

## HPLC Analysis of Mesembrine-Type Alkaloids in *Sceletium* Plant Material Used as An African Traditional Medicine

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**Abstract – Purpose.** *Sceletium* plant species have been reported to contain psychoactive alkaloids, specifically belonging to mesembrine-type alkaloids. *Sceletium* is presently marketed through health shops and on the internet as dried plant powder and as pharmaceutical dosage forms and purported to be useful in the treatment of psychological disorders. However, there are no validated analytical methods and reference standards of the relevant alkaloids are not commercially available for use in the analysis and quality control of *Sceletium* products and dosage forms. Hence, the objective of this research was to isolate and characterize appropriate analytical markers for use in the assay and as well as markers for fingerprinting by high performance liquid chromatography (HPLC). **Methods.** The separation of the relevant alkaloids was carried out on a C18 column and detected at a UV wavelength of 228nm. The method was validated and used to assay the mesembrine-type alkaloids namely  $\Delta^7$ mesembrenone, mesembranol, mesembrenone, mesembrine and epimesembranol. **Results.** The calibration curves were found to be linear over the entire concentration range of 400-60,000 ng/ml with correlation coefficients  $>0.99$ . The accuracies of the relevant alkaloids were found to be between 94.8 and 103.6% with an inter-day relative standard deviation (RSD) of less than 2.8%. The precision studies showed inter-day RSD's of less than 3%. The recoveries were all within the range of 95 and 105% (RSD $<4.5\%$ ) and the limits of quantitation (LOQ) and detection (LOD) were found to be 100 and 200 ng/ml respectively using the respective S/N ratios of 3 and 10. **Conclusions.** An HPLC method for the quantitative analysis of  $\Delta^7$ mesembrenone, mesembranol, mesembrenone, mesembrine and epimesembranol in *Sceletium* plant material has been developed and validated. This assay method can be applied for the quality control of *Sceletium* plant material which is used as an African Traditional Medicine for the treatment of psychological disorders.

### INTRODUCTION

Evaluation of natural products for their chemical components is challenging due to the inherent diversity of their chemical composition. Separation techniques and their application to evaluate specific chemical components of natural products is an important aspect which permits accurate characterization and quantification. It has been reported that almost 80% of known natural substances are non-volatile and thermolabile, which makes high performance liquid chromatography (HPLC) with ultra violet (UV) detection the most commonly used analytical technique for natural products (1). Photo diode array (PDA) detectors enhance the versatility of the analysis by allowing multi-wavelength detection of compounds based on their distinct chromophore active regions. PDA is thus particularly useful for multi-component sample analysis since one sample can usually provide sufficient data to assess the purity of the individual components (2). There are many

reports on the use of HPLC for qualitative analysis of herbal medicines, more importantly for the fingerprinting of South African traditional medicines (3). In addition, chromatographic fingerprinting has been widely accepted and recommended by various regulatory authorities such as WHO (4), US-FDA (5), EMEA (6) and MHRA (7) to assess the consistency of batch to batch dosage forms containing phytochemical components of the harvested plants.

In the current international regulatory scenario, qualitative and quantitative analytical methods are considered mandatory. Even though *Sceletium* plants have been relatively well-researched for their chemical constituents, validated methods for quantitative analysis of the various alkaloid components found in *Sceletium* species have not yet been reported in the published literature.

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Smith *et al.* (8) reported a qualitative method using gas chromatography-mass spectrometry (GC-MS) attached to a nitrogen-phosphorus detector (NPD). The GC-MS analysis was carried out on a DB-5 capillary column of 0.25 mm internal diameter (i.d), which was temperature programmed from 230°C to 260°C at 1°C per minute. The detector was maintained at 350°C. The method described the separation of 4'-O-demethylmesembrenol, mesembrine and mesembrenone.

Gericke *et al.* (9), in their patent application, reported three methods using GC and one using a HPLC method. In all three methods, the samples were prepared by dissolving the extracts in a minimum quantity of methanol. The first method is mentioned as a "fast system" for "large number of samples". The method uses a DB-1 fused silica capillary column 30 m x 0.25 mm i.d, with helium as carrier gas. The column temperature was maintained isothermally at 200°C for 15 minutes and then programmed at 100°C/min to 300°C. The injector and the flame ionization detector were maintained at 230°C and 300°C respectively (9). A second method is mentioned as "high resolution analyses" for "selected samples". The method also uses a DB-1 column and helium as carrier gas, similar to the first method. The column temperature was isothermally maintained at 150°C for 15 minutes and then programmed at 60°C/min to 320°C. The injector and the NPD were maintained at 230°C and 300°C respectively (9). A third method mentioned is for GC-MS analysis, with the column, carrier gas and the temperatures maintained similar to the second method. The MS conditions were not described (9). The patent also includes an HPLC method using a reversed phase system (C<sub>18</sub> column). The mobile phase comprised 30% trimethylamine solution (1% in water) and 70% acetonitrile solution (60% in water) at a flow rate of 1 ml per minute. The PDA detector was set for two channels, one at 280±30 nm (Channel A) and the other at 292±10 nm (Channel B). The method suggests that "for concentrations below 0.05 mg per ml, channel A is more accurate" (9).

The South African National Biodiversity Institute (SANBI), South African Medical Research Council (MRC) and the University of Western Cape (UWC) published a collaborative paper describing another HPLC method on the internet as a part of a monograph project for herbal plants. It was suggested that the method could be applied to perform fingerprinting analysis of medicinal plant extracts. The HPLC

method which was obtained by personal communication from the authors, uses methanol and 1% acetic acid in water mixed over a 25 minute gradient program, passing through a C<sub>18</sub> column and dual channel detection at 280 and 325nm. A typical HPLC chromatogram of a methanol extract and the retention times of the major compounds purported to elute at 2.38, 3.15, 4.89 and 7.84 was displayed on the chromatogram (10).

No validation data were reported in any of the analytical methods described for *Sceletium*. Furthermore, the use of UV wavelengths in the range of 280 nm and 325 nm for detection are not optimum for most of the *Sceletium* alkaloids where, except for  $\Delta^7$ mesembrenone ( $\lambda_{\text{max}}$  =290nm), their  $\lambda_{\text{max}}$  is at 228nm.

Hence the main objective was to develop a simple, rapid, accurate, precise and reproducible HPLC method that can be applied for the qualitative and quantitative analysis for the quality control of *Sceletium* plant material and associated products containing such alkaloids.

## METHODS

### Reagents and Materials

HPLC grade methanol, acetonitrile (ACN) (Romil Ltd. Cambridge, Great Britain) and lab grade ammonium hydroxide 25% solution (Associated Chemical Enterprises (Pty) Ltd. Southdale, South Africa) was used. Water was purified in a Milli-Q® system and Millex HV® hydrophilic PVDF 0.45 µm membrane filters (Millipore, Bedford, USA) were used for filtration. *Sceletium* reference compounds mesembrine, mesembrenone,  $\Delta^7$ mesembrenone were isolated from plant material and the compounds mesembranol and epimesembranol were prepared by hydrogenation of mesembrine. Mesembrine hydrochloride was prepared from mesembrine base. All compounds were qualified by <sup>1</sup>H, <sup>13</sup>C and 2D NMR and mass spectroscopy. *Sceletium* plant samples were obtained as gift samples from cultivators.

