conv. (%)</th>
<th>Mn</th>
<th>Mw/Mn</th>
</tr>
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<tbody>
<tr>
<td>1:40</td>
<td>50</td>
<td>2.00</td>
<td>92</td>
<td>6240</td>
<td>1.21</td>
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<tr>
<td></td>
<td>150</td>
<td>0.67</td>
<td>74</td>
<td>5560</td>
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<tr>
<td></td>
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<td>0.33</td>
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<td>29</td>
<td>3300</td>
<td>1.27</td>
</tr>
<tr>
<td></td>
<td>500</td>
<td>0.20</td>
<td>17</td>
<td>2770</td>
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</tr>
<tr>
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<td>2.00</td>
<td>62</td>
<td>12740</td>
<td>1.26</td>
</tr>
</tbody>
</table>

1.4.2.2 Reversible Addition-fragmentation Chain Transfer Radical (RAFT) Polymerization Using Flow Microreactor Systems

Reversible addition-fragmentation chain transfer radical polymerization (RAFT) [192-194] is known for its compatibility with a wide range of monomers, temperature, and impurities, as compared to other living radical polymerizations such as ATRP and NMP.

RAFT polymerization in miniemulsion has been carried out in a tubular reactor. Emulsion is prepared in a batch reactor using sodium dodecyl sulfate (SDS, surfactant), Triton X-405 (surfactant), styrene (monomer), hexadecane (costabilizer), and 1-phenylethyl phenyldithioacetate (PEPDTA, RAFT agent), and is pumped into a continuous sonication vessel to obtain miniemulsion (Figure 25) [212]. The miniemulsion is introduced to tube reactors where the polymerization takes place. The polymerization in the tube reactor behaved kinetically similar to the batch polymerization. However, the tubular reactor produces polymer with a slightly higher molecular weight distribution than that for polymer produced in a batch reactor, presumably because back-mixing or axial dispersion effects in the tubular reactor that would broaden the residence time distribution of particles within the reactor. The present systems can be extended for the synthesis of block copolymer of polystyrene and poly(butyl acrylate) [213].
RAFT polymerizations of *N*-isopropylacrylamide (NIPAM) as monomer and a trithiocarbonate as chain transfer agent has been carried out using a flow microreactor under homogeneous conditions (Figure 26) [214]. In flow process, an increase in the inner diameter of the tube results in slightly lower conversions and wider molecular weight distributions. Polymerization rates in a flow microreactor are considerably higher in comparison with those of batch polymerization because of uniform heating (Table 6).

Then, RAFT polymerizations of various monomers including acrylamides, acrylates, and vinyl acetate has been studied by Hornung *et al* [215]. Polymers of narrow molecular weight distribution and average molecular weights similar to those of batch polymerizations are obtained on a multigram scale.

### 1.4.2.3 Nitroxide-mediated Radical Polymerization (NMP) Using Flow Microreactor Systems
Nitroxide-mediated radical polymerization (NMP) [195-197] can be applied to a wider range of monomers such as styrenes, acrylates, acrylamides, acrylonitrile, and 1,3-dienes. Acyclic nitroxides such as 2,2,5-trimethyl-4-phenyl-3-azahexane-3-nitroxide (TIPNO) or \( N\)-tert-butyl-\(N\)-(1-diethyl phosphono-2,2-dimethylpropyl) nitroxide (DEPN) enables the reversible termination of the growing polymer chains.

Nitroxide-mediated radical polymerization (NMP) of styrene and \(n\)-butyl acrylate at 140 °C has been performed in a continuous-flow microtubular reactor (Figure 27) consisting of a stainless steel tube reactor and a back pressure cartridge [216]. In the case of styrene polymerization, there is no difference between batch reactors and flow microreactors. However, for \(n\)-butyl acrylate, a better control of the polymerization has been observed in the flow microreactor (Mw/Mn = 1.80 (batch reactor), Mw/Mn = 1.44 (flow microreactor)). Moreover, consumption of the monomer is much faster using the flow microreactor (Figure 28).

