STUDY OF THE PHYSICAL PROPERTIES, SAFETY AND EFFECTIVENESS OF

AN ORGANIC OINTMENT

Abstract

**Introduction:** Women have always used cosmetics to maintain their hair in order to become

pretty.

Aim: To help this category of people with non-harmful ointments, anointment made from

palm kernel oil and shea butter was developed.

Method: In the presence of water, the palm kernel paste was boiled at 100°C to bring the oil

to the surface of the water. The supernatant is collected in a kettle, and after cooling and

filtration is collected in a bottle. The shea butter (used as an excipient) is melted in a bain-

marie, then the required quantity is added to the corresponding palm kernel oil in a container.

The mixture is homogenized until a single phase is obtained, then poured directly into 150mL

jars. The jars were then left to stand at a temperature below 30°C until the contents solidified.

**Results:** Physical, chemical, microbiological and other tests showed that the ointment was

suitable use.

Conclusion: The organic ointment showed better performance in hair growth in rabbits and

could be a fundamental element in the formulation of ointments for human hair growth.

**Key words**: Oil, Palm kernel, Shea, Ointment.

**INTRODUCTION** 

Hair is one of the elements of human hair found on the head. Hair is made up of keratin. As a

whole, hair forms a protective barrier against external aggressions (shocks, sunrays, rain...). It

also acts as a thermal insulator, with the scalp's sweat glands absorbing excess moisture to

maintain the surface temperature of the area it covers.[1]. In addition to its aesthetic character,

1

hair is a symbol of strength and virility for men, and femininity and seduction for women[2]. In these conditions, hair loss is always a source of concern for both men and women, especially when the problem persists. Many people affected by early hair loss try to hide it by any means possible, such as wearing a wig, weaving or wearing a hat, especially as these people no longer fully benefit from its protective action, but also from psychological and social suffering. Men no longer feel virile, while women feel ugly and have lost their femininity. These people are often subject to a masked form of mockery, stigmatization and discrimination in society. Generally speaking, the consequences of hair loss are: loss of selfesteem, difficulty in withstanding the gaze of others, and isolation. [1]. However, there are several solutions for combating premature hair loss, such as taking certain pills, using hair lotions, hair transplants and hair ointments. With regard to ointments, authors have formulated a dermal ointment based on *Mitracarpusscaber* extracts [3]. Formulation of an analgesic and anti-inflammatory ointment based on the same medicinal plantwas initiated [4]. Another team of researchershaveformulated an antimicrobial ointment based on hydroethanolic extract from Bersama abyssinica Fresen to combat skin ailments [5]. It is in this context that plant species such as *Elaeisguineensis* Jacq caught our attention because in several regions of Ivory Costpharmacopea it is commonly used by rural populations for hair maintenance and growth.,

### I.MATERIAL AND METHODS

### I.1.MATERIAL

### **Plant material**

The plant material consisted of *Elaeisguineensis* kernels from the Arecaceae family (Figure 1), obtained after removing the pulp from a few households. Originally from tropical Africa, oil palm is widely cultivated in tropical zones, particularly in Asia. 70% of these plantations

are owned by village planters. These plantations are located in the south of the country, along the coast up to almost 6° north latitude.



Figure 1 Oil palm tree

# Technical equipment

Laboratory equipment and materials

The main laboratory equipment and materials used in this work are:

- Precision balance, Precisa® JB 12000C (capacity = 2100 g sensitivity = 0.1 g),
   (Switzerland);
- - autoclave;
- - Memmertoven (Germany);
- - HANNA® HI 8010 pH meter;

- Glassware: sight tubes, Petri dishes, 1 ml pipettes, 100 mL graduated burette,
   beakers;
- Retsch SK 100 milling machine.
- Culture media and reagent
- PCA (Plate Count Agar);
- BEA (Bile EsculinAzide);
- Sabouraud;
- VRBG (Glucosée Biliée au CristalvioletandAurouge Neutre).

### I.2.METHOD

### **I.2.1.** Production of palm kernel oil

The processing stages are: reception, drying (for a week) and crushing of the pulped seeds, isolation and grouping of the kernels in a container. The kernels are then crushed (using a grinder) and finally the oil is extracted.

