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Research Article

Ethnomedicinal Knowledge of Plants Used in Nonconventional Medicine in the Management of Diabetes Mellitus in Kinshasa (Democratic Republic of the Congo)

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Background. People with diabetes, herbalists, and traditional medicine practitioners (TMPs) from Kinshasa use plants to treat diabetes, but no study has inventoried the plants used by these populations. The present study was conducted to identify the plants used in Kinshasa to treat diabetes mellitus. Methods. The survey conducted in the form of a semistructured interview between March 2005 and August 2006 made it possible to collect ethnobotanical information from people with diabetes (n = 126), herbalists (n = 80), and TMPs (n = 120). Results. The 326 subjects consulted (sex ratio M/F = 0.6, age 51 ± 7 years, and experience: 17 ± 5 years) provided information on 71 plants, most of which are trees (35%), belonging to 38 families dominated by Fabaceae (19.7%) and indicated in 51 other cases of consultation dominated by malaria (12%). From these 71 plants derived, 86 antidiabetic recipes were administered orally, where the leaf is the most used part (>50%) and the decoction (>46%) is the most common mode of preparation. This study reports for the first time the antidiabetic use of 11 species, among which Tephrosia vogelii^X (0.08), Chromolaena corymbosa^X (0.06), and Baphia capparidifolia^X (0.06) present the highest consensus indexes (CI) and Marsdenia latifolia^W (UVp = 0.08) and Rauvolfia mannii^X (UVp = 0.06) present the highest UVs. Conclusion. The results show that Kinshasa people treat diabetes using several plants. Some are specific to the ecological environment; others are used in other regions. Pharmacological studies are underway to assess the therapeutic efficacy of these plants.

1. Introduction

Diabetes mellitus (DM) is a metabolic and chronic disease involving inappropriately elevated blood glucose levels (hyperglycemia). Hyperglycemia alone can impair pancreatic beta-cell function and contributes to impaired insulin secretion. The body cannot produce and secrete sufficient insulin hormone or use it effectively. This insulin deficiency leads to elevated blood glucose levels and decreases carbohydrate and protein metabolism [1, 2].

Worldwide, 537 million adults (20–79 years old) live with diabetes. This number is expected to rise to 643 million by 2030 and 783 million by 2045. More than 3 out of 4 adults with diabetes live in low-income countries like DR Congo, and diabetes has been responsible for 6.7 million deaths in 2021, i.e., one end every 5 seconds [1]. Cardiac, vascular, neurological, and renal damage and neuropathy may occur without appropriate treatment. Treatment includes diet, exercise, and medication [3]. Given that most cases occur in low-income countries, where a significant fraction of the

population resorts to traditional medicine, which essentially uses plant resources, medicinal plants constitute a credible alternative in the fight against diabetes mellitus.

Natural products have been shown to play an essential role in regulating pathophysiological signaling pathways, particularly in diabetes [4]. Over 800 plant species showing hypoglycemic activities can be vital sources in discovering and developing new types of antidiabetic molecules [5].

Medicinal plants have a high potential to treat various ailments due to the presence of their significant bioactive phytoconstituents. Certain plants are rich sources of compounds reputed to be antidiabetic such as flavonoids, alkaloids, phenolic compounds, and tannins, improving pancreatic tissues' efficiency by increasing insulin secretion or decreasing intestinal glucose absorption [6].

The World Health Organization insists that scientists conduct ethnomedicinal, ethnobotanical, and ethnopharmacological investigations to record and preserve traditional knowledge, create databases, and validate scientifically traditional claims from the perspective of developing improved medication [7].

In the Democratic Republic of the Congo, a country with similarities to other developing countries, data on the prevalence of diabetes mellitus and the use rate of medicinal plants in their therapeutic load is not sufficiently available. Although less systematic, some studies have reported presumed antidiabetic plants in Kinshasa [8, 9].

This study completes the data reported in these studies while highlighting plants used by TMPs, diabetics, and herbalists in Kinshasa.

2. Methods

2.1. Experimental Framework. The city-province of Kinshasa is located between 4°18' and 4°25' south latitude and between 15°18′ and 15°22′ east longitude. It is bounded to the north and east by the province of Kwilu, to the south by the province of Congo Central, and to the west by the Republic of the Congo, with an average altitude of 300 m above the sea. Kinshasa city's climate is tropical. It is characterized by a long rainy season lasting for 8 months (October-July), discontinuous between January and February, followed by August and September in favor of the short dry season. The vegetation of Kinshasa consists of degraded primitive forests, savannahs, and aquatic and semiaquatic formations in the valleys and Pool Malebo. It belongs to the Guinean-Congolese region, the Congolese basin domain, and the Congolese-Zambezian transition sector [10].

2.2. Ethnobotanical Data Collection. This cross-sectional descriptive ethnobotanical study was conducted between March 2005 and August 2006 through semistructured interviews based on a questionnaire. The discussion focused on knowledge about plants used to manage diabetes mellitus. Three groups of subjects were consulted: people with diabetes, herbalists, and TMPs from Kinshasa.

People with diabetes were met in 4 health zones of the city of Kinshasa: Bumbu, Kalamu, Limete, and Makala, covering the biomedical centers for the care of people with diabetes in Kinshasa supervised by BDOM (diocesan office of biomedical works). The reference health centers (RHCs) Libundi, Bondeko, Saint Clément, 2e Rue, the Bondeko clinic, and the Saint Joseph General Reference Hospital were concerned.

The herbalists consulted during this study were met in 5 popular markets in Kinshasa, the central market, the Selembao market, the Matadi Kibala market, the Mariano market, and the UPN market.

The TMPs were met in 4 communes of Kinshasa (Bumbu, Kalamu, Limete, and Makala). These municipalities are covered by the health zones selected during the survey of people with diabetes. TMPs were reached by snowballing from the TMPs provided by the population in each area concerned, and the investigations were carried out in Lingala. The listed plant species were collected and placed in herbariums, then compared to the reference herbariums from Kisantu (National Agricultural Study and Research Institute), to identify the scientific names. These were then formatted according to the Plants of the World online database: Plants of the World Online/Kew Science (https://powo.science.kew.org/) or African Plant Database (https://africanplantdatabase.ch/) or the World Flora Online (https://www.worldfloraonline.org/).

2.3. Data Analysis. Three ethnobotanical indexes were calculated to assess the significant species: the usual value (UV), the fidelity index (FI), and the consensus index (CI). The following formula determined the usual value (VU): $VU = (\sum Ui)/Ni$, where Ui is the number of uses mentioned by an informant for a plant (organ) and Ni = the number of informants who cited the plant. The following formula calculated the consensus index on the plant (organ) (CI): CI= Np/N, where Np is the number of people who cited the plant (organ) and N is the number of people who were consulted in the study. The fidelity index of the recipe was calculated by the formula FI = nr/Np, where nr = the number of people who cited the recipeand Np is the number of people who cited the plant. Apart from the 3 ethnobotanical indexes mentioned above, the relative citation frequency (RCF) was also determined. It was calculated by the formula $F_{\rm CR} = n \times 100/N$ with *n* being the number of occurrences of the factor examined and N = total number of the population concerned. This factor was used to quantify various factors analyzed in this study, except those involved by the ethnobotanical indexes.

In this study, UV makes it possible to evaluate the medicinal importance of a plant in the environment; CI makes it possible to identify the level of consensus of the population on the use of a species in the management of diabetes mellitus; and FI makes it possible to establish the level of loyalty that emerges among informants on an antidiabetic recipe.

3. Results

3.1. Ethnobotanical Profile of Inventoried Plants. Seventy-one plants were inventoried during this study. These plants were informed by 3 types of informants, herbalists (class Y), diabetics (class X), and TMPs (class W). The coexistence of these 3 types of informants gave rise to two other classes, the class of plants common to the 3 sources (class Z) and plants familiar to herbalists and TMPs (class V). As such, the results show that 16 plants are from diabetics, 7 plants are from herbalists, 22 are from TMPs, 6 are from both herbalists and TMP, and 20 are from TMPs, herbalists, and diabetics (Table 1). It should be noted, however, that the TMP class provided the largest most significant number of species, i.e., 31% (Figures 1(c) and 1(d)).

Each of these plants is named in one of the 15 Congolese ethnic groups reported in this study, of which Kikongo (50.7%) and Lingala (11.3%) occupy the first two places (Figure 2). These plants belong to 38 families dominated by the Fabaceae (19.7%), followed, respectively, by Rubiaceae Juss (7.0%), Apocynaceae Juss (5.6%), Asteraceae Bercht and J. Presl (5.6%), and Phyllanthaceae Martinov with 5.6% (Figure 3). These plant taxa are mostly trees (35%) or perennial herbs (31%) endemic to tropical Africa (Figures 1(a) and 1(b)). Among the 71 inventoried species, only Persea americana Mill (30.67%), Senna alata (L.) Roxb (30.67%), Garcinia kola Heckel (18.40%), Piliostigma reticulatum (D.C.) Hochst (17.79%), and Vachellia karroo (Hayne) Banfi and Galasso (17.79%) showed a relative citation frequency >15% (Table 1).

85% of identified medicinal plants in the present study have been previously used in local medicines as antidiabetic plants (13%). However, 72% have demonstrated an antihyperglycemic effect. Nevertheless, this study highlights 11 plant species never reported before as antidiabetic plants, among which *Tephrosia vogelii*^X ($F_{\rm CR}$ = 7.98), *Chromolaena corymbosa*^X ($F_{\rm CR}$ = 6.44), and *Baphia capparidifolia*^X ($F_{\rm CR}$ = 5.52) are the most cited (Table 1).

3.2. Ethnomedical Profile of Inventoried Plants. The 71 plants inventoried during this study are used in 86 antidiabetic recipes, of which 80 recipes use one plant, and 6 combine two plants (Table 2). The administration of the two types of recipes is essentially done orally, the leaf is the most used organ (50 and 58.3%), and decoction is the most predominant mode of preparation with 46.3 and 66.7% (Figure 4).

Figure 4 indicates that *Baphia capparidifolia*, *Chromolaena corymbosa*, *Marsdenia latifolia*, *Rauvolfia mannii*, and *Tephrosia vogelii* are among the critical plant species, according to their consensus index and use value.

