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# ORIGINAL RESEARCH ARTICLES

# Ethnobotany and preliminary bioactivity investigation on hepatoprotective medicinal plants from the Mouhoun Region of Burkina Faso

Jotham Yhi-Pênê N'DO1\*, André TIBIRI1,2, Ernest Nogma SOMBIE1, Tata Kadiatou TRAORE3, Noufou OUEDRAOGO<sup>2</sup>, Adama HILOU<sup>1</sup>, Pierre Innocent GUISSOU<sup>2,3</sup> and Odile Germaine NACOULMA<sup>1</sup>

#### **Abstract**

An ethno botanical survey of medicinal plants used to treat hepatitis in the Mouhoun region of Burkina Faso was undertaken. The extracts of the most quoted species were then evaluated for their phytochemistry and their antioxidant activity:

The anti-radical activity (by the method of the discoloration of the ABTS cation radical) and that of the antioxidant (by the method of the reduction of the iron ion, FRAP) were evaluated. Then the phenolic content of the aqueous extracts was determined and a correlation was studied between these two parameters.

The antioxidant tests showed that the plants counted have a good antioxidant power. The three most active extracts are those of the trunk and root bark of Pseudocedrela kostchyi, and the trunk bark of Sterculia setigera. The extract of the bark of the trunk of Sterculia setigera showed the highest total tannin and phenolic content, while that of the leaves of Piliostigma reticulatum showed the highest content of flavonoids. The analyses showed that there is a relationship between the total phenolic contents and the antioxidant capacities of all the extracts (R2 = 0.82).

The extracts of the trunk and root barks of Pseudocedrela kostchyi, and that of the trunk bark of Sterculia setigera showed the best antioxidant properties. They could be good candidates for the search for liver protective molecules.

Keywords: hepatoprotection; Medicinal plants; antioxidant activity; Mouhoun region; Burkina Faso

# Introduction

Hepatitis is an inflammation of the liver. Two billion people worldwide have already been in contact with the hepatitis B virus with 800,000 deaths a year, [1]. In many developing countries there is no real program for the prevention and management of viral hepatitis [2]. According to WHO in 2004, more

than 3,300 deaths were recorded in Burkina Faso. The standard treatment of hepatitis is based on use of alpha interferon associated with nucleotide analogues and it is very expensive and not always effective in Burkina Faso villages [3]. The chronicity of liver infections causes an increase in toxic free radicals [4]. These free radicals play an important role in altering the hepatocyte membrane. That causes lysis of the liver cell membrane, producing inflammation. The antioxidant compounds of plant extracts having the ability to neutralize free radicals can serve to protect the integrity of hepatocytes. In this sense, an ethnob-

\*Correspondence: yhipene2005@yahoo.fr

<sup>1</sup>Laboratory of Biochemistry, Applied Chemistry (LABIOCA), University of Ouagadougou, 03 BP 848 Ouagadougou 03,, Burkina Faso

Full list of author information is available at the end of the article.

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otanical survey in the Mouhoun region of Burkina Faso has identified medicinal plants used against liver diseases and retained the six most widely used plants: Cassia mimosoides L. (Caesalpiniaceae), Piliostigma reticulatum (DC.) Hochst. (Caesalpiniaceae), Pseudocedrela kostchyi Harms (Meliaceae), Sterculia setigera Del. (Sterculiaceae), Prosopis chilensis (Molina) Stuntz emend. Burkart var (Mimosaceae) and Trametes versicolor (L.) Lloyd (Polyporaceae). The extracts of these species were evaluated for their antioxidant capacity, awell as the content of phenolic compounds.

# **Material and Methods**

#### Ethnobotany survey

#### Ethnobotanical investigation area

The survey was conducted in Mouhoun region (with an area of 34497 km² which represents 12% of the national territory, [5]. Located in north-west of Burkina Faso, the chief town in the region is Dedougou, occupies about. Region of Mouhoun haspedoclimatic conditions propitious to the diversity of the vegetation [6]. 90% of the population has agriculture as a mainsource of income [7]. The diversity of the vegetation is an indisputable asset for traditional practitioners and herbalists who are constantly consulted by the population for their health problem [8].

