Determination of Phosphodiesterase Type 5 Enzyme (PDE-5) Inhibitors and Analogues as Adulterants in Selected Herbal Products using Gas Chromatography–Electron Impact-Mass Spectrometer (GC-EI-MS)

Nur Azzalia Kamaruzaman*, Mazlin Mohideen2, Yin-Hui Leong1, Azaharudin Awang Ahmad1, Norjuliana Mohd Noor1

1National Poison Centre, Universiti Sains Malaysia, 11800 Penang, Malaysia
2Faculty of Pharmacy and Health Sciences, Universiti Kuala Lumpur-Royal College of Medicine Perak (UniKL-RCMP), 30450 Ipoh, Perak, Malaysia
*Corresponding author: azzalia@usm.my, mazlin.mohideen@unikl.edu.my; yhleong@usm.my; azaharudin@usm.my; julianamnoor@usm.my

ORCID ID: https://orcid.org/0000-0002-6846-1642*, https://orcid.org/0000-0003-3483-7493; https://orcid.org/0000-0001-6767-6425; https://orcid.org/0000-0002-0882-5117; https://orcid.org/0000-0002-1171-1639

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Abstract:
Sildenafil, tadalafil, and vardenafil are phosphodiesterase type 5 enzyme (PDE-5) inhibitors used for the treatment of male erectile dysfunction. This present study aims to investigate 55 herbal products indicated for men’s sexual health from the Malaysian market for adulteration of PDE-5 inhibitors and analogues. The screening and identification of 20 PDE-5 inhibitors and analogues in herbal supplements of various forms (powder, capsules, tablets, and pastels) were conducted using gas chromatography–electron impact-mass spectrometer (GC-EI-MS). The analysis has shown that 19 herbal products were adulterated with PDE-5 inhibitors and analogues. Unique ion fragmentations and the presence of molecular ion serve as potential markers, and the limit of detection (LOD) was 0.1-5.0 µg/g. Ten PDE-5 inhibitors and analogues were randomly selected and successfully validated for simultaneous quantification, whereby the limit of quantitation (LOQ) was 5.0–50.0 µg/g with recoveries at 86.1-118.9%. The developed method of GC-EI-MS was shown to be simple, throughput, economical and had sufficient sensitivity to screen, identify and quantify PDE-5 inhibitors and analogues in herbal matrices.

Key words: Adulteration, Chromatography–electron impact-mass spectrometer (GC-EI-MS), Herbal products, Men’s sexual health, Phosphodiesterase type 5 enzyme (PDE-5) inhibitors.

Introduction:
Adulteration of phosphodiesterase type 5 enzyme (PDE-5) inhibitors in herbal supplements and food products has been a public health concern in many countries around the world (1-4), including Malaysia. For surveillance and monitoring, Control of Drugs and Cosmetics Regulations 1984 was enacted by the Malaysian government to enforce all herbal products that are manufactured, imported and sold in the country to be tested for the presence of adulterants, before registration with the Drug Control Authority of Malaysia (DCA). Unfortunately, even with law enforcement, adulteration of PDE-5 inhibitors in herbal preparations is still prevalent in the Malaysian market (5). As of currently, sildenafil, tadalafil, vardenafil, and udenafil are the only synthetic PDE-5 inhibitors approved by the Ministry of Health Malaysia (MOH) for the treatment of male erectile dysfunction (5). The prescription of PDE-5 inhibitors is fully controlled due to the harmful side-effects from the vasodilation nature of PDE-5 inhibitors on the pulmonary vessels. The use of PDE-5 inhibitors is also contraindicated in patients prescribed with antihypertensive such as nitrates as well as untreated patients with hypotension (6).