In these recipes (R81–R86), we can observe the associations of leaves-leaves, roots-roots, and leaves-roots, representing half of these associations. Only 12 of the 71 plants are involved in the recipes combining two plants. For these recipes of plants in association, the treatment duration varies between 30 and 45 days, and it is the

combination of *Albizia adianthifolia* (leaves) and *Annona senegalensis* (roots) with an RCF of 4.6 which is the most cited: R85 (Table 3).

The consensus indexes of the identified antidiabetic recipes (ICR) varied between 0.01 and 0.31, with the highest value observed for the recipes R61, based on *Persea americana*^X leaves, and R72, based on leaves of *Senna alata*^X.

For the 11 plants reported for the first time by this study as antidiabetics, the ICRs vary between 0.08 and 0.01, with *Tephrosia vogelii*^X (0.08), *Chromolaena corymbosa*^X (0.06), and *Baphia capparidifolia*^X (0.06) showing the highest values. Usual plant UVp values vary between 0.02 and 0.12; the highest values were observed in *Annona senegalensis*^W, *Nauclea latifolia*^Y, and *Erythrina abyssinica*^W with an UVp = 0.12 each. Note also that in the group of 11 taxa reported for prayer times by this study, UVp varies between 0.02 and 0.08; these are *Marsdenia latifolia*^W (UVp = 0.08) and *Rauvolfia mannii*^X (UVp = 0.06) which presented the highest values in this class (Table 2).

The 71 plants inventoried are involved in 51 other causes of consultation, including malaria (12%), cough (9%), diarrhea (7%), wounds (6%), and sexual weakness, which occupy the first 5 places. The other pathologies are below the 5% mark (Figure 5).

3.3. Sociodemographic Profile of Subjects Consulted. The surveyed and consulted during this study were either diabetics (38.7%) or herbalists (24.5%) or practitioners of traditional medicine (36.8%), primarily women (62.3%) whose majority age is between 48 and 58 years old. Still, the extremes are 18 and 70 years old. They mainly live either in the municipality of Kalamu (29.8%) or in Makala (29.4%), and their level of education is, on average, secondary (47.2%). They mainly exercise 4 types of profession, the most representative of which is either commerce (28.5%) or liberal work (24.2%). In most cases, they have experienced more than 11 years of use of medicinal plants in managing diabetes (Table 4).

We also sought to determine the correlation between a few variables that characterize the subjects consulted within the framework of this study. The results show a positive linear correlation (R = 0.95, y = 0.4134x-6.5581) between the age of diabetic subjects and the number of years they have lived with diabetes. Similarly, a positive linear correlation is also observed between the age of herbalists and their years of experience (y = 0.5148x - 13.198, $R^2 = 0.8372$). In the same way, we noted a positive between the TMP age of the TMPs and the profession's expertise (y = 0.4076x - 8.3606, $R^2 = 0.807$) and their age with the number of patients they receive per quarter, y = 0.4525x - 10.394, $R^2 = 0.9482$ (Figure 6).

4. Discussion

The Congolese, particularly the flora of Kinshasa, have plants that traditional medicine practitioners and the general population use to manage diabetes. This study was interested in inventorying these plants and presenting their

Table 1: Botanical characteristics and previous knowledge of the plants inventoried.

Moench ^X chum ^W nach.) W. Wight ^W nach. and Thonn.)	Malvaceae	Dongodongo	Lingala		[17] 20 000 000 100 000 000 000 000 000 000
	Eupliotolaceae	Kabobo	bemba	Fr [11] Nr	1n vivo rat: 100–200 mg/kg [12] Nr
	Zingiberaceae	Mundongo	Lingala	Tb	In vivo rat: $150-400 \text{ mg/kg}$ [13] In vivo rat: $200-400 \text{ mg/kg}$ [14]
	Fabaceae	Mulu	Kongo	Lv [15]	In vivo guinea pig: 500 mg/kg [16]
	Euphorbiaceae	Mambunzila	Kongo	Lv [17]	In vivo rat: 200-600 mg/kg [18]
	Asphodelaceae	Langa	Ngwaka	Lv [19]	In vivo rat: 100–200 mg/kg [20]
Anacardium occidentale L.* 0.92	Anacardiaceae	Nkasu	Kongo	Lv [23] Sh	In vivo rat: 100 mg/kg [24] In vivo rat: 100 mg/kg [24] In vivo rat: 200 mg/kg [75]
•	Annonaceae	Kilolo	Kongo	30 Lv [8]	In vivo rat: 100–200 mg/kg [25] In vivo rat: 100–200 mg/kg [26]
Tul.	Phyllanthaceae	Misengo	Kongo	Sb [15, 27]	Nr.
Artemisia afra Jacq. ex Willd.* Asparaous racemosus Willd.*	Asteraceae Asparagaceae	Lengana Lutwa	Bemba Mbala	Ap [28] Wp [30]	<i>In vivo</i> rat: 500–1000 mg/kg [29] <i>In vitro</i> amvlase model [31]
	Meliaceae	Marumaru	Shi	Sq [8]	In vivo rat: 500 mg/kg [32]
Roublin contraviditalin BalowX	Fahacasa	Mumburo	Mahandi	Ko Y	111 V1VO rat:100–800 mg/kg [33]
Bidens pilosa L. ² 5.52	Asteraceae	Kokoyalimo	Lokele	Wp [15]	<i>In vivo</i> rat: 250 mg/kg [34]
Boerhavia diffusa L. ^Y	Nyctaginaceae	Ndemba	Ngwaka	L [35] Ro	In vivo rat: 500 mg/kg [36] In vivo rat: 50–300 mg/kg [37]
				0	In vivo rat: 250–500 mg/kg [37]
Brassica juncea (L.) Czern. ^W 3.68	Brassicaceae	Nkofi	Kongo	Sd [38]	<i>In vivo</i> mice: 100 mg/kg [40]
				Lv	In vitro and in vivo rat: 500 mg/kg [41]
Bridelia ferruginea Benth. ^Z 3.37	Phyllanthaceae	Kimwindu kinseke	Kongo	Kb [42] Lv	In vivo rat: 50 mg/kg [4 <i>5</i>] In vivo rat: 200–800 mg/kg [44]
Brillantaisia owariensis P. Beauv. Y	Acanthaceae	Lembalemba	Lingala	Nr [45]	In vivo rat: 500-2,000 mg/kg [46].
	Fabaceae	Ngoliolio F:-1-1	Ngwaka	Lv [35]	In vivo rat: $400-600 \text{ mg/kg} [47]$
Catharanthus roseus (L.) G. Don 1.25	Apocynaceae	Fulele	Ngwaka	1b [48]	<i>In wwo</i> rat: 37.5–150 mg/kg [49]
Chromowena corymbosa (Aubi.) K. M. Milg and 6.44 H. Rob. ^X	Asteraceae	Mpala kasakula	Kongo	$ m N_{r}$	Nr
rantium L. ^w	Rutaceae	Mala	Kongo	Fr [15]	In vivo rat: 200-300 mg/kg [50]
d K. Schum. ^Y	Costaceae	Ngo nkini	Kongo	Lv	<i>In vivo</i> rat: 100–200 mg/kg [51]
	Costaceae	Mafulungu	Kongo	Lv [52]	r.
Crinum ornatum (Aiton) Herb. 1.84	Amaryllidaceae	Munselebende	Kongo	Nr [38]	$ m N_{r}$
Crossopter) x febrifuga (Afzet. ex G. Don) 4.91 Benth. W	Rubiaceae	Mvala	Kongo	Sb [15]	<i>In vivo</i> rat: 500–1500 mg/kg [53]
	Fabaceae	Nkeka zango	Yanzi	$^{ m N}$	$N_{ m r}$
$ ilde{rapf}_{ ilde{\mathbf{v}}}^{W}$	Poaceae	Sinda	Lingala	Wp [8]	In vivo rat: 2500-5000 mg/kg [54]
h) Pax^	Dioscoreaceae	Ngamba	Kongo	Bb [55]	<i>In vivo</i> rat: 100 mg/kg [56]
	Ebenaceae	Lutwa lundomba	Kongo	Ko [52]	$\frac{Nr}{L^2}$ injector and $\frac{100}{r}$ 400 mm and $\frac{1}{r}$ [57]
Dysphania ambrosioides (L.)* Erythrina abyssinica Lam.**	Amaranthaceae Fabaceae	Kulamoka Kikumbu	Yaka Kongo	Lv [7] Rb [8]	<i>In vivo</i> rat: 100–400 mg/kg [57] <i>In vivo</i> guinea pig: 500 mg/kg [16]

TABLE 1: Continued.

