Comparative Study of Different Column Types for the Separation of Polar Basic Hallucinogenic Alkaloids


Abstract
The number of hallucinogenic compounds that have been separated simultaneously by liquid chromatography is limited. This research aimed to identify a column(s) that can allow for separation of several hallucinogens. The extent of separation of seven polar hallucinogenic tryptamine and phenethylamine derived alkaloids containing a basic N atom that becomes protonated at low pH values were investigated on five reverse-based columns and one hydrophilic interaction liquid chromatography (HILIC) column. An RP-phenyl and a negatively charged fused core HILIC were identified and recommended as effective columns in this regard. This research is the first to introduce a HILIC column in separation of hallucinogenic alkaloids and to simultaneously study the separation of the two principal psychoactive agents, muscimol and psilocin. The results of this study showed that better separation in HILIC columns can be achieved if an aprotic diluent of relatively lower polarity is used to introduce analytes into a mobile phase of higher polarity. Tetrahydrofuran was recommended where acetonitrile is used as the organic component of the mobile phase.

Keywords
Hallucinogen, polar compounds, psilocin, muscimol, HILIC, tetrahydrofuran.

1. Introduction
Hallucinogens are pharmacologically psychoactive agents that have mind-altering effects and result in non-ordinary psychological experiences.1 A hallucinogen can be psychedelic, dissociative or deliriant depending on its subjective and behavioral effects. Common are the psychedelics, which are structurally classified as tryptamine or phenethylamine derivatives. Mushrooms in the *Amanita* and *Psilocybe* genera are known to have hallucinogenic properties due to the presence of ibotenic acid and psilocybin as well as their main respective metabolites, muscimol and psilocin, which are considered the principal psychoactive agents.2–4 The toxicity and psychological consequences of the main psychoactive agents are well documented.1,3–11

Almost all hallucinogens are alkaloids and contain an aromatic-alkyl-N moiety that is basic at the N atom, except salvinorin A and tetrahydrocannabinol. The pentafluorophenylpropyl (PFP) stationary phase has been used successfully in separating some phenethylamine alkaloids.36–37 It has also been used on a combined mixture of eight tryptamine and phenethylamine derivatives in *A. rigidula* and its dietary extracts with only two analytes co-eluting.34 When a C18 column was used, five of the eight analytes could not be baseline separated. PFP is itself a reverse phase approach and has shown great separating ability for basic N-based compounds. Brandt and Martins38 have replaced particle-packed reverse phase columns with porous monolithic packing in C4 column in separating 24 N, N-dialkylated tryptamines using liquid chromatography-electrospray ionization-ion trap mass spectrometry/mass spectrometry.

Even though the two psychoactive agents have a minimal chance of coexisting in mushrooms, an approach that allows for their simultaneous quantification in body fluids of victims or culprits of neuro-excitatory mushroom consumption will help in saving time towards proving prior exposure to a specific mushroom genus. Our research thus aimed to identify column types that can enable a successful separation and quantification of hallucinogenic alkaloids using a liquid chromatographic approach. A successful concurrent study of the hallucinogenic alkaloids can help maximize the number of these alkaloids that can be analyzed in biological samples. The results of our study can also help in affording an informed starting point for the separation of related compounds by liquid chromatography. Muscimol and psilocin and their precursors, tryptophan and tryptamine, were used as model diamine alkaloid compounds. Included was the monoamine phenethylamine and its two derivatives, 3,4-dimethoxyphenethylamine and hordenine. Despite having different chemical structures, these tryptamine...
and phenylalkylamine derivatives all contain a substituted ethylamine group that is attached to a ring and also produce remarkably similar subjective effects in the human body as neurotransmitter agonists.\(^1,2,6,18,20,38,39\) Only muscimol has a methylamine group connected to a five-membered ring. The alkylamine group in hallucinogens found in mushrooms is connected to a five-membered ring while the Cactus-sourced have this branch connected to a six-membered ring (Fig. 1).