Certain herbal products are marketed to be ‘natural’ sexual enhancers for men. Demand for these products is high as the public mistakenly associate them as safer than prescription medicine due to claims of all-natural ingredients (4,7). In
addition, some consumers may feel embarrassed about their health situation and feel more comfortable with the purchase of these herbal products which are easily more accessible rather than consulting with a medical professional (5). To keep up with the demand for ‘natural’ aphrodisiacs and to avoid regulatory inspection, errant manufacturers and suppliers have begun synthesizing PDE-5 inhibitor analogues of sildenafil, tadalafil and vardenafil (5,8). The analogues are synthesized using various chemical modifications of the parent compounds, thus resulting in the unpredictability of potency, side effects, and toxicity (9). Many reports have shown that some commercially available herbal products are adulterated with approved PDE-5 inhibitors as well as unapproved analogues (7,9).

This study aims to investigate the adulteration of 20 PDE-5 inhibitors and analogues in 55 herbal products of various preparations (powder, tablets, capsules, and pastels) from the Malaysian market, between the duration of 1st January 2016 and 11th October 2017 received by the Malaysia National Poison Centre (NPC). Gas chromatograph–electron impact-mass spectrometer (GC-EI-MS) was used to screen and identify these PDE-5 drugs and analogues. The potential use of GC-EI-MS for quantification assay was further confirmed by using ten randomly selected PDE-5 inhibitors and their analogues.

Materials and Methods:
All certified PDE-5 inhibitors were purchased from TLC PharmChem, Canada: Acetylildenafil (98.6%), Acetildenafil (99.8%), Aminotadalafil (99.9%), Carbodenafil (99.0%), Desmethylcarbodenafil (99.7%), Dimethylsildenafil (99.9%), Gendenafil (98.9%), Homosildenafil (99.6%), N-octyl-nortadalafil (99.5%), Nornesildenafil (99.7%), Nortadalafil (100%), N-desmethylsildenafil (99.9%), Piperiacetildenafil (99.1%), Pseudovardenafil (99.8%), Sildenafil (99.8%), Tadalafil (100.0%), Thiosildenafil (98.9%), Thiohomosildenafil (100.0%), Thiodimethylsildenafil (99.9%), Vardenafil (99.5%) and \(d^8\)-sildenafil (99.7%). Other chemicals were purchased from Sigma Aldrich (UK) (6α-Methylprednisolone), Fischer Scientific (UK) (methanol) and James Burrough, FAD-UK (ethanol).

Standard and sample preparations
Screening, and identification
Stock solutions containing 1.0 mg/mL of 20 PDE-5 drugs and their analogues, standard solutions (0.1 mg/ml), mixture of the drugs (40.0 µg/mL), internal standard (IS), \(d^8\)-sildenafil (50 µg/ml), and external standard (ES), 6α-methylprednisolone (0.25 mg/ml) were prepared in methanol. Samples of 0.1 g were dissolved in 0.5 mL methanol through 7 min sonication, followed by 4 min vortex and 3 min centrifugation at 1000 rpm. A volume of 0.25 mL supernatant was added with 25 µL IS and dried over nitrogen gas flow at 40 °C. The dried extract was reconstituted with 50 µL of ES before GC-EI-MS analysis. Validation samples (20 µg/g), each consisting 10 different tablets, capsules, pastels and powdered herbs were prepared by spiking 2 µg drugs in each 0.1g blank herbs and subjected to above preparations. Blank herb matrices were tested according to the procedure, and found negative before being spiked and treated as validation samples.

Quantification
Standard mixtures containing 0.001, 0.010 and 0.100 mg/mL of sildenafil, vardenafil, gendenafil, norneosildenafil, pseudovardenafil, thiosildenafil, thiodimethylsildenafil, acetiildenafil, piperiacetildenafil and nortadalafil were prepared in ethanol. They were used in the daily preparation of the calibration curves from to 1 – 40 µg/mL or equivalent to 5 – 200 µg/g drugs.