Taxon	$F_{\rm CR}$ $(n = 326)$	Family	Local name	Ethnicity	A.u.	A.D.
Ficus benghalensis L. ^Y Ficus exasperata Vahl ^Y Garcinia kola Herkel ^V	1.53	Moraceae Moraceae	Nsanda Kikuya Ngadiadia	Suku Kongo	Sb [8] Lv [38] Sd [60]	In vitro: AC $_{50} = 84.44 \pm 1.65 \mu g/mL$ [58] In vivo rat: 150 mg/kg [59] In vivo rat: 200.800 mg/kg [61]
Gladiolus gregarius Welw. ex Baker ^W	0.61	Iridaceae	Litungulu ya Zamba	Lingala	Bb [15]	Nr
Gymnanthemum amygdalinum (Delile) Sch. Bip. ^V	3.07	Asteraceae	Mukari kari	Kongo	Lv [62]	In vitro- α -glucosidase: IC ₅₀ 47.29 ± 1.12 μ g/mL [63]
Vernonia amygdalina Delile (Synonym)	27.6	Hamino	Mfrimii	, of M	I [15]	Is with mot. EAD may led [64]
Heinsia crinita (Wennberg) G. Taylor ^Z	2.76	rrypencaceae Rubiaceae Iuss.	Ivaku	Mongo	Lv [13] Lv [65]	In vivo rat: 300 mg/kg [64] In vivo rat: 400 mg/kg [66]
Hymenocardia acida Tul.X	4.29	Phyllanthaceae	Kigeti	Kongo	Lv [15]	F: <i>in vivo</i> rat: 250–1000 mg/kg [67]
Lantana camara L. ^w	0.61	Verbenaceae	Ngbanyamando	Ngwaka	Lv [15] Fr	In vivo rat: 200–400 mg/kg [68] In vivo rat: 50–2000 mg/kg [69]
Lippia multiflora Moldenke ^Y	1.23	Verbenaceae	Bulukutu	Kongo	Lv [52]	Nr
Marsdenia latifolia (Benth.) K. Schum." Goneronema latifolium Benth. (Svnonym)	0.92	Apocynaceae	Lolango	Kongo	Nr [70]	<i>In vivo</i> rat: [71]
Millettia drastica Welw. ex Baker ^W	0.92	Fabaceae	Mwengzti	Kongo	Ro [52]	Nr
Mitragyna stipulosa (DC.) Kuntze ^W	1.23	Rubiaceae	Liluku	Lingala	Nr [72]	Nr.
Monodora myristica (Gaertn.) Dunal	2.45	Annonaceae	Mpei	Kongo	Sd [73]	<i>In vivo</i> rat: 50–200 mg/kg [74]
Morinda lucida Benth. ^Y	1.23	Rubiaceae	Nkisi	Kongo	Lv [8] Sb	In vitro-α-glucosidase: [75] In vivo rat: 125–500 mg/kg [76]
Musanga cecropioides R. Br. ex Tedlie ^W	1.23	Urticaceae	Nsanga	Kongo	Lv [52] Sb	In vivo rat: 10-40 mg/kg [77] In vitro α-glucosidase [78]
Nauclea latifolia Sm. Y	1.53	Rubiaceae	Kilolo	Kongo	Lv [79]	In vivo rat: 100-200 mg/kg [80]
Ocimum gratissimum L. ^Y	1.23	Lamiaceae	Dinsusu-nsusu	Kongo	Lv [81]	In vitro a-glucosidase [78]
Palisota schweinfurthii C. B. Clarke ^X	8.28	Commelinaceae Mirb	Mabongubongu	Yanzi	Lv [52] Wp [52]	Z, Z
Pentaclethra macrophylla Benth. ^X	1.84	Fabaceae	Mutunzama	Kongo	Sb [45]	In vivo rat: 200-400 mg/kg [82]
Persea americana Mill. ^X	30.67	Lauraceae	Savoka	Yanzi	Lv [83] Fr	In vivo rat: $100 \text{ mg/kg} [84]$ In vitro α -elucosidase [85]
Phaseolus vulgaris L. ^W	1.53	Fabaceae	Madesu	Lingala	[98] ps	In vivo rat: 1–5 mg/kg [87]
Phyllanthus niruri L. ^Y	2.45	Phyllanthaceae	Kahungahunga	Luba-kas	Ro [88] Ap	In vivo rat: 200–400 mg/kg [89] In vitro α-elucosidase: [90]
Piliostigma reticulatum (DC.) Hochst. ^X	17.79	Fabaceae	Falabuta	Suku	Lv [91]	In vivo rat: $250-1000 \text{ mg/kg}$ [92]
Piliostigma thonningii (Schumach.) Y	1.53	Fabaceae	Tshifumbe	Luba-kas	Sb [15]	In vivo rat: 500 mg/kg [93]
Piper guineense Schumach. and Thonn.W	0.61	Piperaceae	Kapinde	Kongo	Lv [52]	In vivo rat: 40–100 mg/kg [94]
Platymitra arborea (Blanco) P. J. A. Kessler	2.45	Annonaceae	Mpei	Kongo	Ņ	$N_{ m r}$
Psidium guajava L. ^w	0.61	Myrtaceae	Mapela	Lingala	Lv [79]	In vivo rat: $40-100 \mathrm{mg/kg} [94]$
Quassia amara L. ^x	0.61	Simaroubaceae	Mupesipesi	Luba-kat	LV, I, KO [95]	In vivo rat: 200–500 mg/kg [96]
Rauvolfia mannii Stapf ^X Shumekia amaican I ^Y	0.92	Apocynaceae	Kilungu Lunzilanzila	Kongo	Nr Wr [15]	Nr In wing wat 600 mayles [07]
Schwenckia americana L.	2.43	Solaliaceae	Lunzhanzha	Nongo	[C P] qw	In VIVO tat: 600 mg/kg [97]

CABLE 1: Continued.

Taxon	$F_{\rm CR}$ $(n=326)$	Family	Local name	Ethnicity	A.u.	A.D.
Senna alata (L.) Roxb. ^X	30.67	Fabaceae	Mbwabwa	Kongo	Lv [98] Fr [100]	In vivo rat: 500–500 mg/kg [99] in vivo: 250–500 mg/kg [101]
Senna occidentalis (L.) Link ^V	2.15	Fabaceae	Ntsambi ntsambi	Kongo	Lv [102]	In vivo rat: supplement at doses of 12.5%, 25%, and 50% [103]
Svzvoium cordatum Hochst ex Kranss ^Y	1 53	Murtaceae	Mugorhe	Shi	Ro Ly [105]	In vivo rat: 400 mg/kg [104] In vivo rat: 3–50 mo/kg [106]
Tephrosia vogelii Hook. f. ^X	7.98	Fabaceae	Uleku	Kongo	Nr [52]	Nr. S. J.
Terminalia mollis M. A. Lawson ^y	2.45	Combretaceae	Mboumbou	Luba-kas	Lv [15] Ro [15]	žž
Vachellia karroo (Hayne) Banfi and Galasso ^x Acacia karroo Havne (<i>Synonym</i>)	17.79	Fabaceae	Munga	Luba-kat	Lv [15]	In vitro a-glucosidase: 25 µg/mL [107]
Voacanga africana Stapf ^W	0.61	Apocynaceae	Kondiankondia	Yaka	Lv [79]	In silico $[108]$
Ziziphus mucronata Willd. ^w	1.23	Rhamnaceae	Sula	Hemba	Lv Sb Ro [111]	In vitro α-glucosidase: 1 mg/mL, A: 70% [109] In vitro α-glucosidase: 62.5 μg/mL, A: 20% [110] Rat: 150–300 mg/kg [112]

A.U.: antidiabetic use, A.D.: previous antidiabetic activity, F_{RC}: relative frequency of citation in percentage, Bb: bulb, Rb: root bark, Sb: stem bark, Lv: leaves, Fr: fruits, Sd: seeds, and Ap: aerial parts.

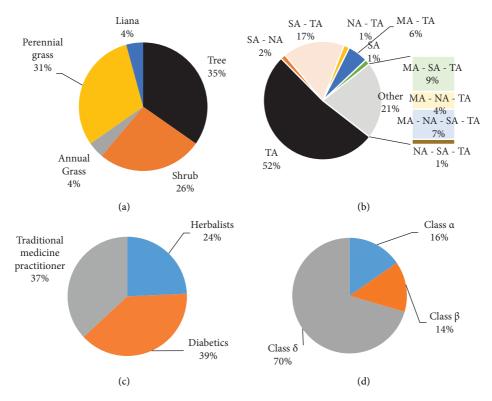


FIGURE 1: Morphological (a) and geographical (b) type of plants, category of informants (c), and situation of the plant with the literature (d). NA: North Africa; CA: Central Africa; TA: tropical Africa; SA: Southern Africa; MA: Madagascar; plants with pharmacological study concerning diabetes: class α ; plants without ethnobotanical or pharmacological study related to diabetes: class β ; plants for ethnobotanical use but without pharmacological research concerning diabetes: class δ .

ethnobotanical and ethnomedicinal profiles. The study reported 71 taxa used in 86 recipes to treat diabetes mellitus.

These 71 plants are primarily trees from the Fabaceae family, endemic to tropical Africa, and are named mainly in Kikongo (Table 1 and Figures 2 and 3). These results are in agreement with the literature. Indeed, Fabaceae is reported to be the most prominent family of trees in Africa's tropical and dry forests [113]. This importance of Fabaceae is observed both within the plant kingdom [114] and within African medicinal plants in particular [115]. The numerical predominance of Fabaceae in sub-Saharan Africa has been attributed to their ability to scavenge atmospheric nitrogen, allowing them to grow in nutrient-poor and rich soils [116]. In our experimental framework, no accessible study has addressed the question of the preponderance of a botanical family over all the taxa used in traditional medicine in Kinshasa. Such a study would be desirable. However, the analysis of some ethnobotanical studies in Kinshasa shows that the Fabaceae constitutes one of the three most evoked families. Some previous studies in the Kinshasa reported the predominance of certain botanical families over others. As an illustration, a study on ethnopharmacological surveys of plants used in female intimate baths [116] mentioned the predominance of Rubiaceae (37%) and Fabaceae (11%). In contrast, another study on plants used to treat the symptoms of tuberculosis [117] reported the predominance of Fabaceae, Apocynaceae, and Lamiaceae, each with an FCR of 8.3%. In the same way, FCR = 47% and FCR = 12.7% were found respectively in the families of Rubiaceae and Fabaceae for a study on plants sold on the Kinshasa market [118], and an FCR of 7.7% in Fabaceae, Rubiaceae and Zingiberaceae for medicinal plants of the Lukunga district [119]. There are also studies done on the management of diabetes in the region. This is the case of a survey carried out in the Kimbanseke and Selembao communes: n = 21, Rubiaceae $F_{\rm CR} = 33.3\%$, Fabaceae $F_{\rm CR} = 19.1\%$ [9], or the study in Kwango and Kinshasa: n = 68, Fabaceae $F_{\rm CR} = 19.1\%$ [8]. It would nevertheless be interesting to carry out an inventory of plants known to be medicinal in the DRC to highlight their specific ethnobotanical characteristics. This concern has not been the subject of this study, which nevertheless contributes to highlighting the need for such data.

The literature review carried out on 71 taxa shows that it is possible to group the plants into 3 classes: species for which no ethnobotanical or pharmacological information is available (class α), those with ethnopharmacological use without any scientific evidence (class δ), and species for which scientific evidence is available (class β) (Figure 1(d)). The fact that 70% of species found are reported to be antihyperglycemic *in vivo* models (using rats) reinforces the credibility of the information obtained from our study. It suggests a high probability of finding those with antihyperglycemic activity among the 30% of the remaining species.