The optimum separation of the seven alkaloids was investigated on six columns. This was achieved by investigating parameters responsible for maximizing separation on two alkyl reverse phase (RP-alkyl) columns, a reverse phase amide (RP-amide), two phenyl based reverse phase columns and a HILIC column.

### 2. Experimental

#### 2.1. Chemicals

All compounds except psilocin (PSI) were purchased from Sigma-Aldrich (Johannesburg, South Africa). PSI in powder form was obtained via the Department of Forensic Sciences, Capetown, South Africa. Muscimol (MUS), tryptophan (TRP), tryptamine (TA) and 3,4-dimethoxyphenethylamine (DMPE) were all purchased in powder form while phenethylamine (PEA) and hordenine (HO) were in liquid form. Ammonium acetate, ammonium formate, acetic acid and formic acid were also purchased from Sigma-Aldrich (Johannesburg, South Africa). HPLC-grade acetonitrile (MeCN), methanol (MeOH), acetone and tetrahydrofuran (THF) were obtained from Merck (Pty) Ltd, Johannesburg, South Africa. Deionized water used to prepare standard solutions and the mobile phase was purified using a Milli-Q-RO4 system (Millipore, Bedford, MA, USA). All six columns investigated were purchased from Sigma-Aldrich (Johannesburg, South Africa).

#### 2.2. Standards

The injection standards were prepared by dissolving appropriate volumes of the stock solutions into deionized water, MeOH, MeCN, THF or acetone depending on the parameter and the column being investigated. Stability issues associated with psilocin and muscimol\(^20,29,40\) and the difficulty in their procurement limited the quantities sufficient enough for this analytical research. Psilocin and muscimol were then injected at 1 and 2 \(\mu\)g mL\(^{-1}\) respectively. The other analytes were injected at 4 \(\mu\)g mL\(^{-1}\).

#### 2.3. Instrumentation

A Bishoff LC-CaDI 22-14 system (Leonberg, Germany) with a Lambda 1010 UV detector set at 280 nm was used for quantification. Elution was done by isocratic means and the analyte injection was done using Rheodyne Series 7725i manual injector with 10 \(\mu\)l sample loop. Prior to elution, each column was equilibrated with the mobile phase for at least 15 min. After each HILIC investigation, the column was first washed with a MeCN:H\(_2\)O mobile phase containing the same organic component composition as the last mobile phase used for elution and then finally with 100 % MeCN for 10 min.

#### 2.4. Column and Hallucinogen Properties

The physicochemical properties of the columns and the compounds used are given in Table 1 and Fig. 1, respectively. X log P\(_3\) is an atom based additive strategy for predicting the octanol/water partition coefficient (log P). Developed by Cheng et al.\(^41\), it estimates the log P value by adding the contributions from individual atoms constituting the compound and some correction factors. The alkyl RP columns are the most used in liquid chromatographic separations of hallucinogenic alkaloids. Active phase-analyte interactions are exclusively hydrophobic. The RP-amide was chosen for the presence of the embedded polar amide group, which is expected to improve retention of polar analytes and also because it has always been viewed as one of the first definite alternative to C\(_{18}\) columns for analysis of nonpolar compounds.

The phenyl-based columns were targeted because of the \(\pi-\pi\) interactions with the cyclic groups of the target compounds. The negatively charged fused-core silica Ascentis Express HILIC choice was based on studies by Chirita et al.\(^42\) and recommendations by McCalley considering that our model compounds have positively charged N atoms at low pH values.\(^42,43\)

#### 2.5. Optimization of Column Parameters

The selection of columns suitable for separation of alkaloids was achieved by optimizing each column parameter. The initial conditions for all parameters investigated were based on the principles of retention for each type of column. The mobile phase was always buffered at pH 3 with formate. Under these conditions all our research compounds were positively charged on the basic N atom, while the \(-\)OH group remained neutral except for TRP whose pKa (OH) value makes it zwitterionic. While the reversed phase principle requires that the analytes be...
in their neutral state for effective retention, the limiting factor was the pKa values (Fig. 1) that lie outside the pH 2–8 working range of most columns. All mobile phases in quantitation of hallucinogenic compounds use acetate and formate buffers.4,12–32 The analytes are charged in the pH working range of these buffer systems.