Weight of powdered herb sample or validation sample (0.05 g) was added with 25 µL IS and mixed in 0.25 mL ethanol through 7 min sonication, followed by 4 min vortex and 3 min centrifugation at 1000 rpm. A total of 15 validation samples at concentration 1 µg/mL or 5 µg/g (n=5), 10 µg/mL or 50 µg/g (n=5) and 40 µg/mL or 200 µg/g (n=5) were prepared daily in blank herbs. The analysis was carried out for three different days.

GC-EI-MS conditions
The GC-EI-MS system consists of HP6890 GC, HP5973 MS (Agilent Technologies) and fused silica capillary column, HP-5MS (15m x 0.25mm i.d. 0.25 µm film thickness). A 1 µL injection using splitless mode was performed on 300 °C injector port. The carrier gas was helium at a flow rate of 2.1 mL/min. The MS interface, ion source and quadrupole temperatures were set at 280 °C, 230 °C and 150 °C, respectively. The electron ionisation was performed at 70 eV with full scan mode at 30-800 amu. The oven ramping temperature was programmed with the initial temperature at 150 °C, raised at 25 °C /min to 290 °C and raised again at 5 °C/min to 320 °C and held for 4.4 min. For quantification, the initial oven temperature was set at 100 °C, raised at 25 °C /min to 320 °C (held for 15 min).
Results and Discussion:
A total of 55 herbal samples were screened for PDE-5 inhibitors and analogues. Analysis has shown that 19 samples were found to be positive for sildenafil (n=7), tadalafil (n=6), aminotadalafil (n=1), desmethylcarbodenafil (n=1), thiodimethylsildenafil (n=1) and mixtures of sildenafil with either tadalafil (n=2) or aminotadalafil (n=1). Currently, the number of PDE-5 inhibitors and analogues reported in the literature has reached 61 (5). For cost-effective purposes, only 20 PDE-5 inhibitors and analogues were chosen for identification, and they are within the capability of detection by GC-EI-MS.

This study found that ethanol was a better solvent than methanol and isopropanol for the extraction of PDE-5 inhibitors and analogues from herbal matrices. However, recovery of analogues was found to be higher by using lower chain alcohol (methanol>ethanol>isopropanol). Therefore, methanol was chosen as a solvent for screening, while ethanol was employed for quantification. Using each of the 100 µg/mL standard solutions of PDE-5 inhibitors and analogues, the GC-EI-MS spectra were produced and added to our database for screening and identification purposes. The MS fragmentations for 20 PDE-5 inhibitors and analogues are shown in Table 1. The molecular ions of the studied compounds remained intact in all of the drugs. Some existed as base ions, thus facilitating the identification process. The GC-EI-MS assay is capable of screening all 20 drugs with good chromatographic separation within 15 min, as shown by the retention index (RI) and retention time (RT) in Table 1. ES was used in order to monitor the RT shift due to the presence of endogenous substances in herbs.

The chemical structures of sildenafil, vardenafil and their analogues consist of three main moieties i.e. alkyl piperazine, sulfonyle and alkoxy-pyrazolopyrimidine (sildenafil) or alkoxy-imidazolotriazine (vardenafil) (Fig.1).

![Figure 1. Structures of: (a) Sildenafil (b) Vardenafil and (c) Tadalafil](image)

Among PDE-5 inhibitors, analogue tadalafil is known to have stereoisomers which are identified using chiral LC column (10). In this developed GC-EI-MS assay, chromatographic separation of the high purity of tadalafil, aminotadalafil and n-octyl-nortadalafil produced two adjacent peaks with similar mass spectrum (Table 1). These stereoisomers were probably derived from GC high injection temperature. The tadalafil analogues contain functional groups; alkyl and amino bonded at N-2 position in piperazinedione ring. No isomer peak was seen for nortadalafil which has no functional group at N-2 position in piperazinedione ring. Tadalafil and its analogues exhibited unique MS fragmentations with high intensity (base ion) of the molecular ions and subsequent loss of benzodioxole ring (-m/z 121). Other characteristic ions are shown in Table 1. MS fragmentation for other analogues was also described in Table 1.

