

SHORT COMMUNICATION

Variation of dietary fibre content and gel-chromatography profile of the fruit pulp of four morphologically different fruit types of palmyrah

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Abstract: Palmyrah (*Borassus flabellifer*) is a common plant in Sri Lanka. Its fruit pulp (PFP) has active constituents, which are steroidal saponins called flabelliferins. Although these flabelliferins may play a minor role in the hypocholesterolaemic effect, it is clear that other important factors are also involved. This study investigates the content of the soluble and insoluble dietary fibre and some characteristics of the soluble dietary fibre from the pulp of four common types of palmyrah. Results showed that dietary fibre content was high (12.3% to 24.3% Dry weight) but type dependent. The pectin and soluble dietary fibre of the four types showed distinct elution patterns on sepharose (CL-2B-300) gel-chromatography indicating different molecular weight profiles.

Keywords: Dietary fibre, palmyrah fruit pulp, pectin, sepharose gel chromatography.

INTRODUCTION

The palmyrah palm (*Borassus flabellifer*) is widely distributed in the arid zones of Sri Lanka, South East Asia and East Africa. Palmyrah fruit pulp (PFP) is consumed widely in Sri Lanka as dried fruit leather, jams, beverages, etc. Palmyrah cultivars can be categorized into four common types according to the morphology of fruits that have been described previously in detail¹. It has been shown that depending on the fruit type, the profiles of bioactive compounds in PFP e.g. flabelliferins (steroidal saponins) vary¹. Past studies have shown that the PFP contains pectins with methoxy value of 3.4². Pectins that are classified as soluble fibres³ are polymers of galactouronic acid together with methoxy/ethoxy derivatives of galactouronic acid and are of commercial value in the food industry. Dietary fibre which comprises soluble dietary fibre (SDF) and insoluble dietary fibre (IDF) is well known as a serum cholesterol lowering agent³ and has many other nutritional benefits such as lowering glycaemic index and the incidence of large bowel disease.

The objectives set for this study were to quantify the soluble and insoluble dietary fibre contents in PFP of the four common morphologically different fruit types and to determine whether there is any variation in their pectin and soluble fibre profiles by subjecting them to sepharose-gel chromatography.

METHODS AND MATERIALS

Palmyrah fruits (n=80) were collected from Kalpitiya in the North-West of Sri Lanka and separated into four common types¹ (n=10-30) given below, bagged separately and transported to Nugegoda (nearly 200km away) where they were stored at -20°C.

Type I: Colour, black, pericarp rough with brown longitudinal striations, distal side black¹;

Type II: Colour black, pericarp smooth and no striations, distal side three orange spots¹;

Type III: Colour black and orange longitudinal stripes, pericarp smooth, distal side black and orange stripes¹;

Type IV: Colour orange, pericarp smooth¹.

Within one week of collection, PFP was extracted from fruits manually⁴ using water in the ratio of 2:1 (v/w) with respect to PFP. The pulp was stored in high-density polyethylene bags at -20°C. The moisture content of PFP was determined in duplicate using the AOAC and Dean and Stark method⁵. The extraction of dietary fibre (insoluble and soluble) was carried out in duplicate using the enzymic method described by Asp *et al.*⁶, from PFP (10 g) of each type, sampled from homogenous samples of 2-3 kg obtained from 10–20 fruits each. Soluble dietary fibre (SDF) was isolated following the same procedure i.e

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by methanol precipitation, after the insoluble dietary fibre was separated. The filtration of SDF precipitate was carried out using filter paper instead of sintered glass crucibles and the incineration step of the Asp⁶ method was omitted since the SDF had to be isolated and could not be burned off. The SDF was freeze dried and weighed. The SDF samples (0.05-0.1g) were dissolved in phosphate buffer (0.1M) in volume of 2ml and introduced into the Sepharose column (CL-2B-300) of 20ml (height = 21cm). The same phosphate buffer was used as the eluting solvent. Fractions were collected and freeze dried. The soluble fibre of each fraction was determined gravimetrically after freeze-drying (subtracting the weight of the buffer blank from each fraction). The freeze-dried fractions dissolved in water (0.25 ml), were then assayed by the carbazole reaction, after primary and secondary hydrolysis with H₂SO₄⁷.

RESULTS AND DISCUSSION

The insoluble and soluble dietary fibers were determined separately and in duplicate. Soluble dietary fiber of type I, II, III and IV were 9.0, 10.6, 5.5 and 8.0% dry weight PFP respectively while the insoluble dietary fiber for the four types were 8.8, 11.9, 6.8 and 16.3% dry weight PFP. There was also a difference in the visual appearance

of SDF of the 4 types: Type I gel-like, Type II soft, cloudy, Type III gel-like, Type IV gel-like, cloudy.

