

# Metabolomics-Guided Isolation of Anti-trypanosomal Metabolites from the Endophytic Fungus *Lasiodiplodia theobromae*

## Authors

Nurkhalida Kamal<sup>1,2</sup>, Christina V. Viegelmann<sup>1</sup>,  
Carol J. Clements<sup>1</sup>, RuAngelie Edrada-Ebel<sup>1</sup>

## Affiliations

- 1 Strathclyde Institute of Pharmacy and Biomedical Sciences, University of Strathclyde, Glasgow, United Kingdom
- 2 Department of Pharmaceutical Sciences, Faculty of Pharmacy, Cyberjaya University College of Medical Sciences (CUCMS), Cyberjaya, Selangor, Malaysia

## Key words

fungal endophytes, *Lasiodiplodia theobromae*, Botryosphaeriaceae, anti-trypanosomal activity, HR-LCMS, NMR, metabolomics, chemometrics

received March 9, 2016

revised Sept. 19, 2016

accepted Sept. 28, 2016

## Bibliography

DOI <http://dx.doi.org/10.1055/s-0042-118601>


Published online October 19, 2016 | *Planta Med* 2017; 83: 565–573 © Georg Thieme Verlag KG Stuttgart · New York | ISSN 0032-0943

## Correspondence

RuAngelie Edrada-Ebel  
Strathclyde Institute of Pharmacy and Biomedical Sciences,  
University of Strathclyde  
161 Cathedral Street, Glasgow G4 0RE, United Kingdom  
Phone: + 44 14 15 48 59 68, Fax: + 44 14 15 52 25 62  
[ruangelie.edrada-ebel@strath.ac.uk](mailto:ruangelie.edrada-ebel@strath.ac.uk)

## Correspondence

Nurkhalida Kamal  
Department of Pharmaceutical Sciences, Faculty of Pharmacy,  
Cyberjaya University College of Medical Sciences (CUCMS),  
No 3410  
Jalan Teknokrat 3, Cyber 4, 63000 Cyberjaya, Selangor,  
Malaysia  
Phone: + 60 3 83 13 73 24, Fax: + 60 3 83 13 70 01  
[nurkhalida@cybermed.edu.my](mailto:nurkhalida@cybermed.edu.my)

 Supporting information available online at  
<http://www.thieme-connect.de/products>

## ABSTRACT

Fungal endophytes offer diverse and unique secondary metabolites, making these organisms potential sources of promising drug leads. The application of high-resolution-liquid chromatography mass spectrometry and nuclear magnetic resonance-based metabolomics to fungal endophytes is practical in terms of dereplication studies and the mining of bioactive compounds. In this paper, we report the application of metabolomics in parallel with anti-trypanosomal assays to determine the ideal conditions for the medium-scale fermentation of the endophyte *Lasiodiplodia theobromae*. The <sup>1</sup>H NMR comparison between the active versus inactive fractions identified several unique chemical fingerprints belonging to the active fractions. Furthermore, by integrating high-resolution-liquid chromatography mass spectrometry data with multivariate data analysis, such as orthogonal partial least squares-discriminant analysis (OPLS-DA) and the bioactivity results of the fractions of *L. theobromae*, the anti-trypanosomal agents were easily discerned. With available databases such as Antibase and Dictionary of Natural Products coupled to MZmine through in-house algorithms optimized in our laboratory, the predicted metabolites were readily identified prior to isolation. Fractionation was performed on the active fractions and three known compounds were isolated, namely, cladospirone B, desmethyl-lasiodiplodin, and *R*-(-)-mellein. Cladospirone B and desmethyl-lasiodiplodin were among the predicted compounds generated by the OPLS-DA S-plot, and these compounds exhibited good activity against *Trypanosoma brucei brucei* with minimum inhibitory concentrations of 17.8 μM and 22.5 μM, respectively.

## Introduction

Despite the increasing disinclination of the pharmaceutical industry to pursue natural products in their pipelines, statistical findings show that natural products still play a major role in drug discovery with more than 50% of FDA-approved drugs were derived from natural products [1]. However, natural products research has been found to be too laborious, time-consuming, and uneconomical, which may have led to the declining trend. Nevertheless, with the emergence of new and more advanced technologies such as genomics, transcriptomics, proteomics, metabolomics, and bioinformatics, natural products research has become more competent in finding promising novel drugs for the pipeline [1–4].

