



Up-Scale of Production and Characterization of Homolog Vivacel from Rice Straw

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Abstract

Up-scale of production of microcrystalline cellulose from rice straw has been done. A total of 1 kg of rice straw powder were made into alpha-cellulose by the method of multistage pulping and hydrolyzing time was 60 minutes with hydrochloric acid 2,5 N and the temperature was 100 °C to produce optimum microcrystalline cellulose. The yield of microcrystalline cellulose which obtained in 60 minutes hydrolyzed was 95 %. Microcrystalline cellulose from rice straw based on the characters test, identification, pH, water soluble substances, solubility, loss on drying, bulk density, and starch test are not significantly different from Vivapur or Avicel PH 102[®], meanwhile for tap density, Carr's index and Hausner ratio are significantly different.

Keywords: microcrystalline cellulose, rice straw, up-scale production, Vivapur or Avicel PH 102.

1. Introduction

Microcrystalline cellulose (MCC) is an important additional ingredient in the pharmaceutical, food, cosmetics, and other industries. In addition, MCC in the pharmaceutical industry is very often used as an excipient in the manufacture of tablets especially for direct compression tablets. Direct compression tablet manufacturing with more and more because it has many advantages such as not using a granulation process, giving a uniform particle size, and make the tablet more stable in a long time, as well as favorable economic terms [1]. In the manufacture of tablets, microcrystalline cellulose (MCC) serves as a binder, filler, and the destroyer. MCC will produce tablets with high hardness, not easily fragile and have destroyed a relatively short time, and can improve the flow properties of the granules [2].

MCC can be produced by reacting cellulose in mineral acid solution boiling for a certain time to limit the degree of polymerization is reached [1]. The process aims to reduce the molecular weight, degree of polymerization, and the long chain to form microcrystalline cellulose. In the process of hydrolysis using hydrochloric acid will decrease the molecular weight of the cellulose from 300000-500000 be 30000-50000. Likewise, the degree of polymerization fell from 2000 - 3000 be 200-300 at MCC, while the length of the cellulose chain dropped from 10000-18000 Armstrong became 1000-1500 Armstrong [2].

Agricultural waste rice straw is available in a number of relatively more who have been wasted which largely burned. In general, rice straw (*Oryza sativa*) and other lignocellulose materials composed of cellulose, hemicellulose and lignin. Cellulose is a polymer of β -glucose with β -1-4 bonds between glucose units. Cellulose found in wood, cotton, hemp and other plants. Cellulose is an organic compound found in the cell



walls together lignin role in cementing the structure of plants. Wood cellulose generally ranges from 40-50 %, whereas in cotton almost 98 % [3].

The purpose of this study was to determine the bioconversion techniques of rice straw into raw materials derived Microcrystalline Cellulose (MCC) i.e. homologous Vivacel or Avicel. Avicel is a pharmaceutical raw material which is very important and widely used as additives or matrix drugs in the pharmaceutical industry, agriculture and packaging. In previous studies we have managed to find a way to convert rice straw into homologous Avicel in common with more than 75 % [4, 5]. This research is still in the laboratory scale. The yield obtained is not optimum, so it needs to be optimized bioconversion processes, in order obtained bioconversion technology which can be upgraded for production on a larger scale. However, MCC has been obtained at the optimal hydrolysis time for 60 minutes using 2.5 N HCl at a temperature of 100 °C from rice straw [6]. In addition, it is also necessary to determine the physical properties, chemical and physicochemical properties of the resulting product, in order to qualify as modern pharmaceutical raw materials [7].

2. Materials and Methods

2.1 Materials

Rice straw powder, distilled water, ethanol (Brataco®), nitric acid (Brataco®), sodium hydroxide (Brataco®), sodium sulfite (Brataco®), sodium nitrite (Brataco®), sodium hypochlorite (Bayclin®), hydrochloric acid (Merck®), zinc chloride (Merck®), potassium iodide (Merck®), hexane (Brataco®), copper(II) sulphate (Merck®), ammonia (Merck®), iodine (Merck®), and Avicel PH 102 (Rettenmeyer®).

2.2 Equipment

Water bath, desiccator, measuring glass, beaker, test tubes, filter paper (Whatmann 42), flask, watch glass, spatula, vaporizer cup, oven, pH meter, analytical balance (Mettler PM200®), round-bottom flask, condenser, electric stove, Jasco FT IR spectrometer 460+, and tap volumeter.

2.3 Procedure

2.3.1 Sample preparation

Rice straw is chopped and washed several times with water, then dried at 60 °C for 24 hours and pulverized in a blender. Rice straw powder was weighed as much as 1 kg, refluxed with a mixture of hexane and ethanol in a ratio of 2:1 for 6 hours, then allowed to cool and filtered. Furthermore, the remaining is dried at room temperature [6].

