

# Heparin Sodium Was Prepared from Pig Intestinal Mucosa by Dialysis and Spray Drying

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## Abstract

A method to extract crude heparin sodium from pig intestinal mucosa by dialysis and spray drying was established. The pig intestinal mucosa was treated in the following steps: enzymolysis, resin exchange adsorption-washing, elution, pressure filtration, dialysis, spray drying. Activity of the product was measured using a heparin anti-IIa factor assay kit. The yield of crude heparin obtained by this method was 2.79% higher than that of oven drying method; the production of 1 kg crude heparin sodium saved 43.4 pigs small intestine. The activity was  $98.48 \pm 2.49$  IU/mg ( $n = 5$ ), 15.18 IU/mg higher than that obtained by oven drying method. The product is pale white powder, attractive color and easy to dissolve.

## Keywords

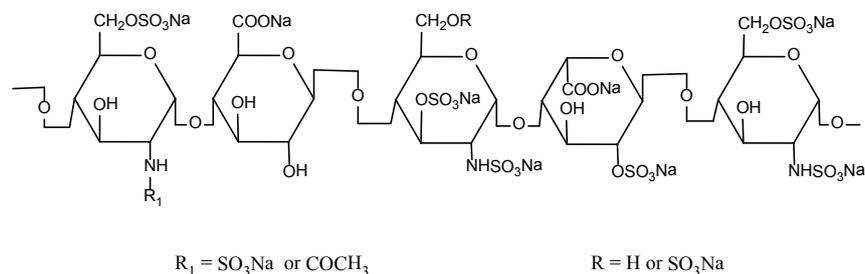
Heparin Sodium, Dialysis and Spray Drying, Oven Drying Method, The Production Process

## 1. Introduction

Heparin sodium is the sodium salt of heparin; it is a kind of natural mucopolysaccharide sulfate ester with complex structure extracted from biological materials. Its basic framework is a polysaccharide chain linked by a glycosylation bond between uronic acid and glucosamine. It generally forms a complex with proteins in animals [1]. The structure of heparin sodium is generally considered as **Figure 1**.

Heparin is a sulphated polysaccharide with a high affinity for antithrombin. When combined with heparin, antithrombin exhibits a fast acting and potent inhibitory activity for coagulant serine esterases: IXa, Xa and thrombin. Low Molecular Weight Heparin (LMWH), and heparin analogues inhibit more efficiently

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**Figure 1.** Heparin sodium structure formula.

Factor Xa than thrombin [2], whilst unfractionated heparin (UFH) inhibits efficiently thrombin and also the other serine esterases. Heparin sodium is widely distributed in the intestinal mucosa, lung and liver of mammals, and its molecular weight is generally between 3000 and 30,000 Da [3]. Heparin sodium is of great medical and biological value [4]. It can prevent and treat arterial and venous thrombosis and pulmonary embolism. It is used as an anticoagulant in artificial heart and lung, peritoneal dialysis or hemodialysis [5] [6] [7].

In the 1980s, heparin research group of the department of biology of Si-Chuan University promoted the salt-hydrolysis process to extract heparin sodium from pigs' small intestine. Generally, it takes 3500 pigs' small intestine to produce 100 million units. In the 1990s, heparin was produced by enzymatic hydrolysis, which typically required 2500 small pigs to produce 100 million units. At present, these two processes are in use, and the most used process is enzymatic hydrolysis/salt hydrolysis-resin adsorption-elution-alcohol precipitation-drying-grinding. After decades of optimization, 100 million units of heparin sodium can be extracted from the 1400 - 1700 pigs' small intestine. The conventional process is simple, but it has many disadvantages, such as poor stability of the product, uneven color, great damage to the product, and a large amount of alcohol is needed to precipitate heparin [8]. Dialysis and spray drying provide the possibility to solve the above problems [9]. Therefore, the two steps of alcohol precipitation and drying in the production process of heparin sodium were improved by dialysis and spray drying in this article.

Anti-IIa assays are then the right methods for measuring the anti-thrombin activity of large heparin molecules [10]. The Heparin Anti-IIa method is a two-stage method based on the inhibition of a constant amount of Thrombin (IIa), by the tested heparin in presence of exogenous antithrombin, then hydrolysis of a Thrombin specific chromogenic substrate, by Thrombin in excess. pNA is then released from the substrate. The amount of pNA released is then a relation of the residual Thrombin activity. There is an inverse relationship between the concentration of heparin and color development, measured at 405 nm.