### Instrumentation

Method validation was performed on a HPLC SpectraSYSTEM P2000 pump connected to an AS1000 auto sampler and UV1000 variable-wavelength UV detector (Thermo Separation Products, Riviera Beach, FL, USA) was used to monitor the compounds at 228nm. The output was interfaced to LCQ software (Finnigan). NMR analysis was performed on a Bruker Advance DRX 400 MHz NMR Spectrophotometer

(Rheinstetten, Germany). The method development was carried out on an Alliance 2690 HPLC connected to a PDA detector 2996 (Waters Corporation, Milford, MA, USA). The alkaloids were separated on two different HPLC columns, Luna® C<sub>18</sub> (2), 5 µm, 150 mm x 4.6 mm i.d. and Hypersil® 150 x 4.6 mm i.d. C<sub>18</sub> column (Phenomenex®, Torrance, CA, USA). Analytical balances, Type AG 135 and a Micro Balance MX-5 (Mettler Toledo, Switzerland) were used for weighing standards and samples. An electronic pipette model 71050XET (Biohit PLC, Helsinki, Finland) was used to transfer standard and sample solutions for dilutions.

### Method development

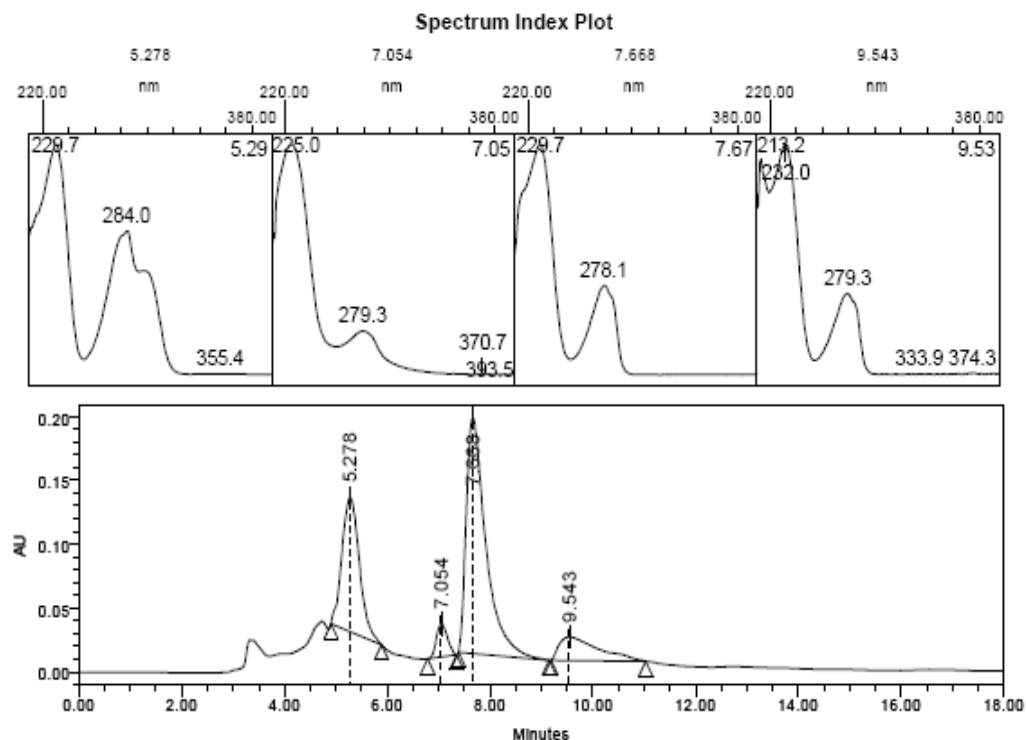
Initially, the method described in the patent literature (9) was used to observe the profiles of *Sceletium* plant extract. The resulting chromatographic profile detected with a PDA detector in the range of 400 to 200 nm indicated many unresolved peaks which eluted within 5 minutes and could not be identified since reference standards for each individual alkaloid were not available. Since the alkaloids are basic molecules, a mobile phase comprising of 100 mM solution of ammonium acetate in water and ACN 70:30 (v:v) was initially used to separate the

alkaloid constituents. The methanol extract showed distinct peaks at RT 5.278, 7.054, 7.668 and 9.543 with UV spectra showing maxima at 229±5 nm, 278±2 nm and 284 nm (Figure 1).

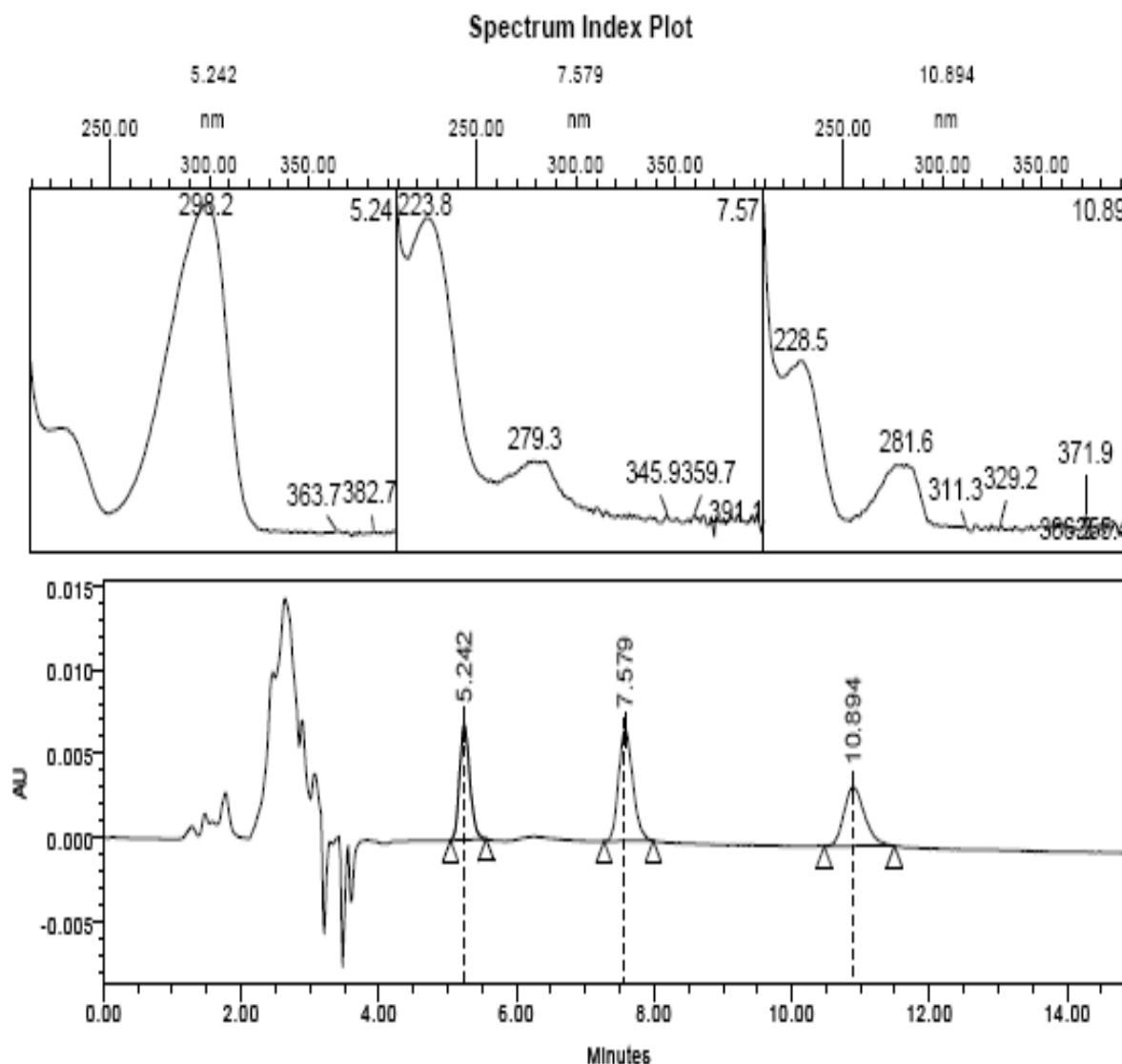
Whereas, using a mobile phase consisting of relatively high salt concentration 100mM ammonium acetate solution facilitated identification of the relevant alkaloid components, the resulting peaks eluted with relatively poor peak shape. Hence, attempts were made to exclude the use of buffer salts and develop a mobile phase using a higher pH to resolve the various *Sceletium* alkaloids and which would be beneficial for use with LC-MS.