The 6 plant species whose antidiabetic ethnomedicinal knowledge is reported for the first time during this study and for which no pharmacological antidiabetic activity survey is registered in the accessible literature to date are *Acalypha*

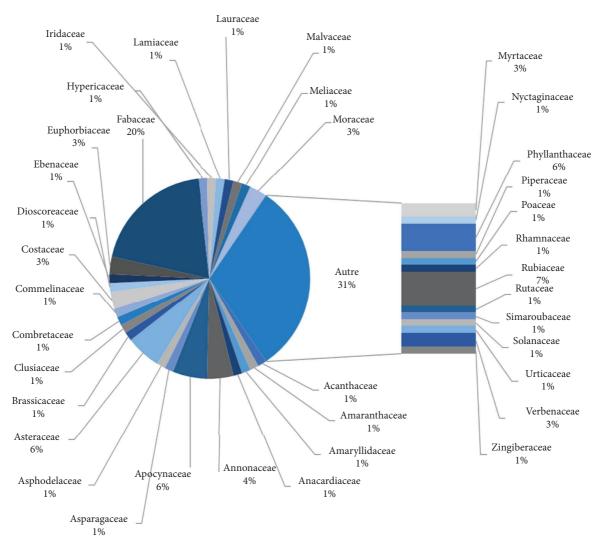


FIGURE 2: Botanical family of listed plant species. Wp: whole plants, Ro: roots, Tb: tubers, and Nr: not reported.

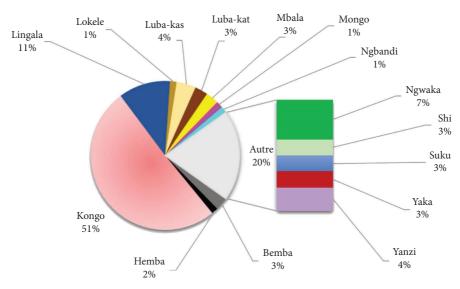


FIGURE 3: Languages of names of inventoried plants.

TABLE 2: Antidiabetic recipes in monophytotherapy and other indications.

Plant species	R_{EC}	UP	Preparation	E_R	Eesp	ICR	ICp	IFR	OAO	UVP	Other indications
Abelmoschus esculentus ^X	R1	Lv	Maceration	2	2	0.01	0.01	1.0	0.04	0.04	Constipation
Acalypha paniculata ^W	R2	Rb	Decoction	4	4	0.01	0.01	1.0	0.04	0.04	Amoeba
Aframomum melegueta ^W	R3	ps	Infusion	7	^	0.02	0.02	1.0	0.04	0.04	Gastroesophageal reflux
Albizia adianthifolia ^W	R4	Lv	Maceration	4	4	0.01	0.01	1.0	90.0	90.0	Sexual weakness and sores
Alchornea cordifolia ^Y	R5	Ro	Decoction	3	3	0.01	0.01	1.0	90.0	90.0	Malaria, stomach aches
Aloe succotrina ^V	R6	Lv	Infusion	2	2	0.02	0.02	1.0	0.08	0.08	Acne, malaria, and wounds
Anacardium occidentale ^X	R7	Lv	Maceration	3	3	0.01	0.01	1.0	0.04	0.04	Warts
Annong conegalousis W	R8	Sp	Decoction	8	10	0.02	0.03	8.0	0.10	0.12	Hepatitis, malaria, sexual impotence, and sores
THEOREM SCHOOLINGS	R9	Ro	Decoction	7		0.01		0.2	0.04		Urogenital infections
Antidesma venosum ^Y	R10	Ro	Decoction	2	2	0.02	0.02	1.0	0.08	0.08	Chest pain, pneumonia, and cough
Artemisia afra ^x	R11	Lv	Maceration	2	2	0.02	0.02	1.0	0.04	0.04	Malaria
Asparagus racemosus ^x	R12	Wp	Decoction	4	4	0.01	0.01	1.0	0.04	0.04	Joint stiffness
Azadirachta indica	R13	Lv	Maceration	7	7	0.01	0.01	1.0	0.04	0.04	Diarrhea, malaria, and stomachaches
Baphia capparidifolia ^x	R14	Γv	Infusion	18	18	90.0	90.0	1.0	0.04	0.04	Inflammation
ı	R15	Ro	Infusion	14	18	0.04	90.0	8.0	0.04	0.08	Cataract
Bidens pilosa ²	R16	Fr	Decoction	7		0.01		0.1	0.04		Glaucoma
;	R17	Lv	Maceration	7		0.01		0.1	0.04		Pyomyositis
Boerhavia diffusa ^Y	R18	Lv	Maceration	∞	8	0.02	0.02	1.0	0.04	0.04	Angina
Brassica juncea ^W	R19	Lv	Infusion	12	12	0.04	0.04	1.0	0.04	0.04	Eczema
Bridelia ferruginea ²	R20	Rb	Decoction	11	11	0.03	0.03	1.0	0.04	0.04	Dislocation
Brillantaisia owariensis ^Y	R21	Rb	Decoction	4	4	0.01	0.01	1.0	0.04	0.04	Epilepsy
Cajanus cajan ^z	R22	Γv	Maceration	18	18	90.0	90.0	1.0	0.04	0.04	Malaria
Catharanthus roseus ^V	R23	Lv	Infusion	4	4	0.01	0.01	1.0	0.04	0.04	High blood pressure
Chromolaena corymbosa $^{ m X}$	R24	Lv	Maceration	21	21	90.0	90.0	1.0	0.04	0.04	Tuberculosis
$Citrus \times aurantium^{W}$	R25	Lv	Infusion	9	9	0.02	0.02	1.0	0.04	0.04	Colic
Costus lucanusianus ²	R26	Γv	Maceration	11	11	0.03	0.03	1.0	0.04	0.04	Peptic ulcer disease
Costus phyllocephalus ^x	R27	Γv	Decoction	4	4	0.01	0.01	1.0	0.04	0.04	Cough
Crinum ornatum ²	R28	Γv	Maceration	9	9	0.02	0.02	1.0	0.04	0.04	Conjunctivitis
Crossopteryx febrifuga ^W	R29	. Ro	Decoction	14	16	0.04	0.05	0.9	90.0	0.06	Diarrhea and malaria
1 / 6 / 6	K30	Υ	Decoction	7		0.01		0.1	90.0		Diarrhea and malaria
Crotalaria medicaginea ²	R31	Wp	Infusion	4	4	0.01	0.01	1.0	0.04	0.04	Cough
Cymbopogon citratus 🖔	R32	Γv	Maceration	15	15	0.02	0.02	1.0	0.04	0.04	Malaria and cough
Dioscorea dumetorum ^x	R33	Tp	Decoction	19	19	90.0	90.0	1.0	90.0	90.0	Diarrhea and wound
Diospyros heudelotii ^w	R34	Ro	Decoction	12	12	0.04	0.04	1.0	0.04	0.04	Contusion
Dysphania ambrosioides ^x	R35	Wp	Decoction	31	31	0.10	0.10	1.0	90.0	90.0	Intestinal bruising and worms
Existence abuscinica W	R36	Rb	Decoction	7	4	0.01	0.01	0.5	90.0	0.12	Sexual weakness and malaria
	R37	Γv	Decoction	7		0.01		0.5	0.08		Cancer, cough, and tuberculosis
Ficus benghalensis ^Y	R38	Sb	Decoction	2	5	0.02	0.02	1.0	90.0	90.0	Eczema and vomiting
Ficus exasperata ^x	R39	Γv	Infusion	4	4	0.01	0.01	1.0	90.0	90.0	Convulsions and peptic ulcer disease
Garcinia kola	R40	ps	Infusion	09	09	0.18	0.18	1.0	0.04	0.04	Glaucoma
Gladiolus gregarius ^W	R41	Bb	Infusion	7	7	0.01	0.01	1.0	90.0	90.0	Angina and amoeba
Gymnanthemum amygdalinum	R42	À Ä	Decoction	10	10	0.03	0.03	1.0	0.10	0.10	High blood pressure, STI, malaria, and cough
Harungana madagascariensis	K43	8	Decoction	×	∞	0.02	0.02	1.0	0.06	0.06	Sickle cell disease and malaria

TABLE 2: Continued.