In quantitation analysis, the limit of detection (LOD) of each drug was found to be between 0.1-1.0 µg/g except for aminotadalafil (4.7
µg/g) and N-desethylsildenafil (1.6 µg/g) in 10 different herbal preparations of powder, capsules, tablets and pastels. Generally, the sensitivity achieved for sildenafil and vardenafil were better than the previously reported GC-MS data (11). Currently, the doses for the intended pharmacological effect are only available for sildenafil (25-100 mg), tadalafil (10-20 mg) and vardenafil (5-20 mg) (12). To the best of our knowledge, no recommended dose is available for the analogues since they are banned substances and their pharmacokinetic effect and safety profile have yet to be determined. For the estimated maximum amount of serving size, 0.3 g (pills, tablets, capsules, pastels) and 20 g powdered herb samples; it may contain at least 1.25-83.33 mg/g sildenafil, 0.50-33.33 mg/g tadalafil and 0.25-16.67 mg/g vardenafil. The current developed GC-EI-MS assay has very low LODs (0.0001-0.0050 mg/g). Hence, the assay has sufficient sensitivity to screen PDE-5 inhibitors and their analogues in herbal matrices for adulteration screening.

Table 1. Retention time, retention index and mass fragmentations of the PDE-5 enzyme inhibitors and analogues analysed by GC-EI-MS