The alkaloids, mesembrine, mesembrenone and  $\Delta^7$ mesembrenone were isolated and their structures confirmed by NMR. The HPLC peaks were identified based on the retention times of the prepared reference compounds.

Subsequently, the optimum conditions involved the use of a mobile phase comprising of water:ACN:ammonium hydroxide solution (25%) mixed in a ratio of 70:30:0.01 (v:v:v) and was tested under isocratic conditions using a Hypersil® C<sub>18</sub> column. The three alkaloids were well resolved with  $\Delta^7$ mesembrenone at RT 5.242, mesembrenone at RT 7.579 and mesembrine at RT 10.894 minutes (Figure 2).



**Figure 1.** HPLC-PDA chromatogram of *Sceletium* extract using a mobile phase comprising of 100 mM solution of ammonium acetate in water and ACN 70:30 (v:v). UV spectra for each relevant peak obtained on-line using a PDA detector are depicted above the chromatogram.



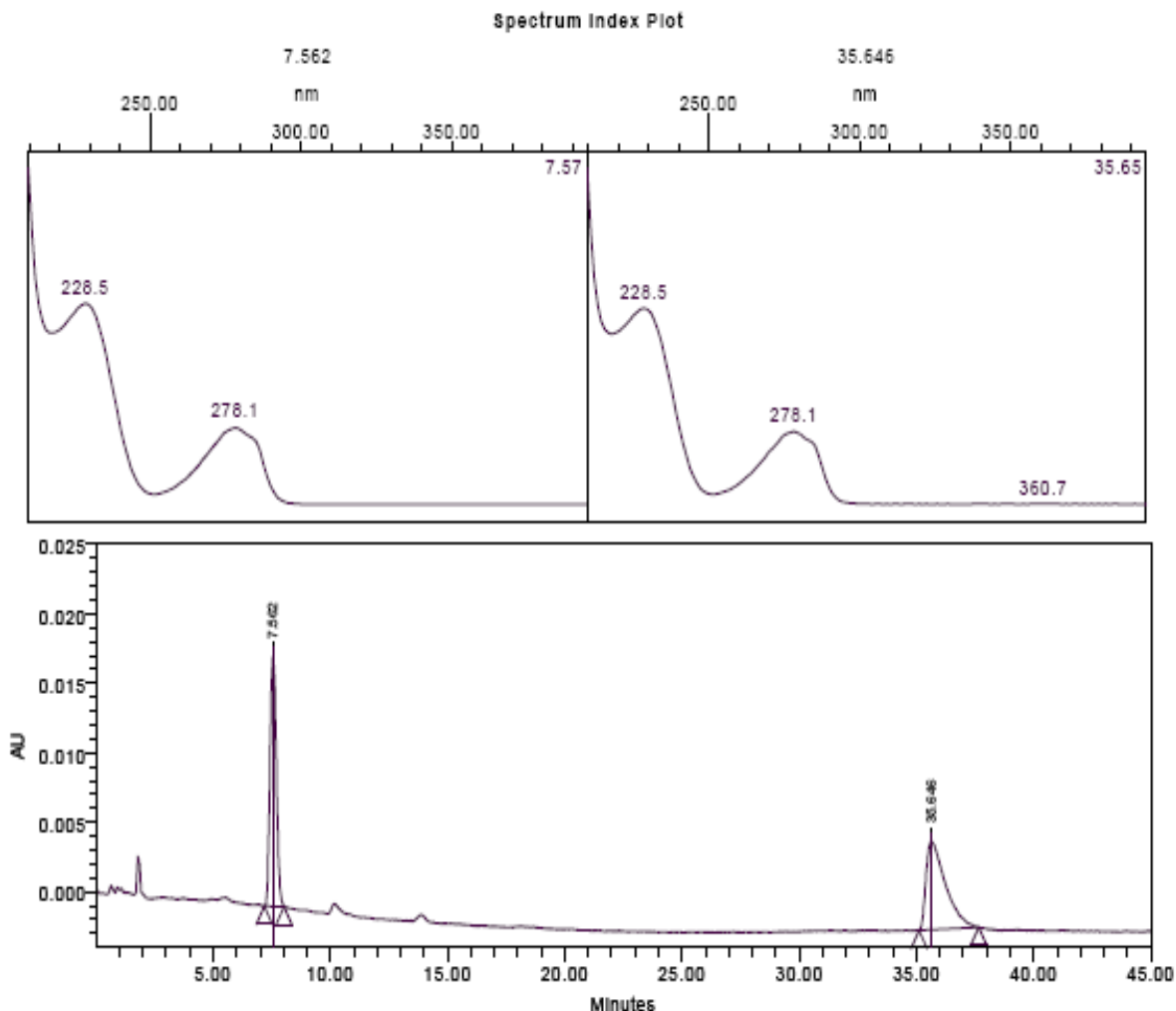
**Figure 2.** HPLC-PDA chromatogram of *Sceletium* standards,  $\Delta^7$ mesembrenone, mesembrenone and mesembrine using a mobile phase comprising of water:ACN:ammonium hydroxide solution mixed in a ratio of 70:30:0.01 (v:v:v). UV spectra for each peak obtained on-line using a PDA detector are depicted above the chromatogram.

The isocratic conditions when applied to the analysis of mesembranol and epimesembranol, posed two problems. Firstly, the epimesembranol peak was retained on the column for an extended period of over 35 minutes (Figure 3) and secondly, separation between mesembranol and mesembrine could only be achieved with an increase of water content to 72 parts. Even though separation was achieved, the resolution and the peak shape were not ideal due to peak tailing (Figure 4).