# The survey

Asemi-structured interview process has been used [9]. with tradi-tional practitioners and herbalists. It allows to record local tradi-tional medicine knowledge dealing with hepatitis treatments. The dialogue for data collection using the questionnaire has been in French and/or local languages (Dafin, Dioula, Moore). The an-swers to the questionnaire provided information on the name of the plant used, the part used, the method of preparation, the route of administration, the other pathologies treated, the type and du-ration of the treatment. After every questionnaire fulfillment a field trip with the respondent allows collection (for the herbar- ium), picturing and the GPS data each plant cited, Herbariums were authenticated by Professor Jeanne MILLOGO, botanist at University de Ouagadougou.

The ethnobotany data were processed by using Sphinx V5 software to obtain the citation frequency of each plant.

The choice of the six more used plants was made on the basis of this survey data and bibliographic research indatabank.

# Antioxidant activities and determination of phenolic compounds Plant material

The whole plants of Cassia *mimosoides* L., trunk barks of *Piliostigma reticulatum* (DC.) Hochst., Trunk and root barks of *Pseudocedrela kostchyi* Harms and *Sterculia setigera* Del., Fruits of *Prosopis chilensis* (Molina) Stuntz emend. Burkart var. was harvested in the Mouhoun region. Seagrass beds were deposited in the UFR / SVT herbarium under the respective identification codes 05ID 16677, 07ID 16679, 04ID 16676, 08ID 16680, 01ID 16673. *Trametes versicolor* (L.) Lloyd was harvested in the same Locality and identified at the Laboratory of phytopathology of the University Ouaga I Pr Joseph KI-ZERBO.

#### Extraction

The fine powder obtained by grinding the dried plant parts was extracted with distilled water according to the maceration process for the root and trunk bark of Sterculia setigera. For other plant organs (Cassia mimosoides, Piliostigma reticulatum, Pseu- docedrela kostchyi, Prosopis chilensis, Trametes versicolor), an aqueous decoction was carried out. Extractions were carried out to conform to the traditional forms in which these plants are used by the respondents. The aqueous maceration was done at 1/10 keeping the powder in contact with the distilled water for 24 hours. For the decoction, it consisted of maintaining 20 g of plant powder of each drug in a flask containing 300 ml of distilled water to boil for 30 minutes. The decoction was done under reflux. The extracts obtained were filtered with a nylon fabric and then centrifuged at 2000 rpm for 10 minutes. The su-pernatant was collected in plastic jars and placed in the freezer for freeze-drying thereafter.

# Solvents and reagents

Ethanol 95% (prolabo); Distilled water; Tris-HCl buffer (50 mM pH 7.4); Methanol 95%. ABTS or 2,2'-azinobis- (3-ethylbenzothiazoline-6-sulfonic acid) (FLUKA Biochimika, Germany); Acetic acid; Ascor- bic Acid, Hydrochloric Acid (SIGMA-ALDRICH, Germany); Sulfuric acid (SDS, France); Trichloroacetic acid (FLUKA Chimika, Switzerland); Pure acetic anhydride (Labosi, France); Sodium carbonate (Labosi, France); Magnesium chloride (SIGMA-ALDRICH, Germany); Sodium hydroxide (Merck, Germany); Potassium persulfate (SIGMA-ALDRICH, Ger- many); 0.2M phosphate buffer pH 6.6; Phosphate buffer 1.15M pH 7.5; Aluminum trichloride (Labosi, France); Gallic acid (Sigma); Ellagic Acid (Sigma); Tanic Acid (Sigma); FeCl3 (Labosi); NaOH (Merck); Quercetin (Sigma).

