



Review 1

Mesembrine alkaloids: Review of their occurrence, chemistry, and pharmacology 2



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ABSTRACT 8

Ethnopharmacological relevance: Mesembrine alkaloids are considered to be the primary active constituents 9 of the South African medicinal plant *Sceletium tortuosum* (L.) N.E.Br. (Aizoaceae), and it is used as the dried or fermented aerial material from the plant, which is known as kanna (aka, channa, kougoed). Traditional regional use ranged from relieving thirst, mild analgesia, and alteration of mood. Current interest has focused primarily on the antidepressant action of preparations based on the plant and commercialization is expanding the recognition and availability of these preparations.

Materials and methods: Searches for the keywords “Sceletium or mesembrine” were performed in “PubMed-NCBI”, “Chemical Abstracts SciFinder” and “Thomson Reuters Web of Science” databases in addition to the inclusion of references cited within prior reviews and scientific reports. Additionally the “SciFinder” database was searched using 3a-phenyl-cis-octahydroindole in the SciFinder Substructure Module (SSM). Plant taxonomy was validated by the database “The Plant List”. 10

Results: This review focuses on the chemistry, analysis, and pharmacology of the mesembrine alkaloids. 11 Despite a long history of medicinal use and research investigation, there has been a renewed interest in the pharmacological properties of the mesembrine alkaloids and much of the pharmacology has only recently been published. The two major active alkaloids mesembrine and mesembrenone are still in the process of being more fully characterized pharmacologically. They are serotonin reuptake inhibitors, which provides a rationale for the plant's traditional use as an antidepressant, but other actions are beginning to appear in the literature. Additionally, mesembrenone has reasonably potent PDE4 inhibitory activity. This review intends to provide an overview of the available literature, summarize the current findings, and put them in perspective with earlier studies and reviews.

1. Introduction 12

Kanna (aka, channa, kougoed) is the dried or fermented aerial material from the plant, *Sceletium tortuosum* (L.) N.E.Br. (Aizoaceae). This succulent species is a long-recognized, medicinal plant whose history and ethnopharmacology has been well reviewed (Gericke and Viljoen, 2008; Smith et al., 1996). The primary recognized active constituents are a group of mesembrine alkaloids; particularly mesembrine (1) and mesembrenone (2) (see Fig. 1). Popular interest in the cognitive and antidepressant actions of kanna (Chiu et al., 2014) and products based on kanna (Terburg et al., 2013) have been growing in recent years. Its abuse in combination with cannabis or other psychoactive agents has attracted the attention of the US Drug Enforcement Administration (Dye, 2013) and a seizure of the material was reported as an intelligence alert the DEA's Microgram Bulletin (DEA, 2008). Interestingly, the retail marketplace, amazon.com, bans kanna sales and includes it in their list of “Drugs & Drug 13

Paraphernalia” as “Products that simulate the effects of any illegal drug” (Amazon.com, 2016). Additionally, due to growing interest in the use of this plant, it is now mostly grown commercially in greenhouses and a 2013 study seeking to survey species in the genus for alkaloid content was unable to source material from the wild and relied on cultivators for their material (Patnala and Kanfer, 2013). In view of the renewed attention on this plant and the mesembrine alkaloids, the purpose of this review is to update the prior reviews of these area (Gericke and Viljoen, 2008; Smith et al., 1996) with a focus on recent analytical and pharmacological reports, and provide a comprehensive listing of the available literature with some commentary on key studies. 14

Some particularly valuable early reviews of the mesembrine alkaloids themselves occur within a couple of extensive chemistry-focused reviews of *Sceletium* alkaloids (Jeffs, 1981; Popelak and Lettenbauer, 1967). Mesembrine alkaloids are a subgroup within the *Sceletium* alkaloids. Additionally, an ongoing series of reviews covering literature reports of *Amaryllidaceae* and *Sceletium* alkaloids in one to several 15

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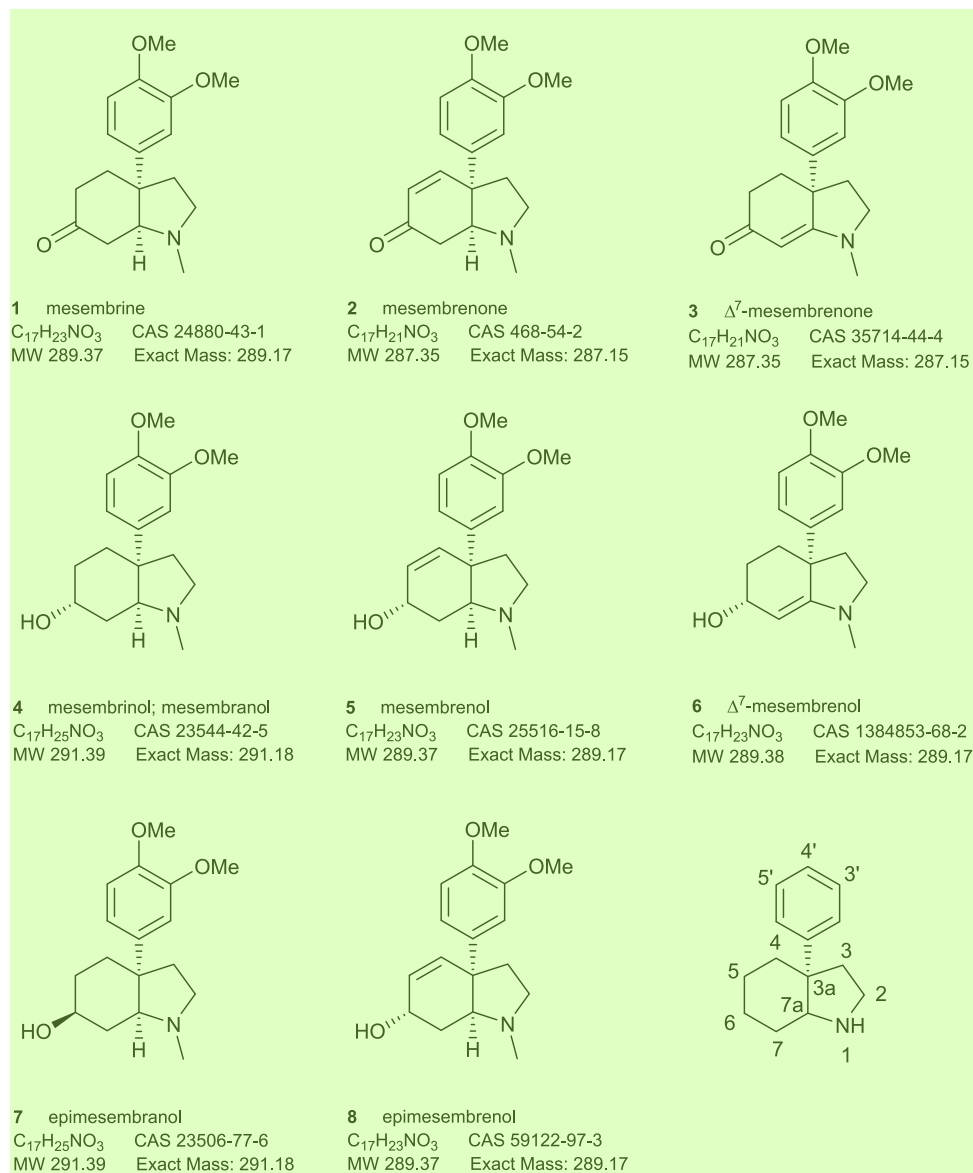


