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Supercritical carbon dioxide systems for sustainable and efficient dissolution of solutes: a review

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Abstract

Green chemistry aims at replacing toxic solvents by safer solvents such as supercritical carbon dioxide (scCO_2), which displays zero surface tension, outstanding transport properties, high diffusivity, and tuned solubilization by changing the pressure and temperature, or adding cosolvents. Nonetheless, the practical application of $\, \text{scCO}_2$ -based technologies under supercritical conditions requires sophisticated systems for optimal operation. Here, we review $\rm scCO_2$ -based systems with focus on dyeing, extraction, chromatographic, and cleaning systems. All $\rm scCO_2$ -based systems consist of a CO_2 supply and pump part, a cosolvent providing part, and a parameters controlling part, yet there are different reaction vessels depending on solute types and practical functions. Developing more normalized, intelligentized, and durable $\sec O_2$ -based systems should render the $\sec O_2$ -based technologies be more economic, controllable, flexible, and suitable for wider applications.

Keywords Apparatus · CO₂ · Equipment · Green solvent · Reactor · Supercritical fluid

Introduction

Since the chemist Anastas (1999) introduced the concept of "green chemistry" in 1990s, avoiding the utilization of toxic reagents and solvents has become the tendency in the manufacture and chemical industries (Cao and Su 2021). Toxic solvents have been widely used in industrial processes despite the high risk of contaminating the environment (Anastas and Eghbali 2010; Aravindhan et al. 2017). However, the solvents cannot be absolutely avoided due to their vital role for dissolving components, promoting mass and heat transfer, and accelerating the reaction process (Mohammad and Inamuddin 2012; Cheng et al. 2023). Accordingly,

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there is an urgent need for more environmentally friendly and efficient solvents (Pacheco-Fernández and Pino 2019).

Supercritical carbon dioxide ($\sec O_2$) as a sustainable and efficient solvent with mild critical temperature of 304.1 K and critical pressure of 7.4 MPa has attracted increasing attention in recent years (Liu et al. 2006; Qi et al. 2018; Ramsey et al. 2009). ScCO₂ with low viscosity, zero surface tension, high gas-like difusivity, and high liquid-like density has becoming an attractive solvent that can be widely used in various felds (Bai et al. 2019; Lebedev et al. 2015). Moreover, $\sec 0₂$ can be completely miscible with the gas solutes, while the solubility of the liquid and solid solutes in $\sec 0₂$ can be adjusted by fexibly varying the temperature and pressure as well as adding cosolvents during the scCO_2 -assisted reaction process (Chauvet et al. 2017; Kang et al. 2020). The use of $\sec O_2$ as the solvent also has the advantages of easy recompression, complete separation with the solutes by simply depressurization, and recycled utilization of $CO₂$ (Machado et al. 2013; Sihvonen et al. 1999). ScCO₂ has been widely applied for dissolving various types of solutes, such as dyes, target solutes in other solvents, component solutes to be separated or purifed, and contaminants to be removed, among which especially for non-polar solutes, based on different $\sec O_2$ -based technologies, i.e., dyeing technology, extraction technology, chromatography technology, and cleaning technology, respectively (Aslanidou et al. 2013; Friedrich and Pryde 1984; Rezaei and Temelli 2000). It is also worth noting that $\sec O_2$ -based technologies are limited since $\sec O_2$ lacks polarity and capacity to form specific solvent–solute interaction, and several studies have found that the addition of a small amount of proper cosolvent can efectively improve its solvent power for part of non-polar solutes (Li et al. 2003; Ting et al. 1993; Güçlü-Üstündağ et al. 2005; Zulkafi et al. 2014; Patil et al. 2018).

Recent reviews on $\sec O_2$ -based technologies mainly focus on the reactions that performed assisted by $\sec O_2$ as the solvent. Gonzalez et al. (2014) comprehensively presented the perspectives of $scCO₂/green$ solvent biphasic systems to offer a good insight into the practical usages and potential applications of the $\frac{\text{cCO}}{\text{g}$ green solvent biphasic systems. Abou et al. (2018) highlighted the great role of scCO₂-based technology in textile dyeing processes, which shows a detailed introduction about $\sec O_2$ -assisted dyeing applications. Moreover, Sharif et al. (2014) introduced the common experimental designs using in the supercritical fuid extraction process and further analyzed the infuential factors. Speybrouck et al. (2016) systematically summarized the advantages of using $\sec O_2$ as the mobile phase, such as short equilibration time, high flow rate, and very low solvent consumption, for analytical and preparative chiral supercritical fuid chromatography. Additionally, Meziani et al. (2017) comprehensively elaborated novel scCO_2 -assisted cleaning applications in the semiconductor cleaning process and introduced the recent advances of the scCO_2 -assisted cleaning applications in the emerging technologies. All of these offer a valuable view in the aspects of $\sec O_2$ as a green solvent for flexible and efficient dissolution of various types of solutes. However, it is worth noting that the practical application of $\sec O_2$ -based technologies under supercritical reaction conditions requires sophisticated systems for optimal operation. Although many diferent types of scCO_{2} -based systems have been built, there is actually no reported review detailing various $\sec O_2$ -based systems used for different $secCO₂$ -based applications.

Herein, compared to most reviews that focused on the reactions that performed assisted by $\sec O_2$ as the solvent, this review summarizes the current research status of the $\sec CO_2$ -based systems for flexible and efficient dissolution of diferent types of solutes, such as dyes, target solutes in other solvents, component solutes to be separated or purifed, and contaminants to be removed, based on different scCO_{2} -based technologies and related systems, i.e., dyeing system, extraction system, chromatograph system, and cleaning system. Additionally, this review also introduces the similarity and diference among the systems which matched with diferent practical production requirements in each application, and further successively points out the perspectives and valuable directions for future development in both fundamental research and commercial applications of the scCO_2 -based systems. The outline of this review is shown in Fig. 1.

Dyeing systems

The traditional printing and dyeing technology are mainly based on water as the solvent, which has the disadvantages of water consumption, energy consumption, and environmental pollution (Banchero 2013). As a consequence, Saus et al. (1993) invented the supercritical fuid dyeing technology. The supercritical fuid dyeing technology has shown excellent characteristics in overcoming the waste of water resources and serious waste pollution, and the most prominent supercritical fuid used for supercritical fuid dyeing is $\sec CO_2$ (Abou et al. 2015; Banchero 2013; Jaxel et al. 2020; Kazarian et al. 1997). ScCO₂ shows extremely high solubility and affinity of dyes; as a result, it could be used as medium to carry dyes to dye the fabrics. After dyeing, $CO₂$ and the unused dyes will be separated and recycled, which avoided the pollution at the source (Abou et al. 2018a, 2020; Banchero 2020).

Moreover, the $\sec O_2$ dyeing technology does not have the washing and drying process that existed in the traditional dyeing technology, and not need to add dispersant, leveling agent, retarder, and other auxiliary agents during the dyeing process, which not only improves the production efficiency, but also reduces energy consumption, thus improving the economic efficiency (Abou et al. 2021; Agrawal 2015; Goñi et al. 2021). Therefore, developing the application and corresponding systems of $\sec O_2$ dyeing technology will inject vitality into the printing and dyeing industry. The critical factors of $\sec 0₂$ dyeing systems are reasonable design of the high-pressure dyeing vessel that can satisfy the dye homogenously dispersed and dissolved in $\sec O_2$, and then uniformly absorbed and difused onto the fabric to realize the dyeing requirements. Meanwhile, separators should be designed for removing excess dye and residue. Actually, there are two types of $\sec O_2$ dyeing methods, namely static dyeing and dynamic dyeing. The systems used in the above two scCO₂ dyeing methods are generally divided into static dyeing system, dynamic beam dyeing system, and dynamic jig dyeing system, respectively, which will be summarized and reviewed separately in the following.

Static dyeing system

Static $\sec O_2$ dyeing systems are typically applied in laboratory-scale $\sec O_2$ dyeing process. Static $\sec O_2$ dyeing systems are simply composed of $CO₂$ supply part, pressure- and temperature-controlling part, high-pressure dyeing vessel, and separator. In the traditional static $\sec O_2$ dyeing system, the fabric, dye, and cosolvent are usually placed together

Fig. 1 Sustainable-solvent-supercritical carbon dioxide (scCO₂)-based systems for flexible and efficient dissolution of solutes. This report reviews scCO₂-based systems for sustainable and efficient dissolution

of different types of solutes, based on different scCO₂-based technologies and associated systems, i.e., dyeing system, extraction system, chromatograph, and cleaning system. BPR: backpressure regulator

into the dyeing vessel, and then, $CO₂$ will be injected by the high-pressure pump. By adjusting the temperature and pressure to achieve the $\sec O_2$ dyeing conditions, the dye will be continuously dissolved and difused onto the fabric to complete the dyeing process. The core of the traditional $\sec CO_2$ dyeing system is the agitation device for promoting the movement of the dyeing solution in order to enhance the dyeing uniformity.