| No | Drugs            | Chemical Formulae | Retention Index ± 30 | RT ± 0.5 min | M+ (m/z)     | Base ion (m/z) | Other major ions (m/z) |
|----|------------------|-------------------|----------------------|--------------|--------------|---------------|----------------------|
| 1  | Gendenafil       | C₁₉H₂₂N₂O₅       | 3096                 | 5.66         | 354 (100%)   | 354           | 339 (47%), 336 (7%), 326 (79%), 321 (22%), 312 (4%), 311 (18%), 297 (15%), 283 (19%), 282 (27%), 193 (11%), 166 (28%), 162 (6%), 136 (46%), 43 (16%) |
| 2  | Piperacutedilenafil| C₂₄H₃₃N₂O₅      | 3661                 | 7.77         | 437 (1%)     | 98            | 312 (1%), 311 (2%), 297 (1%), 283 (1%), 282 (1%), 166 (1%), 136 (2%), 70 (3%), 55 (4%), 42 (4%), 30 (1%) |
| 3  | Desmethylcarbodenafil | C₂₅H₂₃N₂O₅  | 3847                 | 8.75         | 438 (10%)    | 70            | 400 (2%), 381 (7%), 368 (7%), 355 (6%), 339 (7%), 312 (1%), 311 (6%), 283 (1%), 282 (1%), 166 (3%), 136 (3%), 99 (15%), 83 (29%), 56 (21), 43 (10%), 30 (2%) |
| 4  | Tadalafil        | C₂₂H₃₃N₂O₄       | 3838                 | 8.7          | 389 (100%)   | 389           | 318 (5%), 317 (8%), 289 (12%), 275 (6%), 268 (20%), 263 (45%), 262 (68%), 254 (2%), 233 (20%), 204 (33%), 169 (21%), 115 (8%), 102 (12%), 44 (7%) |
| 5  | Acetildenafil    | C₂₃H₃₅N₂O₅       | 3826                 | 8.63         | 466 (4%)     | 127           | 312 (1%), 311 (1%), 283 (1%), 282 (0.6%), 166 (1%), 136 (2%), 112 (4%), 98 (3%), 84 (16%), 70 (17%), 56 (6), 42 (7%), 30 (1%) |
| 6  | Acetylvardenafil | C₂₃H₃₆N₂O₃       | 3859                 | 8.83         | 466 (3%)     | 127           | 326 (1%), 312 (1%), 311 (1%), 283 (1%), 282 (1%), 133 (1%), 112 (2%), 98 (3%), 84 (14%), 70 (15%), 56 (4), 42 (6%) |
| 7  | Carbodenafil     | C₂₄H₃₄N₂O₃       | 3882                 | 9.07         | 452 (9%)     | 84            | 381 (10%), 339 (9%), 312 (3%), 311 (8%), 283 (1%), 282 (1%), 281 (2%), 166 (3%), 136 (4%), 113 (8%), 97 (21%), 70 (13%), 56 (14%), 42 (10%), 30 (1%) |
| 8  | Pseudovardenafil| C₂₃H₂₃N₂O₅S      | 3931                 | 9.26         | 459 (54%)    | 431           | 444 (14%), 416 (3%), 312 (4%), 311 (5%), 296 (3%), 284 (15%), 283 (15%), 282 (10%), 254 (5%), 215 (5%), 166 (3%), 136 (2%), 135 (6%), 94 (7%), 84 (14%), 67 (17%), 55 (11%), 42 (18%), 30 (2%) |
| 9  | Aminotadalafil  | C₂₃H₁₈N₂O₄       | 3950                 | 9.38         | 390 (100%)   | 390           | 374 (14%), 331 (8%), 318 (11%), 317 (9%), 289 (38%), 274 (5%), 269 (10%), 263 (33%), 262 (51%), 252 (4%), 233 (17%), 204 (36%), 169 (18%), 115 (12%), 102 (12%), 44 (4%) |
| 10 | Sildenafil       | C₂₃H₈N₂O₅S       | 3955                 | 9.41         | 474 (1%)     | 99            | 404 (33%), 312 (3%), 311 (3%), 283 (3%), 282 (1%), 254 (1%), 166 (1%), 136 (2%), 84 (1%), 70 |
| Compound            | Chemical Formula | C: H: N: O: S | Molecular Weight | MS Detection |
|---------------------|------------------|--------------|-----------------|--------------|
| 11 Dimethylsildenafil | C23H32N6O4S      | 3959         | 488 (0.3%)      | 9.43         |
| 12 Nortadalafil     | C23H32N6O4S      | 4008         | 375 (100%)      | 9.71         |
| 13 Homosildenafil    | C23H32N6O4S      | 4074         | 488 (1%)        | 10.15        |
| 14 Vardenafil       | C23H32N6O4S      | 4105         | 488 (6%)        | 10.36        |
| 15 Thiosildenafil    | C23H32N6O4S      | 4268         | 490 (5%)        | 11.5         |
| 16 N-desmethylsildenafil | C23H32N6O4S  | 4111         | 460 (2%)        | 10.4         |
| 17 Thiodimethylsildenafil | C23H32N6O4S  | 4282         | 504 (2%)        | 11.6         |
| 18 Thiohomosildenafil | C23H32N6O4S   | 4274         | 504 (6%)        | 11.54        |
| 19 N-octyl-nortadalafil | C23H32N6O4S  | 4435         | 487 (100%)      | 12.83        |
| 20 Norneosildenafil  | C22H29N5O4S      | 4619         | 459 (100%)      | 14.75        |