Fig. 1. Structures and identifying information for the main mesembrine alkaloids of current commercial and pharmacological interest. The numbering system is shown for the 3a-phenyl-*cis*-octahydroindole core that is characteristic of the mesembrine subgroup of *Sceletium* alkaloids. The average (MW) and monoisotopic (Exact Mass) molecular weights are given along with the molecular formula and Chemical Abstracts Service (CAS) number for each compound.

year segments has been published in Natural Product Reports (Jin, 2003, 2005, 2007, 2009, 2011, 2013; Jin et al., 2002; Lewis, 1990, 1992, 1993, 1994, 1995, 1996, 1997, 1998, 1999, 2000, 2001, 2002).

2. Occurrence of mesembrine alkaloids

Sceletium is currently the only known genus having species with relatively high levels of mesembrine. Other related plants or other genera, such as *Narcissus*, contain significantly smaller amounts of the alkaloid, if at all (Berkov et al., 2014; Pigni et al., 2013; Smith et al., 1998). Early work used the basonym *Mesembryanthemum*, but the genus was abandoned. Earlier publications list species of plant that contain mesembrine or other *Sceletium* alkaloids; however, some of these species listed may be currently classified under different names (Jefferies, 1981; Smith et al., 1996). For example, a 1967 review mentions this genus reclassification and gives four species names as being accepted at the time (Popelak and Lettenbauer, 1967). As of 2016 “The Plant List” for the *Sceletium* genus currently lists 23 species with 19 species accepted (The Plant List, 2013). Of these, *Sceletium namaquense*, a reported mesembrine containing species (Popelak and Lettenbauer, 1967), is one of the four considered unresolved and

“in review” by the World Checklist of Selected Plant Families (WCSP) (The Plant List, 2013). Many, but not all of *Sceletium* species obtained from commercial sources, contain mesembrine (Patnala and Kanfer, 2013). This fairly recent publication equates *Sceletium tortuosum* with *S. joubertii* and *S. namaquense*, which is consistent with Gerbaulet, 1996 despite “The Plant List” considering the former as an accepted separate species and the latter as unresolved (Gerbaulet, 1996; The Plant List, 2013). Therefore, this review focuses on the key alkaloids of current pharmacological interest as much as possible without regard to their source. When alkaloid isolation is discussed, some of the differences in individual alkaloid levels will be noted and the species identification is cited from the paper where available. This review focuses on the mesembrine subgroup of the *Sceletium* alkaloids, characterized by the 3a-phenyl-*cis*-octahydroindole ring system, because they are the subject of the majority of recent commercial and scientific interest. For these reasons, other related *Sceletium* alkaloids belonging to different subgroups and having different core structures than those listed in Fig. 1, such as *Sceletium* A4 or tortuosamine, have not been included in this review.

Because of the commercial importance of preparations containing mesembrine alkaloids, there have been efforts to distinguish *Sceletium*

Table 1
Current *Sceletium* species and status (The Plant List, 2013). For status, “A” = accepted, moderate confidence level (source: Missouri Botanical Garden, Tropicos); “U” = unresolved, low confidence level (WCSP (in review). Mesembrine presence as reported in the literature with reference given or reference where it was not detected. If multiple references cite its presence, then a “+” is added after the citation.

Name	Status	Mesembrine	Reference	
<i>Sceletium albanense</i> L. Bolus	A	Yes – possible	Footnote 1	
<i>Sceletium anatomicum</i> L. Bolus	A	Yes – confirmed	Rimington and Roets, 1937	+
<i>Sceletium archeri</i> L. Bolus	A	No – possible	Footnote 2	
<i>Sceletium boreale</i> L. Bolus	A	Yes – possible	Footnote 3	
<i>Sceletium compactum</i> L. Bolus	A	Yes – possible	Footnote 3	
<i>Sceletium concavum</i> Schwantes	A	Yes – possible	Footnote 3	
<i>Sceletium crassicaule</i> L. Bolus	A	Yes – confirmed	Shikanga et al., 2013	
<i>Sceletium dejagerae</i> L. Bolus	A	No – possible	Footnote 4	
<i>Sceletium emarcidum</i> L. Bolus	A	No – confirmed	Patnala and Kanfer, 2013	
<i>Sceletium exalatum</i> Gerbaulet	U	No – confirmed	Patnala and Kanfer, 2013	
<i>Sceletium expansum</i> L. Bolus	A	Yes – confirmed	Hartwich and Zwicky, 1914	+
<i>Sceletium framesii</i> L. Bolus	A	Yes – possible	Footnote 3	
<i>Sceletium gracile</i> L. Bolus	A	Yes – possible	Footnote 3	
<i>Sceletium joubertii</i> L. Bolus	A	No – confirmed	Arndt and Kruger, 1970	
<i>Sceletium namaquense</i> L.Bolus	U	Yes – confirmed	Popelak and Lettenbauer, 1967	+
<i>Sceletium ovatum</i> L. Bolus	A	Yes – possible	Footnote 3	
<i>Sceletium regium</i> L. Bolus	A	Yes – possible	Footnote 5	
<i>Sceletium rigidum</i> L. Bolus	A	No – confirmed	Patnala and Kanfer, 2013	
<i>Sceletium strictum</i> L. Bolus	A	Yes – confirmed	Jeffs et al., 1971	+
<i>Sceletium subvelutinum</i> L.Bolus	U	No – confirmed	Herbert and Kattah, 1990	
<i>Sceletium tortuosum</i> (L.) N.E. Br.	A	Yes – confirmed	Hartwich and Zwicky, 1914	+
<i>Sceletium tugwelliae</i> L. Bolus	A	Yes – possible	Footnote 3	
<i>Sceletium varians</i> L. Bolus	A	No – possible	Footnote 6	