The typical static scCO₂ dyeing system is shown in Fig. 2 (Penthala et al. 2019). Here, the fabric, dye, and cosolvent are usually placed together into the dyeing vessel, and then, $CO₂$ will be injected by the high-pressure pump. By adjusting the temperature and pressure to achieve the $\sec O_2$ dyeing conditions, the dye will be continuously dissolved and difused onto the fabric to complete the dyeing process. The dyeing vessel usually equips with an agitation device to promote the movement of the dyeing solution in order to enhance the uniformity. After the reaction, the excess dye will be collected into the residual dye trap and $CO₂$ will also be recycled. Moreover, horizontal stirring of the fabric can overcome the infuence of gravity during the dyeing process than the vertical stirring and thus further improve the color presentation uniformity; also, the small volume and easy operation of the systems are also important factors for the practical applications of the $\mathrm{s}\mathrm{c}\mathrm{C}\mathrm{O}_2$ dyeing systems. Aiming at the issue, Mo et al. (2020) designed an integrated static $\rm scCO$ ₂ desizing system (Fig. S1), where the high-pressure autoclave is frstly pre-cooled down and then the fabric will be put in the autoclave. Subsequently, the autoclave is heated by the oil bath until it reached the target temperature and stirred continuously until realizing the dyeing process. Finally, the cooling water circulation will be turned on until the pressure of the autoclave decreases to the atmosphere pressure. Compared with the traditional static $\sec O_2$ dyeing system, this system is equipped with an integrated heating and cooling tube, which can control the temperature and pressure more accurately. There are also two *Z*-shaped stirring shafts at both ends of the autoclave, and therefore, it is more conducive to promote the movement of dyeing solution than the vertical stirring and thus to make the dyeing process more homogeneous and efficiency.

The static $\sec O_2$ dyeing system has been widely used in laboratory-scale dyeing experiments to study the efect of parameters, such as temperature, pressure, and time, on the dyeing of different fabrics during the scCO_2 dyeing process. However, due to the absence of fuid circulation during the

Fig. 2 Typical static supercritical carbon dioxide (scCO_2) dyeing system: 1. $CO₂$ cylinder, 2. circulated cooling bath, 3. high-pressure pump, 4. heating bath, 5. temperature controller, 6. dyeing vessel, 7. stirrer, 8. pressure regulator, and 9. separator. The fabric, dye, and cosolvent are usually placed together into the dyeing vessel, and then,

static $\sec 0$, dyeing process, the diffusion rate of dyeing solution and the dyeing efficiency is poor, even if the stirring device is used. Therefore, scientists have developed the dynamic $\sec O_2$ dyeing technology, detailed in the next section.

Dynamic dyeing system

Static $\sec O_2$ dyeing system is lack of dyeing fluid circulation resulting in limited difusion rate of dyeing solution and poor dyeing efficiency, even if the stirring device is used. Aiming at the issue, dyeing solution circulation device is designed in dynamic $\sec O_2$ dyeing system for continuously circulating the dye between the dyeing vessel and the dyestuff vessel, ensuring that the dye enters the high-pressure

 $CO₂$ will be injected by the high-pressure pump. The dye will be continuously dissolved and difused onto the fabric to complete the dyeing process. Reproduced with permission of Elsevier from Penthala et al. (2019)

dyeing vessel with a saturated state. The continuous circulation of dyeing solution can strengthen the mass transfer process, thus improving the dyeing efficiency. Compared with the static scCO₂ dyeing system, the dynamic scCO₂ dyeing system equips with a dyeing solution circulation device (Fig. 3) (Penthala et al. 2020). During the dynamic $\sec O_2$ dyeing process, the dispersed solid dye is packed into the dyestuff vessel and dissolved by $\sec O_2$ and then circulated by the syringe pump passing through the flter into the dyeing vessel. The dyeing solution circulation can continuously circulate the dye between the dyeing vessel and the dyestuf vessel, ensuring that the dye enters the high-pressure dyeing vessel with a saturated state. The continuous circulation of dyeing solution strengthens the mass transfer process, thus improving the dyeing efficiency.

Fig. 3 Dynamic supercritical carbon dioxide ($scCO₂$) dyeing system. During the dynamic $\sec CO₂$ dyeing process, the dispersed solid dye will be packed into the dyestuff vessel and dissolved by $\sec O_2$ and then circulated by the syringe pump passing through the flter into the dyeing vessel. The dyeing solution circulation can continuously circulate the dye between the dyeing vessel and the dyestuff vessel, ensuring that the dye enters the high-pressure dyeing vessel with a saturated state. Reproduced with permission of Elsevier from Penthala et al. (2020)

According to whether the fabric has reciprocating motion in the dyeing vessel, dynamic $\sec O_2$ dyeing systems can be divided into two types, i.e., $\sec O_2$ beam dyeing system and \rm{scCO}_{2} jig dyeing system, which will be introduced separately in the following.

Dynamic beam dyeing system

The dynamic $\sec O_2$ beam dyeing system is one of the most widely used dynamic $\sec O_2$ dyeing systems. The critical factors of dynamic $\sec O_2$ beam dyeing systems are porous beam and associated pipe connections in the high-pressure dyeing vessel. Porous beam is designed with a barrel with evenly distributed holes for uniformly forcing the dyeing fluid into the fabric under supercritical conditions. Meanwhile, the associated pipe connections can ensure the continuous circulation of the dyeing fluid during the $scCO₂$ dyeing process.

The basic structure and dyeing process is shown in Fig. 4 (Kim et al. 2018; Reji et al. 2019). Before dyeing, the fabric will be wrapped around the dye beam and then placed in the center of the dye vessel, which is connected with the inlet of the dyeing solution. The dyeing solution fows through the fabric by the holes on both sides of the dye beam and circulates by virtue of the circulating pump. Moreover, the flowing direction of the dyeing solution can be changed by the reversing device to ensure the dyeing uniformity of the inner and outer layers of the fabric. The dynamic $\sec O_2$ beam dyeing system has the characteristics of small dyeing vessel, simple structure, and convenient loading, which is beneficial for dyeing cloth and yarn. However, it also has certain limitations. With the increased amount of dyed fabric, the radius

of the dyed fabric wound on the dye beam will also increase, thus enhancing the resistance of the dyeing solution passing through the fabric. As a result, the dyeing speed will slow down and the uniformity will be infuenced.

The dyeing efficiency of the traditional dynamic $\sec O_2$ beam dyeing systems is restricted by the gravity and the limited volume of the dyeing vessel. Designing horizontal beam dyeing vessel by employing the rotational drive shaft can efficiently avoid the gravity influence, and increasing the dyeing vessels to realize simultaneous multiple dyeing process leads to improved dyeing efficiency. Aiming at the issue, Gao et al. (2015) designed a high-efficiency scCO₂ beam dyeing system with two beam dyeing vessels, which can dye two batches of fabrics independently and simultaneously. The horizontal beam dyeing vessel equips with the rotational drive shaft to ensure the continuous rotation of the wrap beam (Fig. S2a and b). This system can overcome the gravity infuence during the dyeing process, thus achieving the high uniformity and production efficiency. The mobile driver section outside the vessel can actuate the movement of the cover, and the hydraulic lock device can fast lock the dyeing vessel cover to keep the air tightness of the whole dyeing process. Moreover, the double dyeing vessel system can overcome the defects that single-vessel system needs repetitive heating and gas venting to anew loading fabrics; as a result, the cost and $CO₂$ gas consumption can be decreased.

It is worth noting that there are various types of textiles in the practical dyeing production, such as fabrics, skeins, garments, and balls. In the above systems, the fabrics are wrapped around the beam to achieve the dyeing process. However, the traditional beam cannot accommodate the loading need of skeins, garments, and balls; for that, they

Fig. 4 Structure and working process of dyeing vessel in the dynamic supercritical carbon dioxide beam dyeing system (Kim et al. 2018). Before dyeing, the fabric will be wrapped around the dye beam and then placed in the center of the dye vessel, which is connected with the inlet of the dyeing solution. The dyeing solution fows through the fabric by the holes on both sides of the dye beam and circulates by virtue of the circulating pump

cannot be efectively wrapped around the beam. Considering the issue, the frame of the traditional beam dyeing vessel should be improved and specially designed for realizing different dyeing requirements. Zheng et al. (2015a) designed a special beam dyeing system to accomplish the uniform dyeing of loose fbers (Fig. S2c and d). The frame was designed as a tubular structure with inner porous cylinder, distributing porous tube and media channel to meet the need of the liquid flow, which made the best use of space in the dyeing vessel. It is worth noting that the system works with two dyeing modes. One is "inner dyeing/outer dyeing" mode that the fluid flows from the top of the dyeing vessel to penetrate fibers inside and then flows out at the bottom of the dyeing vessel, and another is "outer dyeing/ inner dyeing" mode that the dyeing solution enters from the bottom, adsorbed and dyed fbers from the outside of the dyeing frame. The alternant working modes can not only decrease the eddy of the dyeing solution but also increase the surface contact area of the dyeing solution and fbers, thus enhancing the dyeing rate and levelness.

Moreover, in the traditional skein dyeing system, it is easy to produce friction between the yarns due to the lack of pores, thus infuencing the dyeing uniformity. For this reason, Zheng et al. (2016) designed a scCO₂ skein dyeing system with double-ply parallel skein frame (Fig. S3a). During the process of $\sec O_2$ skein dyeing, one part of the dyeing fuid fows through the beam to the skein frame and penetrates the skein from the middle to both sides, while another part enters the skein frame from the $CO₂$ overflow port, and then penetrates the skein from the bottom to top, thereby efectively improving the leveling performance of the skein. Besides, the dyeing demand of wool ball is large in the industrial dyeing systems but cannot be wrapped around the beam as ordinary fabrics, and the gravity factor has a great infuence on the uniformity of wool ball dyeing. In order to change the gravity distribution and improve the dyeing uniformity of wool ball, a dynamic $\sec O_2$ wool ball dyeing system was constructed (Fig. S3b) (Zheng et al. 2017a). There are four limit covers in the vessel, among which the frst limit cover is used to control the amount of total wool balls and the other three limit covers are used to stratify the whole wool balls. The limit covers can be layered inside the dyeing vessel as needed in practical production. In addition, there are many micropores distributed on the limit cover, which can not only load the wool balls, but also ensure the circulation of dyeing liquid. The diferent dyeing vessel can be chosen according to actual dyeing needs of the fabric for efficient dyeing.