## 2. Materials and Methods

### 2.1. Materials and Reagents

Pig intestinal mucosa (*C.P. Food (Xiangyang) Co. LTD.*), sodium chloride

(*Tianjin Beilian Fine Chemicals Development Co. LTD.*), alkaline protease (*Solarbio*), sodium hydroxide (*Chengdu Kelong Chemical Reagent Plant*), absolute ethyl alcohol (*Tianjin Yongda Chemical Reagent Co. LTD.*), heparin sodium standard (*Scientan*), 2000 Da ultrafiltration membrane (*Solarbio*), heparin anti-IIa factor assay kit (*Hyphen BioMed*), D254 resin (*The Dow Chemical Co.*).

R1 (Human Antithrombin, lyophilized vial containing about 1.25 IU/mL), R2 (Purified human Thrombin, mainly in the form, lyophilized vial containing about 120 IU/mL), R3 (Chromogenic substrate specific for Thrombin, lyophilized vial of about 6.25  $\mu$ mol, in presence of mannitol), R4 (Assay reaction buffer Tris 0.05 mol/L, NaCl 0.175 mol/L, EDTA 0.0075 mol/L, at pH 8.40, containing Bovine Serum Albumin (BSA) at 0.2% and sodium azide as preservative), R5 (Citric Acid at 20 g/L).

## 2.2. Instrument and Equipment

1/1000 balance (*OHAUS*), spray dryer (*YAMATO*), constant temperature drying oven (*Sartorius*), 200  $\mu$ L - 1 mL pipettor (*Sartorius*), magnetic stirrer (*IKA*).

## 2.3. Method

### 2.3.1. Heparin Sodium Extraction Steps

1) The volume of fresh porcine intestinal mucosa was taken and added to the reaction pot. Added salt to the reaction pot to make the solution salt concentration of 5%. The pH was adjusted to 8.5 - 9.5 with 5 mol/L sodium hydroxide. The temperature was stirred up to 55°C. The solution was hydrolyzed by stirring for 1 h. If the salt concentration or pH value decreased, the salt and sodium hydroxide solution were added to adjust. Four over ten thousand of the protease was added, stirred and dissolved, kept for 2 hours, heated to 90°C, and stopped heating. The decomposition solution was filtered with 120 mesh filter cloth while it was hot.

2) The filtrate was cooled to about 50°C and pH was adjusted to 8.5 - 9.5. The treated resin was combined with the filtrate and the resin content was 6% of the filtrate weight. Stirred for 6 - 8 hours, stirring should not be too fast or too slow to make the resin up and down, otherwise the adsorption effect is poor. After adsorption, filtrate the resin with 120 mesh filter cloth.

3) The filtered resin was rinsed 2 - 3 times with 38°C - 42°C warm water of twice resin volume to remove fat and impurities adsorbed on the resin surface. The resin was then stirred and washed for 30 minutes with 5% salt solution of the same volume as the resin to remove low molecular weight heparin and some proteins that were not tightly bound to the resin. The resin was then filtered out with 120 mesh filter cloth.

4) For the first elution, the filtered resin was mixed with 22% salt solution in a volume ratio of 1:1, stirred and eluted for 2 hours, then the resin was filtered out to collect the filtrate. For the second elution, the concentration of salt solution was 18%, stirred for 1 hour, the filtrate was combined with the first elution filtrate.

5) The collected filtrate passed through the membrane of 4.5  $\mu$ m to remove

particulate impurities and precipitates, then collected the filtrate.

6) The filtrate was collected into the dialysis bag, dialyzed for 16 hours, changed water every 2 hours to remove the impurities with molecular weight less than 2000.

7) The obtained dialysate was spray dried and crude heparin sodium was obtained.

### 2.3.2. Activity Determination of Crude Heparin Sodium

Into the micro well or the plastic test tube, incubated at 37°C, introduce:

1) 40 µL reference material or tested sample and 40 µL *R1* was added to the micropore respectively, then mixed and incubated at 37°C, for 2 minutes.

2) 40 µL *R2* was added to the micropore, mixed and incubated at 37°C, for exactly 2 minutes.

3) 40 µL *R3* was added to the micropore, mixed and incubated at 37°C, for exactly 1 minute.

4) 80 µL *R5* was added to the micropore, mixed and measured the absorbance at 405 nm against the corresponding blank.

## 3. Result and Discussion

### 3.1. Effect of Dialysis and Spray Drying on the Appearance of Crude Heparin Sodium

As shown in **Figure 2**, **Figure 2(a)** shows directly dried samples, while **Figure 2(b)** shows spray dried samples. The directly dried samples are plate shaped and brown in color. The spray-dried samples are powder and pale white in color. Compared with both of them, the latter is more beautiful in terms of traits and colors. And the step of grinding the samples of former into powder, the product obtained by spraying has a higher degree of powder and is more conducive to product dissolution.