The oil extraction process involved boiling the palm kernel pulp in the presence of water. At the boiling temperature of the water (100°C), the oil begins to appear on the surface of the water. All the supernatant is collected in a jar.

The jar is brought to the boil until all the water has evaporated, leaving only the oil which, after cooling, is filtered and collected in a bottle.

## **I.2.2.** Ointment formulation

The ointment has been formulated using palm kernel oil as the active ingredient. To prepare it, shea butter (used as an excipient) is melted in a bain-marie, then the required quantity is added to the corresponding palm kernel oil in a container. The mixture is homogenized until a single phase is obtained, then poured directly into 150mL jars. The jars are then left to stand at a temperature below 30°C until the contents solidify (Figure 2).



Figure 2: Ointment in tins

- I.2.3. Carrying out quality control, sterility and efficacy tests onointment
- Ointmentquality control
- - Color and odor
- The color and odor of the ointment are determined by a panel of ten (10) people.
- - Consistency
- Consistency is determined by touching the ointment.
- - Stability
- The ointment is subjected to various temperature conditions, followed by an evaluation of its melting point.
- - Homogeneity

Homogeneity is checked by observation with the naked eye, after spreading the
ointment thinly on a flat surface with a spatula. The even or uneven distribution of
extracts in the excipient was noted.

## - pH measurement

To measure pH, 50 g of the ointment was melted at 40°C in a water bath, then, when cold, pH was determined using a HANNA HI 8010 pH meter, by inserting the probe directly into the samples. Under the same conditions, the pH of shea butter and palm kernel oil was measured.

## Ointment sterility test

## - Sampling

After melting in a water bath, a 20mL sample of the ointment was taken.

## - Microbiological analysis

Germ identification and enumeration were carried out using the method based on standard procedures [6]. Enumeration included total aerobic mesophilic flora, total coliforms, yeasts and molds, and Streptococcus.

## - Preparation of culture media

Culture media were prepared according to manufacturers' recommendations. Table 1 summarizes the quantities sampled and the germs tested for each culture medium.

Table 1: Germs sought according to culture medium

Prepared	Quantitiescollected	Distilled	Boiling time	Sterilization time	Indication
media		water			
		quantity			
BEA	5.6g			121°C during 20	Streptococci

PCA	2.05g			min	FMAT
Sabouraud	4.2g	100mL	5 min		Fungalgerms
VRBG	3.8g			Not autoclavable	Total
					coliformbacteria

## I.2.4.Inoculation of culture medium for germ detection and enumeration

• Surface seeding was carried out for each culture medium. To do this, 0.1mL of the ointment sample was removed and aseptically deposited on the surface of the agar contained in the Petri dish. Then, using a sterile rake, the inoculum is spread over the agar surface. The plates were incubated at 30°C in the oven, then observed after 24 hours for BEA, PCA and VRBG media, and after 48 hours for Sabouraud medium. Colony enumeration

According to the standard [6], Petri dishes with more than 30 colonies will be retained. The number of microorganisms present in a given product sample is obtained by the following formula [7]:

$$N= \frac{\sum C}{V (n_1+0,1 \ n_2) \ d}$$
 (germes/mL)

- N: number of germs
- $\sum C$ : colonysums
- V: inoculum volume n1: number of Petri dishes counted at 1st dilution
- n2: number of Petri dishes counted at 2nd dilution

• d: dilution rate at first dilution

## **I.2.4.1.** Microbiological standard

The number of germs per mL or g of sample was calculated for each germ tested, then compared to the normative reference of microbiological criteria for herbal medicinal products [8].