The utilization of metabolomics in natural products research is increasingly powerful in several perspectives. Metabolomics is defined as a global study of all or a subset of chemical entities including either or both primary and secondary metabolites that are present in living organisms (cells or tissues) under certain processes [5–7]. Metabolomics, or metabolome mining, in natural products research has been used for dereplication studies of both known and new compounds in crude plant, marine, or microbial extracts [8–18], in differentiating biologically active natural products (NPs) from non-active fractions [19–23], optimizing the production of bioactive secondary metabolites, as well as in developing cultivation processes for large-scale fermentation and understanding their biosynthetic pathways [7, 24].

Endophytic fungi are microorganisms that mutually live inside plant tissues without causing any immediate negative effects towards the host plant for at least a part of the fungal life cycle [25]. The total population of endophytic fungi species has been estimated to be up to 1.3 million [26]. Another study places this estimate at approximately 1.5 million [27]. However, as of the year 2000, only 75 000 fungal species have been identified; the remainder are still untapped and unexplored [28]. Bioactive NPs derived from endophytic fungi display wide ranges of activities, such as (–)-oxysporidinone, (2,6-dihydroxyphenyl)pentan-1-one, and (Z)-1-(2-(2-butyl-3-hydroxyphenoxy)-6-hydroxyphenyl)-3-hydroxybut-2-en-1-one, which exhibited antimicrobial activity [29, 30], pullularin A and hinnuliquinone displayed antiviral activity [31, 32], and spiropreussione A and 9-deacetoxyfumigaclavine C showed anticancer activity [33, 34]. Compounds like cochlioquinone A, isocochlioquinone A, and cercosporin exhibited activity against neglected tropical diseases [35, 36].

Human African trypanosomiasis, or sleeping sickness, is a fatal vector-borne parasitic disease caused by *Trypanosoma brucei brucei* transmitted by the tsetse fly (*Glossina* spp.). This neglected tropical disease occurs only in rural areas of sub-Saharan Africa [37]. To date, only a few drugs have been approved for the treatment of human African trypanosomiasis. These include suramin, pentamidine, melarsoprol, eflornithine, and the combination of nifurtomox/eflornithine. Most of the drugs are old, having been discovered in the 1940s and 1950s, and have adverse effects such as nausea, vomiting, fatigue, seizures, fever, diarrhea, hypoglycemia, abdominal cramping, peripheral neuropathy, hypertension, heart damage, and neutropenia on the patients. For this reason,

► **Table 1** Anti-trypanosomal activity of *L. theobromae* extracts derived from *V. pinnata* in different types of media and incubation periods. MIC was only determined for the bioactive extracts.

Sample	<i>T. b. brucei</i> 20 µg/mL % of viability	<i>T. b. brucei</i> MIC average ± SD (n = 4)
LT-LC-7	112.3	Not tested
LT-LC-15	106.3	Not tested
LT-LC-30	126.0	Not tested
LT-RC-7	103.9	Not tested
LT-RC-15	94.3	Not tested
LT-RC-30	1.4	25 ± 1.3 µg/mL
Suramin	Not tested	0.1 ± 0 µM