2.3.2 Extraction of alpha-cellulose with multistage pulping method

One kg rice straw powder was mixed with 13.4 L 3.5% nitric acid (containing 40 mg of sodium nitrite) in a beaker. The mixture in the container was then placed in a water bath for 2 hours at a temperature of 90 °C. The next part of the insoluble was separated by filtration and the residue obtained was washed with distilled water. The residue was immersed in a 10 L solution containing sodium hydroxide and sodium sulfite respectively of 2 % w/v at 50 °C for 1 hour. Then the mixture was filtered and washed again as described above to obtain a residue. The residue was bleached by mixing them into 8 L mixture of water and sodium hypochlorite 3.5 % w/v (the ratio of water and 3.5 % sodium hypochlorite solution is 1:1), then boil for 10 minutes, followed by filtration and washing. The residue obtained from filtering was mixed with 8 L of sodium hydroxide 17.5 % w/v and heated at a temperature of 80 °C for 30 minutes. Then the mixture was filtered and washed. The residue obtained is alpha-cellulose. The extraction process was continued by mixing alpha-cellulose into 4 L mixture of water and sodium hypochlorite 3.5 % w/v (the ratio of water and 3.5 % sodium hypochlorite solution is 1:1), and heated at 100 °C for 5 minutes. This mixture was filtered and washed with distilled water to obtain a residue which was clean. The residue is then dried at a temperature of 60 °C in order to obtain alpha-cellulose [6].



2.3.3 Production of Microcrystalline Cellulose (MCC)

A total of 500 g of alpha-cellulose was introduced into the beaker and hydrolyzed by heating in 2.5 N HCl (12 L) for 60 minutes. The hot mixture was poured into cold water while stirring strong by using a spatula and allowed to stand overnight. Microcrystalline cellulose formed was then neutralized again by washing it in water, then filtered, dried in an oven at a temperature of 57-60 °C for 60 minutes and then crushed. MCC obtained was stored in at room temperature in a desiccator [6].

2.3.4 Characterization of Microcrystalline Cellulose

Characterization of microcrystalline cellulose include organoleptic examination, identification, measurement of pH, determination of substances soluble in water, measuring the solubility in a mixture of ammonia-copper (II) sulphate, determination of drying shrinkage, determination of real density and compressed density, determination of Carr's index and Hausner ratios, starch test, and determination of FT IR spectra [1, 8, 9].

3. Results and Discussion

Up-scale of microcrystalline cellulose manufacture was done by hydrolyzing alpha-cellulose with 2.5 N hydrochloric acid solution and then be hydrolysis for 60 min. Alpha-cellulose was obtained as much as 560 grams (56 %) of 1 kg of dry rice straw. Results of up-scale production of alpha-cellulose were obtained a yield of MCC 96 % (see Table 1).

Table 1: Results of microcrystalline cellulose from rice straw

Total of rice straw powder (kg)	Results	
	Alpha-cellulose (g)	Microcrystalline cellulose (MCC) of alpha-cellulose
1	560 g (56 %)	96 %

Selection of rice straw to make microcrystalline cellulose was because rice straw contains a lot of cellulose in the stalks. In addition, rice straw was also easily found and many discarded as agricultural waste. Rice straw was used only as fodder. The sample size of rice straw affects the acquisition of microcrystalline cellulose. The greater the amount of rice straw the better microcrystalline cellulose obtained. Conversely the smaller the rice straw it would be a little too microcrystalline cellulose obtained. This is because the finer the sample the more easily cellulose disconnected at the time of multistage pulping so more depolymerization of cellulose to form monosaccharide.

The result of up-scale of alpha-cellulose hydrolysis into MCC has been compared with Avicel PH 102®. Statistically the average measurements of the sample were tested by t-test and non-parametric test of independent samples to see the difference between MCC and Avicel PH 102. The MCC produced meets the requirements of the United States Pharmacopoeia and British Pharmacopoeia.

Comparisons of characters ranging from organoleptic inspection include shape, color, smell, and taste. The results obtained showed similarities to one another as well as meeting the requirements of the Pharmacopoeia (see Table 2). On the results of the identification with iodinated zinc chloride solution, MCC showed positive results. The reagent was a reagent specific to microcrystalline cellulose. Avicel PH 102® also gave similar results (Table 2). Measurement of pH of MCC has a value of eligible British Pharmacopoeia. The pH test results do not differ much from the comparison Avicel pH 102®. Substances dissolved in water, the maximum residue weight was not more than 0.25 % of tests performed [9]. Measurement of solubility of MCC and Avicel PH 102® in a solution of copper ammonium sulfate tetramine was also performed. The results were in accordance with the requirements that do not leave residual dissolved (see Table 2) [9].