*Sceletium* plant powder was analyzed using the above mentioned isocratic conditions which

separated  $\Delta^7$ mesembrenone (RT 5.981 min), an unknown compound (RT 6.421 min), mesembrenone (RT 8.774 min), mesembranol (RT 10.502 min) and mesembrine (12.577 min) (Figure 5).

Even though the developed isocratic system resolved the alkaloidal components, the RTs and the peak shape prompted changes to be made to the mobile phase which involved the use of gradient system. In this manner the RTs of the identified alkaloids could be manipulated to optimize peak shapes and also to reduce the RT of epimesembranol.



**Figure 3.** HPLC-PDA chromatogram of mesembranol and epimesembranol using a mobile phase comprising of water:ACN:ammonium hydroxide solution mixed in a ratio of 70:30:0.01 (v:v:v). UV spectra for each peak obtained on-line using a PDA detector are depicted above the chromatogram.

A binary gradient elution system made up of 0.1% ammonium hydroxide solution in water mixed with ACN was developed with an initial 80% aqueous phase changing to 50% over a period of 15 minutes on a Luna<sup>®</sup> C<sub>18</sub>(2) HPLC column. The analysis resulted in satisfactory resolution and retention times for all the reference standards (Figure 6) and related components in *Sceletium* raw material with a total run time of 16 minutes (Figure 7).

## METHOD VALIDATION

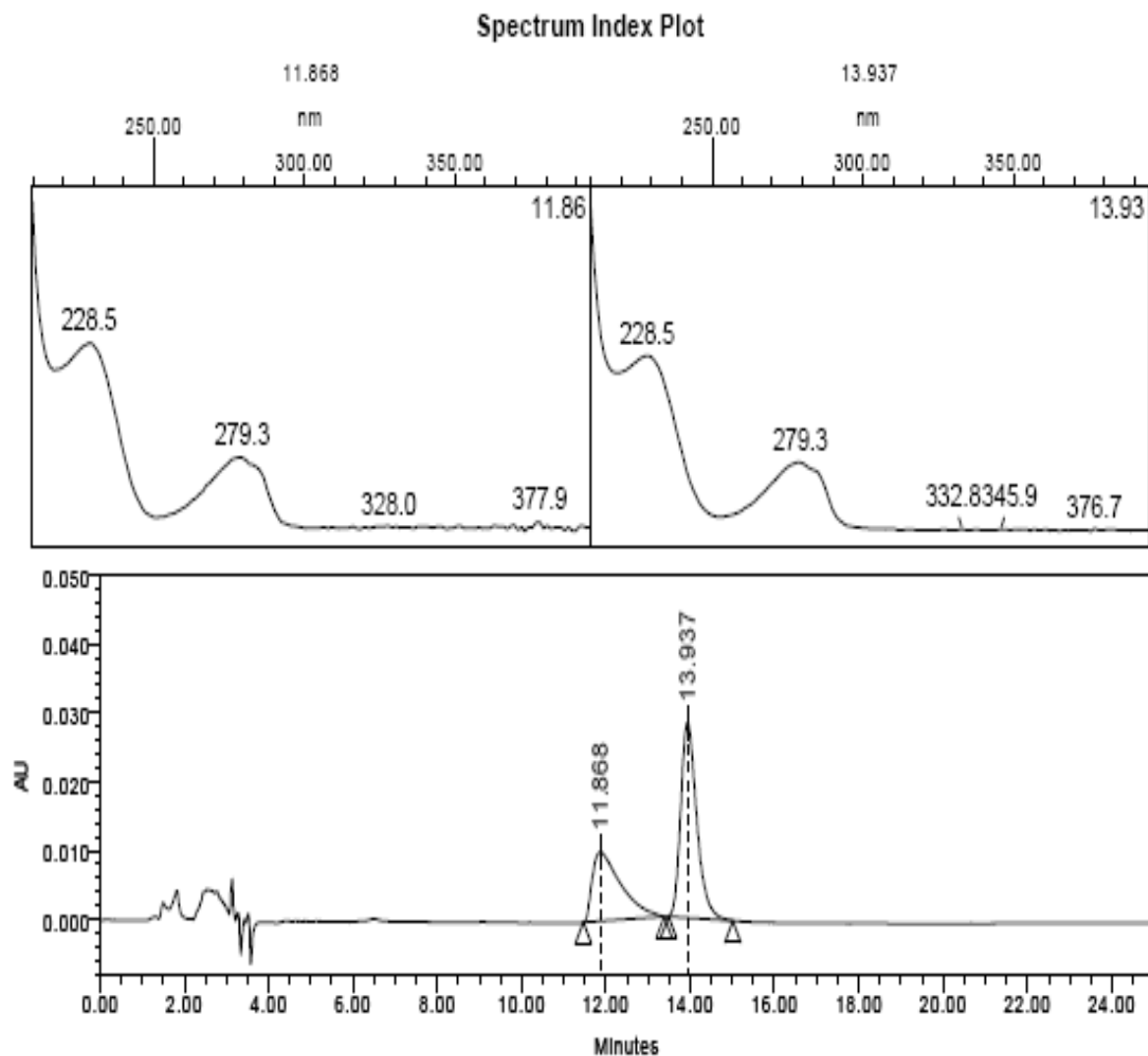
### Linearity

Standard stock solutions (1mg/ml) of  $\Delta^7$ mesembrenone, mesembranol, mesembrenone,

mesembrine hydrochloride and epimesembranol were prepared in methanol. A working stock solution was mixed and diluted appropriately to obtain a concentration of 100  $\mu$ g/ml of each alkaloid. Standard solutions comprising a set of eleven calibrators in a concentration range of 400-60,000 ng/ml were prepared.

### Accuracy

Accuracy studies were performed by separately preparing standard solutions and appropriate dilutions to obtain final concentrations of 4000, 8000 and 10,000 ng/ml for use as QC standards.



**Figure 4.** HPLC-PDA chromatogram of mesembranol and mesembrine using a mobile phase comprising of water:ACN:ammonium hydroxide solution mixed in a ratio of 72:28:0.01 (v:v:v). UV spectra for each peak obtained on-line using a PDA detector are depicted above the chromatogram.

The selected *Sceletium* plant powder was prepared in methanol containing 7.5 mg, 15 mg and 25 mg/ml in triplicate to provide the low, medium and high samples by sonication for 20 minutes and filtered through 0.45  $\mu$ m PVDF membrane filter.