Even though GC-EI-MS is useful for qualitative assay, it may not be suitable for quantification of specific PDE-5 inhibitors and their analogues. Formation of diastereoisomers of tadalafil and its amino- or alkyl- tadalafil limit the assay for screening and identification only. Degradation of parent compounds may occur in certain drugs analysed by GC-EI-MS. Extensive degradation may lead to poor recovery or high quantification limit of the parent drugs. It was observed that sildenafil, vardenafil and their analogues degraded into multiple related derivatives at a high concentration (1 mg/mL) as seen in Table 2. Extensive degradation was recorded among thioke-toke analogues as only 15-30% of parent drugs were detected. Approximately 76% of N-desmethysildenafil was detected. The majority (80%) of PDE-5 inhibitors and their analogues had minor degradations (0-14%). Nevertheless, no degradation was observed at a concentration of 0.1 mg/mL standard solutions. Thus GC-EI-MS had potential application in the quantification of these drugs in diluted samples.
Validation was successfully performed using ten randomly selected PDE-5 inhibitors and analogues as shown in Table 3. Calibration curves for acetildenafil, piperiacetildenafil and nortadalafil were plotted from 10 to 40 µg/mL (or 50 to 200 µg/g). The calibration range for other drugs was plotted from a lower point, i.e. 1 µg/mL or 5 µg/g. The calibration curves were linear with an average correlation coefficient, R² ≥ 0.996 (Table 3). The within- and between-assays precision and accuracy of all drugs were within the acceptable range with an excellent recovery at 91.28-118.85 %. This data proved that PDE-5 inhibitors and analogues could be quantified in herbal preparations by GC-EI-MS. More than 85% thio ketone analogues (thiosildenafil and thiodimethylsildenafil) was recovered at a concentration of 5 µg/g. However, low recovery (49-61%) was recorded at higher concentrations which may be due to the degradation in matrices and the use of a weak organic solvent (ethanol). Nevertheless, a study had reported a recovery of ≥ 73.6 % for 38 PDE-5 inhibitors and analogues using LC-MS/MS assay (12). Limit of quantitation for the assay was 1 µg/mL or 5 µg/g for all drugs except acetildenafil, piperiacetildenafil and nortadalafil (10 µg/g).

The validated GC-EI-MS assay was successfully tested on real samples, qualitatively and quantitatively. However, GC-EI-MS may not be able to replace or provide for an alternative technique to other more superior techniques such as LC-MS (13,14) and LC-MS/MS (15,16), its simplicity, sensitivity, selectivity and economical are sufficient for compound identification and as a complementary technique to support the findings of other conventional methods such as HPLC (15), ELISA (17) and thin-layer chromatography (18).

A Malaysian study conducted between 2014-2016 which screened 61 PDE-5 inhibitors and analogues showed that 82% of unregistered herbal and food products were adulterated with at least one PDE-5 inhibitor or analogue (5). In this study, 35% of herbal products which were dispatched to the NPC from 2016 to 2017 were proven to contain at least one PDE-5 inhibitor or analogue (5). In this study, 35% of herbal products which were dispatched to the NPC from 2016 to 2017 were proven to contain at least one PDE-5 inhibitor or analogue (5). In this study, 35% of herbal products which were dispatched to the NPC from 2016 to 2017 were proven to contain at least one PDE-5 inhibitor or analogue (5). In this study, 35% of herbal products which were dispatched to the NPC from 2016 to 2017 were proven to contain at least one PDE-5 inhibitor or analogue (5). In this study, 35% of herbal products which were dispatched to the NPC from 2016 to 2017 were proven to contain at least one PDE-5 inhibitor or analogue (5). In this study, 35% of herbal products which were dispatched to the NPC from 2016 to 2017 were proven to contain at least one PDE-5 inhibitor or analogue (5). In this study, 35% of herbal products which were dispatched to the NPC from 2016 to 2017 were proven to contain at least one PDE-5 inhibitor or analogue (5). In this study, 35% of herbal products which were dispatched to the NPC from 2016 to 2017 were proven to contain at least one PDE-5 inhibitor or analogue (5). In this study, 35% of herbal products which were dispatched to the NPC from 2016 to 2017 were proven to contain at least one PDE-5 inhibitor or analogue (5).
Table 3. The within assay, between assay precisions and accuracies and recovery of spiked 1, 10, 40 μg/mL (or 5, 50, 200 μg/g) PDE-5 inhibitors and analogues analysed by GC-EI-MS