- Ointmentquality control
- The following parameters were determined:
- Color and odor (organoleptic)
- Appreciation of the color and odor of the ointment by a panel of ten (10) people. The ointment is yellowish in color. The odor is that of attenuated shea butter.
- Consistency
- The ointment has a semi-solid consistency. It feels slightly hard to the touch and softens on contact with the skin when set.
- - Stability
- The ointment is unstable at temperatures above 30°C. Above this temperature, it begins to melt.
- The following observations have been made concerning changes over time:
- Three batches containing the ointment were left to stand at room temperature and examined after 0 days, 1 week, 2 weeks and 1 month of storage:
- Batch permanently open: no change;
- Batch permanently closed until the end of the experiment: the ointment was kept intact;
- Batch opened and tested at each control: no change observed.
- - Homogeneity

- The homogeneity of the ointment was checked by spreading it out in a thin layer on a flat surface using a spatula. In addition, as the oils are miscible with each other, very good homogeneity was achieved (even distribution of the extract).
- - pH measurement

The pH measurementyielded:

Palm kernel oil (pH=4.16), Shea butter (pH=3.85) and formulated ointment (pH=3.93),

## **I.2.4.2.** Cutaneous tolerability test for ointment

Application of the ointment to the flank of an albino rabbit showed no skin reaction on different days. The Primary Irritation Index (PII) was zero after 24 and 72 hours of ointment application to the rabbits' flanks. Even to see how the skin reacts to the ointment, but also to note the length of the ointment.

### **I.2.4.3.**Ointment sterility test

This test is a microbiological analysis carried out in agar plates, in which the ointment was surface-seeded on the respective media (PCA, BEA, Sabouraud and VRBG). The test showed that the ointment was free from contamination after 24 hours incubation at 30°C. The results of the microbiological analysis are reported in Table 2.

Table 2: Results of microbiological tests on ointment

	Number of germs / mL sample (CFU/mL or CFU/g)						
Samples	Yeasts andmolds	Streptococci	Germsmesophilic	Total			
				coliformbacteria			
Ointment	Absence	Absence	Absence	Absence			

### II. RESULTS AND DISCUSSION

Vegetable oil from palm trees provided a very interesting formulation for this study. The use of the traditional extraction method described by [9] enabled palm kernel oil to be extracted with a yield of 26%, providing sufficient palm kernel oil.

Quality control results obtained with the ointment show good macroscopic parameters with very good homogeneity. The attenuated shea butter smell of the ointment is a very pleasant fragrance, insofar as shea butter itself is a preferred excipient over others because of its smell [10], and it also releases the active ingredient better [11].

The pH values were determined using a HANNA HI 8010 pH meter, an electronic device providing precise pH values. The pH values obtained for palm kernel oil (pH=4.16), shea butter (pH=3.85) and formulated ointment (pH=3.93) show that palm kernel oil, shea butter and formulated ointment are acidic.

Furthermore, these different values indicate that the ointment has a pH value closer to that of the skin [11].

At temperatures above 30°C, ointments begin to melt, so we recommend keeping them in a cool place (temperature below 30°C).

The results of the skin tolerability test indicated skin tolerance in rabbits. Researchers have always been more interested in using these animals for their work. In fact, these animals are relatively well suited to the task, as they are phylogenetically closer to humans and are easy to handle. For this study, rabbits were chosen because of the sensitivity of their skin. Furthermore, for ethical reasons, *in vivo* tests cannot be carried out directly on humans. They have to be carried out on rodents such as mice or rats, or on other animals [12] such as rabbits in this case, whose metabolism is far from resembling that of humans. The test consisted in investigating possible skin irritation reactions due to the ointment by determining the primary

irritation index according to the Draize scoring system. No skin irritation reactions were observed.

So, according to the Draize scale, the ointment was considered non-irritating to the skin. This is explained by the fact that the shea butter (excipient) is not aggressive to the skin, since it has been applied for ages.

The result of the microbiological analysis of the ointments showed an absence of germs, which can be explained by the respect of hygiene measures adopted during the formulation of the ointment; This allows us to say that the ointment was produced according to Good Manufacturing Practice (GMP) standards. In addition, as the ointment consists entirely of fats, it limits contamination and the development of microorganisms.

In terms of efficacy tests, the performance of the ointments was better in male rabbits than in females. In fact, in males, the performance of the ointment can be seen from D16, with hair growth of 21 mm at the end of 28 days, whereas it was lower than in the control. We can appreciate the effectiveness of palm kernel oil, used for ages by rural populations for hair care ([9],[14],[13]). In females, on the other hand, at the end of 28 days of experimentation, hair grew faster without the ointments (in the control), reflecting a slowdown in hair growth in rabbits in the presence of the ointments.