Table 1 Footnotes: 1: \equiv *S. crassicaule* (Haw.) L.Bolus (Gerbaulet, 1996) indicating the possible presence of mesembrine; 2: \equiv *S. rigidum* L.Bolus (Gerbaulet, 1996) indicating a possible lack of mesembrine; 3: \equiv *S. tortuosum* (L.) N.E.Br. (Gerbaulet, 1996) indicating the possible presence of mesembrine; 4: \equiv *S. emarcidum* (Thunb.) L.Bolus ex H.J.Jacobsen (Gerbaulet, 1996) indicating a possible lack of mesembrine; 5: \equiv *S. expasum* (L.) L.Bolus (Gerbaulet, 1996) indicating the possible presence of mesembrine; 6: *Sceletium varians* (Haw.) Gerbaulet comb.nov. \equiv *Sceletium subvelutinum* (L.) Bolus forma *subvelutinum* (Gerbaulet, 1996) indicating a possible lack of mesembrine.

species by imaging techniques (Shikanga et al., 2013) and to quantify the alkaloid components within samples from commercial preparations of *Sceletium tortuosum* (Shikanga et al., 2012a, 2012b, 2012c).

Table 1 lists the current 23 species from “The Plant List” indicating the current status of their classification. The earliest reference for the identification of mesembrine in a particular species is given. If a species had been evaluated, but mesembrine had not been detected in significant amounts, then this is indicated as “No – confirmed”. Nevertheless, in these cases, other mesembrine alkaloids were typically present (Patnala and Kanfer, 2013). Species lacking a reference have not been evaluated in the literature reviewed so the presence of mesembrine is uncertain. Footnotes in the table point to species equivalencies based on Gerbaulet, 1996, which may indicate the likelihood of the presence of mesembrine for species not otherwise cited in the literature as having been examined for the presence of mesembrine and mesembrine alkaloids (Gerbaulet, 1996). The basis for drawing these tentative conclusions stems from the alkaloid patterns observed for the species subsets within *Sceletium* (Patnala and Kanfer, 2013). Thus, it can be seen that mesembrine is present in many, but not all species of the *Sceletium* genus. From a commercial standpoint, studies directed at the variability of mesembrine alkaloid composition with regard to species, strain, time of harvest, and growing conditions would be of great importance.

3. Chemistry

3.1. Structure and properties

The absolute stereochemistry for the mesembrine alkaloids as shown in Fig. 1 was established by x-ray crystallography of 6-epimesembranol methiodide and later that of (-)-mesembrine HCl (Capps et al., 1977; Coggon et al., 1970). It was in agreement with the assignment of stereochemistry of mesembrine based on CD and NMR studies (Jeffs et al., 1969). It should be noted that a number of papers and internet sources depict incorrect stereochemistry for some of the alkaloids, so it is important to carefully note and establish the correct stereochemistry within any source. The structures depicted in this review reflect the stereochemistry as established by Jeffs’ studies noted above.

3.1.1. Mesembrine (1): (3a*S*,7a*S*)-3a-(3,4-dimethoxyphenyl)-1-methylhexahydro-1*H*-indol-6(2*H*)-one

This alkaloid is the most studied of the mesembrine alkaloids due to its relative abundance in *Sceletium tortuosum* and biological activity. Mesembrine was initially partially isolated, characterized, and named by Zwicky (Hartwich and Zwicky, 1914; Zwicky, 1915). Following on that report, mesembrine was purified and crystallized as the picrate (mp 193–194 °C) and the platinichloride (mp 181 °C) (Rimington and Roets, 1937). They also reported some color tests performed on the non-crystalline hydrochloride salt.

In 1957, mesembrine was isolated and crystallized as the hydrochloride salt from isopropanol-ethanol (mp 199–201 °C) and reported the IR spectrum (Bodendorf and Krieger, 1957). They correctly identified that the material contained two methoxy groups and an N-methyl group, correcting earlier reports claiming mesembrine to be a phenolic base, which it is not.

In 1960, Popelak, et al. correctly reported the structure of mesembrine, physical data, and characterized a variety of derivatives through chemical modification of the isolated alkaloid (Popelak et al., 1960a, 1960b). It should be noted that mesembrine HCl can be recrystallized from 2-propanol; however, attempts at crystallizing this salt from ethanol produces an ethyl enol ether from the keto group that can be converted back to mesembrine HCl by treatment with 0.1 N aq. HCl. Enol formation does not occur with the free base in ethanol (Popelak and Lettenbauer, 1967).