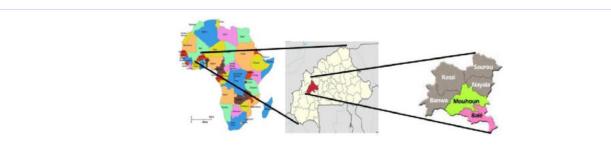


Figure 1 Location of the inventory sites

#### Antioxidant activities

Reduction of the radical cation ABTS +.

The TEAC (Trolox Equivalent Antioxidant Capacity) test was determined using the ABTS •+ radical [10]. ABTS was dissolved in water at the concentration of 7 mM (stock solution). The ABTS (ABTS +) cation radical was produced by reacting the stock solution of ABTS with 2.45 mM potassium persulfate (final concentration) and the mixture placed in the dark at room temperature for 12-16h before use. The reaction mixture consists of 1 ml of diluted ABTS + solution and 10  $\mu$ L of extract (1-25  $\mu$ g/ml) or Trolox as standard (0-15  $\mu$ M) in ethanol or the appropriate solvent. The mixture is incubated for 30 min at room temperature and the absorbance is read at 734 nm. The percent inhibition of absorbance at 734 nm was calculated according to formula:

% Inhibition = 
$$\frac{A_0 - A_1}{A_0} \times 100$$
  $ARP = \frac{1}{IC_{50}} \times 100$   $TEAC = \frac{ARP_{produit}}{ARP_{Troxlox}}$ 

Where  $A_0$  is the absorbance of the control and  $A_1$  is the absorbance of the sample or standard. The curve of inhibition of the absorbance as a function of the concentration of the extract or trolox was established for the determination of the 50% inhibitory concentration (IC 50). The Anti-Radical Power (ARP), trolox Equivalent Antioxidant Capacity (TEAC) has been determined through the formulas [11].

# Reducing power of iron

The reducing power of the samples was evaluated according to the spectrophotometric method described by [12] . 1 ml of the extract (0.02 g / ml was mixed with 2.5 ml of phosphate buffer (0.2M, pH 6.6) and 2.5 ml of 1% potassium ferricyanide (v/v). The mixture was incubated at 50° C for 20 min. 2.5 ml of 10% (v/v) trichloroacetic acid (TCA) were then added and the mixture centrifuged for 10 min. 2.5 ml of the obtained supernatant were then removed and mixed with 2.5 ml of distilled water and ml of 0.1% (v/v) ferric chloride (FeCl<sub>3</sub>). The absorbance was read at 700 nm. The AAE (Ascorbate Acid Equivalent) value is inversely proportional to the reducing power.

## Determination of total phenolics

The total phenolics were assayed according to the Singleton method [13]. The phenolic compounds react with the folinciocalteu reagent (FCR) in an alkaline medium. Molybdate reduced forms a blue complex whose maximum absorption is at 760 nm. The mixture consisted of 1 ml of extract, 1 ml of 2N FCR and 3 ml of 20% sodium carbonate. The absorbance was measured after forty minutes at room temperature with an agilent spectrophotometer (Agilent 8453) equipped with UV-visible and ChemStation software. The standard curve was plotted with tannic acid (0.1 to 1 mg/ml).

#### Determination of tannins

The assay is performed according to the method described by Tibiri [14]. 100mg of polyvinyl polypyrrolidone (pvpp) were used to precipitate 2mg of total phenolics. Such a mixture is vortexed, kept at  $4^{\circ}$  C for 15 minutes and then centrifuged for ten minutes. After forty minutes, the assay tube is centrifuged and the absorbance of the supernatant is measured at 760 nm. The tannin content is determined by differentiating the total phenolics (including tannins) and the second missing value of the tannins. The standard curve was plotted with tannic acid (0.1 to 1 mg/ml) with  $R^2 = 0.99$ .