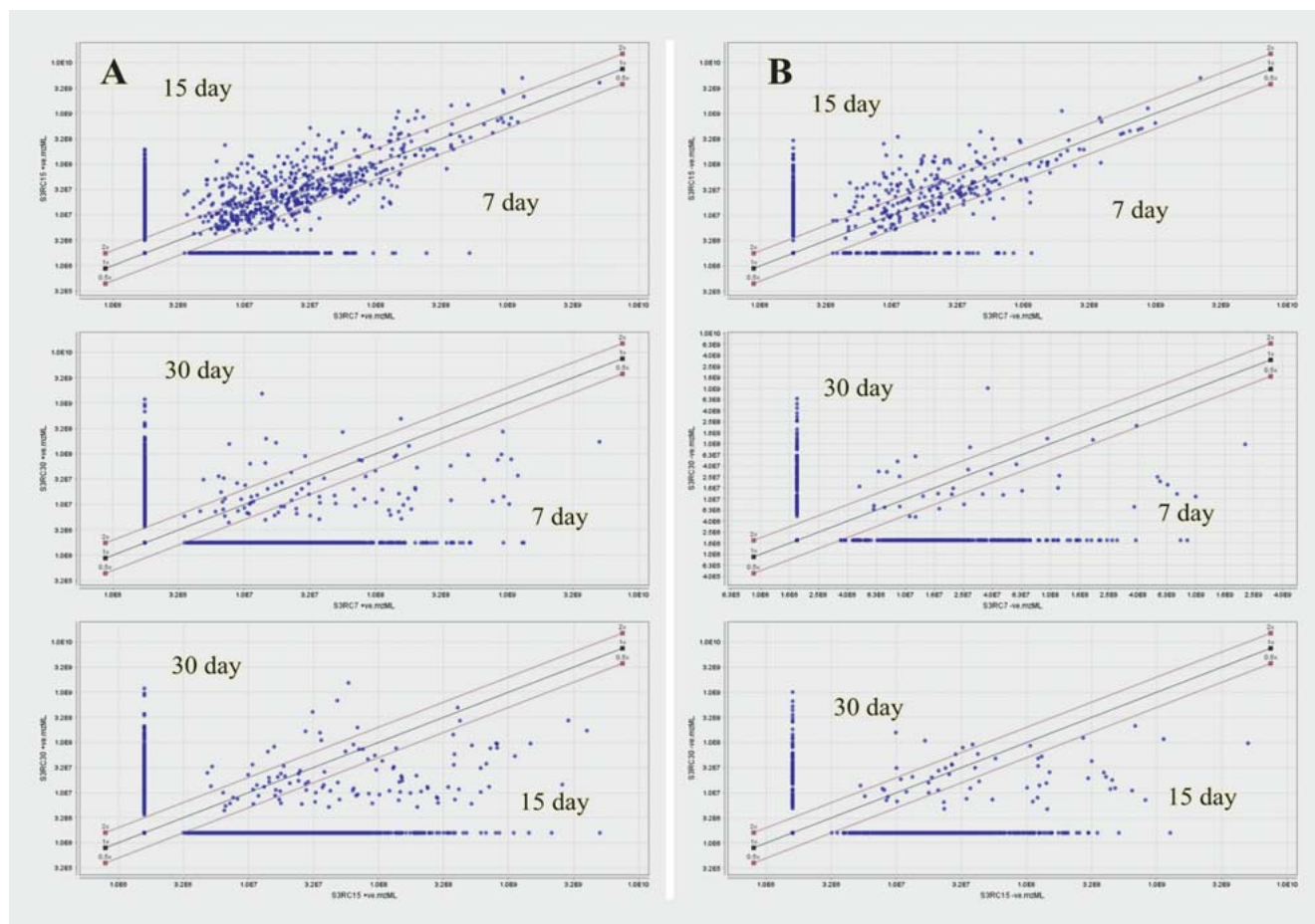
LT = *L. theobromae*.

mining and developing new human African trypanosomiasis drugs from natural products is crucial and essential because various natural sources including plants, microorganisms, animals, and marine organisms offer a high number of NPs with diverse chemical structures and novel pharmacological mechanism of action [38].

The aim of this study is to adopt an untargeted HR-LCMS and NMR-based metabolomics approach to determine the optimal fermentation conditions of *Lasiodiplodia theobromae* for medium scale-up, and also to capture and trace the production of active anti-trypanosomal metabolites by using statistical multivariate data analysis of generated HR-LCMS data, such as principal component analysis (PCA) and orthogonal partial least squares-discriminant analysis (OPLS-DA). To verify the dereplication results obtained from HR-LCMS data, 1D and 2D <sup>1</sup>H NMR data were utilized. Finally, the isolation of active metabolites was performed based on the outcome of HR-LCMS and NMR-based metabolomics profile data.

## Results and Discussion

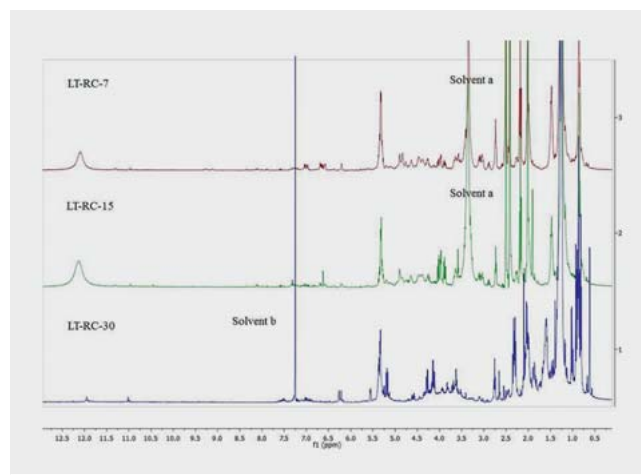
In search of the best condition for scaling up the culture of the endophyte *L. theobromae* obtained from the leaves of *Vitex pinnata*, HR-LCMS and NMR-based metabolomics along with the bioassay data were utilized. The fungus was grown in solid rice cultures and liquid Wickerham cultures for 7, 15, and 30 days, after which the metabolites were extracted. Three different incubation times were chosen based on the fungal life cycle [39]. In this case, the first 7 days represent the germination phase, while the 15-day and 30-day cultures cover the hyphal growth stage and sporing phase, respectively, of *L. theobromae*. The production of secondary metabolites was monitored using HR-LCMS and NMR at each of the growth phases parallel to the bioassay results. Based on the bioassay results (► **Table 1**), the 30-day rice culture extract exhibited the strongest activity against *T. b. brucei* with a minimum inhibitory concentration (MIC) of less than 25 µg/mL. The HR-LCMS raw data was processed using MZMine 2.10 [40]. The results of the assay were reproducible between the scale-up batches.