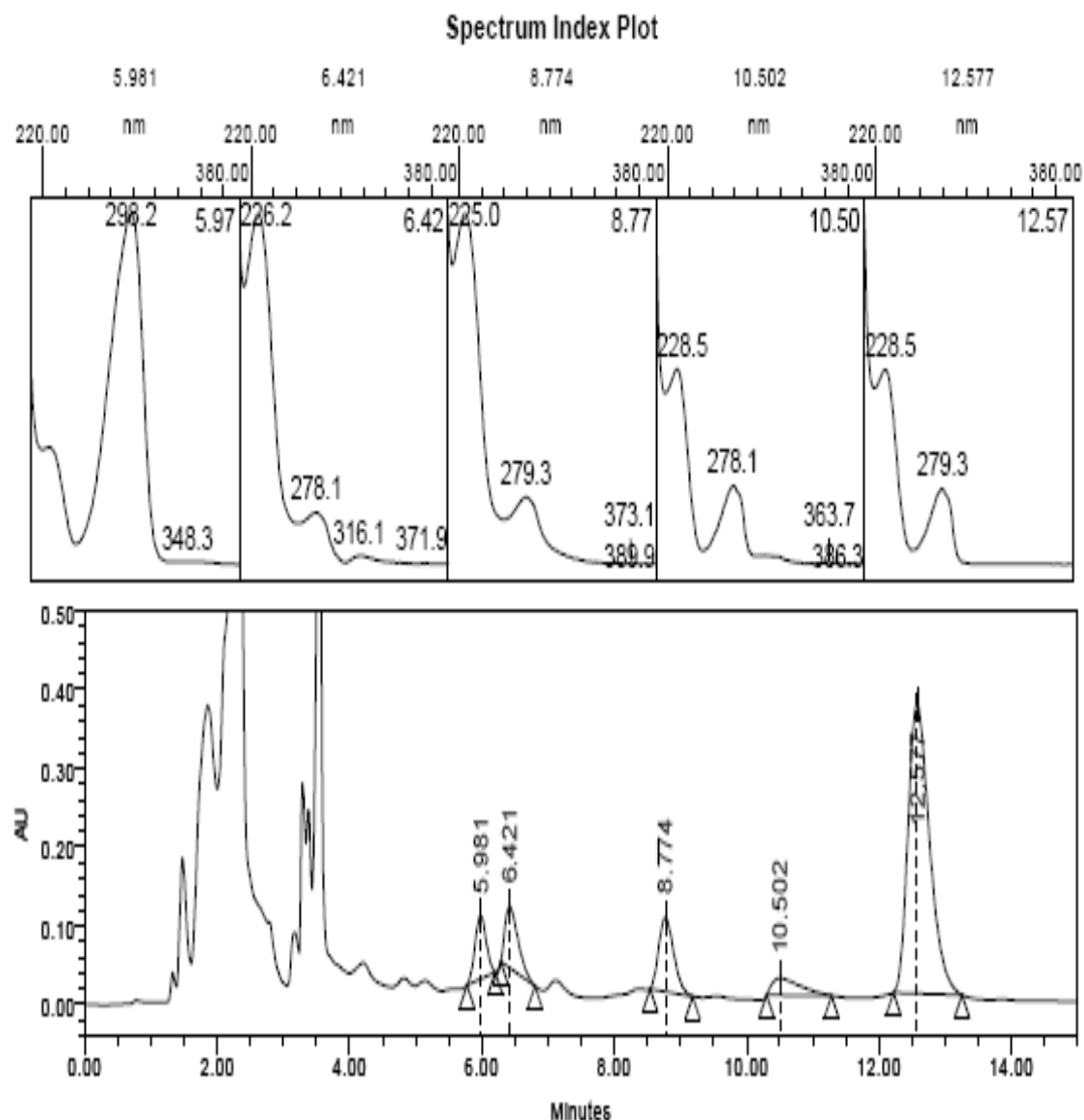
#### Recovery

Recovery studies were carried out by preparing in triplicate, three individual concentrations of 10 mg/ml of plant material. Standard stock solutions

of each of the five alkaloids were added to each of the triplicate sample solutions to result in concentrations of 2000 ng, 4000 ng and 10,000 ng/ml of each of the alkaloids/sample.

#### Limit of Detection and Limit of Quantitation

Standard stock solutions were diluted appropriately to obtain concentrations for the estimation of the limit of detection (LoD) and limit of quantitation (LoQ) according to a signal to noise (S/N) ratio of 3:1 and 10:1, respectively.



**Figure 5.** HPLC-PDA chromatogram of Sceletium plant extract using a mobile phase comprising of water:ACN:ammonium hydroxide solution mixed in a ratio of 72:28:0.01 (v:v:v). UV spectra for each relevant peak obtained on-line using a PDA detector are depicted above the chromatogram.

## RESULTS

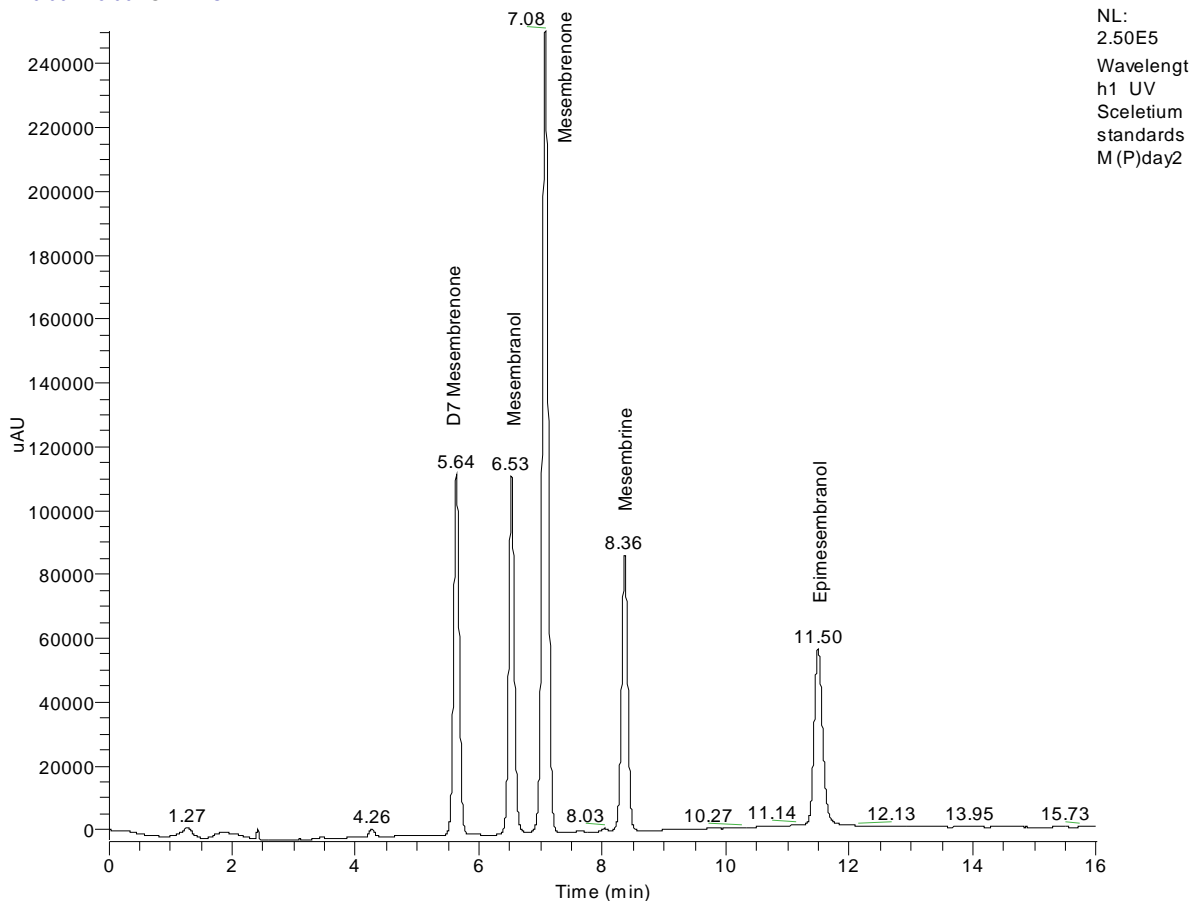
### Confirmation of peak identity

The identity of each relevant alkaloidal peak in the standard chromatogram (Figure 6) was confirmed by the respective spectral data ( $^1\text{H}$ ,  $^{13}\text{C}$  and 2D NMR and mass spectroscopy as well as x-ray crystallography for the crystalline compounds, mesembrine hydrochloride and mesembranol, the others being semi-solid/liquids) obtained for each of the mesembrine-alkaloids used as reference standards.