#### Dosage of flavonoids

The flavonoid assay was carried out according to the method described by Abdel-Hameed [15].  $100~\mu\text{L}$  of extract of concentration 10~mg/mL in methanol are mixed with  $100~\mu\text{L}$  of 2% aluminum trichlorid and a drop of acid Acetic acid then the volume is brought to 5 ml. The absorbance is measured at 415 nm after 40 minutes. The control tube consists of  $100~\mu\text{L}$  of extract and one drop of acetic acid. The amount of flavonoids in the extract was determined by reading Agilent (Agilent 8453) spectrophotometer equipped with UV-visible and ChemStation software against a standard quercetin curve (R2 = 0.999).

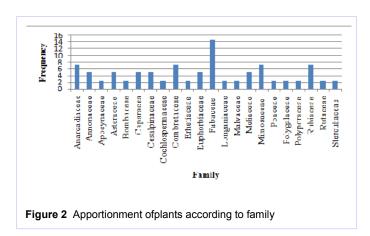
#### Statistical analysis

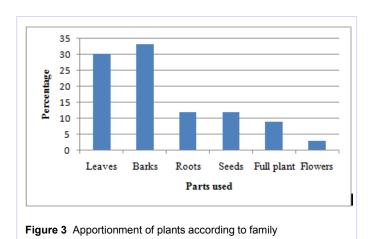
For statistical analyzes, Microsoft Excel has been used to obtain averages and standard deviations. One-way ANOVA followed by the Turkey test were used to measure the degree of statistical significance of the results using the XL stat module. A significant difference is considered for p < 0.05.

# **Results and Discussion**

#### Results

The majority of respondents (65%) were male (71%). The plants counted with 41 species belong to 22 families (Table 1). The most represented families were fabaceae, mimosaceae, combretaceae, rubiaceae and anarcadiaceae with a citation frequency greater than 7% (Figure 1). Trunk barks and leaves were the most frequently used parts with frequencies above 30% (Figure 2). The most common method of preparation used by the respondents was the decoction with 52.9% (Figure 3).





Evaluation of the antioxidant capabilities through the ABTS method showed that the bark extract of the root of *Pseudoce-drela kostchyi* had the highest antioxidant capacity  $(0.53 \pm 0.03)$  as the TEAC value, while the most Low capacity was observed with the whole plant extract of Cassia mimoso ides (Table 2) The FRAP method showed that the iron ion reduction capacities ranged from 13.63 to 134.52 ESA / ml. Extracts from the trunk bark of *Pseudocedrela kostchyi* showed the greatest reduction in iron ion compared to other extracts (Table 2).

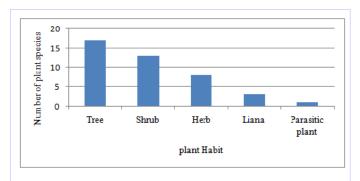
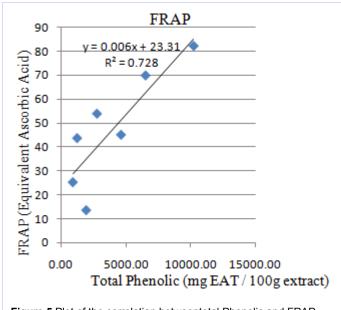


Figure 4 Apportionment according to plant habit



 $\textbf{Figure 5} \ \textbf{Plot} \ \textbf{of the correlation between total Phenolic and FRAP}$ 

The total phenolics content of the extracts was between 2.76 and 17.33 g EAT / 100 g of extract. The high content was obtained with the extract of the trunk bark of *Sterculia setigera* (Table 2). The tannin content of the extracts varied from 0.71 to 15.03 g EAT / 100 g. Extracts from the trunk bark of *Sterculia setigera* and *Pseudocedrela kostchyi* showed the highest levels of tannins. The flavonoid content was higher in the leaves of *Piliostigma reticulatum* with 739.67  $\pm$  0.01mg EQ/100g compared with the rest of the extracts. The flavonol content was greater than 300mg EQ/100g in the extract of the trunk bark of *Sterculia setigera* and *Pseudocedrela kostchyi* (Table 2).