The formate buffering system was prepared only when needed by mixing appropriate quantities to create a specific pH value buffering system. The reported formate concentrations were the final concentrations in the mobile phase. Because of its low viscosity44–48, MeCN was used as the organic component of the mobile phase. The chromatographic results presented in this study were for the optimized parameters and for those columns where a separation was observed.

The parameters investigated for alkyl columns included the mobile phase composition and salt concentration of the buffer, flow rate and analyte injection solvent at ambient temperatures. Chirita et al. has suggested the use of HILIC columns for separation of polar analytes.42 While the essential parameters in HILIC separation are known42,44,49–51, our research further investigated the effect of analyte injection diluents. Water, methanol, acetonitrile, acetone and tetrahydrofuran were investigated as potential analyte diluents. These solvents were chosen based on polarity, eluotropic order and their involvement in other procedures of the research.

3. Results and Discussion

3.1. Alkyl Columns

The composition of the mobile phase was altered up to Buffer: MeCN 80:20 % (v/v) 100 mM formate, beyond which it was apparent that the seven compounds could not be separated with most eluting in the column void volume and all detected within 2 min of \( t_m \) at a 0.6 mL min \(^{-1} \) flow rate. The THF diluent had a comparably superior retention power even though all analytes eluted before the THF solvent, a situation also observed by Loeser and Drumm.52 Solvophilic effects due to high polarity and the charged state of these basic analytes might explain the reduced retention behaviour.53

3.2. Ascentis Hexyl Phenyl Column

Better retention and separation was observed when the above alkyl column conditions were applied. This was attributed to the presence of the \( \pi-\pi \) interactions, hydrophobic, \( \pi-\pi \) interactions and HILIC H-bonding. The structural position of the benzyl group in TA, in addition to the hydrophobic nature of the ethyl connecting group, might have explained its position in the elution order (Fig. 2).

TRP has a similar positional benzyl group, but TRP is also an amino acid. Both PSI and HO have a phenolic ring and their X log P3 values are the same, but PSI also has a pyrrole ring joining the phenolic group and the ethyl-amino group. The electron-withdrawing methoxy group in DMPE might explain its retention behaviour compared to PEA. Generally, the three indolealkyltryptamine derivatives (TA, TRP and PSI) could be separated based on type of substituents on the indole ring. This seemed to be the case too with the phenethylamine alkaloids. HO, TRP and DMPE were less retained by the hexyl phenyl column resulting in co-elution.

3.3. Ascentis Phenyl Column

Better retention and separation compared to the alkyl and the alkyl phenyl columns was observed. The elution order between TA and PSI was reversed, PEA and DMPE co-eluted while TRP, HO and DMPE became clearly resolved. Compared to the com-
peting effect of the hydrophobic hexyl spacer in hexyl phenyl column, the butyl linking group in the phenyl column allowed for stronger $p-p$ electron interactions between the phenyl group and the cyclic groups of the analyte. The shorter linking group in the phenyl column also allows for some kind of hydrophilic interaction (HILIC) that might take advantage of the polarity of the analytes. The presence of the O atom as a substituent on the benzyl group might have improved the HILIC effects of the column hence the observed behaviours of PSI, HO and DMPE compared to the other RP columns. The mobile phase was buffer (100 mM formate) and MeCN in a ratio 87:13 % (v/v) at 0.5 mL min$^{-1}$ beyond which no further separation could be achieved.

3.4. Ascentis RP-Amide Column

Compared to alkyl columns, better retention and selectivity for RP-amide was achieved at Buffer:MeCN 35:65 % (v/v), 100 mM formate. This was expected because, in addition to the hydrophobic interactions, the RP-amide has the H-bond accepting amide carbonyl group that interacted with the –OH and uncharged –NH groups of the analytes. The presence of two H-bond donating groups in PSI might have been responsible for its position in the elution order while HO only has –OH group. Such groups are absent in PEA implying that its retention was based on the hydrophobic alkyl group while the nucleofugic methoxy group in DMPE could have acted as an H-bond acceptor. However, the sp$^2$-hybridized O atom of the isoxazole in MUS cannot be an H-bond acceptor. The amide column was the only RP mode that retained MUS and the elution order was different from the phenyl columns. The observed reduction in selectivity of the RP-amide compared to phenyl columns was attributed to the charged state of these basic analytes. Retention was low and separation less achieved at low organic solvent content of the mobile phase.