Lit. (Popelak and Lettenbauer, 1967)	
mesembrine (free base)	$[\alpha]_D^{20} = -55^\circ$ (MeOH) bp _{0.05} 138 °C
mesembrine HCl	$[\alpha]_D^{20} = -8.4^\circ$ (MeOH) mp 205–

206 °C **1**

Lit. (Taber and He, 2005)

mesembrine, colorless oil $[\alpha]_D^{20} = -55.4^\circ$ (c 0.54, MeOH);IR (neat, cm^{-1}): 2944 (m), 1716 (s), 1520 (s), 1454 (m), 1253 (s), 1148 (m), 1027 (m); ^1H NMR (400 MHz, CDCl_3) δ : 6.93 (dd, $J = 8.4, 2.4$ Hz, 1 H), 6.90 (d, $J = 2.0$ Hz, 1 H), 6.84 (d, $J = 8.4$ Hz, 1 H), 3.90 (s, 3 H), 3.88 (s, 3 H), 3.13–3.17 (m, 1 H), 2.96 (t, $J = 3.6$ Hz, 1 H), 2.61 (m, 2 H), 2.35–2.46 (m, 2 H), 2.33 (s, 3 H), 2.04–2.26 (m, 5 H); ^{13}C NMR (CDCl_3 , 90 MHz) δ u: 211.5, 149.3, 147.8, 140.4, 55.1, 47.8, 40.8, 39.1, 36.4, 35.5; d: 118.2, 111.3, 110.3, 70.6, 56.2, 56.1, 40.3

Racemic mesembrine, which is the product of several total synthesis routes (Chavan et al., 2004; Gelas-Mialhe et al., 1992; Gramain and Remuson, 1985; Hackett and Livinghouse, 1986; Honda et al., 2012; Hoshino et al., 1978, 1987; Jeffs et al., 1983; Kochhar and Pinnick, 1983; Kulkarni et al., 2002; Martin et al., 1979; Michael et al., 1992; Ohishi and Kugita, 1968, 1970; Parkinson and Pinhey, 1991; Rajagopalan, 1997; Rigby and Dong, 2000; Roe et al., 2008; Shamma and Rodriguez, 1965, 1968; Shono et al., 1989; Spittler et al., 2016; Stevens et al., 1975; Takano et al., 1981a; Zhao et al., 2009), can be resolved to the natural *l*-isomer through fractional crystallization using α -bromo-D-camphor- π -sulfonic acid (Popelak and Lettenbauer, 1962).

3.1.2. Mesembrenone (**2**): (3*aR*,7*aS*)-3*a*-(3,4-dimethoxyphenyl)-1-methyl-3,3*a*,7,7*a*-tetrahydro-1*H*-indol-6(2*H*)-one

The material readily racemizes (e.g., ethanolic solution for 17 h at 25 °C or 5 min at 50 °C) via acid or base catalyzed reversible β -elimination to **9** (Fig. 2) (Jeffs et al., 1970).

Lit. (Jeffs et al., 1970)

Mesembrenone, CD $[\theta]_{395} 0^\circ$, $[\theta]_{334} -4060^\circ$, $[\theta]_{310} 0^\circ$ (c an oil 1.072 mg/mL, ethanol)Mesembrenone mp 204–207 °C
HCl

3.1.3. mesembrinol; mesembranol (**4**): (3*aS*,6*R*,7*aS*)-3*a*-(3,4-dimethoxyphenyl)-1-methyloctahydro-1*H*-indol-6-ol

Much of the historical literature uses the mesembranol name. This alkaloid differs from epimesembranol (**7**) only in the stereochemistry at the 6 position alcohol. It can be separated from its epi-counterpart by crystallization, since the epi-form does not crystallize, or by chromatography (Popelak and Lettenbauer, 1967).

Lit. (Popelak and Lettenbauer, 1967)

Mesembrinol $[\alpha] = -32^\circ$ (CHCl_3); mp 144–145 °C

Lit. (Chida et al., 1994)

Mesembrinol $[\alpha] = -28^\circ$ (c 0.2, CHCl_3); mp 146–147 °C

3.1.4. mesembrenol (**5**): (3*aR*,6*S*,7*aS*)-3*a*-(3,4-dimethoxyphenyl)-1-methyl-2,3,3*a*,6,7,7*a*-hexahydro-1*H*-indol-6-ol

Jeffs, et al. first isolated, crystallized, and identified this alkaloid in 1970 as the major alkaloid component of *Sceletium strictum* (L.) Bolus (Aizoaceae) (Jeffs et al., 1970). The material used in this study had been grown from seed in a greenhouse for three years and was identified by L. Bolus, Bolus Herbarium, University of Cape Town, South Africa.

Lit. (Jeffs et al., 1970)

Mesembrenol $[\alpha] = +91^\circ$ (c 0.0176, CHCl_3) mp 140 °C (recrys. from EtOAc)
mp 122–124.5 °C (recrys. from acetone) ^1H NMR (100 MHz) δ : 6.84 (m, 3 H, ArH), 5.70 (q of t, $J_{4,5} = 10.5$ Hz, $J_{4,6} = 1.8$ Hz, $J_{5,6} = J_{5,7a} = J_{4,7a} = 1.5$ Hz, 2 H, $-\text{CH}=\text{CH}-$), 4.30 (m, 1 H, C6H), 3.76 (s, 3 H, $-\text{OCH}_3$), 3.80 (s, 3 H, $-\text{OCH}_3$), 2.30 (s, 3 H, NCH_3).

3.2. Biosynthesis **13**

The biosynthetic pathway depicted in Fig. 3, suggested by the work involving radiolabeled precursor materials, proposes that the aromatic group of mesembrine comes from phenylalanine and the perhydroindole portion comes from tyrosine (Jeffs et al., 1971). From phenylalanine, cinnamic acid intermediates are produced, but the 3'-aryl oxygen substituent is introduced late in the pathway as evidenced by the late steps involving sceletenone (CAS 51934-31-7), 4'-O-demethyl-mesembrenone, and mesembrenone (Jeffs et al., 1978).