In the $\sec O_2$ beam dyeing system, the dyeing uniformity of the fabric can be improved by reversing the fow direction of the dyeing liquid. However, reversing the fow direction of the dyeing liquid required the dyeing system having the reversing pipeline of high-pressure fuid, which increases the risk of the dyeing process. As a result, scientists have developed the dynamic $\sec O_2$ jig dyeing system, detailed in the next section.

Dynamic jig dyeing system

In the $\sec O_2$ beam dyeing system, the dyeing uniformity of the fabric can be improved by reversing the fow direction of the dyeing liquid. However, reversing the fow direction of the dyeing liquid required the dyeing system having the reversing pipeline of high-pressure fuid, which increases the risk of the dyeing process. Meanwhile, forcing the fabric to have reciprocating motion in the high-pressure dyeing vessel can effectively improve the dyeing uniformity and efficiency. Dynamic $\sec O_2$ jig dyeing system was improved on the basis of the beam dyeing system, which added the fabric roller, guide roller, and the driver device in the dyeing vessel to realize the reciprocating motion of the fabric (Fig. 5). The fabric is wound around multiple guide rollers from one fabric roller and then connects to another fabric roller. While the supercritical fuid is dynamically circulated in the dyeing system, the fabric wound around the roller is also reciprocated in the dyeing vessel. After the dye dissolves in the supercritical fuid, the dyeing liquid will pass through and penetrate the fabric, which truly realizes the dynamic dyeing between the fabric and the dyeing solution. Generally speaking, the structure of the dynamic $\sec O_2$ jig dyeing system is compact and reasonable, and the dynamic $\sec O_2$ jig dyeing process combines the properties of supercritical fuid dyeing process and the characteristics of traditional water dyeing process, thus greatly improving the dyeing quality and efficiency. Nevertheless, the dynamic $\sec O_2$ jig dyeing system has the disadvantages that the process of fabric wrapping and unloading is inconvenient and therefore needs to be further improved.

Fig. 5 Dynamic supercritical carbon dioxide ($\sec O_2$) jig dyeing system. The fabric is wound around multiple guide rollers from one fabric roller and then connect to another fabric roller. While the supercritical fuid is dynamically circulated in the dyeing system, the fabric wound around the roller is also reciprocated in the dyeing vessel

Zhang et al. (2020) designed a horizontal scCO_2 jig dyeing system as shown in Fig. S4. This system equips with a supercritical fuid distributor between the guide rollers, from which the fabric can be penetrated evenly when it passes by. There are also two motors to drive the fabric rollers to unwind and rewind, respectively, as a constant speed during the dyeing process. After dyeing, the shell of the dyeing valve will slide along the guide rails, so that the fabric can be unloaded easily. Gao et al. (2022) further improved the system as shown in Fig. S5. The position of the guide rollers was changed and the dyeing liquid distributor was designed as the shape of conveyor belt, which make this system can be used in both ordinary jig dyeing and $\sec O_2$ jig dyeing. For the ordinary dyeing, the fabric in the closed valve will not be polluted by the external debris, thus creating a cleaner dyeing environment. Furthermore, in the $\sec O_2$ dyeing system, the unique valve space distribution and compact wrapping structure make the contact between the fabric and the dyeing liquid more sufficient, so that the dyeing uniformity and efficiency are improved.

For better contact between supercritical fuid and the fabric, Wang et al. (2014) improved the traditional scCO₂ jig dyeing system with spray nozzle in the dyeing vessel (Fig. S6). During the dyeing process, while the fabric is moving on the guide roller, $\sec O_2$ dissolved with the dye will be sprayed out by the nozzle. After the rapid decompression and expansion, the dyeing liquid will pass through the fabric to accomplish the dyeing process, and then, $\sec O_2$ with the remaining dyes will enter the recovery device and can be reused after the separation. Compared with the system where the dyeing liquid is controlled by the distributor, the spray nozzle can dye the fabric more precisely and in detail. Moreover, according to the specifc dyeing requirements, the positions of the spray nozzle and the recycler can be adjusted, so as to realize single-side or double-side dyeing of the fabric. The $\sec O_2$ spray dyeing system has the characteristics with simple structures, safe and convenient to operate, good dyeing levelness and continuity, and so on. However, the $\rm{s}\rm{c}\rm{CO}_{2}$ spray dyeing system generally relies on the high-pressure dyeing nozzle, so when $\sec O_2$ is directly sprayed on the fabric without any protection measures, the fabric will be easily damaged by the high-pressure fuid jet. In order to overcome the disadvantages, Wang et al. (2012) designed a new type of $\sec O_2$ spray dyeing system (Fig. S7). The system consists of squirrel-cage dyeing vessel and rolling dyeing rollers, which can realize dynamic contact dyeing between $\sec O_2$ and the fabric. Moreover, the dyeing fabric is supported by the rolling dyeing rollers, which effectively eliminates the deformation and tension damage caused by the high-pressure fuid jet and traction force. Furthermore, for the light and deformable fabric dyeing, Yang et al. (2014) improved the $\sec O_2$ spray dyeing system by combining the advantages of jig dyeing system and spray dyeing system

(Fig. $S8$). This scCO₂ spray dyeing system drives and circulates the fabric by the guide cable and eliminates the complex nozzle systems that avoids damage of the light fabrics. This $\sec O_2$ spray dyeing system not only increases the contact area between the dyeing solution and the fabric, but also reduces the energy consumption of the circulation pump, especially suitable for high-volume dyeing production of deformable fabrics.

In a summary, the $\sec O_2$ dyeing systems are generally divided into static dyeing system, dynamic beam dyeing system, and dynamic jig dyeing system, according to whether the dyeing liquid and the fabric are dynamically circulated during the dyeing process. Static dyeing system is mainly consisted of $CO₂$ cylinder and pump, temperature and pressure controller, and the dyeing vessel. Static dyeing system is easy to operate and suitable for using in the laboratoryscale dyeing experiments to study the effect of parameters on the dyeing of different fabrics during the $\sec O_2$ dyeing process. Dynamic dyeing system additionally adds a circulation system and a recycling device on the basis of the static dyeing system. Moreover, the dynamic beam dyeing system and the dynamic jig dyeing system show diference on the dyeing vessel, which matched with the specifc dyeing requirements. Dynamic dyeing system has higher dyeing efficiency and is more suitable for large-scale practical industrial production. Diferent types of fabric dyed in different $\sec O_2$ dyeing systems and the corresponding dyeing conditions are summarized in Table 1.

Moreover, it is worth noting that $\sec O_2$ dyeing process may afect the properties of the dyeing fabric, which generates the resulting need for the design and operation of the $\sec CO_2$ dyeing system. The $\sec CO_2$ dyeing process contains the rapid pressurization and depressurization step, and the depressurization rate turns into an important factor affecting the properties of the dyeing fabric. Zheng et al. (2017b) investigated the infuence of the depressurization rate on the properties of yak hairs and found that the breaking strength and the elongation at break of the yak hairs both increased with the increasing depressurization rate. The result may attribute to the rapid air fow during the pressurization step and depressurization step during the $\sec O_2$ dyeing process. The pressurization process is realized by operating the fxedfrequency high-pressure pump, while the depressurization process is achieved by the valve control leading to relatively irregular pressure decreasing rate. Consequently, the torsion, bending, and shearing action for the yak hair strengthens when the depressurization rate increases, especially in the depressurization step during the scCO_2 dyeing process.

Meanwhile, the surface fber morphology may also be afected during the high-pressure process, of which the surface of the dyeing fabric may be altered to be rougher with the appearance of stripes and wrinkles. Zheng et al. (2015b) observed the same conclusion when studying the infuence

Table 1 Fabric dyed in different scCO₂ dyeing systems and the associated dyeing conditions

Fabric	Type of dyeing system	Temperature $(^{\circ}C)$	Pressure (Mpa)	Time (min)	References
Cotton	Static	$30 - 120$	$12 - 30$	$30 - 120$	Zhang et al. (2017)
Cotton	Dynamic	$60 - 120$	20	$60 - 180$	Yang et al. (2017)
Cotton, wool, silk	Static	110	26	Not reported	Liu et al. (2018)
Cotton, wool, silk	Dynamic	130	20	Not reported	Yan et al. (2020)
Wool	Static	120	20	90	Zhang et al. (2016)
Wool	Dynamic	$70 - 120$	$15 - 35$	$30 - 180$	Zheng et al. $(2017a)$
Yak hair	Dynamic	$80 - 120$	$21 - 25$	$40 - 80$	Zheng et al. (2017b)
Polyester, cotton	Dynamic	80	18	60	Gao et al. (2015)
Polyester	Dynamic	$60 - 140$	$15 - 35$	$20 - 100$	Xiong et al. (2017)
Polyester	Static	120	25	60	Abate et al. (2020)
Polyester	Static/Dynamic	$100 - 120$	24	60	Zheng et al. (2016)
Nylon 6, Polyester	Dynamic	$100 - 130$	$15 - 25$	$60 - 120$	Abou et al. $(2018b)$
Nylon 6, Polyester	Static	$100 - 130$	$15 - 25$	$60 - 180$	Abou et al. $(2019a)$
Nylon 6	Dynamic	$80 - 120$	$5 - 15$	$60 - 180$	Abou et al. (2015)
Polypropylene	Dynamic	$100 - 130$	$15 - 25$	$60 - 180$	Abou et al. $(2018c)$
Polypropylene	Static	$110 - 130$	$15 - 25$	$60 - 120$	Abou et al. $(2019b)$
Zipper tape	Dynamic	90	23	60	Huang et al. (2017)
Zipper tape	Dynamic	$70 - 110$	$14 - 25$	$20 - 65$	Huang et al. (2019)
Ultra-high-molecular- weight polyethylene fiber	Static	120	28.85	60	Zhou et al. (2023)
Wood	Static	$45 - 100$	$10 - 30$	900	Jaxel et al. (2020)
Ramie fiber	Static	$70 - 130$	$12 - 20$	90	Liu et al. (2006)

of depressurization rate on the morphology of the wool fber, and the results investigated that the fber damage and brightness variation would be higher with the increased depressurization rate. The phenomenon can be attributed to the burst of water droplets by the fast pressure release, which can be efectively overcome by precisely controlling and slowing down the depressurization rate. As a consequent, for avoiding the deterioration of mechanical and physical properties of the dyeing fabric, the $\sec O_2$ dyeing system should be equipped with depressurization rate controlling device to precisely release the pressure with a fxed frequency in control. Moreover, the depressurization rate should be set at diferent values when dyeing diferent fabrics for obtaining optimized properties.