Table 2: Examination of characterization of microcrystalline cellulose and Avicel PH 102

No	Examination	MCC	Avicel PH 102
1	Organoleptic [8]	Fine powder, white, odorless, tasteless	Fine powder, white, odorless, tasteless
2	Identification [7]	Blue violet	Blue violet
3	pH [9]	7.15 ± 0.06	7.15 ± 0.03
4	Water soluble substances (%) [9]	0.095 ± 0.017	0.126 ± 0.014
5	Drying shrinkage (%) [9]	4.490 ± 1.028	5.287 ± 0.090
6	Solubility in ammonia-copper sulfate [9]	Dissolved	Dissolved
7	Real specific gravity	0.397 ± 0.046	0.397 ± 0.005
8	Specific gravity of compressed	0.526 ± 0.041	0.582 ± 0.010
9	Carr's index [1]	24.425 ± 2.219	31.751 ± 1.704
10	Hausner ratio [1]	1.314 ± 0.035	1.465 ± 0.037
11	Starch test [9]	Negative	Negative

MCC drying shrinkage value is still below the limit values allowed on British Pharmacopeia, which is 6 %. When compared with Avicel PH 102®, it can be concluded that there is no significant difference in drying shrinkage MCC. On the measurement of the real specific gravity of MCC and Avicel PH 102® there is no significant difference in the real specific gravity between MCC and Avicel PH 102. While the specific gravity of compressed there is a real difference between the MCC and Avicel PH 102.

Hausner ratio and Carr's index is an indirect way of measuring to see the flow properties of powders and compression properties. Hausner ratio indicates the friction between the particles, while Carr's index showed the ability of the powder to reduce its volume or referred to as compressibility. In general, the ratio of Hausner was > 1.25 shows the flow properties were poor. Carr's index-value $< 16\%$ indicates good flow properties and $> 35\%$ showed the strength and cohesion of particles [10].

The compressed nature of microcrystalline cellulose can be caused by hydrogen bonds between the hydroxyl groups plastically changing its form around the particles. During compression, MCC may be suspected because deformation stress release by some mechanism that produces hard tablets with low tensile force [11].

Indeed pure cellulose does not contain starch in it. This can be tested by the presence of starch in cellulose by reacting cellulose with iodine. According to the requirements, the results obtained in the test color, starch is not formed in blue. This occurs because the non-occurrence of the absorption of iodine by the cellulose so no blue coloration as in starch [9]. The test results showed that MCC and Avicel PH 102® meet these requirements.

Test results show the FT IR spectrum of MCC lies in wave numbers which are almost the same as Vivapur 101C. The fingerprint region that is located on the wave number $1600 - 900 \text{ cm}^{-1}$ have similarities peak between MCC and Vivapur 101C (see Figure 1-2). Thus the MCC has the same functional group with Vivapur 101C (see Table 3).