![Figure 27](image_url)

**Figure 27** A continuous flow microreactor system for nitroxide-mediated radical polymerization (NMP) of poly(styrene) or poly(\(n\)-butyl acrylate). R: microtube reactor.

![Figure 28](image_url)

**Figure 28** Conversion vs residence time of nitroxide-mediated radical polymerization of styrene and \(n\)-butyl acrylate without or with acetic anhydride.

Nitroxide-mediated radical polymerization of styrene in a miniemulsion can be also performed in a tubular reactor [217]. In the first step, a macroinitiator is prepared by bulk polymerization in a batch reactor and the subsequent miniemulsion polymerization is carried out in a tubular reactor. The
polymerization kinetics in the tubular reactor is similar to those in a batch reactor. It is also noteworthy that both the preparation of a macroinitiator and miniemulsion polymerization can be achieved in a continuous tubular reactor to obtain polystyrene-block-poly(n-butyl acrylate) diblock and polystyrene-block-poly(n-butyl acrylate)-block polystyrene triblock copolymers [218].

Continuous nitroxide-mediated block copolymerization of n-butyl acrylate (first monomer) and styrene (second monomer) can be performed using two serial 900 µm inner diameter stainless steel microtube reactors (Figure 29) [219]. For the second polymerization process, the influence of mixing was examined by changing micromixers. The use of a high pressure interdigital multilamination micromixer (HPIMM) provided by the IMM (Mainz, Germany), can significantly reduce the polydispersity index (Mw/Mn = 1.36, 120 °C) compared with that obtained in a batch macro reactor (Mw/Mn = 1.74, 120 °C). Efficient mixing of a viscous solution of poly(n-butyl acrylate) and a solution of styrene by virtue of small diffusion paths caused by small lamination widths seems to be responsible. Conversions, molecular weights, and molecular weight distributions are significantly influenced by the nature of micromixers [220]. The results obtained with three micromixers, i.e. two HPIMMs with different lamination widths and a slit plate micromixer (LH2) manufactured by Ehrfeld Mikrotechnik BTS (Wendelsheim, Germany) are summarized in Table 7. Molecular weight distribution and molecular weight strongly depend on the factor F (F = 1/N(Wc + Wl) (N: number of channels per inlet, Wc: channel width, Wl: slit or aperture width)). The relationship between molecular weight distribution or molecular weight and F is linear, enabling prediction of copolymer features.

<table>
<thead>
<tr>
<th>micromixer</th>
<th>HPIMM</th>
<th>HPIMM</th>
<th>LH2</th>
</tr>
</thead>
<tbody>
<tr>
<td>number of channels per inlet N</td>
<td>16</td>
<td>15</td>
<td>10</td>
</tr>
<tr>
<td>channel width Wc/µm</td>
<td>45</td>
<td>20</td>
<td>50</td>
</tr>
<tr>
<td>microstructure thickness/µm</td>
<td>250</td>
<td>100</td>
<td>300</td>
</tr>
<tr>
<td>Slit or aperture width Wl/µm</td>
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<td>60</td>
<td>50</td>
</tr>
<tr>
<td>Form factor F/mm³</td>
<td>0.59</td>
<td>0.83</td>
<td>1.0</td>
</tr>
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</table>
Figure 29 A continuous flow microreactor system for nitroxide-mediated radical block copolymerization of \( n \)-butyl acrylate and styrene. \( \textbf{R1}, \textbf{R2}: \) microtube reactor.

Very recently, Ryu and Studer \textit{et al.} also reported the use of highly sterically hindered amines [221,222] for the nitroxide-mediated radical block copolymerization (NMP) of styrene and butyl acrylate in a flow microreactor [223]. The polymerization in the flow microreactor is faster than that in the batch system, although polymers with slightly smaller molecular are obtained in flow.