In addition, particles formation on the dyeing fabric during the $\sec O_2$ dyeing process has also been observed in several research. Gao et al. (2015) found the particles presented on the surface of the dyeing fabric after the $\sec O_2$ dyeing process attributed to the deposition of active compounds in the dye solution. Abate et al. (2019) reported the particles formed on dyeing fbers due to the oligomer crystals produced by the migration of these molecules to the surface, which was caused by $CO₂$ penetration and high-temperature process. The same phenomenon was also observed in nicotinamide dyeing for viscose fabric and azo-based dispersed dyes for dyeing wool and cotton fabrics. However, it is worth noting that no obvious presence of the formed particles was detected when using the gallic acid with higher difusion rate due to its smaller size than other dyes in the same conditions. Enhancing the fber swelling by suitable operation of the $\sec O_2$ dyeing system can lead to a better penetration of the dye into fabric instead of deposition at the surface of the fabric. Consequently, for avoiding the particle deposition of the high molecular weight compounds on the dyeing fabric during the $\sec O_2$ dyeing process, the $\sec O_2$ dyeing system should be designed with depressurization controlling device to precisely release the pressure in the dyeing vessel for avoiding the nucleation, or operating suitable supercritical fuid fow patterns in the dyeing vessel to enhance the difusion rate of the dyes into the fabric. Moreover, it is also important to design the $\sec O_2$ dyeing system with recirculation flter for fltration of the recirculation current to efectively eliminate the suspended particles, as well as improve the $CO₂/dye$ separators for better separation efficiency of $CO₂$ and the dye.

Overall, $\sec 0₂$ dyeing technology has been identified as an attractive alternative to water-based dyeing. The low mass transfer resistance and high difusion rates observed in \rm{scCO}_{2} assist dye penetration into the fibers causing a reduction on dyeing times. Besides, $CO₂$ completely vaporizes by depressurization, and therefore, no subsequent drying or solvent removal operations are required. The rational design

of scCO₂ dyeing system is the key of scCO₂ dyeing technology applied in practical production. However, current $\sec O₂$ dyeing systems have the problems that diferent types of fabric and dyes, the geometric design of the dyeing vessel, and the flow conditions of $\sec O_2$ will all influence the dyeing uniformity during the $\sec O_2$ dyeing process. Therefore, carrying out the computational simulation of fuid dynamics of $\sec O_2$ flow in different types of $\sec O_2$ dyeing systems is necessary for analyzing the efect of operation conditions on the dyeing uniformity during the $\sec O_2$ dyeing process, as well as for contrapuntally designing the dyeing conditions of diferent types of fabrics. Moreover, the residual dye particles in the pipelines of the systems during the $\sec 0₂$ dyeing process will pollute the fabric after the dye changes in the subsequent dyeing process. Designing novel separator that can provide lower pressure drop corresponding to higher separation efficiency is crucial for solving the $CO₂$ recontamination and color reproducibility problems. In addition, visualization of the $\sec O_2$ dyeing process is crucial for judging the color presentation degree and dyeing quality, which requires visual $\sec O_2$ dyeing systems should be more focused and developed for the future improvement of the $\sec O_2$ dyeing systems.

Extraction systems

With the increasing demand of natural products, more selective and efective extraction technology is required to extract the ingredients from raw materials (Reverchon and Senatore 1992). The supercritical fuid extraction (SFE) technology has been developed as a substitute for traditional solventbased techniques (Sahena et al. 2009). SFE is referred to a process in which the extracted component is separated or dissolved from the raw materials by the supercritical fuid according to their dissolution properties (Spanos et al. 1993). The solubility of the extracted components can be maintained by varying the temperature and pressure above the critical point (Zougagh et al. 2004). The SFE technology has the advantages of selective extraction, high extraction efficiencies, easy separation, short processing time, and low impact on the environment, which paves the way to selectively and mildly extract the ingredients from the natural materials (Attard et al. 2016). When $\sec O_2$ is used, many components can be extracted in a short time, and there is no residual solvent after the extraction process (Herrero et al. 2010). During the scCO₂ extraction process, the liquid-like density provides a good solvating power, while the gas-like high diffusivity and low viscosity lead to the deep penetration into the matrix. As a result, the rate of mass transfer into $\sec O_2$ is higher than that into the liquid (Chassagnez-Mendez et al. 2000; Silva et al. 2017; Vladić et al. 2021). In addition, $\sec O_2$ can be applied in the extraction of both polar compounds and non-polar compounds. $SCO₂$ as nonpolar compound is a good solvent for non-polar compounds extraction, such as essential oils, hydrocarbons, and organic pesticides (Chassagnez-Mendez et al. 2000; Fornari et al. 2012; Mouahid et al. 2013; Tyśkiewicz et al. 2018). Moreover, the addition of polar cosolvents, such as ethanol and methanol, can efectively increase the solubility of polar compounds in $\sec O_2$, thus overcoming the disadvantages of low afnity with polar compounds (Devi and Khanam 2019; Fujii et al. 2018; Soh et al. 2019). With more and more extensive application of $\sec O_2$ extraction, the corresponding systems are also improved continuously.

 $SCO₂$ extraction process typically requires continuous production proceeded under high-pressure conditions, and therefore, the rational design of scCO_2 extraction system has become the research focus in recent years. $ScCO₂$ dyeing systems should satisfy the requirements that the components to be extracted can be effectively dissolved into $\sec O_2$ under desired constant temperature and pressure conditions, and scCO₂ should be evenly distributed and fully contacted with the extract at any cross section of the extraction vessel to ensure the extract purity, which can be realized by equipping backpressure regulator (BPR) and high-pressure sealed screw feed device, respectively. $SCO₂$ dyeing systems can be divided into batch extraction system and semicontinuous extraction system. Moreover, multiple-separator extraction systems were designed to realize multiple extraction of the same or diferent components for obtaining improved extraction efficiency. Different types of scCO₂ dyeing systems will be introduced separately in the following.

Batch extraction system

 $SCO₂$ batch extraction systems are the traditional extraction systems that most widely used in laboratory-scale extraction of various types of components. $SCO₂$ batch extraction systems should be designed with high-pressure vessel, temperature- and pressure-controlling device, and the separator to collect the extracted components. Moreover, when adding the raw materials into the extraction vessel and taking out of the extracted components from the extraction vessel, the extraction vessel needs to be opened and closed frequently. In order to realize facilitate operation and shorten operation time, the extraction vessel should equip with quick-opening device. $SCO₂$ dynamic extraction systems are simply designed and easy to operate and can maintain the constant extraction conditions in the extraction vessel, which is beneficial to measure the extraction rate and extraction influence of parameters, such as extraction temperature, pressure, $\sec O_2$ flow rate, concentration, and particle size of raw materials.

Naikoo et al. (2021) used the typical batch scCO₂ extraction system to extract fat-soluble vitamin A from retinol (Fig. 6). The system is composed with $CO₂$ tank, refrigerator, high-pressure pump, heating device, extractor, and cosolvent tank that can be set according to actual needs. The scCO₂ extraction process is as follows. Liquid CO₂ is frstly cooled at low temperature by the refrigerator and mixed with the cosolvent of methanol, and then, $CO₂$ and the cosolvent are further mixed through the spiral tube. Before entering the extractor, $CO₂$ will be preheated to the desired extraction temperature and then enters the extractor to react with retinol for the extraction of fat-soluble vitamin A. This typical $\sec O_2$ extraction system has been widely used due to its properties of simple, extensively applicable and easy to control. Mouahid et al. (2013) used this scCO₂ extraction system to study the extraction modeling curves of four diferent microalgae, which aims to verify that Sovová equation could be applied to all microalgal extraction cases. Khaw et al. (2019) used the system for scCO_2 extraction of freeze-dried leaf juice from Carica papaya and further investigated the maximal extract activity at the best smallscale extraction parameters. Győri et al. (2019) extracted the aroma compounds from herbs and spices by this static $\sec O_2$ extraction system. It was found that the extraction efficiency, the number of extracted compounds, and the quality of product from $\mathrm{s}\mathrm{c}\mathrm{C}\mathrm{O}_2$ extraction are all higher than those from classic maceration. Chassagnez-Mendez et al. (2000) studied the infuence of temperature and cosolvent on turmeric extraction by this typical scCO_2 extraction system. Moreover, diferent types of components extracted in batch extraction system and the corresponding extraction conditions are summarized in Table 2.