### Linearity

Calibration curves were constructed by plotting the peak area of each alkaloid versus the concentration corresponding to that alkaloid on each of three days. The curves obtained were found to be linear with determination coefficients better than 0.99 and in one instance, 0.9858 (Table 1).

RT: 0.00 - 16.00 SM: 11G



**Figure 6.** HPLC chromatogram of standard Sceletium alkaloids.

### Accuracy

The studies were performed using QC samples that were prepared separately on each day of analysis. The accuracy of the method was found to be between 96.9%-103% for all five compounds. The inter-day relative standard deviation (RSD) values obtained were less than 3% (Tables 2a and 2b).

### Precision

Studies were carried out on *Sceletium* plant material by preparing three sets of low, medium and high sample concentrations on each day of analysis. The RSD's for all the precision data were found to be less than 6% (Table 3).

### Recovery

The values obtained for this experiment which involved spiking plant material with the relevant alkaloids, are shown in Tables 4a and 4b where the recoveries of all five identified alkaloids were seen to be in the range of 95%-105% for each of

the added compounds with inter-day RSDs of less than 5%.

### Limit of Detection and Limit of Quantitation

The LoD and LoQ of each of the 5 alkaloids were found to be 100 and 200 ng/ml respectively using the respective S/N ratios of 3 and 10.

### DISCUSSION

The *Sceletium* plant is currently receiving great deal of attention in view of its purported potential as natural medicine for use in common mental health and wellness conditions including anxiety, depression and stress. A website has recently been launched whereby the South African company HGH Pharmaceuticals Pty Limited has been granted the country's first ever integrated export and bioprospecting permit to produce Zembrin®, which is patented standardized *Sceletium* extract.



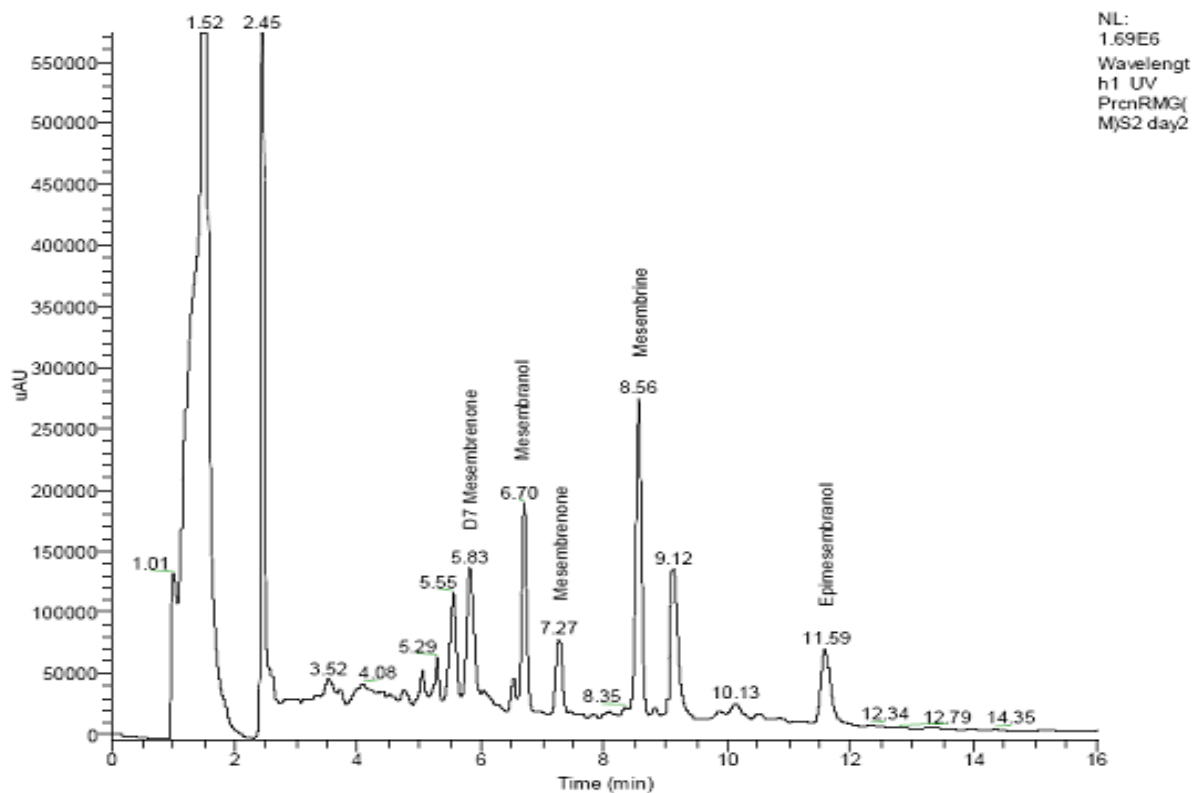


Figure 7. HPLC chromatogram of Sceletium plant material.

Table 1. Linear ranges and coefficients of determination			
Name of the compound	Day	y = mx + c linear model	Determination coefficient (R <sup>2</sup> )
$\Delta^7$ Mesembrenone	Day 1	y = 48.968x + 23904	R <sup>2</sup> = 0.9947
	Day 2	y = 49.425x + 25244	R <sup>2</sup> = 0.9961
	Day 3	y = 47.453x + 25688	R <sup>2</sup> = 0.9960
Mesembranol	Day 1	y = 39.347x + 12817	R <sup>2</sup> = 0.9944
	Day 2	y = 39.748x + 9961.1	R <sup>2</sup> = 0.9960
	Day 3	y = 39.120x + 13789	R <sup>2</sup> = 0.9973
Mesembrenone	Day 1	y = 94.156x + 54025	R <sup>2</sup> = 0.9926
	Day 2	y = 93.704x + 50233	R <sup>2</sup> = 0.9974
	Day 3	y = 91.179x + 41683	R <sup>2</sup> = 0.9977
Mesembrine HCl	Day 1	y = 34.676x + 24050	R <sup>2</sup> = 0.9858
	Day 2	y = 33.674x + 23121	R <sup>2</sup> = 0.9956
	Day 3	y = 33.070x + 23720	R <sup>2</sup> = 0.9945
Epimesembranol	Day 1	y = 32.126x + 11333	R <sup>2</sup> = 0.9925
	Day 2	y = 29.952x + 15173	R <sup>2</sup> = 0.9921
	Day 3	y = 32.581x + 13702	R <sup>2</sup> = 0.9964