3.3. Isolation **15**

Many alkaloid isolations using varied techniques and different species of plant have been reported (Arndt and Kruger, 1970; Bastida et al., 1989; Doepke et al., 1981; Jeffs et al., 1970, 1982; Jeffs and Capps, 1979; Nieuwenhuis et al., 1981; Rimington and Roets, 1937;

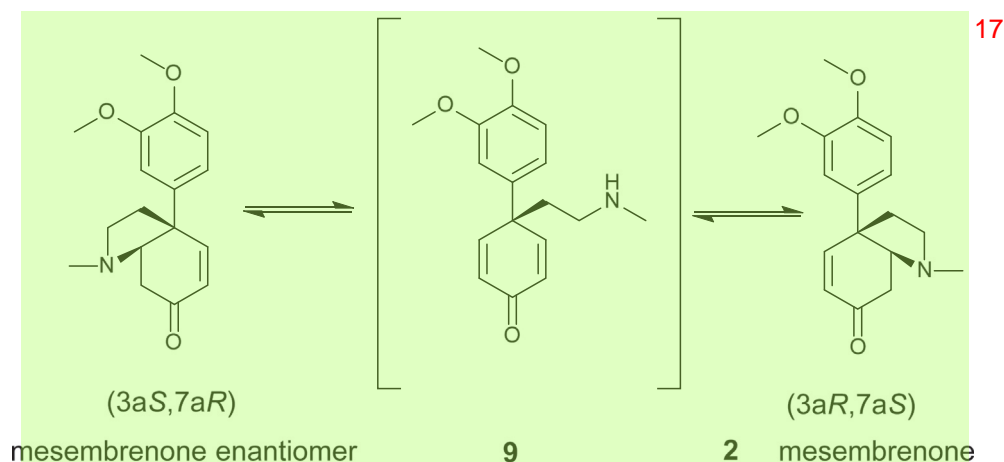


Fig. 2. Racemization of mesembrenone via reversible β -elimination (Jeffs et al., 1970).

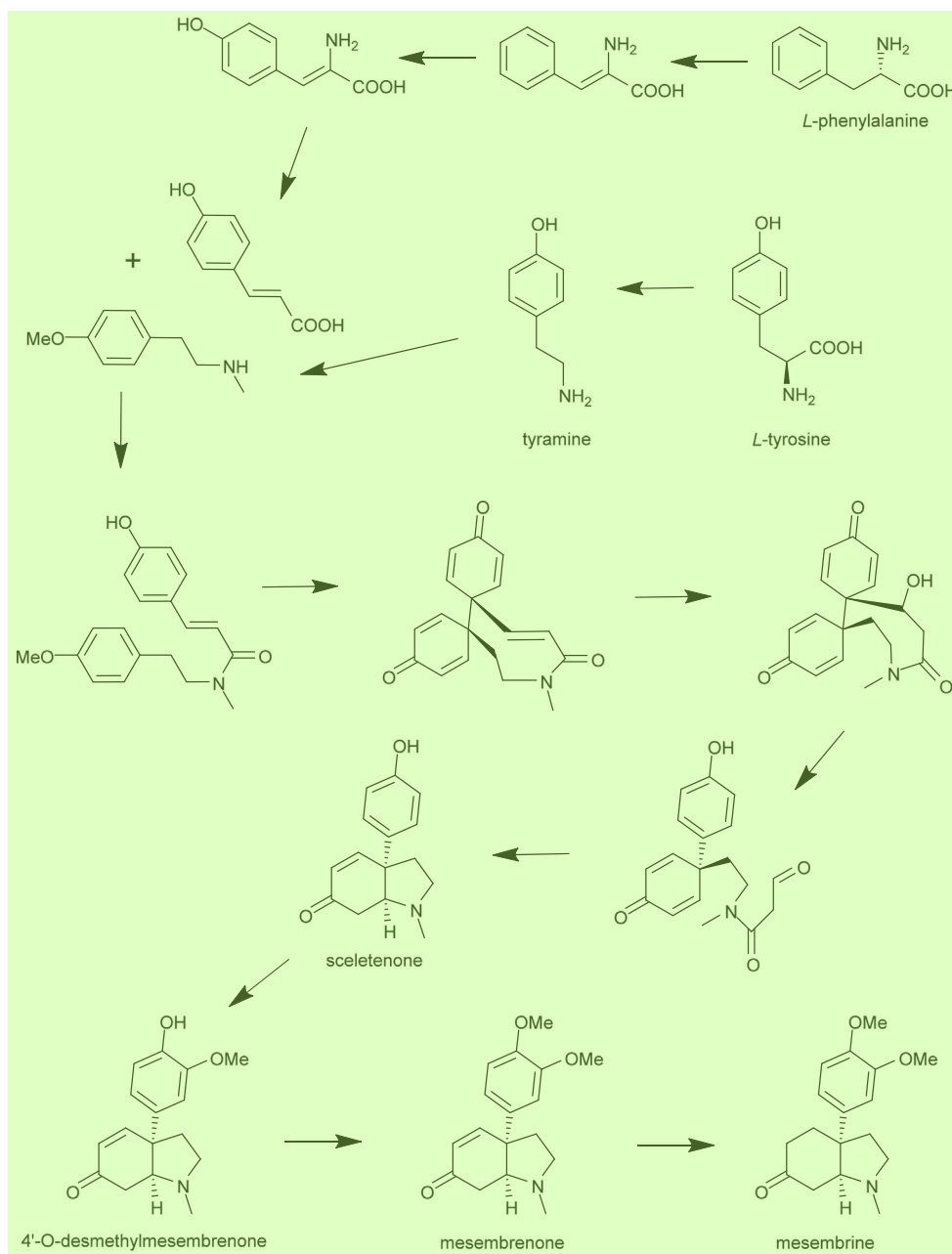


Fig. 3. Proposed biosynthetic pathway (Jeffs, 1981). Some arrows represent more than one step in the pathway.

Shikanga et al., 2011). A couple of examples are presented below with the isolated yield from each. Note that the isolated yield is not only dependent on the efficiency of the method, but also on the abundance of the alkaloids in the starting plant material used.

3.3.1. Alcohol extraction with neutral alumina column

Fresh 3 year old plants of *Sceletium strictum* were homogenized with 95% ethanol, heated and filtered (Jeffs et al., 1971). The material used in this study had been grown from seed in a greenhouse for three years and was identified by L. Bolus, Bolus Herbarium, University of Cape Town, South Africa. The solid was further Soxhlet extracted with methanol. The combined extracts were concentrated, treated with sodium carbonate and extracted with chloroform. Evaporation of the chloroform left the crude alkaloid (2.5% w/w). The crude alkaloid fraction was eluted on neutral alumina column using a linear gradient of benzene against ethyl acetate followed by ethyl acetate/methanol and finally methanol. Fractions were analysed by GC-FID. The follow-

ing materials were obtained from the fractions in the following amounts: mesembrine (1) 0.067%; mesembrenone (2) 0.025%; mesembranol (4) 0.058%; mesembrenol (5) 0.71%; (See Fig. 1 for structures). Note that the major alkaloid from this plant species was mesembrenol.