Moreover, according to empirical knowledge and the literature, palm kernel oil, the active ingredient in the ointment, is a product used mainly by women for hair maintenance and growth. This contradiction between test results in rabbits and information from empirical knowledge and the literature could be explained by the fact that rabbits and humans do nothave the same metabolism [5].

Also, the low density of hair observed in males in the treated area may be due to the mechanical action exerted during ointment application.

#### **CONCLUSION**

The galenic formulation enabled us to develop a cosmetic ointment based on palm kernel oil and shea butter as an excipient. The results of tests carried out on rabbit hair show that these ointments stimulate hair growth, especially in males. The ointment contains appreciable physical, chemical and microbiological properties that encourage its use.

### **ACKNOWLEDGMENTS**

At the end this study, financial support to research and pedagogy unity of biochemical of pharmacodynamy, the national center of floristics from department of University Felix HOUPHOUET BOIGNY in Cocody-Abidjan (Ivory Coast) and unit of fundamental Medical Biochemistry of Pasteur institute (in Cocody-Abidjan; Ivory Coast) are gratefully acknowledged.

#### REFERENCES

[1]. **Mélanie LR**, (2016)The consequences of alopecia in women treated with chemotherapy for cancer. End-of-study work for the state nursing diploma. Nursing training institute, Caen - France. p 1-26.

- [2]. Thomas D, (2015) Alopecia in young people. Doctoral thesis, Université de Picardi Jule Verne, France ; 96p.
- [3]. Sanogo R, Konipo A, Maiga A, Arama AE, Diall D, Bougoudogo, F, (2006)
  Formulation of a dermal ointment based on *Mitracarpusscaber*(zucc) extracts. Pharm. Med,
  Vol. XIV. pp. 159-177.
- [4]. Dembélé DL, (2011)Formulation of an analgesic and anti-inflammatory ointment based on *Securidacalongepedunculata*Fresen (Polygalaceae). Doctoral thesis from the Faculty of Medicine, Pharmacy and Odontostomatology. University of Bamako, Mali. 143 p.
- [5]. Bene K, Camara D, Soumahoro I.A, Kanga Y, Zirihi GN, (2017) Galenical formulation of an antimicrobial ointment based on a hydroalcoholic extract of *Bersamaabyssinica* Fresen. *Ethnopharmacologia*, n°58, p 61-69
- [6]. **AFNOR** (1996)Food microbiology. Enumeration of thermo-tolerant coliforms by counting colonies obtained at 44°C, NF V 08 60, Paris, AFNOR, 2p.
- [7]. **AFNOR** (2001)Food microbiology. General rule for microbiological examinations, NF ISO 7218 / A1, Paris, AFNOR, 12p.
- [8]. European Pharmacopoeia (2014)Microbiological quality of pharmaceutical preparations, p 567 569.
- [9]. Ouattara H, Meite A, Dally T, Kati CS, (2016) Study of the properties of the fruits of three Arecaceae: *Elaeisguineensis* jacq., *Cocos nucifera* L., *Borassus flabellifer* Var. *J.Appl. Biosci.* p 10157-10169.
- [10]. Mboui O SE, (2003)Traditional palm kernel oil. Doctor of Pharmacy thesis from the University of Cheikh Anta Diop. 93p.

- [11]. Eyang EM,(2007)Study of the phytochemistry and antibacterial and antifungal activities of five medicinal plants used in the traditional treatment of dermatoses in Mali,Doctoral Thesis from the Faculty of Medicine, Pharmacy and Odontostomatology, University of Bamako, Mali, 175 p.
- [12]. Kamanzi AK, (2002)Medicinal plants from Côte d'Ivoire: phytochemical investigations guided by bioassays,State Doctorate in Science, Plant Biology, Cocody University,

  Abidjan, Ivory Coast, 176 p
- [13]. Sijelmassi A, (1991)Medicinal plants of Morocco. Ricin, Casablanca: The Fennec. Pp 217.
- [14]. Sadok D, Bentounes FZ, (2016)Study of the insecticidal activity of Ricinus communis and Mentha piperita leaf extracts against *Aphis spiraecola* green citrus aphid (Hemiptera: Aphididae). Master's thesis from Abdelhamid Ibn Badis University-Mostaganem Algeria; 100p.