$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Flant species	R_{EC}	UP	Preparation	E_R	Eesp	ICR	ICp	IFR	UVO	UVP	Other indications
R45 Ro Decoction 14 14 0.04 0.04 1.0 0.04 0.04 R46 Fr Decoction 12 12 0.04 0.04 1.0 0.06 0.06 R48 Ro Decoction 3 3 0.01 1.0 0.08 0.06 R49 Ro Decoction 3 3 0.01 0.01 1.0 0.08 0.08 R50 Rb Decoction 4 0.01 0.01 1.0 0.04 0.04 R52 Lv Maceration 2 0.01 0.01 0.01 0.01 0.04 0.04 R53 Fr Infusion 2 0.01 0.01 0.01 0.04 0.04 0.04 R54 Lv Maceration 4 4 0.01 0.01 0.04 0.04 0.04 0.04 0.04 0.04 0.04 0.04 0.04 0.04 0.04 0.04 <td< td=""><td>Heinsia crinita^Z</td><td>R44</td><td>Lv</td><td>Decoction</td><td>6</td><td>6</td><td>0.03</td><td>0.03</td><td>1.0</td><td>90.0</td><td>90.0</td><td>Cough and HIV</td></td<>	Heinsia crinita ^Z	R44	Lv	Decoction	6	6	0.03	0.03	1.0	90.0	90.0	Cough and HIV
R46 Fr Decoction 12 12 0.04 0.04 1.0 0.06 0.06 R47 Lv Maceration 4 4 0.01 0.01 1.0 0.08 0.08 R48 R0 Decoction 3 3 0.01 0.01 1.0 0.08 0.08 R50 R6 Decoction 4 4 0.01 0.01 1.0 0.04 0.04 R51 S4 Decoction 5 8 0.02 0.02 0.02 0.04 0.04 0.04 R53 Fr Infusion 2 0.01 0.01 1.0 0.04 0.04 0.04 R54 Lv Maceration 4 4 0.01 0.01 1.0 0.04 0.04 0.04 R55 Lv Maceration 4 4 0.01 0.01 1.0 0.04 0.04 0.04 0.04 0.04 0.04 0.04 0.04 <td>Hymenocardia acida$^{ ext{ iny X}}$</td> <td>R45</td> <td>Ro</td> <td>Decoction</td> <td>14</td> <td>14</td> <td>0.04</td> <td>0.04</td> <td>1.0</td> <td>0.04</td> <td>0.04</td> <td>Sickle cell disease</td>	Hymenocardia acida $^{ ext{ iny X}}$	R45	Ro	Decoction	14	14	0.04	0.04	1.0	0.04	0.04	Sickle cell disease
R47 Ly Maceration 4 4 0.01 0.01 1.0 0.08 0.08 R48 Ro Decoction 3 3 0.01 1.0 0.04 0.04 R48 Ro Decoction 3 3 0.01 1.0 0.04 0.04 R50 Rb Decoction 5 8 0.02 0.02 0.05 0.04 0.04 R51 Sd Decoction 2 8 0.02 0.02 0.05 0.04 0.04 R52 Lv Maceration 4 4 0.01 0.01 1.0 0.04 0.04 R55 Lv Maceration 4 4 4 0.01 0.01 1.0 0.04 R6W Lv Decoction 3 5 0.01 0.01 1.0 0.04 0.04 R8W Lv Decoction 3 5 0.01 0.01 0.04 0.04	Lantana camara ^w	R46	Fr	Decoction	12	12	0.04	0.04	1.0	90.0	90.0	Malaria and cough
R48 Ro Decoction 3 3 0.01 0.01 1.0 0.08 0.08 R49 Ro Decoction 3 3 0.01 0.01 1.0 0.04 0.04 R51 Sd Decoction 5 3 0.01 1.0 0.04 0.04 R52 Lv Maceration 2 8 0.01 0.01 1.0 0.04 0.04 R53 Lv Maceration 4 4 0.01 1.0 0.04 0.04 R54 Lv Maceration 4 4 0.01 0.01 0.01 0.01 0.04 0.08 R85 Lv Decoction 3 5 0.01 0.01 0.01 0.01 0.04 0.04 R85 Lv Maceration 4 4 4 0.01 0.01 0.01 0.01 0.04 0.04 0.04 0.04 0.04 0.04 0.04 0.04	Lippia multiflora ^Y	R47	Lv	Maceration	4	4	0.01	0.01	1.0	0.08	0.08	Angina, asthma, and sexual weakness
R49 Ro Decoction 3 3 0.01 0.01 1.0 0.04 0.04 R50 Rb Decoction 4 4 4 0.01 0.01 1.0 0.04 0.04 R51 SA Decoction 2 8 0.02 0.02 0.02 0.04 0.04 0.04 R53 Fr Infusion 1 0.00 0.01 0.01 0.01 0.04 0.04 R54 LV Maceration 4 4 4 0.01 0.02 0.02 0.02 0.03 0.04 0.08 0.08 0.08 0.08 0.08 0.08 0.08 0.08 0.08 0.08 0.08 0.08 0.08 0.08 0.08	Marsdenia latifolia ^W	R48	Ro	Decoction	3	3	0.01	0.01	1.0	80.0	0.08	Constipation, cough, and intestinal worms
R50 Rb Decoction 4 4 0.01 0.01 1.0 0.04 0.04 R51 Sd Decoction 5 8 0.02 0.02 0.02 0.04 0.08 R52 Fr Infusion 1 0.00 0.01 0.04 0.04 R53 Fr Infusion 1 0.00 0.0 0.04 0.04 R54 Lv Maceration 4 4 0.01 0.01 1.0 0.04 0.04 R54 Lv Maceration 4 4 0.01 0.01 1.0 0.04 0.04 R58 Lv Decoction 2 0.01 0.01 1.0 0.04 0.04 R84 Lv Maceration 2 0.02 0.02 0.02 0.05 0.04 0.04 R84 R6 R6 R6 6 6 0.02 0.02 0.05 0.04 0.04 <	Millettia drastica ^W	R49	Ro	Decoction	3	3	0.01	0.01	1.0	0.04	0.04	Sickle cell disease
R51 Sd Decoction 5 8 0.02 0.02 0.02 0.02 0.03 0.04 0.08 R52 Lv Maceration 1 0.00 0.1 0.04 0.04 R54 Lv Maceration 4 4 0.01 0.01 1.0 0.04 0.04 R55 Lv Maceration 4 4 0.01 0.01 1.0 0.04 0.04 R55 Lv Maceration 2 0.01 0.01 1.0 0.04 0.08 R85 Lv Decoction 3 5 0.01 0.01 1.0 0.04 0.08 R87 Rb Decoction 2 0.01 0.01 1.0 0.04 0.04 R84 Lv Maceration 6 8 0.02 0.02 0.02 0.05 0.04 0.04 R85 Sb Infusion 5 5 0.02 0.02 0.02 <td></td> <td>R50</td> <td>Rb</td> <td>Decoction</td> <td>4</td> <td>4</td> <td>0.01</td> <td>0.01</td> <td>1.0</td> <td>0.04</td> <td>0.04</td> <td>Urogenital infections</td>		R50	Rb	Decoction	4	4	0.01	0.01	1.0	0.04	0.04	Urogenital infections
R52 Lv Maceration 2 0.01 0.03 0.04 R53 Fr Infusion 1 0.00 0.1 0.04 R54 Lv Maceration 4 4 4 0.01 1.0 0.04 0.04 R55 Lv Maceration 3 5 0.01 1.0 0.04 0.04 0.04 R55 Lv Decoction 3 5 0.01 1.0 0.04 0.05 0.05 0.05 0.05 0.02 0.05 0.05 0.02 0.05 0.05 0.02 0.05 0.05 0.05 0.05 0.05 0.05 0.05 0.05 0.05 0.05 0.06 <td>Washing chitical</td> <td>R51</td> <td>PS</td> <td>Decoction</td> <td>2</td> <td>8</td> <td>0.02</td> <td>0.02</td> <td>9.0</td> <td>0.04</td> <td>0.08</td> <td>High blood pressure</td>	Washing chitical	R51	PS	Decoction	2	8	0.02	0.02	9.0	0.04	0.08	High blood pressure
R53 Fr Infusion 1 0.00 0.01 0.04 R54 Ly Maceration 4 4 0.01 0.01 0.09 0.08 0.08 R55 Ly Decoction 2 0.01 0.01 0.06 0.08 0.08 R57 Ro Decoction 2 0.01 0.01 0.04 0.08 0.08 R58 Ly Decoction 2 0.01 0.01 0.04 0.06 0.08 R59 Ly Decoction 2 0.01 0.01 0.01 0.01 0.09 0.08 0.08 R64 Rb Decoction 5 5 0.02 0.02 0.02 0.06 0.06 R64 Ap Infusion 2 0	minagyna supaiosa	R52	Lv	Maceration	7		0.01		0.3	0.04		Anthrax
R54 Lv Maceration 4 4 0.01 0.01 0.09 0.04 0.04 R55 Lv Maceration 4 4 0.01 0.01 0.08 0.08 R56 Lv Decoction 3 5 0.01 0.01 0.09 0.08 0.08 R58 Lv Decoction 4 4 0.01 0.01 0.04 0.08 0.08 R59 Lv Maceration 6 6 0.02 0.02 1.0 0.04 0.04 R61 Lv Maceration 6 6 0.02 0.02 0.04 0.04 R62 Sd Infusion 5 5 0.02 1.0 0.04 0.04 R63 Sb Maceration 6 8 0.02 0.0 0.0 0.0 0.04 0.04 R64 Ap Infusion 2 2 0.02 0.0 0.0 0.0 <t< td=""><td></td><td>R53</td><td>Fr</td><td>Infusion</td><td>1</td><td></td><td>0.00</td><td></td><td>0.1</td><td>0.04</td><td></td><td>Cancer</td></t<>		R53	Fr	Infusion	1		0.00		0.1	0.04		Cancer
R55 Lv Maceration 4 4 0.01 0.01 1.0 0.08 0.08 R56 Ly Decoction 3 5 0.01 0.02 0.6 0.08 0.01 R57 Ly Decoction 4 4 4 0.01 1.0 0.08 0.08 0.08 0.08 0.08 0.08 0.08 0.08 0.09	Morinda lucida ^Y	R54	Lv	Maceration	4	4	0.01	0.01	1.0	0; 04	0.04	Cancer
R56 Ly Decoction 3 5 0.01 0.02 0.6 0.08 0.12 R57 Ro Decoction 2 0.01 0.01 0.4 0.06 R58 Ly Decoction 4 4 0.01 0.01 1.0 0.08 0.08 R59 Ly Maceration 6 6 0.02 1.0 0.04 0.04 R61 Ly Maceration 6 6 0.02 1.0 0.04 0.04 R63 SA Infusion 5 5 0.02 0.0 0.03 0.04 0.04 R64 Ap Infusion 2 0.01 0.01 0.01 0.02 0.02 0.0 0.06 0.06 R65 Rb Decoction 5 5 0.02 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0	$\mathit{Musanga}$ cecropioides $^{\mathrm{W}}$	R55	Γv	Maceration	4	4	0.01	0.01	1.0	0.08	0.08	High blood Pressure, cough, and tuberculosis
R57 Ro Decoction 2 0.01 0.04 0.06 R58 Ly Decoction 4 4 0.01 0.01 1.0 0.08 0.08 R89 Ly Maceration 27 27 0.08 0.08 1.0 0.06 0.06 R61 Ly Maceration 6 6 0.02 0.02 1.0 0.04 0.04 R62 Sb Infusion 5 0.02 0.02 1.0 0.04 0.04 R64 Ap Infusion 2 0.01 0.01 0.01 0.01 0.03 0.06 0.06 R65 Rb Decoction 5 5 0.02 0.02 1.0 0.04 0.08 R67 Fr Infusion 2 2 0.01 0.01 1.0 0.04 0.04 R68 Fr Maceration 2 2 0.01 0.01 0.01 0.01 0.01	Mangles latifolish	R56	Lv	Decoction	8	5	0.01	0.02	9.0	0.08	0.12	Malaria, toothache, and sore
R58 Ly Decoction 4 4 0.01 0.01 1.0 0.08 0.08 a ^W R60 Rb Decoction 6 6 0.02 0.02 1.0 0.04 0.04 R61 Ly Maceration 6 6 0.02 0.02 1.0 0.04 0.04 R62 Sd Infusion 5 5 0.02 0.02 1.0 0.04 0.04 R63 Sb Maceration 6 8 0.02 0.02 1.0 0.04 0.04 R64 Ap Infusion 2 0.01 0.01 0.02 0.02 0.02 0.06 0.06 R64 Ap Infusion 2 0.01 0.01 0.01 0.03 0.04 0.08 R65 Rb Decoction 5 5 0.02 0.02 0.02 0.03 0.04 0.04 R68 Fr Infusion 2	induciea tatifotta	R57	Ro	Decoction	7		0.01		0.4	90.0		Bronchitis and dysentery
R59 Lv Maceration 27 27 0.08 0.08 1.0 0.06 0.06 Ad R60 Rb Decoction 6 6 0.02 0.02 1.0 0.04 0.04 R61 Lv Maceration 100 100 0.31 0.31 1.0 0.04 0.04 R62 Sd Infusion 5 5 0.02 0.02 1.0 0.04 0.04 R63 Sb Maceration 6 8 0.02 0.02 0.02 0.02 0.06 0.06 R64 Ap Infusion 2 2 0.01 0.01 0.01 0.08 0.08 R65 Fr Infusion 2 2 0.01 1.0 0.04 0.04 R69 Lv Maceration 3 3 0.01 1.0 0.02 0.02 0.02 0.03 0.04 0.08 R69 Lv Infusion	Ocimum gratissimum ^Y	R58	Lv	Decoction	4	4	0.01	0.01	1.0	0.08	0.08	Diarrhea, wound, and sinusitis