3.3.2. High performance countercurrent chromatography (HPCCC)

An isolation process that started with an acid extraction of the dried *Sceletium tortuosum* material giving 1.9% by weight of crude alkaloid extract has been reported (Shikanga et al., 2011). This plant material used in the study was collected in April 2009 from the Western Cape South Africa with a voucher specimen kept at the Dept. of Pharmaceutical Sciences, Tshwane University of Technology. This was fractionated on a silica gel column to give one fraction containing alkaloids 1 and 2 and another fraction containing 4 and 5. Using a high-performance countercurrent chromatography instrument on the two fractions, they obtained each of the four alkaloids in > 95% purity.

The overall isolated amounts from dried plant material were **1** 0.21%; **2** 0.24%; **4** 0.025%; **5** 0.16%.

3.4. Analysis **2**

Both preclinical, clinical, and forensic study of mesembrine alkaloid-containing products requires sensitive validated assays for the qualitative and quantitative analysis of those alkaloids or the metabolites. The recent literature provides a variety of such methods, which greatly enable a broad range future studies.

3.4.1. High performance thin layer chromatography (HPTLC)-densitometry **4**

HPTLC is a robust method for the analysis and characterization of botanical products (Toniole et al., 2014). Using Silica gel 60 F254 HPTLC plates eluted with dichloromethane–methanol–10% ammonia (90:10:0.1, v/v) (Shikanga et al., 2012b). Densitometric quantification was based on absorbance at 280 nm. Limit of Detection (LOD) was in the range of 18–31 ng per band and limit of quantitation (LOQ) was 44–95 ng per band for the four main mesembrine alkaloids. The validated method was reported useful for raw plant material and commercial products containing *Sceletium tortuosum* alkaloids.

3.4.2. Capillary electrophoresis methods (CE) **6**

A nonaqueous capillary electrophoresis-mass spectrometry (NACE-MS) method for forensic analysis of *Sceletium*-containing products and to analyze products the fermentation process of *Sceletium tortuosum* has been reported (Roscher et al., 2012). The species-identified plant materials used in this study were obtained from two commercial sources. The method, originally developed for the analysis of kratom alkaloids, has a mean dynamic range is 0.1–500 μ M, based on *Harmala* alkaloid standards.

A capillary zone electrophoresis method for the analysis of mesembrine in *Sceletium tortuosum* products has been reported with an LOQ and LOD of 2.5 and 1.5 μ g/mL, respectively (Patnala and Kanfer, 2008).

3.4.3. Gas chromatography-mass spectrometry (GC-MS) **9**

GC-MS combined with other methods have been used to provide a detailed metabolic analysis of the mesembrine alkaloids from rat urine (Meyer et al., 2015), and GC-MS alone has been used to develop a method that has a LOD of 12–23 μ g/mL and LOQ of 35–68 μ g/mL for the four main mesembrine alkaloids.

3.4.4. High performance liquid chromatography (HPLC) **11**

Along with the GC-MS method, a more sensitive reverse phase-ultra high performance liquid chromatography-photodiode array (RP-UHPLC-PDA) method was developed with a LOD of 0.9–1.2 μ g/mL and LOQ of 2.7–3.7 μ g/mL (Shikanga et al., 2012a). A sensitive HPLC method has been published that is linear in the concentration range of 0.4–60 mg/mL with a LOD of 100 ng/mL and a LOQ of 200 ng/mL (Patnala and Kanfer, 2010). Finally, liquid chromatography-linear ion trap high resolution mass spectrometry (LC-HR-MSn) was used as a component of metabolism studies discussed in more detail below (Meyer et al., 2015).

3.4.5. Direct analysis in real time ionization coupled with high resolution time-of-flight mass spectrometry (DART-HRTOFMS) **13**

Forensic analysis of kanna from a variety of suppliers using DART-HRTOFMS was performed in order to assess the presence of the plant materials as well as adulterants (Lesiak et al., 2016). Using five different samples, the major alkaloids were identified, providing evidence for the presence of the advertised plant material, and in one case, adulteration with a banned chemical was identified, establishing this method as useful in identifying misrepresented and potentially unsafe products.

3.5. Total synthesis **15**

As early as 1962, a synthetic route for mesembrine and mesembrine analogs was patented (Popelak and Lettenbauer, 1962). Since then, numerous total syntheses of both the racemic (Chavan et al., 2004; Gelas-Mialhe et al., 1992; Gramain and Remuson, 1985; Hackett and Livinghouse, 1986; Honda et al., 2012; Hoshino et al., 1978, 1987; Jeffs et al., 1983; Kochhar and Pinnick, 1983; Kulkarni et al., 2002; Martin et al., 1979; Michael et al., 1992; Ohishi and Kugita, 1968, 1970; Parkinson and Pinhey, 1991; Rajagopalan, 1997; Rigby and Dong, 2000; Roe et al., 2008; Shamma and Rodriguez, 1965, 1968; Shono et al., 1989; Spittler et al., 2016; Stevens et al., 1975; Takano et al., 1981a; Zhao et al., 2009) and both individual stereoisomers (Arns et al., 2007; Dalko et al., 1998; Denmark and Marcin, 1997; Geoghegan and Evans, 2013; Ilardi et al., 2009; Kosugi et al., 1993; Meyers et al., 1985; Mori et al., 1997a, 1997b; Nemoto et al., 1994, 1995; Nunokawa et al., 2015; Otani and Yamada, 1973; Ozaki and Kobayashi, 2015; Paul et al., 2006; Taber and Neubert, 2001; Taber and He, 2005; Takano et al., 1981b, 1990; Yamada and Otani, 1971; Yokomatsu et al., 1992; Yoshimitsu and Ogasawara, 1996; Zhang et al., 2012) have been reported and much of this work has been presented in a Japanese review covering 31 published routes (Zhao et al., 2010). While the paper is in Japanese, which presents a barrier for those who cannot translate the article, the many synthetic schemes are in English and having them together in one reference is valuable. Despite these extensive synthetic efforts and the availability of commercially grown *Sceletium tortuosum* there is currently no commercially available source for the mesembrine alkaloids in pure form. Commercial availability of pure materials at a reasonable price would be of considerable value enabling pharmacologists worldwide to expand our understanding of these materials.