| Analyte                  | Curve (μg/mL) | R²  | Nominal (μg/mL) | Recovery (%) | Observed (mean ± SD) (μg/mL) | Precision (%) | Accuracy (%) | Observed (mean ± SD) (μg/mL) | Precision (%) | Accuracy (%) |
|--------------------------|--------------|-----|-----------------|-------------|------------------------------|--------------|-------------|------------------------------|--------------|-------------|
| Gendenaﬁl               | 1-40         | 0.998 | 118.85          | 100.0       | 1.034 ± 0.071                | 6.83         | 3.38        | 1.086 ± 0.045                | 8.32         | 5.87        |
|                          |              | 10   | 105.04          | 96.57       | 9.022 ± 1.136                | 12.57        | 9.78        | 9.320 ± 0.635                | 11.19        | 7.13        |
|                          |              | 40   | 92.23           | 89.26       | 40.748 ± 5.625               | 13.80        | 1.87        | 40.728 ± 3.248               | 8.12         | 7.08        |
| Poperiacetildenafil      | 10-40        | 0.998 | 105.32          | 100.0       | 8.563 ± 1.111                | 12.98        | 14.37       | 9.206 ± 0.561                | 6.03         | 7.94        |
|                          |              | 40   | 92.14           | 90.0        | 37.110 ± 4.764               | 12.80        | 7.20        | 39.116 ± 2.747               | 7.02         | 5.95        |
| Acetildenafil            | 10-40        | 0.997 | 92.77           | 100.0       | 10.367 ± 1.070               | 10.37        | 3.67        | 10.337 ± 1.270               | 12.31        | 4.95        |
|                          |              | 40   | 91.28           | 97.04       | 38.921 ± 5.416               | 13.92        | 2.70        | 38.333 ± 3.418               | 9.02         | 7.08        |
|                          |              | 1    | 92.18           | 100.0       | 1.115 ± 0.130                | 11.68        | 11.46       | 0.988 ± 0.116                | 13.20        | 8.84        |
| Noreosildenafil          | 10-40        | 0.997 | 98.35           | 100.0       | 10.105 ± 0.876               | 8.67         | 1.05        | 9.777 ± 0.350                | 7.65         | 2.93        |
|                          |              | 40   | 102.18          | 100.0       | 39.763 ± 2.150               | 5.41         | 0.59        | 40.185 ± 0.368               | 7.53         | 8.86        |
|                          |              | 1    | 111.73          | 100.0       | 1.054 ± 0.011                | 1.03         | 5.46        | 1.050 ± 0.060                | 5.75         | 9.29        |
| Pseudovardenafil        | 10-40        | 0.999 | 91.95           | 100.0       | 10.173 ± 0.938               | 9.23         | 1.73        | 10.054 ± 0.726               | 7.15         | 2.76        |
|                          |              | 40   | 96.39           | 99.71       | 38.413 ± 2.877               | 7.49         | 3.97        | 39.236 ± 3.282               | 8.36         | 1.91        |
| Sildenafil               | 10-40        | 0.996 | 103.63          | 100.0       | 9.702 ± 0.970                | 10.00        | 2.93        | 10.287 ± 0.579               | 6.57         | 4.86        |
|                          |              | 40   | 99.47           | 100.0       | 39.579 ± 2.148               | 5.43         | 1.05        | 39.345 ± 0.941               | 5.18         | 1.64        |
| Nortadalafil            | 10-40        | 0.997 | 114.06          | 100.0       | 8.846 ± 0.839                | 9.48         | 11.54       | 9.311 ± 0.410                | 4.41         | 6.87        |
|                          |              | 40   | 96.54           | 100.0       | 38.228 ± 3.610               | 9.44         | 4.43        | 38.564 ± 0.408               | 1.06         | 3.59        |
|                          |              | 1    | 110.53          | 100.0       | 1.009 ± 0.063                | 6.21         | 0.89        | 1.100 ± 0.080                | 8.55         | 10.02       |
| Vardenafil              | 1-10         | 0.998 | 105.60          | 100.0       | 10.002 ± 0.808               | 8.08         | 0.02        | 10.035 ± 1.037               | 9.74         | 0.35        |
|                          |              | 40   | 103.99          | 100.0       | 40.112 ± 1.124               | 2.80         | 0.28        | 38.192 ± 2.516               | 6.57         | 4.52        |
| Thiosildenafil          | 10-40        | 0.999 | 87.12           | 100.0       | 1.121 ± 0.110                | 9.82         | 12.12       | 1.061 ± 0.139                | 14.95        | 12.64       |
|                          |              | 40   | 76.45           | 76.45       | 8.526 ± 1.166                | 13.67        | 14.74       | 8.926 ± 0.914                | 12.57        | 10.74       |
| Thiodimethylsildenafil  | 1-40         | 0.998 | 66.70           | 100.0       | 8.850 ± 1.174                | 13.26        | 11.50       | 9.064 ± 0.340                | 10.41        | 9.36        |
|                          |              | 40   | 48.69           | 100.0       | 35.634 ± 4.458               | 12.51        | 10.91       | 36.934 ± 1.802               | 8.57         | 7.67        |