taw R60 Rb Decoction 6 6 0.02 0.02 1.0 0.04 0.04 R61 Lv Maceration 100 100 0.31 0.31 1.0 0.04 0.04 R62 Sd Infusion 5 5 0.02 0.02 1.0 0.04 0.04 R63 Sb Maceration 6 8 0.02 0.02 0.02 0.09 0.09 0.04 R64 Ap Infusion 2 0.01 0.01 0.10 0.01 0.01 0.01 0.03 0.06 0.08 0.09 0.09 0.09 0.09 0.09 0.09 </td <td>Palisota schweinfurthii^X</td> <td>R59</td> <td>Lv</td> <td>Maceration</td> <td>27</td> <td>27</td> <td>0.08</td> <td>0.08</td> <td>1.0</td> <td>90.0</td> <td>90.0</td> <td>Snakebite and rheumatism</td>	Palisota schweinfurthii ^X	R59	Lv	Maceration	27	27	0.08	0.08	1.0	90.0	90.0	Snakebite and rheumatism
R61 Lv Maceration 100 100 0.31 0.31 1.0 0.04 0.04 R62 Sd Infusion 5 5 0.02 0.02 1.0 0.06 0.06 R63 Sb Maceration 6 8 0.02 0.0 0.0 0.06 0.06 R64 Ap Infusion 2 0.01 0.01 0.03 0.06 0.08 R65 Rb Decoction 5 5 0.02 0.02 0.02 0.08 0.08 R65 Fr Infusion 2 2 0.01 0.02 0.02 0.02 0.02 0.02 0.02 0.02 0.03	Pentaclethra macrophylla $^{ m W}$	R60	Rb	Decoction	9	9	0.02	0.02	1.0	0.04	0.04	Amoeba
R62 Sd Infusion 5 5 0.02 0.02 1.0 0.06 0.06 R63 Sb Maceration 6 8 0.02 0.03 0.04 0.08 R64 Ap Infusion 2 0.01 0.03 0.06 0.08 R65 Rb Decoction 58 58 0.18 1.0 0.08 0.08 R66 Rb Decoction 2 2 0.01 1.0 0.04 0.04 R68 Fr Infusion 2 2 0.01 1.0 0.04 0.04 R69 Lv Maceration 2 2 0.01 1.0 0.01 0.01 0.01 0.01 0.01 0.01 0.00 0.08 0.08 R74 Ro Decoction 3 3 0.01 0.01 1.0 0.06 0.06 R71 Wp Decoction 3 3 0.01 0.01	Persea americana ^X	R61	Lv	Maceration	100	100	0.31	0.31	1.0	0.04	0.04	High blood pressure
R63 Sb Maceration 6 8 0.02 0.03 0.04 0.08 R64 Ap Infusion 2 0.01 0.3 0.06 0.08 R65 Rb Decoction 58 58 0.18 1.0 0.08 0.08 R66 Rb Decoction 5 5 0.02 1.0 0.08 0.08 R67 Fr Infusion 2 2 0.01 1.0 0.04 0.04 R69 Lv Maceration 2 2 0.01 1.0 0.01 0.01 R70 Lv Maceration 2 2 0.01 1.0 0.08 0.08 R71 Wp Decoction 3 3 0.01 1.0 0.01 0.01 0.01 0.06 0.06 R72 Lv Maceration 3 3 0.01 1.0 0.04 0.04 0.04 R75 Lv	Phaseolus vulgaris ^W	R62	ps	Infusion	2	2	0.02	0.02	1.0	90.0	90.0	Wounds and ulcer
K64 Ap Infusion 2 0.01 0.3 0.06 R65 Rb Decoction 58 58 0.18 1.0 0.08 0.08 R66 Rb Decoction 5 5 0.02 1.0 0.08 0.08 R67 Fr Infusion 2 2 0.01 1.0 0.01 0.04 R68 Fr Maceration 2 2 0.01 1.0 0.04 0.04 R74 Ro Decoction 2 2 0.01 0.01 0.01 0.01 0.01 0.00 0.08 R70 Lv Maceration 2 2 0.01 0.01 0.01 0.01 0.01 0.06 0.06 R71 Wp Decoction 8 8 0.02 0.02 0.0 0.06 0.06 R73 Lv Infusion 7 7 0.02 0.0 0.0 0.06 0.06	Dhyllanthus nimi	R63	Sp	Maceration	9	8	0.02	0.02	8.0	0.04	0.08	Wounds
K65 Rb Decoction 58 58 0.18 1.0 0.08 0.08 R66 Rb Decoction 5 5 0.02 0.02 1.0 0.08 0.08 R67 Fr Infusion 2 2 0.01 1.0 0.04 0.04 R68 Fr Maceration 2 2 0.01 1.0 0.02 0.02 R74 Ro Decoction 2 2 0.01 0.01 0.01 0.01 0.00 0.08 R70 Lv Maceration 2 2 0.01 0.01 0.01 0.00 0.06 0.06 R71 Wp Decoction 8 8 0.02 0.02 1.0 0.06 0.06 R73 Lv Infusion 7 7 0.02 1.0 0.06 0.06 R75 Lv Infusion 26 26 0.08 0.02 1.0 0.08 <	1 17 1111 1111 111 111 111 1	R64	Ap	Infusion	7		0.01		0.3	90.0		Urogenital infections and ulcers
R66 Rb Decoction 5 5 0.02 0.02 1.0 0.08 0.08 R67 Fr Infusion 2 2 0.01 0.01 1.0 0.04 0.04 R68 Fr Maceration 2 2 0.01 1.0 0.02 0.02 R69 Lv Maceration 2 2 0.01 1.0 0.01 0.08 R74 Ro Decoction 3 3 0.01 0.01 1.0 0.10 0.10 0.10 R71 Wp Decoction 8 8 0.02 1.0 0.06 0.06 R72 Lv Infusion 7 7 0.02 1.0 0.04 0.04 R75 Lv Infusion 5 5 0.02 1.0 0.06 0.06 R77 Lv Infusion 26 26 0.08 0.02 1.0 0.08 0.08 0.08	Piliostigma reticulatum ^x	R65	Rb	Decoction	28	28	0.18	0.18	1.0	80.0	0.08	Bronchitis, headaches, and rheumatism
R67 Fr Infusion 2 2 0.01 0.01 1.0 0.04 0.04 R68 Fr Maceration 8 8 0.02 1.0 0.02 0.02 R69 Lv Maceration 2 2 0.01 1.0 0.10 0.08 R74 Ro Decoction 2 2 0.01 1.0 0.10 0.10 R70 Lv Maceration 3 3 0.01 0.01 1.0 0.10 0.10 R72 Lv Maceration 100 100 0.31 1.0 0.04 0.04 R73 Lv Infusion 7 7 0.02 1.0 0.06 0.06 R75 Lv Infusion 26 26 0.08 0.02 1.0 0.06 0.06 R77 Lv Maceration 8 8 0.02 1.0 0.08 0.08 R78 R0	Piliostigma thonningii ^x	R66	Rb	Decoction	2	2	0.02	0.02	1.0	0.08	0.08	Diarrhea, wound, and ulcer
R68 Fr Maceration 8 8 0.02 0.02 1.0 0.02 0.02 R69 Lv Maceration 2 2 0.01 0.01 1.0 0.08 0.08 R74 Ro Decoction 2 2 0.01 0.01 1.0 0.10 0.10 0.10 R70 Lv Maceration 8 8 0.02 1.0 0.06 0.06 R72 Lv Infusion 7 7 0.02 0.02 1.0 0.04 0.04 R75 Lv Infusion 5 5 0.02 1.0 0.06 0.06 R76 Lv Infusion 26 26 0.08 0.02 1.0 0.08 0.08 R77 Lv Maceration 8 8 0.02 1.0 0.08 0.08 R78 R0 Decoction 58 58 0.18 1.0 0.08 0.08	Piper guineense ^w	R67	Η̈́	Infusion	7	2	0.01	0.01	1.0	0.04	0.04	Asthma
R69 Lv Maceration 2 2 0.01 0.01 1.0 0.08 0.08 R74 Ro Decoction 2 2 0.01 0.01 1.0 0.10 0.10 0.10 R70 Lv Maceration 3 3 0.01 0.01 1.0 0.06 0.06 R71 Wp Decoction 8 8 0.02 1.0 0.06 0.06 R73 Lv Infusion 7 7 0.02 1.0 0.06 0.06 R75 Lv Maceration 5 5 0.02 1.0 0.06 0.06 R77 Lv Maceration 8 8 0.02 1.0 0.08 0.08 R78 R0 Decoction 58 58 0.18 1.0 0.08 0.08	Platymitra arborea ^Y	R68	Ŧ	Maceration	∞	8	0.02	0.02	1.0	0.02	0.02	NR
R74 Ro Decoction 2 2 0.01 0.01 1.0 0.10 0.10 0.10 0.10 0.10 0.10 0.10 0.10 0.10 0.10 0.10 0.10 0.01 0.01 0.01 0.01 0.01 0.02 0.02 0.02 0.02 0.02 0.06 0.06 0.06 0.06 0.04 0.04 0.04 0.04 0.04 0.04 0.04 0.04 0.04 0.04 0.04 0.04 0.04 0.06	Psidium guajava ^W	R69	ΓΛ	Maceration	7	7	0.01	0.01	1.0	0.08	0.08	Diarrhea, hepatitis, and malaria
R70 Lv Maceration 3 3 0.01 0.01 1.0 0.06 0.06 R71 Wp Decoction 8 8 0.02 0.02 1.0 0.06 0.06 R72 Lv Infusion 7 7 0.02 0.0 1.0 0.04 0.04 R75 Lv Infusion 5 5 0.02 1.0 0.06 0.06 R76 Lv Infusion 26 26 0.08 0.03 1.0 0.02 0.02 R77 Lv Maceration 8 8 0.02 1.0 0.08 0.08 R78 R0 Decoction 58 58 0.18 1.1 0.08 0.08 0.08	Quassia amara ^X	R74	Ro	Decoction	7	7	0.01	0.01	1.0	0.10	0.10	Malaria, sexual weakness, typhoid, and rheumatism
R71 Wp Decoction 8 8 0.02 0.02 1.0 0.06 0.06 R72 Lv Maceration 100 100 0.31 0.31 1.0 0.04 0.04 R73 Lv Infusion 7 7 0.02 0.02 1.0 0.06 0.06 R75 Lv Infusion 26 26 0.08 0.08 1.0 0.02 0.02 R77 Lv Maceration 8 8 0.02 1.0 0.08 0.08 0.08 R78 Ro Decoction 58 58 0.18 1.1 0.08 0.08 0.08	Rauvolfia mannii ^x	R70	Ľ	Maceration	Э	С	0.01	0.01	1.0	90.0	90.0	Jaundice and female infertility
R72 Lv Maceration 100 100 0.31 0.31 1.0 0.04 0.04 R73 Lv Infusion 7 7 0.02 0.02 1.0 0.06 0.06 R75 Lv Maceration 5 5 0.02 1.0 0.06 0.06 R77 Lv Infusion 26 26 0.08 0.02 1.0 0.02 0.02 R78 Ro Decoction 58 58 0.18 1.0 0.08 0.08 0.08	Schwenckia americana ^x	R71	Wp	Decoction	%	∞	0.02	0.02	1.0	90.0	90.0	Diarrhea and cholera
R73 Lv Infusion 7 7 0.02 0.02 1.0 0.06 0.06 R75 Lv Maceration 5 5 0.02 0.02 1.0 0.06 0.06 R76 Lv Infusion 26 26 0.08 0.08 1.0 0.02 0.02 R77 Lv Maceration 8 8 0.02 1.0 0.08 0.08 R78 Ro Decoction 58 58 0.18 1.0 0.08 0.08	Senna alata ^x	R72	Γv	Maceration	100	100	0.31	0.31	1.0	0.04	0.04	Constipation
R75 Lv Maceration 5 5 0.02 0.02 1.0 0.06 0.06 R76 Lv Infusion 26 26 0.08 0.08 1.0 0.02 0.02 R77 Lv Maceration 8 8 0.02 1.0 0.08 0.08 R78 Ro Decoction 58 58 0.18 1.0 0.08 0.08	Senna occidentalis ^V	R73	Lv	Infusion	^	7	0.02	0.02	1.0	90.0	90.0	Sexual weakness and malaria
R76 Lv Infusion 26 26 0.08 0.08 1.0 0.02 0.02 R77 Lv Maceration 8 8 0.02 0.02 1.0 0.08 0.08 0 R78 Ro Decoction 58 58 0.18 1.0 0.08 0.08	Syzygium cordatum ^Y	R75	ΓΛ	Maceration	5	5	0.02	0.02	1.0	90.0	90.0	Diarrhea and cough
R77 Lv Maceration 8 8 0.02 0.02 1.0 0.08 0.08 0.08 0.08 0.08 0.08 0.08	Tephrosia vogelii ^X	R76	Lv	Infusion	56	56	80.0	0.08	1.0	0.02	0.02	NR
R78 Ro Decoction 58 58 0.18 0.18 1.0 0.08 0.08	Terminalia mollis ^Y	R77	Lv	Maceration	∞	8	0.02	0.02	1.0	0.08	0.08	Cough ulcers, measles, and gastroduodenal diseases
	Vachellia karroo ^x	R78	Ro	Decoction	28	28	0.18	0.18	1.0	0.08	0.08	Constipation, diarrhea, and cough
R79 Lv Maceration 2 2 0.01 0.01 1.0 0.06 0.06	Voacanga africana ^w	R79	ΓΛ	Maceration	7	2	0.01	0.01	1.0	90.0	90.0	Sexual weakness and high blood pressure
V R80 Rb Decoction 4 4 0.01 0.01 1.0 0.06 0.06	Ziziphus mucronata ^w	R80	Rb	Decoction	4	4	0.01	0.01	1.0	90.0	90.0	Angina and sickle cell disease