The total synthesis of analogs based on a mesembrine core structure reported modest analgesic activity in a mouse hot plate test for one of the analogs, 3-((3aS,7aS)-1-methyloctahydro-3aH-indol-3-yl)phenol (Langlois et al., 1971, 1972).

Biovitrum submitted a patent application covering substituted mesembrine analogs as melanin concentrating hormone receptor (MCH1R) antagonists for the modulation of appetite (Browning et al., 2005). They present a total synthesis of a large number of analogs. Example 70 in the application is a 6-substituted mesembrine analog with a p-trifluoromethoxyphenylurea group having a K_i of 52 nM at the MCH1R.

4. Biology **19**

4.1. Metabolism **20**

The metabolism of isolated and purified mesembrine and mesembrenone *in vivo* in the rat, and *in vitro* using a human liver preparation has been reported (Meyer et al., 2015). O- and N-demethylations were observed for both alkaloids. Reduction (dihydro metabolite) was observed and hydroxylation was observed at different positions. The resulting phenolic metabolites were partly excreted as glucuronides or sulfates. For mesembrine the most abundant metabolites were the acetylated demethyl-dihydro and hydroxy metabolites in rat urine. For mesembrenone the demethyl-dihydro were the most abundant metabolites in the rat urine.

4.2. Pharmacology **22**

Despite the long history of the medicinal actions of kanna and the depth of the literature on isolation and total synthesis of mesembrine and related alkaloids, there has been surprisingly little published on the pharmacology of the individual alkaloids, which is likely partly due to the lack of commercially available pure alkaloids. Nonetheless, as more is learned about the pharmacology of these materials the number of

patents and applications for the use of mesembrine alkaloid preparations has been gaining pace (Davies, 2016; Gericke et al., 2015, 2016; Gericke and Van Wyk, 2001; Harvey et al., 2010; Holford, 2002). A report of three clinical cases using *Sceletium tortuosum* tablets has previously been reviewed pointing to potential therapeutic utility of such a preparation (Gericke and Viljoen, 2008). Controlled human trials mentioned below are beginning to appear and are the first step in providing evidence for wide acceptance and use of these materials.

4.2.1. Receptor screening 2

In 2011, an attempt at a comprehensive pharmacological screen, began with a commercial material, Zembrin®, which is a standardized extract of *Sceletium tortuosum* (Harvey et al., 2011). The preparation consists of 0.35–0.45% total alkaloids, with mesembrenone and mesembrenol comprising ≥60%, and mesembrine contributing < 20% (Murbach et al., 2014). This mixture was screened in 77 radioligand binding assays at 0.75 mg/mL and a panel of phosphodiesterases. The extract gave > 80% inhibition at GABA receptors (non-selective), serotonin (5-HT) transporter, δ_2 - and μ -opioid receptors, the cholecystikinin-1 receptor, and PDEs 3 and 4 (Harvey et al., 2011). Following up on these results, the potency of the isolated alkaloids at the 5-HT transporter and with PDE4 were reported, as described separately in the following Sections 4.2.2 and 4.2.3.

4.2.2. Serotonin reuptake inhibition 4

Individual alkaloids isolated from *Sceletium tortuosum* extract were tested against the 5-HT transporter (SERT) and the K_s for mesembrine, mesembrenone, and mesembrenol were 1.4, 27, and 63 nM, respectively (Harvey et al., 2011). These values are comparable to the marketed SSRI, fluoxetine (Tavoulari et al., 2009) and significantly more potent than values seen for related *Amaryllidaceae* alkaloids, such as buphanidrine or distichamine with SERT K_s of 312 and 868 μ M, respectively, which derive from another traditional anxiety treating plant *Boophone disticha* (L.f.) Herb (Amaryllidaceae) (Sandager et al., 2005).

4.2.3. PDE4 inhibition 6

The isolated alkaloids mesembrine, mesembrenone, and mesembrenol inhibited PDE4B with IC_{50} s of 7.8, 0.47, and 16 μ M, respectively (Harvey et al., 2011). The prototypical PDE4 inhibitor rolipram is at least 3-times more potent with an IC_{50} for PDE4B of 0.13 μ M (MacKenzie and Houslay, 2000). Related to this mechanism of action, one *in vivo* study correlated the responses seen by Zembrin® extract and rolipram as interpreted as an electropharmacogram, which is a frequency analysis of the field potentials recorded from the frontal cortex, hippocampus, striatum and reticular formation in the presence of drugs in freely moving rats (Dimpfel et al., 2016b).

Since the identification of PDE inhibition for these alkaloids, a number of patents, where a PDE inhibitor would have utility alone or in combination with other substances, have included the mesembrine alkaloids among a list of other PDE inhibitors (Collins et al., 2014; Hammock et al., 2011; Kielian, 2014; Lefebvre and DeMaeyer, 2013).

4.2.4. Increased VAMT-2 expression 9

The effects of a high mesembrine-content standardized extract, Trimesemine™, (3% mesembrine (w/w)) on human astrocytes and mouse hippocampal cells in comparison to the effect elicited by the racemic mixture SSRI, citalopram (Coetzee et al., 2016). The mesembrine extract caused less down-regulation of expression of SERT than citalopram and a greater upregulation of expression of the vesicular monoaminetransporter-2 (VAMT-2) than citalopram. These results were interpreted by the authors as meaning that the primary mechanism of action of this extract is through monoamine release and that serotonin reuptake inhibition is secondary to this release effect.