**Table2a. Accuracy studies**

<b>Table2a. Accuracy studies</b>													
<b><math>\Delta^7</math> Mesembrenone</b>					<b>Mesembranol</b>				<b>Mesembrenone</b>				
	<b>Day</b>	<b>Actual (ng/ml)</b>	<b>Observed (ng/ml)</b>	<b>Accuracy (%)</b>	<b>RSD</b>	<b>Actual (ng/ml)</b>	<b>Observed (ng/ml)</b>	<b>Accuracy (%)</b>	<b>RSD</b>	<b>Actual (ng/ml)</b>	<b>Observed (ng/ml)</b>	<b>Accuracy (%)</b>	<b>RSD</b>
<b>Low spike</b>	1	3520.00	3432.40	97.50		4240.00	4239.11	101.25		4320.00	4293.20	99.40	
	2	3360.00	3296.85	98.12	1.71	4160.00	4196.00	100.86	0.99	4160.00	4150.55	99.77	1.64
	3	3840.00	3866.44	100.68		4320.00	4292.60	99.37		4400.00	4385.91	99.68	
<b>Medium Spike</b>	1	7040.00	6847.90	97.23		8480.00	8038.42	94.79		8640.00	8408.45	97.32	
	2	6720.00	6569.00	97.75	0.30	8320.00	8036.60	96.59	1.10	8320.00	8213.10	98.71	1.57
	3	7680.00	7511.50	97.81		8640.00	8365.60	96.80		8800.00	8417.23	95.65	
<b>High spike</b>	1	8800.00	8802.40	100.03		10600.00	10711.20	101.05		10800.00	10860.40	100.6	
	2	8400.00	8546.85	101.75	2.17	10400.00	10395.51	99.96	1.02	10400.00	10473.80	100.71	2.30
	3	9600.00	9352.00	97.45		10800.00	10689.50	99.00		11000.00	10633.00	96.70	

**Table2b Accuracy studies (continued)**

<b>Table2b Accuracy studies (continued)</b>									
<b>Mesembrine</b>					<b>Epimesembranol</b>				
	<b>Day</b>	<b>Actual (ng/ml)</b>	<b>Observed (ng/ml)</b>	<b>% Accuracy</b>	<b>RSD</b>	<b>Actual (ng/ml)</b>	<b>Observed (ng/ml)</b>	<b>% Accuracy</b>	<b>RSD</b>
<b>Low spike</b>	1	4240.00	4192.53	98.9		4160.00	4109.82	98.79	
	2	4080.00	4068.10	99.71	1.60	3680.00	3777.44	102.64	2.09
	3	3840.00	3710.60	96.6		4160.00	4129.68	99.27	
<b>Medium Spike</b>	1	8480.00	8229.90	97.05		8320.00	8382.05	100.75	
	2	8160.00	8264.74	101.28	2.30	7360.00	7545.04	102.51	2.05
	3	7680.00	7723.68	100.57		8320.00	8185.72	98.39	
<b>High spike</b>	1	10600.00	10303.93	97.21		10400.00	10715.70	103.04	
	2	10200.00	10459.70	102.55	2.80	9200.00	9535.00	103.64	2.60
	3	9600.00	9455.46	98.49		10400.00	10264.00	98.70	

<b>Table 3 Precision studies</b>				
<b>Content in µg/100mg of plant material</b>				
<b>Compound</b>	<b>Day1</b>	<b>Day2</b>	<b>Day3</b>	<b>Inter-day % RSD (n=3)</b>
<b>Δ<sup>7</sup>Mesembrenone</b>				
Low	20.00 (±1.00)	20.33 (±0.58)	21.33 (±0.58)	3.37
Medium	23.67 (±0.58)	23.00 (±1.00)	24.33 (±1.15)	2.81
High	23.00 (±1.73)	23.67 (±1.53)	23.00 (±2.00)	1.67
<b>Mesembranol</b>				
Low	91.33 (±7.50)	94.67 (±9.30)	97.30 (±2.52)	3.17
Medium	97.67 (±3.22)	96.33 (±5.70)	91.30 (±5.03)	3.53
High	88.67 (±1.12)	93.00 (±2.65)	88.67 (±5.13)	2.77
<b>Mesembrenone</b>				
Low	12.10 (±1.87)	12.87 (±0.84)	12.67 (±1.40)	3.20
Medium	12.87 (±0.38)	13.20 (±0.38)	13.03 (±1.72)	1.27
High	13.37 (±0.71)	13.43 (±0.64)	13.83 (±0.47)	1.85
<b>Mesembrine</b>				
Low	144.50 (±0.12)	147.90 (±1.10)	149.40 (±2.51)	1.70
Medium	165.40 (±1.63)	166.03 (±5.65)	163.30 (±4.54)	0.87
High	165.30 (±3.15)	167.24 (±10.16)	162.50 (±5.91)	1.44
<b>Epimesembranol</b>				
Low	55.00 (±2.56)	60.56 (±7.51)	56.03 (±3.47)	5.17
Medium	61.13 (±3.75)	64.80 (±5.60)	64.00 (±3.80)	3.05
High	62.25 (±2.74)	68.43 (±2.92)	62.14 (±0.86)	5.60
(±SD) = Standard deviation				
Low = 7.5 mg/ml (n=3), Medium = 15 mg/ml (n=3), High = 25 mg/ml (n=3); Total samples n=9 each day				

The pharmacological actions of Zembrin® are described in granted international patents and new patent applications. Validated analytical methods to assay *Sceletium* plant material and dosage forms for relevant alkaloidal content have not hitherto been reported in the published scientific literature. Published analytical procedures have focused only on qualitative determinations.

Since the identified markers have closely related structures, of which two alkaloids, mesembranol and epimesembranol are epimers, the main objective of this investigation was to develop an efficient HPLC method for the separation and quantitative analysis of the relevant alkaloid components in *Sceletium*, and also to reduce, in particular the RT of epimesembranol, which was too long (> 30 min) under isocratic conditions. Use of a PDA detector was extremely valuable for peak identification and homogeneity testing during the initial method development.

A simple, accurate, precise, rapid and reproducible HPLC method was developed for the identification and quantitative analysis of 5 relevant *Sceletium* alkaloids which has been successfully applied for the assay and QC of *Sceletium* plant material. Furthermore, this method was found to efficiently separate the alkaloidal markers from complex components present in plant material using a simple methanol sample extraction procedure. This method has also been successfully used to study chemotaxonomy of some *Sceletium* species and has provided impetus for the future development of quality monographs for plant and dosage forms containing *Sceletium*.

#### ACKNOWLEDGEMENT

Funding from the South African Medical Research Council (MRC) and Rhodes University is gratefully acknowledged.