Semicontinuous extraction system

In the traditional batch extraction systems, high $CO₂$ flow rate can increase mass transfer driving force, thus

Fig. 6 Batch supercritical carbon dioxide ($scCO₂$) extraction system designed by Naikoo et al (2021) . Liquid CO₂ is firstly cooled at low temperature by the refrigerator and mixed with the cosolvent, and then, $CO₂$ and the cosolvent are further mixed through the spiral pipe. Before entering the extractor, CO₂ will be preheated to the desired extraction temperature and then enters the extractor to react with retinol for the extraction of fat-soluble vitamin A

improving the extraction rate and reducing the extraction time. However, high $CO₂$ flow rate also leads to inadequate contact between the extract components and scCO_2 , which greatly affects the extraction efficiency. Meanwhile, the extraction vessel needs to be opened and closed frequently to load the raw materials during the static extraction process, thus leading to complex extraction operation and long extraction time. Aiming at the issue, semicontinuous extraction systems are improved on the basis of batch extraction system by adding BPR and $CO₂$ recycling device to realize static and dynamic alternating extraction. The critical factors of semicontinuous extraction systems are rational design of the extraction vessel for simultaneously realizing maintaining constant high pressure in the extraction vessel and also ensuring the continuous fow of the raw materials and extracted components into and out of the extraction vessel, which can be realized by utilizing high-pressure sealed screw feed device for extracting components from solid extracts and by employing highpressure injector to extract liquid components.

The traditional batch extraction system has been improved by adding BPR and a $CO₂$ recycling device to allow smooth alternating of static extraction and dynamic extraction, and to realize $CO₂$ recycle (Fig. S9) (Ghoreishi et al. 2009). During the scCO_2 extraction process, after CO_2 is injected into the extraction column, the BPR will be set to the appropriate working pressure. Simultaneously, the temperature will be maintained through the coil preheater and oven. When the required pressure and temperature are reached in the extraction column, the pump and the valve will be disconnected. Subsequently, \secO_2 can dissolve the ingredients during the static extraction time, and then fow at the set rate after extraction. The dissolved ingredients are trapped in the collection vessel and $CO₂$ is recycled by the circulation device. The alternating static extraction and dynamic extraction

process can increase the scCO_2 -assisted extraction time, thus improving the extraction rate and efficiency.

Soh et al. (2019) used the semicontinuous extraction system to study how the pressure, temperature, and flow rate influence the $\sec O_2$ extraction of patchouli oil. It was found that higher pressure and fow rate would increase the extraction rate, while higher temperature had an inverse effect for the extraction process. Fujii et al. (2018) developed an analogous system, in which the cosolvent was replaced with liquid solution to form the liquid– $CO₂$ emulsion in the micro-mixing device, thus increasing the mass transfer from liquid to $\sec O_2$. This system could fast and continuously extract vanillin from aqueous vanillin solution and also could be applied in other aqueous liquids containing hydrophobic compounds. Moreover, Cornelio-Santiago et al. (2017) measured the $\sec O_2$ extraction solubility of oil from green coffee by the semicontinuous extraction system. Fornari et al. (2012) extracted the essential oil from diferent herbs and plants using the semicontinuous extraction system. Devi and Khanam (2019) utilized the semicontinuous extraction system for the extraction of ω-3 α-linolenic and ω-6 linoleic acids from hemp seed in $\sec O_2$ and further optimized the process parameters. Andersen and Bruno (2003) used this semicontinuous extraction system to extract metal β-diketonate complexes from aqueous solutions, which overcame the limited mass transfer between the CO_2 -soluble chelating agent and the water-soluble metal ion.

Multiple‑separator extraction system

The volume of the extraction vessel also limits the extraction efficiency in the practical production process. $ScCO₂$ extraction systems are mainly small or medium sized; for that, big volume of the extraction vessel requires high $CO₂$ consumption and long pressurization time and also increases the operation difficulties. Aiming at the issue, multiple extraction vessels and separators can be employed to further improve the extraction efficiency. Multiple extraction vessels can be connected in series to extract the same components repeatedly for enhancing the extraction purity and also can be connected in parallel to extract the same components for improving the extraction rate or extract diferent

components in diferent extraction vessels simultaneously. Moreover, multiple separators are equipped connected with the multiple extraction vessels to separate the same or diferent components correspondingly and also can evaluate the separation extent and extraction efficiency.

Rochfort et al. (2020) and Srinivas et al. (2017) further improved the above system to achieve multicomponent separation and evaluate the separation extent (Figs. 7 and S10). The improved system is equipped with an automatic backpressure regulator (ABPR) for automatically regulating the temperature and pressure in the extraction vessel and also added three independent-heated cyclone separators for further separation of the ingredients. Moreover, the temperature and pressure in three cyclone separators are independently and accurately controlled by three manually adjusted BPR, which not only could measure the separation extent of different categories in each cyclonic separator, but also could evaluate the extraction efficiency of the same category by combining the ingredients together.

In addition to the continuous improvement of the extraction process, the design of the extraction reactor is also critical. The original design of the Nottingham reactor is based on a syringe compressor that can convert liquid $CO₂$ to $\sec CO_2$ (Fig. 8a) (Qamar et al. 2021). $\sec CO_2$ enters from the top of the extraction chamber where glass wool and stainless steel flter are used to separate the ingredients from the raw materials. However, the disadvantage of this extraction reactor is that a large amount of ingredients will remain on the glass wool which reduces the extraction yield seriously, and there is also a problem of raw material pollution

Fig. 7 Supercritical carbon dioxide ($\sec O_2$) extraction system with independent separators (Srinivas et al. 2017). The temperature and pressure in three cyclone separators are independently and accurately controlled by three manually adjusted backpressure regulator (BPR), which not only could measure the separation extent of diferent categories in each cyclonic separator, but also could evaluate the extraction efficiency of the same category by combining the ingredients together

to be resolved. In order to solve the above problems, after the extraction is completed, the extraction chamber will be removed from preconditioning chamber and inverted to depressurize with the help of bracket (Fig. 8b). Because the extraction chamber is completely depressurized at 0 bar, the ingredients can be separated with a larger extent, thus improving the extraction efficiency. However, the problem of serious raw material pollution has not been completely resolved. Therefore, the helix reactor with low-pressure regulation chamber was designed (Fig. 8c) (Qamar et al. 2021), where the backpressure is directly regulated by the liquid $CO₂$ cylinder and the preconditioning chamber. During the \rm{scCO}_{2} extraction process, \rm{CO}_{2} enters and extracts from the bottom of the extracting chamber and then the extracted components are separated from the raw materials by the built-in standard flter at the top of the extracting chamber. After the extraction is completed, $CO₂$ dissolved the matrix will enter the separating chamber, where the pressure is about 50 bar to avoid the piston efect of dry ice. Finally, the ingredients will be collected in the extract collecting glass tube. The helix reactor can not only improve the extraction efficiency, but also avoid the raw material contamination problems, which is suitable for extraction requirements with high quality and high extraction rate.

To sum up, the scCO_2 -assisted extraction systems are split into typical static extraction system, static and dynamic alternating extraction system, and multiple-separator extraction system. The typical static extraction system is consisted of $CO₂$ supply and pump device, temperature- and pressure-controlling device, and the extraction vessel. Static and dynamic alternating extraction system is improved on the typical static extraction system by adding BPR and $CO₂$ recycling device. Static and dynamic alternating extraction system has high extraction efficiency and is widely used for most of the extraction requirements. The multiple-separator extraction system is added with several independent-heated cyclone separators on the basis of the typical static extraction system and can realize multicomponent separation and evaluate the separation extent.

Overall, $\sec O_2$ extraction technology has obtained increasing attention in the felds of fractionation of liquid mixtures and solid material extraction due to its fascinating properties of high density, low viscosity, zero surface tension, and adjustable properties by manipulating the pressure and temperature. Now, more and more attention has been paid on the development of $\sec O_2$ extraction processes and corresponding systems and also attracted interests of industries on scCO_2 extraction techniques. However, current $\sec 0$, extraction systems also face several challenges to be further solved and thus more suitable to apply in the practical industry production. In $\sec O_2$ extraction process, the particle size of raw materials will affect the extraction efficiency, of which the contact area between the extracted

Fig. 8 a Supercritical fuid extraction Nottingham reactor. Supercritical carbon dioxide ($\sec O_2$) enters from the top of the extraction chamber where glass wool and stainless steel flter are used to separate the ingredients from the raw materials. **b** Sample-containing chamber inverted and placed on stirrer for the depressurization. After the extraction is completed, the extraction chamber will be removed

from preconditioning chamber and inverted to depressurize with the help of bracket and **c** schematic representation of the Helix reactor (Oamar et al. 2021). After the extraction is completed, $CO₂$ dissolved the matrix will enter the separating chamber, and the ingredients will be collected in the extract collecting glass tube

component and $\sec O_2$ will increase when the particle size of raw materials decreases, thus leading to enhanced extraction efficiency. However, small particle size of raw materials will not only result in serious blockage of the sieve pore, but also cause the plug of the flter at the exit of the extraction vessel. Meanwhile, when the extracted components are powder materials, the pressed powder under high pressure and temperature will seriously wear the seals of the high-pressure extraction vessel, which requires the seals of the high-pressure extraction vessel possessing strong wear resistance to maintain the stable pressure during the whole $\sec CO_2$ extraction process. Moreover, in $\sec CO_2$ extraction process, the water cannot be thoroughly removed, thus

accumulating more and more water with the cycle, which may influence the extract purity and efficiency. The existing $\sec O_2$ extraction system usually equips with drain hole or relies on adding desiccant, which has the disadvantages of poor water removal efect and inconvenient replacement of the desiccant. Improving the $\sec O_2$ extraction system with unique water removal device will effectively enhance the extract purity and efficiency. It is also worth noting that dynamic $\sec O_2$ extraction system can improve the extraction efficiency due to the circulation of $\sec O_2$ compared with static $\mathrm{s}\mathrm{c}\mathrm{CO}_2$ extraction system. However, the dynamic $\rm scCO$ ₂ extraction system has the problems that the pressure fuctuation in the high-pressure extraction vessel will be

caused by $CO₂$ circulation process. Therefore, improving the dynamic $\sec O_2$ extraction system with pressure precisely controlling device to maintain the constant pressure during the whole extraction process is necessary.