### 3.2. Effect of Dialysis and Spray Drying on the Yield of Crude Heparin Sodium

The crude heparin sodium was extracted with 10 L intestinal mucosa by drying



**Figure 2.** The appearance of crude heparin sodium obtained by directly drying and spraying.

method and spray drying method, and the yield of crude heparin sodium was as shown in **Table 1**. Generally, 2 pig bismall intestines can produce 10 L intestinal mucosa fluid, and 1kg crude heparin sodium obtained by spray drying method is less 43.4 pig small intestines than that obtained by drying method. This may be due to the fact that heparin sodium is dissolved in the eluent during alcohol precipitation without complete precipitation, while the crude heparin sodium is completely dried into a powder by spray drying.

### 3.3. Effects of Dialysis and Spray Drying on the Potency of Crude Heparin Sodium

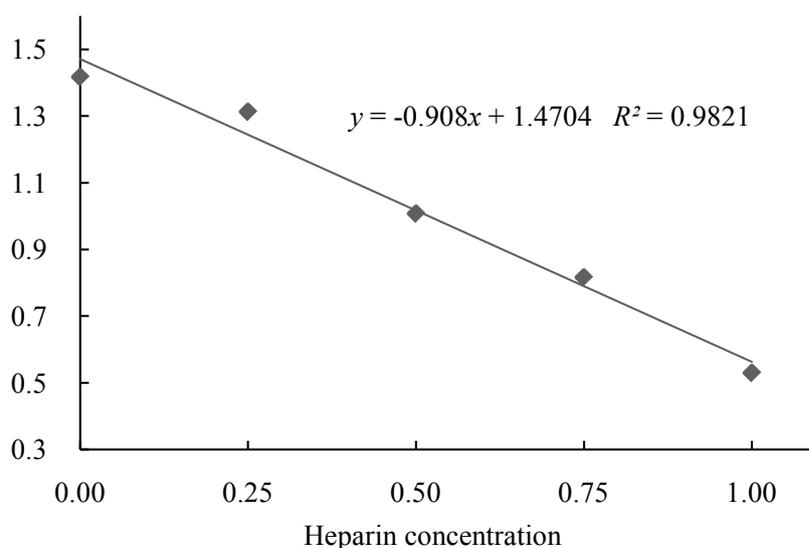
The standard heparin sodium was diluted into a series of gradient, and the absorbance value was measured using the heparin anti-iiia factor determination kit, and the standard curve was drawn as shown in **Figure 3**.

The absorbance of crude heparin sodium obtained by alcohol precipitation drying method and spray drying method was taken into the standard curve and the potency obtained is shown in **Table 2**. The titer of crude heparin sodium obtained by alcohol precipitation drying method was 83.3 IU/mg. The relative standard deviation was 1.36% ( $n = 5$ ), and the titer of crude heparin sodium

**Table 1.** Effect of dialysis and spray drying on the yield of crude heparin sodium.

Technology	Number of small intestine	Volume of intestinal mucosa fluid/L	Weight of crude heparin sodium/g	Number of small intestine to produce 1 kg crude heparin sodium
Alcohol sedimentation	2	10	1.253 ± 0.025	1596.8
Spray drying	2	10	1.288 ± 0.038	1553.4

Absorbance values/OD<sub>405</sub>



**Figure 3.** The potency of standard heparin sodium.

**Table 2.** The titer of crude heparin sodium obtained by alcohol precipitation drying method and spray drying method.

Method	Potency (IU/mg)	RSD/%
Alcohol precipitation drying method	83.3	1.36
Spray drying method	98.48	2.53

obtained by spray drying method was 98.48 IU/mg. The relative standard deviation was 2.53% ( $n = 5$ ). The activity of the heparin sodium is 18.2 percentage points higher than that of alcohol precipitation drying method. This may be due to the removal of small molecular impurities with a molecular weight less than 2000 D during dialysis.

#### 4. Conclusion

This method breaks down the traditional process of preparing crude heparin sodium. First, the collected eluent passed through the membrane of 4.5  $\mu\text{m}$ . Then the filtrate was dialyzed. Finally, spray drying was carried out. The yield of crude heparin was 2.79% higher than that of oven drying method; the production of 1 kg crude heparin sodium saved 43.4 pigs small intestine. The activity of crude heparin sodium was  $98.48 \pm 2.49$  IU/mg ( $n = 5$ ), 15.18 IU/mg higher than that obtained by oven drying method. The product is pale white powder, attractive color and easy to dissolve.

#### Conflicts of Interest

The authors declare no conflicts of interest regarding the publication of this paper.

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