#### Discussion

The listed species belonged to families of which the most represented is the fabaceae. Hailement [16]. found that fabaceae were the most widely used in Ethiopia against malaria. It should be

Table 1 Diseases treat andform of plant used

Scientific name and family	parts used	mode of préparation	Mode Of administration	Associated plants	Number of quotation	Diseases	
Adansonia digitata L. (Bombacaceae)	Fragment leafy and stem bark	Calcination and decoction	Oral and purging	Cassia nigricens and Entada africana	3		
Boscia angustifolia A. Rich (Capparaceae)	Roots	Decoction	Drink and bath	Cassia nigricans	2	Asthma, haemorrhoids, Hepatitis	
Carica papaya L.	leaves	Maceration	Drink, purgation and	Manhhot utilissima, Citrus	4	fever, stomach ache, Hepatitis	
(Annonaceae)	icaves	Maccration	bath	limon and Cordia africana		rever, stomach ache, riepatitis	
Cassia mimosoïdes L. (Caesalpiniaceae)	Full plant	Decoction	Drink	Cassia tora	2	Jaundice	
Parkia biglobosa (Jacq.) R.Br. exG.Don	leaves	Calcination	Digest	Acacia nilotica	1	Hepatitis	
(Mimosaceae)		Culcination	2.8	Treating milestrea			
Chrysantellum americanum L. (Asteraceae)	Full plant	Decoction	Drink	Gardenia erubescens	6	Malaria, Hepatitis	
Citrus limon (L.) Burm. F. (Rutaceae)	leaves	Maceration	Drink	Manhhot utilissima, Carica	2	Hepatitis, Jaundice	
, , , , , , , , , , , , , , , , , , , ,	CONTRACTOR OF THE PARTY OF THE	ABBERT CONTROL CONTROL	Visitable	papaya, Acacia nilotica			
Combretum micranthum G. Don (Combretaceae)	leaves	Decoction	Drink and bath	Manhhot utilissima, Citrus	2	Fever, Malaria, Jaundice	
		222333335		limon and Carica papaya			
Detarium microcarpum Guill. & Perr.	Stem bark	Decoction	Drink	Guiera senegalensis	2	Hepatitis	
(Caesalpiniaceae) Guiera senegalensis J.F. Gmcl. (Combretaceae)	Fragment leafy	Decoction	bath	Saba senegalensis	2	hemorrhoids, abdominal pain,	
2000-14 Province - 15 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	174 - 174 - 174	Manager .	- Control of the Cont			malaria	
Hibiscus sabdariffa Mendonça & Torret	Fragment leafy	Infusion	Boisson		1	Hepatitis	
(Malvaceae)	Water Carl				i i	N	
Manhhot utilissima F. lancifolia Roberty	leaves	Infusion and	Drink and bath		2	Jaundice, hepatitis	
(Euphorbiaceae)		maceration				-07-5000 Gr 6000	
Phyllanthus maderaspatensis L.	Full plant	Decoction	Bath and drink	Chrysantellum americanum	4	Malaria, hepatitis	
(Euphorbiaceae)		D	D. d.		-	0.11	
Piliostigma reticulatum (DC.) Hochst.(Fabaceae	Leaves	Decoction	Bath		3	Cold, cough, bronchitis,	
Prosopis chilensis (Molina) Stuntz emend.	Saade rinanina	Infusion	Bath	1	1	headache, hepatitis Hepatitis	
Burkart var (Milosaceae)	Seeds ripening	imusion	Datii		1	ricpatitis	
Pseudocedrela kostchyi Harms (Meliaceae)	Bark of stem and	Maceration	Drink and bath		4	Hepatitis	
r seudoceureu kosicnyi Hariis (Menaceae)	root	Macciation	Drink and bath			ricpatitis	
Saba senegalensis (A. DC.) Pichon	Fragment leafy	Infusion	Inhalation ant bath		2	Hepatitis	
(Apocynaceae)							
Securidaca longepedunculata (Polygalaceae)	Fragment leafy	Decoction	Inhalation et bain		2	Stomach ache, Hepatitis	
Senna obtusifolia (L.) (Caesalpiniaceae)	Leaves	Infusion	Drink and bath	Cassia mimosoïdes	2	Jaundice	
Sterculia setigera Del.	Bark of stem and	Decoction and	inhalation	Coclospermum planchonii	5	heart palpitations, low blood	
(Sterculiaceae)	root	Maceration				pressure and hepatitis	