3.5. Ascentis HILIC Column

All the model analytes were separated when the analytes were dissolved in THF and eluted at 0.4 mL min$^{-1}$ with MeCN:Buffer 90:10 % (v/v) 10 mM formate. Fig. 3 compares separation when THF and MeCN diluents were used. Retention, and hence separation, was minimal when analytes were diluted with either H$_2$O or MeOH. A general reversal in the elution order compared to that of the RP approach was observed. In HILIC, a relatively hydrophobic high organic content mobile phase is used to elute analytes through a hydrophilic column. The mechanism is the extent of partitioning of analytes between a pseudostagnant aqua layer formed at the surface of the stationary phase and the mobile phase, as confirmed by McCalley and Neue, and Dinh et al. The result is that retention favours the more hydrophilic analytes as confirmed by the negative slope of Fig. 4.

3.5.1. The HILIC Complexity

Our results confirmed that if analyte class is ignored, the general principles of HILIC separation cannot explain the behaviour of some analytes as shown by log k vs X log P$^3$ in Fig. 4. This has also been observed by other researchers. This is due to presence of secondary interactions that come into effect based on the structural composition and sterical orientation of the analyte, and the salt concentration of the mobile phase. Such polar basic interactions have been identified as electrostatic, ion exchange, H-bonding and some weak hydrophobic interactions depending on stationary phase, mobile phase and analyte conditions.

For analytes where the backbone is similar, the electrostatic effect seemed to be more important as observed with each of the two alkaloid types. The presence of alkyl substituents on the basic N resulted in a reduction in retention times, implying that the basic N was involved in these electrostatic interactions. Alkylation was deemed responsible for neutralizing the positive charge on the N atom. This also allowed for elution order to follow protocol. For example, if phenethylamine or tryptamine derivatives only were compared in Fig. 4, $R^2$ values were 0.9882 and 0.9938, respectively. The resonance-stabilized negative
charge on the carboxylic group of TRP enhanced the electrostatic interaction in addition to hydrophilic partitioning hence its observed behaviour together with peak broadening. General peak broadening for those compounds that retain more, as shown by the peak for TRP, has been reported before.\textsuperscript{42,59}