4.2.5. CB1 receptor binding 11

Mesembrine has been shown to bind to cloned human cannabinoid CB1 receptors *in vitro* with about 1% the potency of the Δ^9 -tetrahydrocannabinol (THC) standard (Lubbe et al., 2010). Due to the low potency, a binding constant was not determined. This result was in agreement with a later receptor screening study (Harvey et al., 2011).

4.2.6. Inhibition of forskolin-induced steroidogenesis 13

An *in vitro* study examining the ability of high mesembrine-content standardized extract, Trimesemine™, to affect steroidogenesis using the human adrenocortical carcinoma cell line H295R was reported (Swart and Smith, 2016). The Trimesemine™ preparation is a lyophilized extract prepared from a proprietary hybrid (DV17) of *S. tortuosum* (L.) N.E. Br. and *S. expansum* (L.) L. Bolus (Aizoaceae) using a proprietary method, consists primarily of mesembrine (1), Δ^7 -mesembrenone (3), and mesembrenone (2), as demonstrated by UPLC-MS (see Fig. 1 for structures). The results demonstrate that Trimesemine™ affects steroid hormone production, possibly through modulation of the catalytic activity of steroidogenic enzymes.

4.2.7. Mouse forced swim test 15

In 2014 Schell reported a study comparing a partially purified alkaloid extract containing the main mesembrine alkaloids to paroxetine in a mouse forced swim test, a rodent model for depression (Schell, 2014). The relative composition of the alkaloid mixture used was mesembrine 9.2%, mesembrenone 8.0%, mesembrenol 67.0%, and mesembrenol 15.8% as determined by GC/MS. At 10 and 80 mg/kg i.p. dose of the alkaloid mixture was compared with a 1 mg/kg i.p. dose of paroxetine, which was calculated to give a comparable level of serotonin reuptake inhibition based on the literature IC_{50} s for mesembrine vs paroxetine. Both the alkaloid mixture and paroxetine had comparable activity in this depression model, while neither demonstrated any locomotor effects at the same dose levels.

4.2.8. Human trials 17

Since 2009, the aforementioned preparation Zembrin® has been studied for safety and cognitive effects in a number of human trials.

The first report was a double-blind, placebo-controlled study performed in 2009 that examined the safety and tolerability at doses of 8 mg and 25 mg once daily over a three month period (Nell et al., 2013). A dozen healthy adults were used in each arm of the study and compared to thirteen subjects receiving placebo (18–55 years of age). Both doses were well tolerated and adverse effects were the same as or less frequent than was seen for the placebo group.

Another double-blind, placebo-controlled cross-over design study in sixteen healthy undergraduate students (18–21 years) used functional magnetic resonance imaging (fMRI) 2 h after a single 25 mg dose of Zembrin® or placebo focusing on amygdala activity in response to an anxiety-related activity and amygdala-hypothalamus coupling in an emotion matching task (Terburg et al., 2013). There results were interpreted as demonstrating anxiolytic potential consistent with the previously proposed serotonin reuptake inhibition (SRI) and PDE4 inhibitory activity (Harvey et al., 2011); however, there were no positive controls to assess the relative effects versus established control SRIs or PDE4 inhibitors.

A randomized double-blind, placebo-controlled, cross-over design clinical trial using 25 mg capsules of Zembrin® in 21 cognitively healthy subjects was reported (Chiu et al., 2014). The dosing was once daily for 3 weeks. Zembrin® was well tolerated and demonstrated improved executive function along with positive changes in mood and sleep.

Recently, a larger double-blind, randomized, placebo-controlled study in 60 healthy adults (40–75 years) was conducted using 25 or 50 mg doses of GMP-manufactured Zembrin® versus placebo using quantitative 17-channel EEG and eye-tracking, two hours after dosing,

while being challenged cognitively and emotionally (Dimpfel et al., 2016a). Psychometric performance improvement suggest a small improvement versus placebo, but the results were not statistically significant. Most significant was the documentation of consistent, dose-related changes in brain activity that could serve as a baseline for future studies.

In summary, these studies appear to give good evidence to safety and tolerability in healthy subjects at the doses tested and over the period of time of the studies; however, the modest size of the studies, lack of positive controls limit the conclusions that can be drawn about the therapeutic utility of the agents.

4.3. Toxicology 3

A toxicological study of Zembrin® in male and female rats was reported (Murbach et al., 2014). No mortality or treatment-related adverse effects in the 14- or 90-day studies at doses ranging from 100 to 5000 mg/kg bw/day.

5. Conclusions 5

While the pharmacology and mechanism(s) of action for the behavioral effects of the individual mesembrine alkaloids and preparations containing mixtures of these alkaloids is not entirely clear, it is evident from *in vitro*, *in vivo*, and clinical studies that the main mesembrine alkaloids are bioavailable CNS-active materials worthy of further study. One current limitation is the lack of commercially available pure substance and reference materials. Despite the broad screening of an mesembrine alkaloid mixture followed by testing of the individual alkaloids against the positive hits from the mixture (Harvey et al., 2011), it would still be of value to broadly screen (affinity and functionality) the individual alkaloids to better understand their functional activity and selectivity in addition to their potency. Seldom are CNS drugs completely selective for one target. From the standpoint of developing these materials or their analogs into therapeutic agents, considerably more study is needed concerning the ultimate safety of these alkaloids and alkaloid mixtures. Additionally a competitive threat exists from non-mesembrine compounds that select for the same or multiple CNS targets, such as the dual serotonin reuptake inhibitors having PDE4 inhibitory activity that have been reported (Cashman and Ghirmai, 2009; Cashman et al., 2009a, 2009b).

However, from an herbal product perspective, where the safe and effective threshold of a pharmaceutical product does not need to be met, the trials using standardized commercial preparations are quite helpful in addressing the issues of safety and tolerability needed for an herbal product. As with all natural product preparations, QA/QC for the delivery of a consistent product with predictable biological effects is a considerable challenge. Another issue for the non-prescription use of these particular preparations is the significant potency as a serotonin reuptake inhibitor. Inappropriate use of the product or use in combination with other antidepressants or herbal products might lead to serious interactions or serotonin syndrome. If these products ever see more widespread marketing, then it would be a priority to address these real possibilities.

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