In general, gel permeation chromatography (GPC) is used to characterized polymers. However, sample preparation and analysis requires long time. The integration of a flow microreactor with analytical devices to monitor the progress of polymerization in real time allows fast screening and optimization. For example, the continuous online rapid size-exclusion chromatography monitoring of polymerizations (CORSEMP) system has been developed by Serra and Hadzioannou \textit{et al} [224]. It consists of automatic samplings, dilutions, and injections every 12 min of polymers synthesized in continuous flow (Figure 30). In addition, this system also includes a viscometer, RI and UV detectors, and a GPC to determine the molecular weights and molecular weight distributions. The nitroxide-mediated block copolymerization of \( n \)-butyl acrylate and styrene can be monitored ‘near real-time’ with good accuracy and repeatability. Very recently, bifurcation analysis and grade transition dynamic optimization for NMP of styrene in a tubular reactor have been demonstrated by Flores-Tlacuahuac \textit{et al} [225, 226].
Figure 30 The continuous online rapid size-exclusion chromatography monitoring of polymerizations (CORSEMP) system for nitroxide-mediated radical block copolymerization of \( n \)-butyl acrylate and styrene.

1.5 Ring Opening Polymerization Using Flow Microreactor Systems

Ring-opening polymerization is categorized into chain-growth polymerization, in which the terminal end of a polymer acts as a reactive center [227]. The reaction of cyclic compounds with a polymer end causes the cleavage of the ring and repeating of this process lead to the formation of high-molecular-weight polymers. The synthesis of polyamides from lactams, polyesters from lactones, and polyethers from cyclic ethers has been widely used.

The most popular approach for the synthesis of polyamides from amino acids [228] is an amino acid \( N \)-carboxyanhydride (NCA) method (Figure 31) [229]. The initiation with a base such as tertiary amine followed by the reaction of another NCA molecule with the resulting activated NCA anion produces a dimer with an electrophilic \( N \)-acyl NCA end group and a nucleophilic carbamate group. Then, \( N \)-carbamic acid is detached as carbon dioxide during the propagation reaction. It is known that deprotonation of NCA and the attack of NCA anion are relatively fast [230]. Therefore, it is important to control these steps. However, in conventional batch systems, the polymerization control would be difficult due to local gradients of the concentrations. In contrast, flow microreactor systems enable such control (Figure 32) [231]. For example, polymerization of \( N \)-benzyloxy carbonyl-\( L \)-lysine using a flow microreactor system consisting of a polydimethylsiloxane-based multi-lamination mixer and a PTFE microtube leads to narrower molecular weight distribution (flow: Mw/Mn = ca 1.2, batch: Mw/Mn = 1.5), although molecular weights are similar (flow Mn = ca. 20,000, batch: Mn = ca. 18,000) [232]. The use of a simple T-shaped connector as a mixer leads to broader molecular weight distribution, indicating the importance of fast micromixing. The present flow microreactor system can be applied to polymerization and copolymerization of other amino acid NCAs to obtain
poly(Glu), copoly(Lys, Ala) and copoly(Lys, Leu) of narrower molecular weight distribution than those obtained from the batch reactor (Table 8).

![Figure 31](image1.png)

**Figure 31** Mechanism of polymerization of amino acid N-carboxyanhydride (NCA) initiated by a base.

![Figure 32](image2.png)

**Figure 32** A flow microreactor system consisting of a polydimethylsiloxane (PDMS) multi-layered laminar micromixer and PTFE microtubes for polymerization of amino acid N-carboxyanhydride (NCA) initiated by triethylamine.