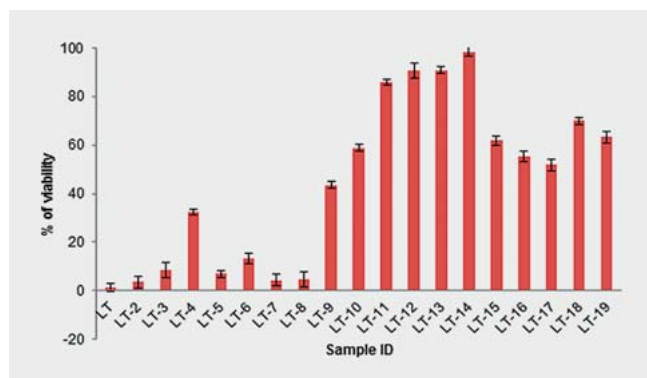
► **Fig. 1** Scatter plot of the ion peaks of the *L. theobromae* extracts from different days (A) in positive ionization and (B) in negative ionization.

Metabolite production and distribution between cultures were analyzed through ion peak scatter plots (► **Fig. 1**). Based on the MS data, the occurrences of the metabolites on the 7th and 15th days were similar, while a decrease in metabolite production was observed on the 30th day. However, the ion chromatogram both in positive and negative modes revealed a different set of metabolites on the 30th day to those of the 7th and 15th day extracts (► **Fig. 1**). Moreover, the  $^1\text{H}$  NMR data revealed findings complementary to the MS data (► **Fig. 2**). Therefore, the bioactive 30-day rice culture condition was chosen for scale-up and further isolation work.

The medium-scale 30-day rice culture extract of *L. theobromae* was fractionated yielding 19 fractions (LT-1 until LT-19). These were submitted for the anti-trypanosomal activity and subjected to MS- $^1\text{H}$  NMR-based metabolomics data profiling. The nonpolar fractions LT-2 to LT-8 exhibited strong bioactivity, except for LT-4, which showed only moderate activity against *T. b. brucei* (► **Fig. 3**). LT-1 was excluded from the bioassay screening because it contained only fatty acids, as indicated by its  $^1\text{H}$  NMR data (► **Fig. 4**). The  $^1\text{H}$  NMR data of the 19 fractions (► **Fig. 4**) were analyzed, and unique chemical fingerprints of the active fractions were detected. Among these active fractions, two distinctive sub-groups, LT-2 to LT-4 and LT-6 to LT-8, clustered together, as they



► **Fig. 2** The  $^1\text{H}$  NMR data of *L. theobromae* extracts obtained from solid rice culture at three different incubation periods (solvent a is  $\text{DMSO-}d_6$ ; b is chloroform- $d$ ). The 30-day rice culture extract could only be fully dissolved in chloroform, indicating the compounds occurring in this extract are semi-nonpolar.



► **Fig. 3** Anti-trypanosomal activity of *L. theobromae* fractions against *T. b. brucei*. LT: *L. theobromae* extract as the positive control; LT2-LT19: *L. theobromae* fractions.

shared similar spectral data. Fraction LT-5 was a mixture of both groups. The  $^1\text{H}$  NMR spectra of fractions LT-2 to LT-5 displayed a pair of *meta*-coupled aromatic protons at  $\delta_{\text{H}}$  6.23 and 6.28 ( $J = 2.7$  Hz) and a multiplet peak at  $\delta_{\text{H}}$  5.15, which may be an olefinic or oxygenated methine. The upfield shift of the *meta*-coupled aromatic protons at the 6 ppm region suggested the presence of an electron withdrawing group such as a hydroxyl or halogen substituent. In fractions LT-5 to LT-8, two *meta*-coupled doublets at  $\delta_{\text{H}}$  6.18 and 6.25 ( $J = 2.6$  Hz) were observed (designated by red arrows on ► **Fig. 4**). The proton signals between 4.2–4.8 ppm revealed the presence of oxygenated methines while proton signals between 6.6 to 7.7 ppm indicated the presence of aromatic compounds in fractions LT-6 to LT-8.