Tamarindus indica L. (Caesalpiniaceae)	Leaves, stem bark	Decoction	Bath and drink	Adansonia digitata, Daniellia oliveri, Crossopterix febrifuga	2	Hepatitis and stomach ach	
Terminalia laxiflora Eng. & Diels	Stem bark	Calcination	Suck	Coclospermum planchonii,	3	Malaria and hepatitis	
(Combretaceae)				Sterculia setigera			
Trichilia emetica subsp. suberosa J.J. de Wilde	Root bark	Maceration	Bath and drink		2	Hepatitis	
(Meliaceae)	Ct t t	December	Tabalada as bad	C.1		M.L. I. I. I.	
Ximenia americana L. (Olacaceae)	Stem bark	Decoction	Inhalation et bath	Saba senegalensis	4	Malaria, Jaundice	
Cochlospermum planchonii	Root bark	Decoction	Buccal way	Tamarindus indica	3	Malaria, liver ailments	
Hook.f.(Cochlospermaceae)	ROOT Dark	Decoction	Duccai way	Tamarmans marca		Maiaria, fiver armients	
Acacia nilotica (L.)Delile(Fabaceae)	Seeds	Infusion	Buccal way and	Citrus limon, Parkia	2	Painful, menstruation, jaundice,	
steach money (E.) Denic(2 noncency	Secus	inteston	purge	biglobosa	-	shingles	
Acacia seyal Delile	Stem bark	Decoction	purge		1	Constipation, hernia, hepatitis	
(Fabaceae)	- A. C.	The state of the s	1				
Cassia nigricansVahl	Full plant	Decoction and	Buccal way and	Boscia angustifolia	3	Hepatitis and heart ailments	
(Fabaceae)		Infusion	purge	Control of the state of the sta			
Cordia africanaLam,	Leaves and stem	Decoction and	Buccal way, bath and	Carica papaya	3	Lack of appetite, infant fever	
(Erhetiaceae)	bark	Infusion	purge			hepatitis and thoraxiales pain	
Daniellia oliveri Benn.	Stem bark	Decoction	Buccal way and	Tamarindus indica	2	Ulcers, hemorrhoids, Hepatitis	
(Fabaceae)			purge	0000			
Entada africana(L.) Merr (Fabaceae)	Bark of stem and root	Decoction and infusion	Buccal way	Adansonia digitata	1	Jaundice, upset stomach cough	
Gardenia erubescensStapf&Hutch(Rubiaceae)	Seeds and stem	Decoction and	Buccal way	Chrysantellum americanum	2	Hepatitis, lunbar rpain and aches	
	bark	infusion	in accommendated to		==		
Lannea velutina (L.) (Anarcadiaceae)	Stem bark	Maceration	Purge and buccal way		1	Fever, abdominal pain, jaundice.	
Maerua angolensisDC. (Capparaceae)	Leaves	Decoction and infusion	Purge and buccal way		2	Hypertension, malaria, jaundice	
Sarcocephalus latifolius(Sm.) E.A.Bruce	Leaves and stem	Decoction or	Buccal way, bath and		3	Stomach aches, skin diseases.	
(Rubiaceae)	bark	Infusion	purge		-	hepatitis	
Cymbopogone giganteus Chiov.(Poaceae)	Trunk and seed	Maceration	Buccal way		3	hepatitis in children, Indigestion,	
To or						colic	
Crossopterix febrifuga (Afzel. ex G. Don) Benth.	Leaves	Infusion	Buccal way	Tamarindus indica	2	Jaundice	
(Rubiaceae) Spondias mombin L. (Anacardiaceae)	Leaves and stem	Decoction	Purge and buccal way		2	Jaundice, Malaria	
The state of the state of the state of	bark	- cocum	. arge and ouccur way			The state of the s	
Annona senegalensis Pers. (Annonaceae)	Young shoots, leaves and stem	Decoction	buccal way	Chrysantellum americanum	2	Enhances the activity of liver and heart	
, and the second						Liver, gargle and buccal diseases	
Scleeocarya birrea (A. Rich) Hochst	bark Seed rupening	Maceration	buccal way		1	Liver, gargle and buccal diseases	
SALAK DI PALAMASA PENGANANGAN SI SERIKATAN KALAK SERIKATAN DI SERIKATAN SALAK SERIKATAN DI SERIKATAN SERIKATAN	bark	Maceration Infusion	buccal way		1 5	Liver, gargle and buccal diseases hepatitis, high blood pressure	