<table>
<thead>
<tr>
<th>polymer</th>
<th>Mn</th>
<th>Mw/Mn</th>
</tr>
</thead>
<tbody>
<tr>
<td>Poly(Glu)</td>
<td>batch 40200</td>
<td>1.56</td>
</tr>
<tr>
<td></td>
<td>microreactor 40000</td>
<td>1.17</td>
</tr>
<tr>
<td>Poly(Lys, Ala)</td>
<td>batch 17700</td>
<td>2.56</td>
</tr>
<tr>
<td></td>
<td>microreactor 18800</td>
<td>1.64</td>
</tr>
<tr>
<td>Poly(Lys, Leu)</td>
<td>batch 18100</td>
<td>2.13</td>
</tr>
<tr>
<td></td>
<td>microreactor 19200</td>
<td>1.54</td>
</tr>
</tbody>
</table>

A silicon-glass based flow microreactor system that is suitable for long periods of use has been developed and applied to the polymerization of amino acid NCAs [233]. The flow microreactor exhibits excellent controllability of the molecular weight distribution. Moreover, a single flow microreactor can produce
100 mg/min of poly(Lys/Leu). It means that more than 200 g of poly(Lys/Leu) acids can be produced in 2 months.

Hyperbranched polymers have attracted much attention because these polymers exhibit different characteristic features, such as a lower viscosity, higher solubility, and higher amount of terminal groups, compared with those of the corresponding linear polymers. Such polymers can be prepared by ring-opening multibranching polymerization. Hyperbranched polyglycerols are popular because they have a large number of hydroxyl groups and exhibit excellent biocompatibility [234]. Continuous micromixer-assisted flow process for the synthesis of hyperbranched polyglycerol by ring-opening multibranching polymerization of glycidol was demonstrated by Wilms et al (Figure 33) [235]. The polymerization was carried out in the presence of trimethylol propane (TMP) as a multifunctional initiator and potassium methoxide necessary to deprotonation 10% of the hydroxyl groups to obtain well-defined hyperbranched polyglycerol with molecular weights up to 1,000 g/mol. However, higher flow rates results in the partial formation of high molecular weight probably because of the formation of “hot spots” where the polymerization proceeds much faster.

Figure 33 A flow microreactor for synthesis of hyperbranched polyglycerol by ring-opening multibranching polymerization of glycidol. M: micromixer.

The continuous-flow microwave-assisted polymerization of 2-ethyl-2-oxazoline was reported by Schubert et al [236]. The flow process overcomes the problems associated with scale-up of the batch process including safety issues.

1.6 Polycondensation Using Flow Microreactor Systems

Polycondensation reactions are important processes for the synthesis of polyesters, polycarbonates, polyamides, and polysiloxanes [237]. Polycondensation reaction is classified into a step-growth polymerization, which involves a stepwise covalent-bond-forming reaction between functional groups of two monomers, between a functional group of a monomer and a polymer end, or between functional groups of two polymer ends. Because the propagating polymers and monomer do not contain an active species, such as cations, radicals, or anions, the polymer chains reach moderately high molecular weight even at
very high conversion. Therefore, in principle, it is difficult to control molecular weight and molecular weight distributions precisely.

Polycondensation of 4,4’-oxydianiline (ODA) and isophthaloyl dichloride (IPA) followed by terminal modification has been carried out in a flow microreactor system (Figure 34) [238]. The polymerization in the flow microreactor is faster than that in the batch system. A higher mixing efficiency of monomer seems to be responsible for the faster reaction. It is also important to note that the molecular weight distribution of the polymer obtained in the flow is slightly narrower than that obtained in batch. Precise control of reaction temperature in the microreactor seems to be responsible.

Figure 34 A flow microreactor system for polycondensation of 4,4’-oxydianiline (ODA) and isophthaloyl dichloride (IPA) followed by terminal modification. M1, M2: micromixer; R1, R2: microtube reactor.

Dendrimers, which have highly branched 3D structure provide a high degree of surface functionality and versatility. Polycondensation reactions are widely used for synthesizing various dendrimers. The synthesis of polyamide dendrons (G1 Dendron, G2 Dendron) and a dendrimer (G1 Dendrimer) as shown in Figure 35 has been carried out using flow microreactors [239]. The use of flow microreactors enables significant reduction of reaction time. In addition, the reaction can be performed at a constant temperature such as 30 °C in flow, whereas low temperature such as 0 °C is necessary for the first mixing to avoid side reactions in batch. Moreover, polyamide dendrons can be deposited onto the functionalized glass surface through amide bond formation in flow.
Reactions of the convergent synthesis of polyamide dendrons and dendrimers.