Chromatographic systems

The availability of efficient analytical separation tools is critical to the pharmaceutical industry; as a result, various chromatographic techniques, such as high performance liquid chromatograph (HPLC), gas chromatograph (GC), and supercritical fuid chromatograph (SFC), are commonly used for the purpose. Among them, SFC has attracted attention in recent years (Tavares and Lanças 2000; Van et al. 2018; Zheng et al. 2020). When the temperature and pressure of the mobile phase in SFC are higher than the critical point, the solvating power of the mobile phase will be changed to achieve the separation of certain components (Desfontaine et al. 2015; Gao et al. 2017; He et al. 2018). The parameters of temperature, pressure, and modifer content during SFC process, compared with GC and HPLC processes, can be controlled independently or cooperatively (Heiland et al. 2019; Lu et al. 2006; Planeta and Roth 2004; Zheng et al. 2020). Moreover, the supercritical fuid with gas-like viscosity and difusivity can still maintain high separation efficiency at high mobile phase velocities, and its liquidlike density and solvating power can also provide fast transfer and good solubility of analytes, which is benefcial for chromatography and purifcation in food or pharmaceutical analysis (Ehlers et al. 1998; Hawthorne and Miller 1986; Taylor 2009). Among the supercritical fluid, scCO₂ with the characteristics of low solvent toxicity, low critical temperature, chemical inertness, and nonfammability has become the preferred mobile phase in SFC technique. Furthermore, the addition of modifers can overcome the limited solubility of polar compounds in non-polar $\sec O_2$, thus greatly expanding the application in pharmaceutical feld (Gere 1983).

SFC utilizes $\sec O_2$ as the mobile phase during the chromatography process and employed porous solid particles in the chromatography column. When the mobile phase carrying the samples continuously impacts the stationary phase, the components to be separated in the samples will be simultaneously subjected to the thrust and gravity of the mobile phase. Due to the combined action of these two forces, the mobile phase carrying the samples will go through "adsorption-elution-adsorption-re-elution" process repeatedly. If a substance in the sample is closely bound to the stationary phase, the repressor force will be high, and the migration rate will be slow. Otherwise, the migration rate will be fast. Then, the speed of each substance movement will be diferent due to the diference of repressor force, thus leading to diferent staying time of each substance in the chromatography column, so as to realize the separation of each component in the samples. SFC systems are gradually developed to become commercialized. In the 1980s, the SFC systems were mainly divided into two types, i.e., open tubular capillary column (GC-like chromatograph) system and packed column (LC-like chromatograph) system, which were exclusively home-made by retroftting GC or LC system for around 20 years (Pyo 2001). In 1992, HP supplier invented a new SFC system with a dynamic compressibility compensation and electronic BPR. Since then, more suppliers ofered diferent SFC systems, such as Shimedzu, Jasco, and Fluitron, which has been commercialized successfully.

Open tubular capillary column system

Novotny et al. designed the open tubular capillary column SFC in 1981, which was a fused-silica capillary tube with 50 μm inner diameter and coated with polymer as the stationary phase. The open tubular capillary column SFC system is equipped with syringe pump for delivering and pressurizing $CO₂$, injection valve for introducing samples, GC-like oven, wall-coated open tubular column, fxed restrictor for maintaining the mobile phase at supercritical state and fame ionization detector (FID) (Fig. 9a) (Saito 2013), and the open tubular capillary column SFC system has been exclusively marketed and patented by Lee Scientifc.

The open tubular capillary column SFC system is equipped with syringe pump, injection valve, GC-like oven, wall-coated open tubular column, fxed restrictor, and fame ionization detector (FID) (Fig. 9a) (Saito 2013), and the open tubular capillary column SFC system has been exclusively marketed and patented by Lee Scientifc. In this system, the backpressure is controlled by the restrictor with a certain flow rate, which can ensure the system pressure above the critical pressure of $CO₂$ to keep supercritical state. However, under the circumstances, pressure as the most important parameter will be restricted by the fow rate, which cannot be changed independently. Therefore, the optimum flow rate cannot be obtained at the optimum pressure. In addition, when the organic modifers are used to increase the polarity of scCO_{2} , excessive baseline background will be produced and thus limit the application of standard FID detectors (Pyo and Lee 1999). Thus, the open tubular capillary column SFC systems declined rapidly in the early 1990s.

Packed column system

It is worth noting that almost all SFC research works were proceeded by utilizing packed column SFC before the open tubular capillary column SFC was introduced. Pack column SFC was LC-like instrument while open tubular capillary column SFC was more like GC. Although open tubular

Fig. 9 a Typical gas chromatograph (GC)-like open tubular column supercritical fuid chromatograph (SFC) system. FID. Flame ionization detector. Reproduced with permission of Elsevier from Saito (2013); **b** typical liquid chromatograph (LC)-like packed column SFC system. PU1. Liquefied $CO₂$ delivery reciprocating pump with chilled pump heads, PU2. Modifer solvent delivery pump, RF. Safety relief valve that prevents over pressure, INJ. Injection valve, PDA.

Photodiode array, PT. Pressure transducer, BR. Backpressure regulator. Reproduced with permission of Elsevier from ref. (Saito 2013); **c** SFC-4000 analytical chromatograph designed by Jasco; **d** 1260 Infnity II analytical SFC system designed by Agilent Technologies; **e** Prep 150 AP preparative SFC system designed by waters; **f** Nexera UC Prep preparative SFC system designed by Shimadzu

capillary column SFC processes longer columns that yield more theoretical plate numbers, pack column SFC renewed the interest of researchers due to its more uniform packing materials, smaller diameter, and the ability to pack narrower fused-silica columns compared with open tubular capillary column SFC. The typical packed column SFC system is developed from HPLC chromatograph by adding photodiode array (PDA) UV detector, pressure transducer, and electronic BPR (Fig. 9b) (Saito 2013).

The typical packed column SFC system is developed from HPLC chromatograph by adding photodiode array (PDA) UV detector, pressure transducer, and electronic BPR (Fig. 9b) (Saito 2013). During the packed column SFC chromatography process, the pressure transducer can monitor the real-time pressure, which will be compared with the set pressure by BPR. Afterward, by adjusting the fow resistance of the regulator, the actual pressure can maintain equal to the set pressure to keep the stability of chromatography process. Comparing with the open tubular capillary column system, the packed column system can control the pressure independently without the restriction of the mobile phase fow rate. Moreover, Saito et al. (1988) designed a new type of electronically controlled BPR with small volume applied in packed column SFC system, realizing efficient chromatography without cross-contamination between different components. Sugiyama et al. (1985) used the packed column SFC system hyphenated with SFE system to evaluate the extraction and chromatography performance of cafeine from the coffee beans. The SFE system is directly connected to the SFC system, which allowed online introduction to the SFC column, and the PDA is used to monitor the signal. Nowadays, various detectors, e.g., mass spectrometer, can connect online with SFC system.

In the recent years, various key system manufacturers, such as Shimadzu, Jasco, Agilent Technologies, Thar Technologies, Thermo Fisher Scientifc, and Waters Corporation, launched their state-of-the-art system, which included both analytical and preparative SFC system.

Analytical supercritical fuid chromatography

As for the analytical SFC system, SFC-4000 analytical chromatograph designed by Jasco is equipped with a temperature-stabilized pump that can ensure the optimum conditions for pumping liquid $CO₂$ (Fig. 9c). This system neither requires complicated pretreatment of $CO₂$, nor needs a chiller to cool the pump heads. There are also automatic stop-valves for controlling the input and output of $CO₂$, as well as the precision BPR for maintaining supercritical conditions with pressure fuctuations. Besides, this system is also equipped with three ovens for diferent temperature zones, and with one diode array for high-sensitivity component identifcation and component identifcation positive confrmation toward enantiomeric separation. These features lead to a signifcant improvement on the detection performance compared with that from the previous SFC system. Moreover, the ACQUITY Ultra Performance Convergence Chromatograph $(UPC²)$ system developed by Waters is improved based on the previous traditional SFC system. In the UPC² system, both the CO_2 chilling and compression take place inside a pump with an integrated device. This system also equipped with an active component to further increase the backpressure and with an independent pump heads cooling device achieved by a Peltier module, realizing accurate control of the pressure and temperature,

respectively. The 1260 Infnity II SFC system designed by Agilent Technologies has presented broad power range to support pressure up to 600 bar at flow rates up to 8.3e – 5 L/s for extended application reach and the patented feed injection technology guarantees outstanding injection precision and LC-like reproducibility (Fig. 9d).