a = 5 replicates, b = 15 replicates

Conclusion:
GC-EI-MS assay was successfully developed and tested on real samples, qualitatively and quantitatively, and it has been proven to be simple, sensitive, selective and economical for the screening and identification of PDE-5 inhibitors and their analogues. From 55 samples received by the NPC between 1st January 2016 and 11th October 2017, 19 samples were found to be adulterated with at least one PDE-5 inhibitor or analogue. Effective strategy must be employed to reduce the alarming increase of adulteration of PDE-5 inhibitors and analogues in Malaysian herbal products, through regular monitoring and surveillance, thus ensuring product quality and safety. In addition, the public should be educated and informed about the risks of using adulterated and tainted products which are claimed to be safe and “natural”.

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تحديد مثبطات ونظائر إنزيم الفوسفودستراز من النوع 5 (PDE-5) باستخدام الكروماتوغرافيا الكهربائية - مطابق تأثير الكثافة الإلكترونية (GC-EI-MS)

نواف عزاليا قمر الزمان1، 2
مازلين محي الدين2
أهراز الدين أوانج أحمد1
تأليف: محمد نور1

1. مركز السموم الوطني ، جامعة سنير ماليزيا ، 11800 بينانغ ، ماليزيا
2. كلية الصيدلة والعلوم الصحية ، جامعة كوالالمبور - الكلية الملكية للطب بيراك ، ماليزيا

(ر)10050 UniKL-RCMP

الخلاصة:

هذه مثبطات إنزيم الفوسفودستراز 5 (PDE-5) وسافنادينول وتادالافينول وسيلة الإنزيم PDE-5، وهو المثبطات الشائعة في المنتجات العشبية التي يتم استخدامها في حالات ضعف الانتصاب عند الذكور. تهدف الدراسة الحالية إلى تحديد وتحديد 20 مثبط يابسة ونظائر PDE-5 لشوكي جزئي لا يمكن تمييزه من الأنواع المضادة للكثير من الأنواع في المنتجات العشبية. تستخدم الكروماتوغرافيا الكهربائية - مطابق تأثير الكثافة الإلكترونية (GC-EI-MS) لتحديد وتحديد 19 منجأ بصارم. يتم تحديد تشخيص النباتات ونظائر PDE-5 بشكل مشترك مع تحديد تشخيص النباتات ونظائر PDE-5. وظيفة إجمالية مفيدة في التحكم في تشخيص النباتات ونظائر PDE-5 لتشخيص التشخيص. وتستخدم الكروماتوغرافيا الكهربائية - مطابق تأثير الكثافة الإلكترونية (GC-EI-MS) لتحديد وتحديد 19 منجأ بصارم. يتم تحديد تشخيص النباتات ونظائر PDE-5 بشكل مشترك مع تحديد تشخيص النباتات ونظائر PDE-5. وظيفة إجمالية مفيدة في التحكم في تشخيص النباتات ونظائر PDE-5 لتشخيص التشخيص.

الكلمات المفتاحية: النبات، الصحافة الجنسية للرجال، مثبطات إنزيم الفوسفودستراز 5 (PDE-5)، المنتجات العشبية، مطابق تأثير الكثافة الإلكترونية