By polycondensation of trifunctional silanols, poly(silsesquioxane)s, which emerge as materials for various applications ranging from low dielectric constant materials [240] to flame-retardant material [241], can be prepared. The polycondensation reaction of methyltrimethoxysilane (MTMS) and various trimethoxysilanes involving p-(chloremethyl)phenylethyltrimethoxysilane (1), dithiobenzoic acid benzyl-(4-ethyltrimethoxysilyl) ester (2), dithiobenzoic acid 1-ethylphenyl-4-(ethyltrimethoxysilyl) ester (3) and N,N-diethylthiocarbamoylethylphenyl(trimethoxy)silane (4) [242, 243] can be successfully achieved in a microreactor to obtain corresponding poly(silsesquioxane)s (Figure 36) [244]. The yields are significantly higher than those in batch. The polydispersity indexes can be smaller than 2, whereas they are usually between 2.2 and 3 in batch. Moreover, molecular weights, which range from Mn = 1900 to 11000, can be controlled by changing the residence time (Table 9).
Figure 36 A flow microreactor system for polycondensation reactions of trialkoxysilanes. M: micromixer; R: microtube reactor.

Table 9 Molecular weight and molecular weight distribution obtained in microreactor synthesis.

<table>
<thead>
<tr>
<th>Initiator</th>
<th>Temperature (°C)</th>
<th>Residence Time (min)</th>
<th>Mn</th>
<th>Mw/Mn</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0</td>
<td>0.22</td>
<td>1900</td>
<td>1.47</td>
</tr>
<tr>
<td></td>
<td>0</td>
<td>0.66</td>
<td>3450</td>
<td>1.49</td>
</tr>
<tr>
<td></td>
<td>0</td>
<td>1.32</td>
<td>4350</td>
<td>1.50</td>
</tr>
<tr>
<td></td>
<td>0</td>
<td>1.98</td>
<td>6250</td>
<td>1.54</td>
</tr>
<tr>
<td></td>
<td>0</td>
<td>2.64</td>
<td>8400</td>
<td>1.59</td>
</tr>
<tr>
<td>1</td>
<td>20</td>
<td>0.22</td>
<td>2700</td>
<td>1.71</td>
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<td>2.64</td>
<td>9550</td>
<td>1.93</td>
</tr>
<tr>
<td>2</td>
<td>0</td>
<td>0.22</td>
<td>2100</td>
<td>1.46</td>
</tr>
<tr>
<td></td>
<td>0</td>
<td>2.64</td>
<td>8200</td>
<td>1.69</td>
</tr>
<tr>
<td>2</td>
<td>20</td>
<td>0.22</td>
<td>3200</td>
<td>1.79</td>
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<td>20</td>
<td>2.64</td>
<td>9700</td>
<td>1.87</td>
</tr>
<tr>
<td>3</td>
<td>0</td>
<td>0.22</td>
<td>2200</td>
<td>1.53</td>
</tr>
<tr>
<td></td>
<td>0</td>
<td>2.64</td>
<td>8900</td>
<td>1.65</td>
</tr>
<tr>
<td>3</td>
<td>20</td>
<td>0.22</td>
<td>3400</td>
<td>1.71</td>
</tr>
<tr>
<td></td>
<td>20</td>
<td>2.64</td>
<td>10100</td>
<td>1.92</td>
</tr>
<tr>
<td>4</td>
<td>0</td>
<td>0.22</td>
<td>2250</td>
<td>1.39</td>
</tr>
<tr>
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<td>2.64</td>
<td>8100</td>
<td>1.61</td>
</tr>
<tr>
<td>4</td>
<td>20</td>
<td>0.22</td>
<td>3700</td>
<td>1.87</td>
</tr>
<tr>
<td></td>
<td>20</td>
<td>2.64</td>
<td>11000</td>
<td>1.95</td>
</tr>
</tbody>
</table>