ER (number of citations per recipe); Eesp (number of citations per plant); CIR (consensus index of the recipe); CIP (consensus index of the plant); RRI (recipe reliability index); UVo (the usual value of the organ); UVp (the number of uses of the plant out of the number of uses of the plant out of the number of uses of the plant out of the number of uses); NR (Nothing to report); Nu(number of uses); N (number of people consulted); Nu = 52; N = 326.

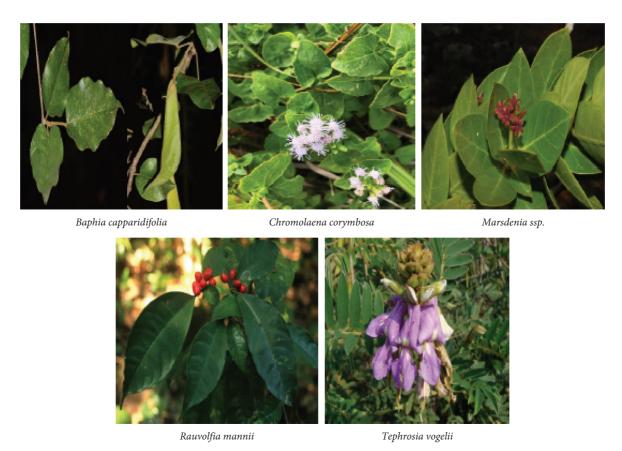


FIGURE 4: Important plant species identified as antidiabetic in Kinshasa (figures collected from https://www.inaturalist.org).

paniculata, Baphia capparidifolia, Chromolaena corymbosa, Crotalaria medicaginea, Platymitra arborea, and Rauvolfia mannii. On the other hand, 11 medicinal plants, including Antidesma venosum, Costus phyllocephalus, Crinum ornatum, Diospyros heudelotii, Gladiolus gregarius, Lippia multiflora, Millettia drastica, Mitragyna stipulosa, Palisota schweinfurthii, Tephrosia vogelii, and Terminalia mollis, have been quoted in traditional medicine as treating diabetes mellitus in various countries. However, they are not yet scientifically validated (Table 1). In both cases, these plants constitute a great richness in the database for future preclinical investigations. However, the antidiabetic evaluation of Costus lucanusianus has been validated after several research attempts; no study has confirmed the local antidiabetic use of this plant.

Previous studies on traditional healers and medicinal plant vendors have identified some species used in Kinshasa city (Figure 7) to manage diabetes mellitus. Thus, Abelmoschus esculentus, Albizia adianthifolia, Alchornea cordifolia, Azadirachta indica, Bridelia ferruginea, Catharanthus roseus, Costus phyllocephalus, Cymbopogon citratus, Dysphania ambrosioides, Erythrina abyssinica, Ficus benghalensis, Gymnanthemum amygdalinum, Lippia multiflora, Morinda lucida, Nauclea latifolia, Phaseolus vulgaris, Phyllanthus niruri, Psidium guajava, Schwenckia americana, Senna alata [8], Annona senegalensis [8, 73, 119], Garcinia kola [73, 119] Gladiolus gregarius [119], Monodora myristica [120], and Persea americana [8, 118] have been cited.

Concerning ethnobotanical indexes, it should be noted that for an ethnobotanical study targeted towards a specific pathology, the relative frequency of citations corresponds to the consensual citation index of the plant, which translates the consensus reached around a particular plant species on targeted use. There is a higher consensus for the 11 taxa not studied on using Tephrosia vogelii leaves (CIp = 0.08) as an antidiabetic than any other species of the same category (Table 2). This precedence which may result from this ethnobotanical index (CIp) is, however, to be put into perspective by the fact that a single category only told the plant of the subjects consulted, namely, people with diabetes, unlike, in particular, the leaves of the Crinum ornatum^Z species which present the weakness of ethnobotanical index whose value is twice lower than that of Tephrosia vogelii (CIp = 0.04), but whose strength lies in the fact that it was informed simultaneously by the three sources: diabetics, TMPs, and herbalists, which reinforces the consensus around its use as an antidiabetic in the study environment.