Preparative supercritical fuid chromatography

The successful commercially available preparative SFC system–Prep 150 AP SFC system from Waters offers column switching, automated sample handling, and tracking and fraction collection in an open-bed format (Fig. 9e). This SFC system utilizes $\sec O_2$ as its main mobile phase in combination with one or more organic solvents, which provides routine compound purifcation with fast separations, high resolution and high throughput for drug discovery labs. The stacked injection capability of system is also available for the purifcation of large amounts of materials delivered by multiple injections and for allowing for multiple peaks collected into separate vessels. In short, the Prep 150 AP SFC system can provide lower pressure drops across the column, faster equilibration, lower cost per sample, and reduced solvent and thus is reproducible and applicable to a wide range of compounds relevant in the pharmaceutical, food and environmental feld, life sciences, and chemical materials. Besides, the semi-preparative SFC system–Nexera UC Prep designed by Shimadzu is equipped with the patented gas–liquid separator (Fig. 9f), which can decrease fow rate without increasing the pipe diameter by splitting fow through multiple channels. $CO₂$ is discharged externally, while the liquid travels along the column and drips directly into the collection vessel without dispersing or spattering the eluate. These features can successfully reduce sample dispersion and carryover and also achieve high recovery rates, so that the high recovery rates can be obtained regardless of fow rate or modifer concentration, even for volatile compounds such as the fragrance linalool. Nowadays, commercialized analytical and preparative SFC systems are developing rapidly and have been applied for analysis and purifcation in various felds, and they will attract more and more interests of scientists in future.

Through the introduction of the foregoing, the SFC systems are mainly divided into open tubular capillary column system and packed column system, which all have the common components of $CO₂$ cylinder, syringe pump, and temperature- and pressure-controlling device, but they present the diference on the reaction column. Moreover, it is worth noting that the SFC systems have been commercialized of both analytical and preparative SFC system.

Overall, SFC has been revived attributed to the introduction of a new generation of columns and instruments from major suppliers of chromatographic instrumentation.

Particularly, using $CO₂$ as the mobile phase in SFC possesses the advantages of short equilibration time, high flow rate, and low solvent consumption. However, SFC also faces the challenges of high consumption of $CO₂$, difficult collection of the solute, and so on. The consumption of CO_2 can be high at high fow rate, such as 3 kg/min for the Supersep 3000 of Novasep supplied by a $CO₂$ tank, and a B50 cylinder can only supply this system for 11 min. Therefore, recycling and reusing $CO₂$ by properly tuning the temperature and pressure of $CO₂$ downstream from the vessel can effectively reduce CO_2 consumption. Consequently, it requires SFC equipped with separate and recovery device to realize good recovery of the solute and pure $CO₂$ without contamination of $CO₂$ by the solute and without dissolved solvent inside. The technique designed by Novasep realized the requirement by cleaning the recycled gaseous $CO₂$ by a flow of liquid $CO₂$. Moreover, preparative SFC also face the challenge of the solute collection. The separation of CO_2 and the modifer dissolved the compound inside is realized by depressurizing the mobile phase in downstream of the BPR, which leads to a 500 times higher expansion in the volume. The signifcant expansion can cause the remixing of the compounds that results in the diference between the fnal fraction purity and the observation with the UV detector. The volume expansion will also cause aerosol spray leading to low recovery of the solute and contamination in the device. Aiming at the issue, using cyclone at high-pressure for fraction collection realized by the centrifugal force to complete the modifer separation can efectively improve the recovery of the solute, which has been used by some suppliers such as Jasco, Novasep, and Sepiatec. Berger also developed a phase separator to control the depressurization process to collect the modifier under low pressure, during which $CO₂$ will depressurize from supercritical fluid state to gas state without aerosol formation. The improved SFC with phase separator has been used by Multigram II and Multigram III devices for batch purifcations and Autoprep for series of compounds purifcation. Moreover, Shaimi designed a new process to collect the solute under atmospheric pressure in a series of vessels, which can realize the purifcation of several compounds consecutively. The process relies on two collection heads, of which one collects the solute and another is fushed by the solvent for cleaning it before collecting the next fraction.

Cleaning systems

The cleaning of metal devices, optical and laser components, precision mechanical parts, and electronic assemblies for contaminants removal are extremely important and necessary in the manufacturing industry (Meziani et al. 2017). The traditional industrial cleaning mainly uses organic

solvent cleaning and water solution cleaning. The organic solvent cleaning usually employs volatile solvents and halogen-containing chlorofuorocarbons as the solvent, which causes serious damage to the ozone layer and the atmosphere (Allafi et al. 2021). Meanwhile, the water solution cleaning has the problems of complex surfactant accession, long drying time, treated metal easy to rust, secondary pollution, and also inevitably increasing the treatment costs. As an alternative, $\sec O_2$ with low viscosity, zero surface tension, high gas-like difusivity, and high liquid-like density has becoming an attractive solvent that can be widely used for cleaning of the precision devices (Matz and Reidy 2005). Moreover, $\sec{CO_2}$ can easily reach the hard-to-reach surfaces of the parts with complex geometry and can be absolutely released without any residue. The solvating power of scCO_2 can be adjusted by varying the temperature and pressure as well as adding the cosolvents (Korzenski et al. 2005; Lawandy and Smuk 2014; Weibel and Ober 2003). More importantly, the $\rm scCO_2$ -assisted cleaning process can be completed simultaneously with drying process, which greatly saves the treatment time and cost in the practical production.

At present, $\sec O_2$ cleaning technology has been applied in several fields, such as precision machinery industry, medical system industry, electronics industry, and optical industry, and therefore, the $\sec O_2$ cleaning systems have also attracted increasing attention. $ScCO₂$ cleaning systems are designed based on the traditional wet-cleaning systems and employ $CO₂$ as the cleaning medium to complete the cleaning process under supercritical conditions, and then, the separation of $CO₂$ and contaminants and the drying of the cleaned samples are realized through decompression process. The critical factors of $\sec O_2$ cleaning systems are reasonable design of the high-pressure cleaning vessel to maintain the contaminants stably dissolves into $\sec O_2$ and not produced repeated pollution due to the parameters fuctuation during the $\sec O_2$ cleaning process, which requires high tightness of the cleaning vessel and precise controlling device of the cleaning conditions. Moreover, ensuring no reattaching of the original dissolved contaminants on the surface of the cleaned samples during the separation and drying process is also important to obtain high cleaning efficiency. In the recent years, $\sec O_2$ cleaning systems at laboratory scale and commercial scale both developed rapidly, which will be introduced separately in the following.

Laboratory cleaning system

The typical laboratory $\sec O_2$ -assisted cleaning system is shown in Fig. 10 (Ventosa et al. 2008), which consisted of gas cabinet, high-pressure pump, heat exchanger, cosolvent delivery part, and reaction chamber. Ventosa et al. (2008) studied $\sec O_2$ -assisted cleaning for removal of the postetch residues and oxidized copper formed in the post-via after **Fig. 10** Typical laboratory supercritical carbon dioxideassisted cleaning system. Reproduced with permission of Elsevier from Ventosa et al. (2008) . CO₂ is firstly pumped and pressurized up to the target pressure, and maintained in the process chamber for the residence time. Then, the cosolvent will be pumped and mixed with CO₂ with a fixed proportion, and the mixture will fow into the chamber containing the sample to be cleaned when the mixture reaches the supercritical condition

SiOC-based porous materials etching and photoresist plasma ashing processes by employing this scCO_2 -assisted cleaning system. During the scCO_2 -assisted cleaning process, CO_2 is frstly pumped and pressurized up to the target pressure and maintained in the process chamber for the residence time. Then, the cosolvent will be pumped and mixed with $CO₂$ with a fixed proportion, and the mixture will flow into the chamber containing the sample to be cleaned when the mixture reaches the supercritical condition. The samples will be continuously rotated with a fxed rotate speed to enhance the mass transfer, thus speeding up the cleaning process. The scCO_2 -assisted cleaning process was divided into two steps, i.e., $\sec O_2$ with cosolvent cleaning process and individual $\secO₂$ flowing process for removing the cosolvent and drying of the samples. Finally, $CO₂$ will be released to the room pressure and drove to the exhaust, and the cosolvent will be collected to the vessel. Ito et al. (2004) used the $\rm scCO_{2}$ -assisted cleaning system for $\rm scCO_{2}$ -assisted cleaning of the air filters and found that the removal efficiency of dioctyl phthalate can highly reached 100%. Wang et al. (1999) explored $\sec O_2$ -assisted cleaning of the residuals of Ni (or Sn/Pb) ions from the surface of electroplated terminations and simultaneously realized batch cleaning and flow cleaning with this scCO_2 -assisted cleaning system by employing the metering valve at the exit of the reaction chamber to control the flow rate of $CO₂$ and also using a BPR to adjust the pressure to the target level. The results showed that $\sec O_2$ -assisted cleaning ability was superior than that of the deionized water cleaning and found that higher temperature and pressure led to noticeably improved cleaning ability of scCO_2 . Moreover, the scCO_2 -assisted

cleaning ability can be further improved through the agitation by ultrasonic transducers set in the cleaning chamber (Kohli 2015).

Lin et al. (2010) further optimized the typical laboratory $\rm{scCO_{2}}$ -assisted cleaning system by adding the boil-off vessel located downstream from the cleaning vessel and the syringe-type high-pressure pump subsystem. Diferent from the typical scCO_2 -assisted cleaning process, after the samples soaked and rotated in the $\mathrm{s}\mathrm{c}\mathrm{CO}_2$ and cosolvent mixture under the supercritical condition, $CO₂$ mixed with contaminants will be fushed out of the cleaning vessel through continuous flow of pure $CO₂$ in a first-in–first-out (FIFO) process, in which pure $CO₂$ is pushed by the syringe-type high-pressure pump subsystem at a constant flow rate. The stabilized temperature and pressure in the cleaning vessel during the FIFO fushing and soaking process make the solvent power maintain unchanged, thus minimizing the contamination leftover. After the cleaning process, the contaminants will be left in a flter subsystem. It is worth noting that the decompression process should be slow enough for guaranteeing the temperature in the cleaning vessel will not be too cold due to the Joule–Thomson efect, and the temperature in the boil-off vessel should always maintain slightly higher than the saturated vapor pressure during the decompression process, thus avoiding the bubbles formed that may stir up the contaminants by gravity at the bottom of the cleaning vessel. As a result, most of the bubbles will be formed in the boil-off vessel. Lin et al. (2010) used this improved $\sec O_2$ -assisted cleaning system to study the cleaning ability of the compressed Martian air mix (consisting of 95% $CO₂$) under supercritical condition for removing the trace amounts of microbial and organic contaminants from spacecraft material surfaces. The results demonstrated that $\rm{scCO_{2}}$ -assisted cleaning technology can achieve the cleanliness levels of 0.01 μ g/cm² or less for the cleaning of hydrophobic contaminants, which shows great potential for future Mars in situ cleaning.