Table 2 Means of determination of phenolic compounds and antioxidant activities

		Dosage of phenolic compounds		Antioxidant activities	
Plant species Phenolic content (g EAT/100g of extract)		Tanin content (g EAT/100g of extract)	Flavonoid content (mg EQ/100g of extract)	ABTS	FRAP
C. mimoso ides (Full plant)	2,76±0,01 <sup>e</sup>	$2,50\pm0,36^{e}$	562,03±0,19°	(TEAC)	(ASE/mI)
S. setigera (Root bark)	1,24±0,02 <sup>g</sup>	1,12±0,01 <sup>fg</sup>	327,31±0,18 <sup>e</sup>	$0,05\pm0,01^d$	53,83±5,6 <sup>bc</sup>
S. setigera (Stem bark)	17,33±0,05 <sup>a</sup>	$15,03\pm0,33^a$	590,84±0,11 <sup>b</sup>	$0,17\pm0,07^{c}$	43,64±3,3°
P. reticulatum (Stem bark)	$4,61\pm0,02^d$	4,13±0,14 <sup>d</sup>	739,67±0,01 <sup>a</sup>	$0,16\pm0,02^{c}$	63,38±0,9 <sup>b</sup>
P. kostchyi (Stem bark)	$6,49\pm0,01^{c}$	$5,63\pm0,30^{c}$	484,50±0,09 <sup>g</sup>	$0,1\pm 0,01^{cd}$	45,05±1,7 <sup>bc</sup>
P. kostchyi (Root bark)	10,17±0,08 <sup>b</sup>	10,02±0,30 <sup>b</sup>	629,31±0,19	$0,53\pm0,03^{ab}$	$69,77\pm0,8^b$
P. chilensis (Seeds)	1,93±0,01	1,40±0,21	$425,19\pm0,22^d$	$0,4\pm0,007^b$	82,06±1,3 <sup>b</sup>
T. versicolor (Full plant)	$0,91\pm0,01^{h}$	0,71±0,21 <sup>fg</sup>	108,44±0,13 <sup>h</sup>	$0,06\pm0,002^d$	13,63±0,8
Trolox Quercetin				$0.08\pm0.001^d$ $1\pm0.08^a$	25,28±3,2 <sup>d</sup> 134,52±2,5 <sup>a</sup>

noted that the pathophysiology of malaria presents a hepatic phase, which could bring about a rapprochement with hepatitis. Among the plants cited appears a fungus Trametes versicolor which is a saprophyte. The virtues of saprophytes used in traditional medicine depend on the host [17] . In the case of our fungus work is the one being harvested on the species *Parkia biglobosa*.

Trunk barks and leaves were the most frequently cited by respondents [18]. while flowers, fruits and roots held low frequencies [19]. The low use of roots and fruits could contribute to the preservation of biodiversity [20]. in the Mouhoun region. The most cited method of preparation is the aqueous decoction, which is corroborated by some authors in the literature [21].