1.7 Ziegler-Natta polymerization Using Flow Microreactor Systems

Ziegler-Natta polymerization [245, 246] is an important method of vinyl polymerization because it allows us to make polymers of specific tacticity. As reported by Santos and Metzger, Ziegler-Natta polymerization can be carried out in a flow microreactor system coupled directly to the ESI source of a Q-TOF mass spectrometer (Figure 37) [247]. In the first micromixer M1, a catalyst (Cp2ZrCl2/MAO) and a monomer solution are mixed continuously to initiate the polymerization. The polymerization occurs in a microtube reactor. The solution thus obtained is introduced to the second micromixer M2, where the polymerization is quenched by acetonitrile. The quenched solution is fed directly into the ESI source. The transient cationic species can be characterized by mass spectrometry. This is the first case where an alkyl zirconium cation intermediate in the homogeneous Ziegler-Natta polymerization of ethylene is detected directly.
1.8 Fabrication of Polymeric Structures inside Microchannels Using Fluid Flow

1.8.1 Polymerization on the Surface of Microchannels

Surface-initiated polymerization is an attractive method for fabricating microfluidic devices [248-251]. For example, Beers et al. reported surface-initiated atom transfer radical polymerization inside microchannels to produce flat gradient and patterned surfaces [252]. A solution of 2-hydroxyethyl methacrylate (HEMA) is introduced into the microchannel (300 mm × 8 mm × 4.5 cm), which was manufactured on a silicon wafer and functionalized with initiator-functionalized self-assembled monolayer (Figure 38). A polymer gradient is created inside the microchannel so that the thickness of the polymer brush decreases linearly from the inlet to the outlet.

Microfluidic system can be applied to generate complex gradient solutions [253-255]. Recently, the generation of the solution gradient in a microfluidic passive mixer [256] is used to synthesize a surface-grafted statistical-copolymer-brush composition gradient via surface-initiated atom transfer radical polymerization (Figure 39) [257]. Using surface-initiated polymerization, the monomer solution gradient of butyl methacrylate (nBMA) and 2-(N,N-dimethylamino)ethyl methacrylate (DMAEMA) is applied to synthesize a statistical-copolymer-brush composition gradient. Burdick et al. also reported a similar approach to fabricating photo-cross-linked hydrogels with gradients of immobilized molecules and crosslinking densities using a microfluidics/photopolymerization process [258].
Figure 39 A schematic drawing of the setup used for the patterning of a surface with a statistical-copolymer-brush gradient.

1.8.2 Fabrication of Polymer Membranes inside Microchannels

Characteristic features of controlled laminar flow in microfluidics devices have been utilized in many applications such as diffusion-based separation and detection solvent extraction, mixing and hydrodynamic focusing [259, 260].

A method of microfabrication based on multicomponent laminar flow inside microchannels was developed by Whitesides and Kenis et al. [261, 262]. Laminar streams of solutions enable a reaction at the interface between streams to make membranes inside the microchannel. For example, the reaction at the interface between two aqueous phases containing oppositely charged polymers (poly(sodium 4-styrenesulfonate) and hexadimethrine bromide) flowing laminarly in parallel produces a polymeric structure (membrane) deposited on glass at the laminar flow interface (Figure 40).

Figure 40 A polymeric structure deposited on glass at the laminar flow interface of two solutions of poly(sodium 4-styrenesulfonate) and hexadimethrine bromide.