Commercial cleaning system

With the increasing attractive attention of $scO₂$ -assisted cleaning technology, various key system manufacturers launched their commercialized $\sec O_2$ -assisted cleaning systems. The TC-SFE-50-1-120S laboratory $\sec O_2$ -assisted cleaning system manufactured by Suofuer is shown in Fig. 11a (Li et al. 2015; Liu et al. 2014). TC-SFE-50-1-120S laboratory $\sec O_2$ -assisted cleaning system is consisted of a 1-L cylindrical stainless steel cleaning vessel with a length of 360 mm and a diameter of 60 mm, which can be operated at temperatures from 15 to 300 °C and pressures up to 50 MPa. The TC-SFE-50-1-120S laboratory scCO_2 -assisted cleaning system also has two series-wound separators used

for the recovery of the cosolvent, contaminants, and $CO₂$. During the scCO_2 -assisted cleaning process, gaseous CO_2 will be transformed into liquid state by condensation and be preheated to the target temperature and then enters the stainless steel cleaning vessel pumped by the high-pressure pump to the pre-set pressure. The temperature in the cleaning vessel is measured by the thermocouple contacted with the fuid inside the vessel to ± 2 °C, and the pressure is measured by the piezometer with an accuracy of \pm 0.2 MPa. The flow rate of $CO₂$ and pressure in the cleaning vessel can be controlled by both the pump and the heated micro-metering valve. Furtherly, the contaminants in the separator can be recovered, recycled, or reused, if suitable, or the contaminants can be disposed as an individual component without any wastewater, solvents, or other residues that may complicate the subsequent treatment process.

The commercial $\sec O_2$ -assisted cleaning system is shown in Fig. 11b. The commercial $\sec O_2$ -assisted cleaning systems have the low operating and waste disposal costs, but installed costs are from \$100,000 for small-capacity (1-L volume) system to several hundred thousand dollars for

Fig. 11 a TC-SFE-50-1-120S laboratory supercritical carbon dioxide ($scCO₂$)-assisted cleaning system manufactured by Suofuer. The TC-SFE-50-1- 120S laboratory $scCO₂$ -assisted cleaning system also has two series-wound separators used for the recovery of the cosolvent, contaminants, and $CO₂$. Reproduced with permission of Elsevier from Liu et al. (2014), and **b** commercial scCO₂-assisted cleaning system designed by Unitech-Annemasse

large-capacity (30-L volume) system, and the installed costs increase with the size, the complexity of the components, and the required automation degree. However, the subsequent contaminants treatment cost in the scCO_2 -assisted cleaning process is lower than that of solvent or water solution cleaning technologies, since the waste residue treated by the $\rm{s}\rm{c}\rm{CO}$ ₂-assisted cleaning process is 100% contaminant without any other residues or solvents. Consequently, for the contaminants readily soluble in $\sec O_2$, the $\sec O_2$ -assisted cleaning is undeniably competitive with the solvent or water solution cleaning technologies (Kohli 2015).

Through the introduction of the foregoing, the $scCO₂$ -assisted cleaning system consists of $CO₂$ cylinder and pump, cosolvent delivery device, temperature and pressure controller, and the cleaning vessel. Moreover, the $\rm scCO_2$ -assisted cleaning system can be further improved by employing the separators which are used for the recovery of the cosolvent, contaminants, and $CO₂$. The scCO₂-assisted cleaning system has been commercialized and widely used for the practical industrial cleaning process. $ScCO₂$ -assisted cleaning technology can meet the contaminants cleaning requirements without compromising the component properties of the devices, and the corresponding scCO_2 -assisted cleaning systems will attract more and more attentions and be applied in practical productions.

Overall, $ScCO₂$ cleaning technology has been considered a precision cleaning technology applied in many diferent industries, especially attractive for cleaning intricate parts with complex geometries due to its fascinating properties of gas-like viscosity, liquid-like density, and low surface tension, leading to high wettability. $SCO₂$ cleaning system has developed rapidly in both laboratory and industry scale. Nevertheless, $ScCO₂$ cleaning system also faces several challenges and problems to be solved and further improved for better applying in the practical production. Firstly, when cleaning a large quantity of particulate contaminants in the $\sec{CO_2}$ cleaning system, the blockage can be easily caused at the valves and pipe joints due to the aggregation of particle contaminants, which can lead to local pressure surges in the pipelines and thus result in serious safe problems. Consequently, installing $CO₂$ flow regulating device to precisely control the $CO₂$ flow rate and particulate contaminants dissolution in $\sec O_2$ can effectively improve the cleaning efficiency. The second challenge of scCO_2 cleaning system is that when separating $\sec O_2$ and the contaminants by reducing the temperature and pressure after the $\sec O_2$ cleaning process, the solubility of contaminants will decrease due to the transformation of $CO₂$ from supercritical state to gas state, which may lead to reattaching of the original dissolved contaminants on the surface of the cleaned samples. Aiming at the issue, low-pressure vessel can be added after the high-pressure cleaning vessel for separating $CO₂$ and the contaminants, while the high-pressure cleaning vessel

can be maintained at constant pressure to avoid reattaching of the contaminants. Moreover, cleaning for large-volume precision instruments still needs to be explored due to the limitations of current volume of the high-pressure cleaning vessel. Developing larger and more flexible $\sec O_2$ cleaning system for cleaning various types of instruments can realize wider applications of $\mathrm{s}\mathrm{c}\mathrm{CO}_2$ cleaning system in the practical industrial production process.

Perspective

 $ScCO₂$ -based technologies with unique advantages have achieved great achievements for dissolving various types of solutes. Although great progress of scCO_2 -based systems has been made, there are still numerous challenges to be conducted, which also means the valuable directions for future development in both fundamental research and commercial applications of scCO_2 -based systems.

First, the standard of various $\sec O_2$ -based systems for diferent applications should be more unitized and normalized. The current $\sec O_2$ -based systems for dissolving different types of solutes have similar components composition but present the diference on the reaction vessel due to the specifc requirements for dissolving diferent types of solutes. More unitized and normalized standard of the systems can realize fexible switching of diferent functions by changing diferent reaction vessel for dissolving diferent types of solutes.

Second, the premiumization and intelligentization of $\sec CO_2$ -based systems are expected to be realized in future. Most of the current $\sec O_2$ -based systems mainly depend on the manual operation, such as setting the amount of cosolvent, controlling the working conditions, and judging the degree of reaction completion. The manual intervention may infuence the accuracy and repeatability of the reaction process, especially under supercritical conditions where minor changes of the temperature/pressure during the reaction process may lead to major variation of the $\sec O_2$ solubility. Therefore, improving the premiumization and intelligentization and reducing the manual intervention of $\sec O_2$ -based systems for more precisely controlling the properties of $\sec CO_2$ during the reaction process are in urgent need.

Third, the long-term stability and durability of $\sec CO_2$ -based systems should be focused and further improved. The $CO₂$ compression process usually accompanied with obvious exothermic process, which may afect the normal operation of the $CO₂$ pressure pump. Meanwhile, the $CO₂$ decompression process is along with endothermic process, thus leading to freezing around the pipeline, which may cause the blockage in pipelines and the damage of reducing valve. Moreover, the scCO_{2} -assisted reactions usually should be maintained at a residence time

under the high-pressure conditions, which greatly requires high air tightness of the reaction vessel. Improving the long-term stability and durability of scCO_{2} -based systems can ensure the stability during the reaction process and reduce the system cost.

Conclusion

Supercritical carbon dioxide (scCO_2) as a sustainable and efficient solvent has been widely applied for dissolving various types of solutes. This review comprehensively and systematically summarized the sustainable-solvent- scCO_2 based systems, including dyeing system, extraction system, chromatograph system, and cleaning system, for fexible and efficient dissolution of different types of solutes, i.e., dyes, target solutes in other solvents, component solutes to be separated or purifed, and contaminants to be removed, respectively. Based on the detailed introduction of scCO_2 -based systems for dissolving diferent solutes, it can be found that $CO₂$ supply and pump part, cosolvent providing part, and parameters controlling part are commonly included in the $\sec O_2$ -based systems. However, the reaction vessel presents diference depending on the specifc dissolving requirements and practical functions. Additionally, this review also introduces the diference between the systems, which matched with diferent production requirements in each application and can be fexibly chosen to apply in the practical production. Through the analysis of the foregoing, focusing and further improving $\sec O_2$ -based systems for flexible and efficient dissolution of various types of solutes based on the application scenarios will render $\sec O_2$ -based technologies be more economic, controllable, fexible, and suitable for wider practical application in future. This review can provide a comprehensive understanding for the construction of scCO_2 -based systems and supply valuable reference for further development and practical industrial applications of $\sec O_2$ -based technologies.

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Declarations

Conflicts of interest The authors declare no confict of interest in this work.

Ethics approval EL declares that he is Chief Editor of Environmental Chemistry Letters.

Consent to participate Not applicable.

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