Pectin content of types I, II, III and IV were 7.2, 7.8, 5.3 and 7.2% dry weight of PFP respectively, showing that pectins comprise the major fractions of soluble dietary fibre, in each case. Figure below shows the profiles of pectins obtained from the sepharose column. As expected with carbohydrate polymers, the pectins showed polydispersity. Type I showed the lowest elution volume fraction and type II and III distinctly shifted to higher elution volume (lower molecular weight). Blue dextran molecular weight about 2 million eluted at 8-10 ml.

Pectin content was 5.3-7.8%. This is more than the amount reported previously (4-6% dry weight²). The better range of sampling adopted in this study, gives a more sound basis for pectin content of the common PFP. This is important, as isolated pectin is commercially valuable in food industry.

Separation of pectin on sepharose column gel chromatography resulted in different elution profiles indicating that there is a varied distribution of molecular weight of pectins in the different types. The more gel like visual appearance could be due to the high molecular

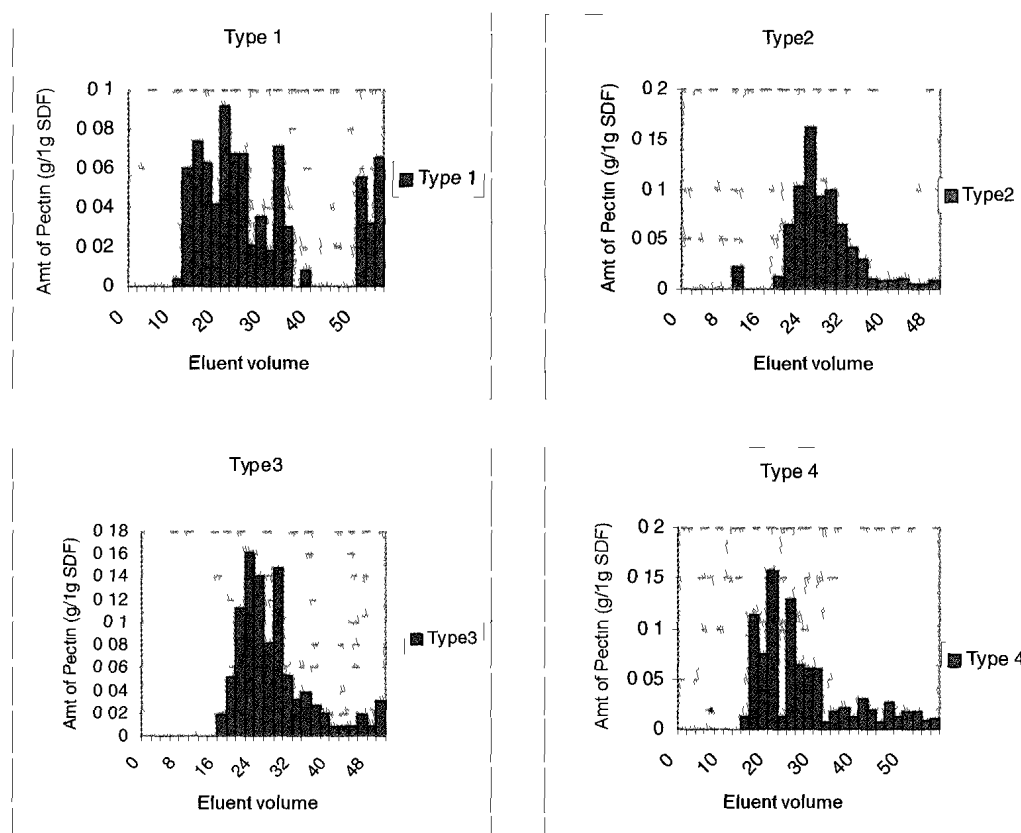


Figure. Pectin profiles of four common types of PFP

weight molecules that are present in type I and type IV that have lower elution volumes and are highly viscous. It was not possible to determine the molecular weight ranges as no monodisperse carbohydrate standards, except for blue dextran (2 million Da) was available. Results indicated that type I had pectin molecules of MW 2 million

It is clear that high dietary fibre (both SDF and IDF) can result in lowering of cholesterol³. This will be significant as PFP is used as a food. Highly viscous pectins also have a major effect on binding bile salts and preventing their reabsorption, a known serum cholesterol lowering effect⁸. The difference observed in SDF, IDF and pectin could, to a certain extent, explain the varied lowering of serum cholesterol in mice fed with 10% PFP of these four types of PFP compared to the controls⁹.

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