Beer et al. demonstrated the formation of a laminar flow interface of immiscible liquids in a microchannel using partial chemical modification of the channel surface [263]. Synthesis of a polyamide nylon membrane by an interfacial
polycondensation reaction of adipoyl chloride in 1,2-dichloroethane and hexamethylenediamine in water can be achieved in the cross-junction of the microchannel. Kitamori et al. also demonstrated design and synthesis of nylon polyamide membrane structures by an interfacial polycondensation reaction of adipoyl chloride in 1,2-dichloroethane and hexamethylenediamine in water (Figure 41) [264]. Uozumi et al. also reported that catalytic membrane-installed microchannel devices prepared from linear polymer ligands and palladium complexes and applied to Suzuki-Miyaura coupling reaction, oxidative cyclization of alkenols, allylic arylation, and hydrodehalogenation [265-269]. In addition, the influence of microfluidic device geometry and flow rate on the membrane formation by the interfacial polymerization was also demonstrated by Gargiuli et al [270].

![Figure 41: Polymer membrane formation under organic/aqueous two-phase flow in an X-shaped microchannel.](image)

Single and parallel dual-membrane structures are successfully prepared by using multilayer flow such as organic/aqueous two layer flow and organic/aqueous/organic three-layer flow inside a microchannel, and this method can be applied to preparing surface modified polymer membranes (Figure 42). For example, horseradish peroxidase is immobilized on one side of the membrane surface, and this enzyme-modified membrane realizes substrate permeation and a subsequent reaction.
Figure 42 Channel patterns and cross-sectional views of the nylon membrane prepared inside a microchannel. (a) Single membrane formed under organic/aqueous two-layer flow, (b) parallel dual membranes formed under organic/aqueous/organic three-layer flow.

Protein-polymeric membrane in a microchannel is prepared by using a concentric laminar flow (Figure 43) [271]. Cross-linking condensation reaction of a cross-linked enzyme aggregate (CLEA) [272] with aldehyde groups, which react with amino groups of the enzyme in a concentric laminar flow results in the formation of the cylindrical enzyme-polymerized membrane on the inner wall of the microtube. The use of this technology for membrane formation in a microchannel can be extended to a broad range of functional proteins.

Figure 43 Schematic illustration of the procedure on preparation of enzyme-membrane in micro-tube.

1.9 Industrial Applications

It is easily anticipated that flow microreactors can enjoy industrial applications by virtue of inherent advantages based on microstructures and flow nature. Significant progress in flow-microreactor-system-controlled polymerization to obtain structurally well-defined polymers has already been
made to meet the demands of the chemical industry. No need of cryogenic conditions for anionic polymerization may enable commercial production. Some pilot plants have already been built and tested to examine the feasibility and durability of polymerization in flow microreactors. For example, a microchemical pilot plant for radical polymerization of MMA has been built by numbering up eight microtube reactors which can be operated continuously for 6 days to produce 4.0 kg of the polymer, indicating that flow microreactor systems can be used for industrial production of polymers (Figure 44) [171]. The information which has been accumulated in laboratories and pilot plants should profit the development of this field to realize commercial plants for making polymers in the future.

![Figure 44](image)

**Figure 44** A pilot plant for radical polymerization.

1.10 Conclusion

The examples shown in this review article demonstrate that a variety of methods for polymer synthesis have been developed in flow microreactors. The continuous flow synthesis enables serial combinatorial synthesis, in which a variety of polymers in a sequential way using a single flow reactor with flow switch. Space integration, which enables the synthesis of structurally well-defined polymers without isolating living polymer ends, also enhances the power and speed of polymer synthesis. Because several test plants for continuous production have already been built, there is no doubt that flow microreactors can contribute to polymer production in industry.

In conclusion, continuous flow polymer synthesis will be an indispensable technology for research in a laboratory and production in industry. Various methods for polymer synthesis by virtue of characteristic features of flow microreactors will be developed, and they will work together to help meet the demanding expectations in polymer chemistry in the future.

References


Because the yield of N-acyliminium ion from precursor is estimated as ca. 80% based on the reactions with various nucleophiles, 1.2 equiv of precursor is used when 1.0 equiv of N-acyliminium ion is needed for polymerization.


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