The study population uses three plant species most because of their highest UV of the series (0.12). These are *Annona senegalensis, Erythrina abyssinica*, and *Nauclea latifolia* (Table 2). This justifies their lower usual value of 0.02 (Table 2). All of its plants belong to the taxa category, whose ethnobotanical and pharmacological knowledge of diabetes has previously been reported. Therefore, the most common plants in the study would not be the most interesting in the context of this study, which aims to enhance the

N°	Taxon 1	Taxon 2	No recipe	PU-ratio	Preparation	D.T. (D)	$F_{\rm CR}$
1	Chromolaena corymbosa ^X	$Dioscorea\ dumetorum^{ m X}$	R81	F-F $(1 \div 2)$	Decoction	30	3.1
2	Senna alata ^X	Alchornea cordifolia ^Y	R82	F-R $(1 \div 1)$	Maceration	45	3.7
3	Morinda lucida ^Y	Nauclea latifolia ^Y	R83	F-R $(2 \div 1)$	Infusion	30	2.5
4	Platymitra arborea ^Y	Schwenckia americana ^Y	R84	R-R $(1 \div 1)$	Decoction	45	3.7
5	Albizia adianthifolia ^W	Annona senegalensis ^W	R85	F-R $(2 \div 1)$	Decoction	45	4.6
6	Garcinia kola ^V	Gymnanthemum amygdalinum ^V	R86	F-F $(2 \div 1)$	Decoction	45	1.8

TABLE 3: Antidiabetic recipes that use two herbs in combination.

PU-ratio: part used and proportion of the mixture; F: leaf; R: root; D.T.: duration of treatment; D: day; the F_{CR} is expressed in % (n = 326); N° recipe-antidiabetic recipe.

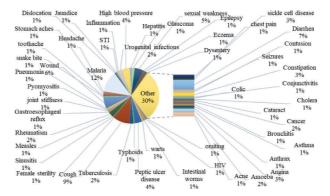


FIGURE 5: Other diseases are cured by the plants inventoried.

TABLE 4: General characteristics of the subjects consulted.

Class	Category	dia	le with betes = 126)		palists = 80)	TMP	(n = 120)	Total % (n = 326)
		E	%	E	%	E	%	
	18-28	3	2.4	6	8	7	5.8	4.9
	28-38	21	16.7	6	8	18	15.0	13.8
Age	38-48	7	5.6	12	15	27	22.5	14.1
C	48-58	35	27.8	41	51	60	50.0	41.7
	>58	60	47.6	15	19	8	6.7	25.4
	1-5	10	7.9	14	18	10	8.3	12.9
	6-10	15	11.9	7	9	58	48.3	24.5
Experience (year)	11-15	20	15.9	40	50	26	21.7	26.4
•	16-20	78	61.9	8	10	20	16.7	32.5
	21-25	3	2.4	11	14	6	5.0	6.1
Profession	Trade	21	16.7	68	85	4	3.3	28.5
Profession	Public sector	3	2.4	0	0	5	4.2	2.5
	Liberal	57	45.2	12	15	10	8.3	24.2
	Housework	45	35.7	0	0	3	2.5	14.7
	TMP	0	0.0	0	0	98	81.7	30.1
C 1	Female	84	66.7	64	80	55	45.8	62.3
Gender	Male	42	33.3	16	20	65	54.2	37.7
Gender	Bumbu	15	11.9	16	20	32	26.7	19.3
	Kalamu	45	35.7	23	29	29	24.2	29.8
Habitation	Lemba	11	8.7	10	13	6	5.0	8.3
	Limete	25	19.8	10	13	8	6.7	13.2
	Makala	30	23.8	21	26	45	37.5	29.4
	None	2	1.6	20	25	6	5.0	8.6
	Primary	21	16.7	15	19	18	15.0	16.6
Level of studies	Professional	7	5.6	3	4	10	8.3	6.1
	Secondary	47	37.3	41	51	66	55.0	47.2
	University	49	38.9	1	1	20	16.7	21.5

TMP: practitioner of traditional medicine; experience in the diabetic category is the number of years since the patient was diagnosed with diabetes.

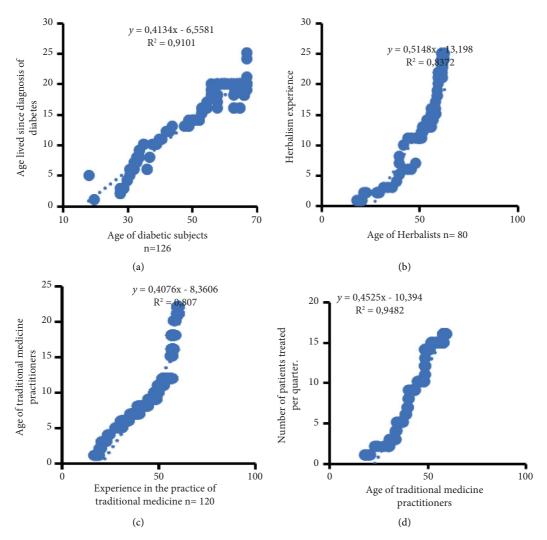


FIGURE 6: Correlations between Y-axis and X-axis variables of the characteristics of the subjects surveyed: Age lived since diagnosis of diabetes and age of diabetic subjects n=126 (a); Herbalism experience and age of herbalists n= 80 (b); Age of traditional medicine and practitioners (c); and Number of patients treated per quarter and age of traditional medicine practitioners (d).

ethnopharmacological knowledge of Kinshasa. If, in most cases, each plant inventoried during this study is also known and used by our resource persons for at least one other pathology, it should be emphasized that this is not the case with *Tephrosia vogelii*^X and *Platymitra arborea*^Y for which our informants only use it in the management of diabetes mellitus.

Malaria (FCR = 12%), cough (FCR = 9%), and diarrhea (FCR = 7%) constituted the 3 first pathologies, apart from diabetes, for which the plants listed during this study are used. These pathologies are among the 10 most deadly pathologies in the DRC [121], suggesting that traditional medicine from Kinshasa will likely contribute to managing specific pathologies of concern in DR Congo. It would be interesting to carry out studies to validate this ethnopharmacological knowledge to rightly understand the assistance of conventional medicine from Kinshasa in managing pathologies such as diabetes.

The 71 inventoried plant taxa made it possible to list 86 antidiabetic recipes, 80 of which use a single plant (Table 2), and 6 recipes combine 2 simultaneous medicinal plants

(Table 3). Nonetheless, considering all previous ethnopharmacological studies conducted in DR Congo [8, 15, 38, 48], this one reports antidiabetic recipes of the plants used in association for the first time.

Of all the antidiabetic recipes using a single plant and reported during this study, the leaf and the decoction constitute the most used organ and mode of preparation. These results (Figure 8) are in agreement with the studies carried out on the plants of Kisangani: leaf (57.6%), decoction (78.8%), and n = 33 [38], and plants of DR Congo gathered in a review article, leaves 39.2%, decoction: 60.5%, and n = 213 [52]. Some disparities are nevertheless observed in a study in South Katanga, where the decoction (62.5%) was found to be the primary way of preparation and the root was found to be the most used organ: 41.3%, n = 95[15]. The same is true in the study carried out simultaneously in Kwango and Kinshasa (n = 68), where the leaves at 65% proved to be the most used organ and maceration with 63% was found to be the most common mode of preparation solicited [8]. The herbal antidiabetic recipes

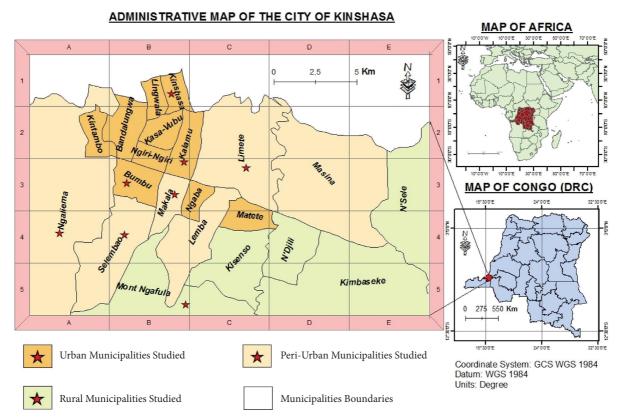


FIGURE 7: Map of Kinshasa city with different sites where data have been collected.

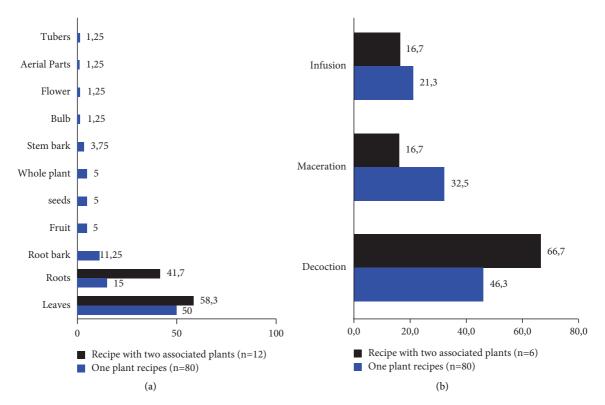


FIGURE 8: Parts used (a) and mode of preparation of antidiabetic recipes (b).

used in Kinshasa align with the national trend, unlike those used in Lubumbashi, the country's second-largest city.

According to the subjects consulted, using a decoction would aim to extract and activate the active ingredient. This is a mixed idea, especially since this extraction procedure is more beneficial than harmful. Indeed, as much as it could facilitate the release of certain active principles often present in the plant in the form of glucosides, it could not only release specific toxic secondary metabolites such as cyanogenic glycosides [122] but also deteriorate the active ingredient when the latter is thermolabile [123–125]. Therefore, this practice remains to be assessed case-by-case, and only experimental work could determine its fair value.

About the various correlations established on specific characteristics of the subjects consulted (Figure 8), this study shows that the older the diabetic issues, the longer they have lived with diabetes, which suggests that most of the subjects consulted would be diabetics from type 1. Similarly, the study reveals that the oldest TMPs, the most experienced and skilled, see more patients. This supposes that traditional medicine requires time to retain patients who would be more confident towards the most experienced. This attitude is also observed in conventional medicine, where rationality would like the patient to be reassured by the doctor's experience before being consulted, especially for specific pathologies.

5. Conclusion

Several plants are used in Kinshasa to manage diabetes mellitus by herbalists as well as by people with diabetes and traditional medicine practitioners. This study highlights not only antidiabetic recipes but also plants reported only in Kinshasa for the management of diabetes. It also evokes their usual values in this environment. It opens the way for a subsequent study to validate the antidiabetic use, particularly of particular species of the environment and those of specific special interest there.

Data Availability

The data used in this study are included within the article.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

Authors' Contributions

BCV designed, formatted, and analyzed the data, developed methods, and drafted the first manuscript. FMK and MMH performed study checks. BCV, ONP, and FMK were responsible for manuscript corrections.

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