The group of phenolic compounds includes a wide range of secondary metabolites including flavonoids, tannins and phenol acids [22]. In scientific journals, phenolic acids, tannins and flavonoids are compounds with several biological activities such as antioxidant activity [23]. In this work, these phenolic compounds have been assayed for their contribution to the antioxidant capacities of the extracts. The extract of the bark of the trunk of Sterculia setigera showed a high total phenolic content  $(17.33 \pm 0.05)$ . In the work of Osemeahon [24], significant total phenol content was found in the vegetative aerial parts of Sterculia setigera. The extracts with the highest tannin content are those of the trunk of Pseudocedrela kostchyi and Sterculia setigera. The tannins have the ability to adopt conformations that allow them to oppose mechanical and biological attacks that may threaten the integrity of hepatocytes [2]. The dosage of the flavonoid extracts showed higher levels in the leaf extract of Piliostigma reticulatum. Flavonoids would have a great ability to protect primary metabolites such as lipids from oxidation, which is a route of attack from liver aggressors.

The antioxidant tests were carried out by two methods, including the evaluation of the capacities of the extracts to stabilize the ABTS + radical and to reduce the iron ion. The extracts showed on the whole an ability to stabilize the ABTS radical. This study demonstrates the ability of the extracts to neutralize the free radicals produced by the different systems that are the source of oxidative stress [25, 26]. This neutralizing capacity suggests their potential for use as therapeutic agents of pathologies such as viral hepatitis that produce radicals in the body [27].

As regards the reduction of the iron ion, this test consisted in determining the capacity of the extracts to transform the fer- ric ion into ferrous ion by the supply of an electron [28]. The extracts all showed an interesting activity, ranging from 13.63  $\pm$  0.8 to 82.06  $\pm$  1.3 ESA / ml. The antioxidant effects of the extracts demonstrate their ability to give electrons to block the chain of production of free radicals caused by the hepatic affections. The bark of the trunk of *Pseudocedrela kostchyi* and *Sterculia setigera* showed the greatest abilities to reduce the iron ion. Ugulu [29]. In evaluating the antioxidant properties of plants found that the leaves and trunk bark exhibited better antioxidant activities.

A correlation curve between total phenolic content and antioxidant activity gave a positive correlation coefficient with  $R^2 = 0.7281$  (Figure 4). The antioxidant activity of the extracts

could be explained by their richness in polyphenolic substances such as tannins, flavonoids [30]. The antioxidant capacity of the extracts observed is a solid argument that could justify their use in traditional medicine especially for these hepatoprotective properties [14]. The literature has often pointed to the close link between these properties and the antioxidant potential of plants [31]. Several authors explain the hepatoprotective effect of plants by their ability to relieve the liver cells of oxidative stress [32,33].

# **Conclusions**

This study is an asset for the preservation of ethnobotanical knowledge that is transmitted orally. The antioxidant properties of the extracts could justify the use of these plants in traditional medicine against stress diseases such as hepatitis. Extracts from the trunk and root bark of *Pseudocedrela kostchyi* and from the trunk bark of *Sterculia setigera* which exhibited the best antiox- idant properties deserve further investigation into their hepato- protection mechanism.

# **Authors' contributions**

All authors had similar contributions regarding the manuscript writing, literature research, review design, literature analysis and final text approval.

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#### **Author details**

<sup>1</sup>Laboratory of Biochemistry, Applied Chemistry (LABIOCA), University of Ouagadougou, 03 BP 848 Ouagadougou 03, Burkina Faso. <sup>2</sup>Department of Medicine and Traditional Pharmacopoeia (MEPHATRA-PH), 03 BP 7192 Ouaga 03, Institute for Research in Health Sciences (IRSS / CNRST, Burkina Faso. <sup>3</sup>Doctoral School of Health, University of Ouagadougou, PO Box 7021, Burkina Faso.

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