3.5.2. The Moderately Polar Aprotic Diluent for Analyte Injection

While the emphasis is on using a non-polar aprotic solvent for a mobile phase, solvents that are predicted to give better results than MeCN are not recommended because of background noise and detection issues proving the irreplaceability of MeCN.\textsuperscript{30,58,60–63} Studies by Li and Huang\textsuperscript{64} showed better separation when MeCN was used as the mobile phase compared to MeOH, THF and isopropanol. Attempts by Dinh et al.\textsuperscript{58} to admixture THF with the MeCN-containing mobile phase reduced the extent of formation of the water enriched layer on the stationary phase. Also, acetone and THF have very high UV-vis absorbance cut-off wavelengths and show reduced signal response in MS detection. While the recommendations guided by the viscous fingering phenomenon are that the injection solvent must be as close to the mobile phase organic composition as possible with a minimal 50 % MeCN composition, such declarations have been based on comparison of MeCN with polar protic solutions like H$_2$O and MeOH.\textsuperscript{50,55,54} It was noted that most publications follow this protocol and rarely report or optimize their injection diluents. Controversies arise considering that the polar protic MeOH and H$_2$O are the solvents of choice for the extraction of polar compounds from biological matrices. This prompted us to try diluting the polar-dissolved stock solutions with the low elution order aprotic solvents. Our research has successfully identified the introduction of a moderately polar aprotic dilution solvent like THF for analyte injection as an essential parameter in MeCN-mobile phase core column HILIC interactions. Compared to MeCN as an injection eluent at the same ratio, better retention and hence separation was observed when THF was used (Fig. 3). While observations by Dinh et al.\textsuperscript{58} might come into effect, THF is aprotic and relatively apolar and if introduced as an analyte solvent will maximize analyte partitioning into the pseudo-stagnant water-enriched layer thus counteracting the admixture effect. It was suggested that THF allows for analyte on-column focussing where the analytes are concentrated or accumulated into the pseudo water layer within a narrow range at the column entrance, so that by the time MeCN mobile phase enters the column the analytes are already retained in the stationary phase. This postulation is based on the review by Jandera et al.\textsuperscript{55} who attributed this on-column focussing to weak eluents. Related to this observation is the use of dichloromethane in place of water for injection of acrylamide by Backe et al.\textsuperscript{35} While both dichloromethane and THF have similar polarity (polarity indices of 3.1 and 4 respectively), THF has the advantage of 100 % solubility in water. This implies that THF has the ability to penetrate the pseudo-stagnant aqua layer on the surface of the HILIC column thereby enhancing the partitioning of the analytes into the aqua layer. In HILIC columns where H-bond interactions are less important, THF is a weaker eluent than MeCN based on polarity. The situation is reversed for silica modified columns where H-bond accepting abilities come into effect.\textsuperscript{40,44} Peaks were unidentified when analytes were diluted with acetone. Acetone’s absorbance cut-off point is 330 nm.

The effect of electrostatic interactions by increasing concentration of formate buffer beyond 10 mM on the extent of separation and retention was inversely minimal and relatively absent for TRP but seemed to improve sharpness and reduce tailing experienced especially with TRP. Increasing salt concentration is expected to act by competing for electrostatic interaction with the stationary phase hence reducing the effects of electrostatic attractions on retention. TRP also contains a negative charge on its acidic group that tends to cancel out this impact, hence its observed elution status. Other studies where the effect has been more pronounced have identified the residual silanols as the major contributors, a scenario typical of non-endcapped columns. Using a high organic content mobile phase, the embedded water layer on the stationary phase increases, thus reducing amount of available residual silanols and their ionizability.\textsuperscript{44,50,67} Also, the low metal impurity implies that potential acidity of the uncapped residual silanols is greatly reduced.\textsuperscript{46,50}

4. Conclusions

For the analysis of hallucinogens and any other set of polar compounds that contain a basic alkyl-N and aromatic substituents, the phenyl and HILIC-based columns can be considered in order to maximize the number of analytes for simultaneous quantification. Phenyl RP columns are promising through $\pi$–$\pi$ interactions and could be a first choice for separation depending on the number of compounds, presence of cyclic groups with conjugated $\pi$-electron systems, the class and functional groups of the compounds to be analyzed. Based on studies by other researchers, the pentafluorophenylpropyl stationary phase is also recommended.\textsuperscript{34,35,37} The HILIC is a promising and almost definite choice for separation of polar basic compounds that show separation problems under reverse based phases. The declaration on HILIC separation was based on testing basic alkyl-N hallucinogenic compounds on a fused core silica column. For compounds that are derivatives of a single precursor and where the basic N has either H or alkyl substituents, polarity as given by X log P5 values can be used to predict the elution order. The presence of other functional groups attached to this N atom, as exemplified by the amino acid tryptophan in this study, brings in secondary interactions. Such interactions lead to the collapse of the polarity-based elution predictions. This effect is also observed when other polar biomolecules of a different class or precursor are considered.

Compared to effect of salt concentration, the amount of salt in the mobile phase was more important in fused core silica HILIC retention and the opposite was true for resolution. The reverse was observed for reversed phase chromatography. Our results have shown that in addition to the major parameters deemed essential for HILIC elution optimizations, the injection diluent for analytes also needs to be considered when dealing with polar compounds. The use of relatively non-polar diluents for injection of analytes into a polar mobile phase in HILIC separations was suggested in this research. THF was recommended as the first choice of consideration where acetonitrile or methanol is used as the organic component of the mobile phase.

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References

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