Table 4a. Recovery studies

	$\Delta^7$ Mesembrenone content ( $\mu\text{g/ml}$ )	Day-1 Observed content ( $\mu\text{g/ml}$ )	% Recovery	% RSD (n=3)	$\Delta^7$ Mesembrenone content ( $\mu\text{g/ml}$ )	Day-2 Observed content ( $\mu\text{g/ml}$ )	% Recovery	% RSD (n=3)	$\Delta^7$ Mesembrenone content ( $\mu\text{g/ml}$ )	Day-3 Observed content ( $\mu\text{g/ml}$ )	% Recovery	% RSD (n=3)	Inter-day Recovery % RSD (n=3)
*Content	2.2				2.2				2.3				
Low	(+1.76) <sup>1</sup> 3.96 <sup>2</sup>	3.89 ( $\pm 0.05$ )	96.20	2.96	(+1.84) 4.04	4.00 ( $\pm 0.07$ )	99.00	1.75	(+1.92) 4.22	4.31 ( $\pm 0.05$ )	102.13	1.22	3.00
Medium	(+3.52) 5.72	5.83 ( $\pm 0.10$ )	101.90	2.85	(+3.68) 5.88	5.72 ( $\pm 0.10$ )	97.30	1.83	(+3.84) 6.14	6.17 ( $\pm 0.11$ )	100.50	1.82	2.36
High	(+8.80) 11.00	10.73 ( $\pm 0.13$ )	97.50	1.50	(+9.20) 11.40	10.92 ( $\pm 0.24$ )	95.80	2.20	(+9.60) 11.90	11.69 ( $\pm 0.12$ )	98.70	1.10	1.50
	Mesembranol content ( $\mu\text{g/ml}$ )	Day-1 Observed content ( $\mu\text{g/ml}$ )	% Recovery	% RSD (n=3)	Mesembranol content ( $\mu\text{g/ml}$ )	Day-2 Observed content ( $\mu\text{g/ml}$ )	% Recovery	% RSD (n=3)	Mesembranol content ( $\mu\text{g/ml}$ )	Day-3 Observed content ( $\mu\text{g/ml}$ )	% Recovery	% RSD (n=3)	Inter-day Recovery % RSD (n=3)
*Content	9.2				9.5				9.2				
Low	(+2.12) 11.32	11.25 ( $\pm 0.04$ )	99.38	2.05	(+2.08) 11.58	11.58 ( $\pm 0.06$ )	100.00	0.51	(+2.08) 11.28	11.38 ( $\pm 0.05$ )	100.90	0.50	0.76
Medium	(+4.24) 13.44	13.14 ( $\pm 0.16$ )	97.16	3.63	(+4.16) 13.66	13.65 ( $\pm 0.15$ )	99.90	1.10	(+4.16) 13.36	13.46 ( $\pm 0.08$ )	100.70	0.60	1.9
High	(+10.60) 19.80	19.87 ( $\pm 0.50$ )	100.40	4.50	(+10.40) 19.90	19.81 ( $\pm 0.34$ )	99.50	1.72	(+10.40) 19.60	19.95 ( $\pm 0.30$ )	101.80	1.47	1.15
	Mesembrenone content ( $\mu\text{g/ml}$ )	Day-1 Observed Content ( $\mu\text{g/ml}$ )	% Recovery	% RSD (n=3)	Mesembrenone content ( $\mu\text{g/ml}$ )	Day-2 Observed content ( $\mu\text{g/ml}$ )	% Recovery	% RSD (n=3)	Mesembrenone content ( $\mu\text{g/ml}$ )	Day-3 Observed content ( $\mu\text{g/ml}$ )	% Recovery	% RSD (n=3)	Inter-day Recovery % RSD (n=3)
*Content	1.30				1.30				1.30				
Low	(+2.16) 3.46	3.33 ( $\pm 0.90$ )	95.40	4.25	(+2.08) 3.38	3.32 ( $\pm 0.07$ )	98.20	2.10	(+2.2) 3.50	3.32 ( $\pm 0.05$ )	95.00	1.44	1.80
Medium	(+4.32) 5.62	5.37 ( $\pm 0.60$ )	95.60	1.53	(+4.16) 5.46	5.31 ( $\pm 0.09$ )	97.30	1.60	(+4.4) 5.70	5.45 ( $\pm 0.04$ )	95.60	0.73	1.02
High	(+10.80) 12.10	12.12 ( $\pm 0.18$ )	100.16	1.70	(+10.40) 11.70	11.44 ( $\pm 0.42$ )	97.80	3.71	(+11.00) 12.30	11.70 ( $\pm 0.22$ )	95.10	1.87	2.60

\*Actual content; <sup>1</sup>spiked amount; <sup>2</sup>represents the total content, i.e. spiked plus original content (follows through the table), ( $\pm$ SD)

Table 4b. Recovery studies (continued)

	Day-1				Day-2				Day-3				Inter-day
	Mesembrine content (µg/ml)	Observed content (µg/ml)	% Recovery	% RSD (n=3)	Mesembrine content (µg/ml)	Observed content (µg/ml)	% Recovery	% RSD (n=3)	Mesembrine content (µg/ml)	Observed content (µg/ml)	% Recovery	% RSD (n=3)	% RSD (n=3)
*Content	15.82				16.00				15.80				
Low	(+1.88) <sup>1</sup> 17.7 <sup>2</sup>	17.64 (±0.09)	99.65	4.65	(+1.81) 17.81	17.88 (±0.30)	100.40	1.62	(+1.85) 17.65	18.23 (±0.08)	103.30	0.42	1.91
Medium	(+3.77) 19.52	19.58 (±0.09)	100.30	2.33	(+3.63) 19.63	19.75 (±0.12)	100.60	0.63	(+3.70) 19.50	20.12 (±0.11)	103.10	0.53	1.51
High	(+9.42) 25.24	25.21 (±0.17)	99.90	1.78	(+9.07) 25.07	25.45 (±0.18)	101.50	0.72	(+9.24) 25.04	25.62 (±0.14)	102.30	0.55	1.21
	Day-1				Day-2				Day-3				Inter-day
	Epimesembranol content (µg/ml)	Observed content (µg/ml)	% Recovery	% RSD (n=3)	Epimesembranol content (µg/ml)	Observed content (µg/ml)	% Recovery	% RSD (n=3)	Epimesembranol content (µg/ml)	Observed content (µg/ml)	% Recovery	% RSD (n=3)	% RSD (n=3)
*Content	5.90				6.20				6.10				
Low	(+2.08) 7.98	8.04 (±0.06)	100.80	3.04	(+1.84) 8.04	7.75 (±0.08)	96.40	1.00	(+2.08) 8.18	8.14 (±0.04)	99.50	3.50	2.30
Medium	(+4.16) 10.06	10.26 (±0.06)	102.00	1.31	(+3.68) 9.88	9.54 (±0.08)	96.50	0.86	(+4.16) 10.26	10.30 (±0.03)	100.40	0.30	2.80
High	(+10.40) 16.30	17.01 (±0.48)	104.40	4.30	(+9.2) 15.40	15.13 (±0.21)	98.20	1.36	(+10.40) 16.50	16.53 (±0.26)	100.60	1.58	0.45

\*Actual content; <sup>1</sup>spiked amount; <sup>2</sup>represents the total content, i.e. spiked plus original content (follows through the table), (±SD)

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