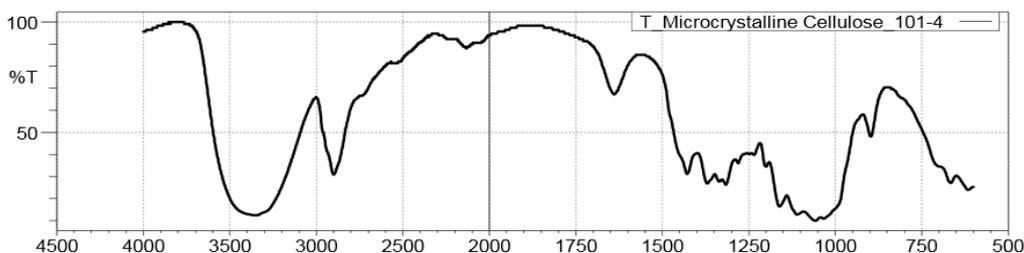


Figure 1: Infrared Spectrum of Vivapur 102

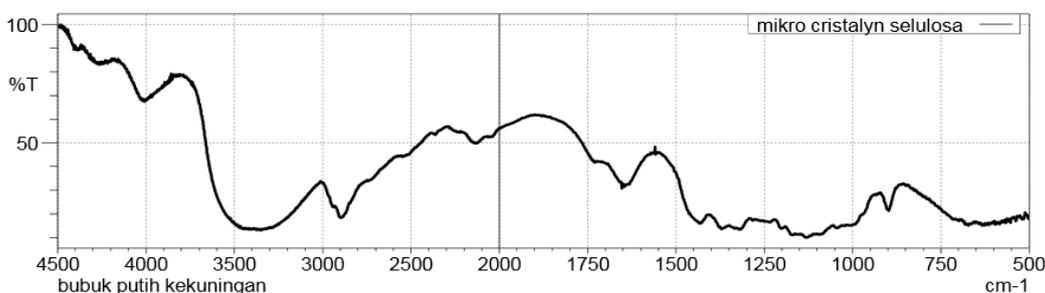


Figure 2: Infrared Spectrum of Microcrystalline Cellulose (MCC) from Rice Straw

Table 3: FT IR Library of microcrystalline cellulose

	Score	Library	Name	Comment
1	840	41 - T_FoodAdditives2	T_Microcrystalline Cellulose_101-4	Microcrystalline Cellulose(Product name;VIVAPUR101CSales origin;TOAKASEI CO.,LTD.)@KBr Wafer
2	814	42 - T_FoodAdditives2	T_Microcrystalline Cellulose_102-4	Microcrystalline Cellulose(Product name;VIVAPUR102CSales origin;TOAKASEI CO.,LTD.)@KBr Wafer
3	812	49 - T_FoodAdditives2	T_Powdered Cellulose-4	Powdered Cellulose(Product name;VITACEL L-600CSales origin;TOAKASEI CO.,LTD.)@KBr Wafer
4	797	20 - T_FoodAdditives2	T_Carboxymethyl Cellulose Calcium-4	Carboxymethyl Cellulose Calcium(Product name;E.C.G-FACSales origin;Gotoku CHEMICAL CO.,LTD.)@KBr Wafer
5	778	43 - T_FoodAdditives2	T_Microfibrillated Cellulose_200L-4	Microfibrillated Cellulose(Product name;CELISH FD-200LSales origin;Daicel Chemical Industries Ltd.)@KBr Wafer

Hydrolysis optimization results have been obtained in the MCC (Figure 3 - 4), which is about 96 % by the hydrolysis time of 60 minutes, using a 2.5 N HCl at a temperature of 100 °C. This shows that the longer hydrolysis time, the more the interruption of the chain of cellulose, whereas cellulose is a polysaccharide that

has a chain length when the hydrolysis time of over 60 minutes the lower the degree of polymerization because the number of polymers or chains are broken off so that the result obtained was can be reduced.

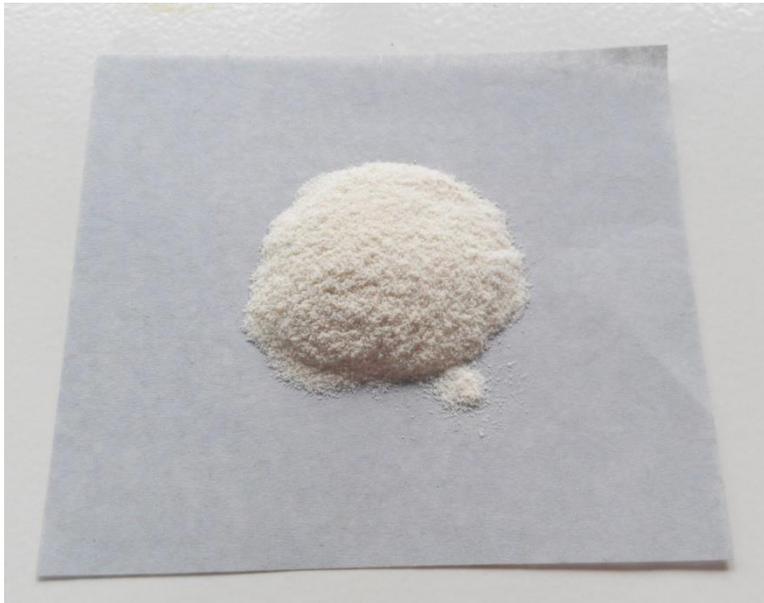


Figure 3: Photo of microcrystalline cellulose (MCC) from rice straw



Figure 4: Microscopic photo of microcrystalline cellulose (MCC) from rice straw

4. Conclusion

Microcrystalline cellulose has been obtained by up-scale production at the optimal hydrolysis time for 60 minutes using 2.5 N HCl at a temperature of 100 °C from rice straw. Microcrystalline cellulose from rice straw is not significantly different from Vivacel by organoleptic inspection, identification, pH, and dissolved



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substances in water, solubility, drying shrinkage, apparent specific gravity, starch test, and have as well as the infrared spectrum. In testing the density of compressed, Hausner ratio and Carr's index there is a noticeable difference between the microcrystalline cellulose from rice straw and Vivacel.

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