Supervised methods of multivariate data analysis were used to analyze the similarity and differences of the data sets between 19 samples. PCA was used in an earlier step to observe an overview of variance between the fractions and metabolites generated from MS data and also to identify any outliers. The distribution difference of the type of metabolites between active vs. inactive fractions of *L. theobromae* against *T. b. brucei* was analyzed by subjecting the data to OPLS-DA. The results of the analysis led to the prediction of compounds that contribute towards the anti-trypanosomal activity of the fractions. For the OPLS-DA model (► **Fig. 5A**), the MS-based metabolomics data set was assigned as the X independent variable, while the fractions' anti-trypanosomal response was the Y dependent variable. The quality of the OPLS-DA model was measured by two parameters, R<sup>2</sup> (goodness of fit) and Q<sup>2</sup> (predictability), and the model's R<sup>2</sup>X was 0.261, R<sup>2</sup>Y was 0.955, and Q<sup>2</sup> was 0.733. These results were interpreted that 26.1% of the X variables could be used to describe 95.5% of the variation between active fractions and inactive fractions, while this model has 73.3% of the average predictability. The value of R<sup>2</sup>Y and Q<sup>2</sup> was greater than 50%, indicating a well-fitted model exhibiting good prediction [41]. The quality and robustness of the OPLS-DA model was validated by a permutation test ( $n = 100$ ). The Q<sup>2</sup> intercept value was  $-0.368$  (below 0.05), showing that the original model is statistically effective (► **Fig. 5B**) [42]. For the OPLS-DA scores plot (► **Fig. 5A**), the active fractions were grouped together versus the inactive ones. Under the active

group, fractions LT-2 to LT-8 were clustered together, indicating a shared set of metabolites, while fractions LT-9 to LT-19 were observed as the inactive group. The generated S-plot (► **Fig. 5D**) determined the “end point” or unique compounds for each of the respective groups, indicating the metabolites that are potentially responsible for the bioactivity against *T. b. brucei*, which discriminated the active from the inactive fractions of *L. theobromae*. Eight metabolites were identified from Antibase as shown in ► **Table 2**. The end point compounds were targeted for bioassay-guided isolation work (► **Fig. 6**).

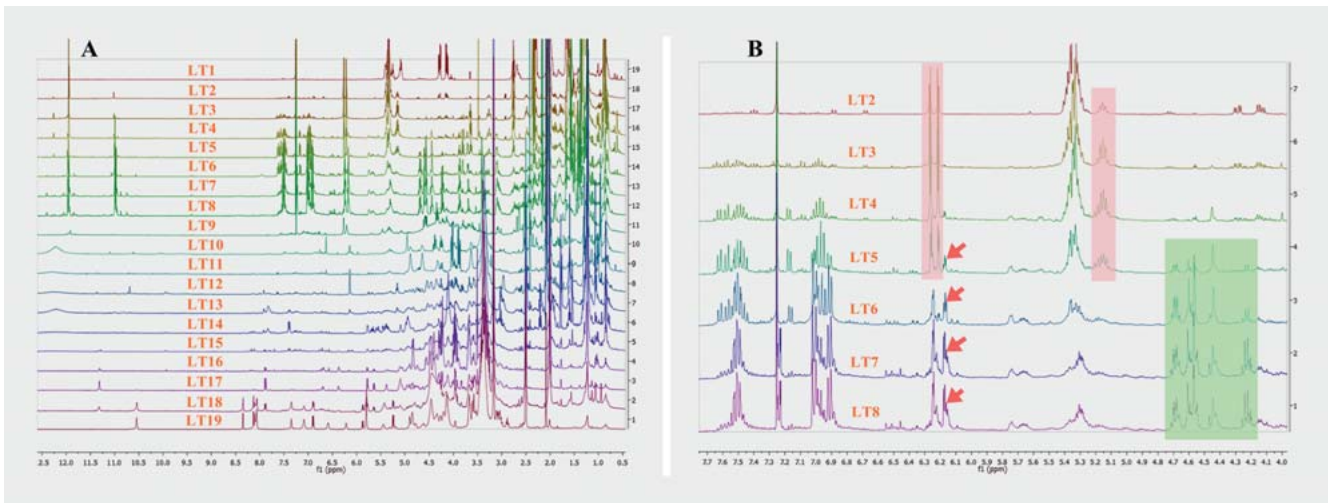
The active metabolites listed from the dereplication step of the HR-LCMS data were identified as palmarumycin CP17 (5), cladospirone B (6), and desmethyl-lasiodiplodin (8). Further analysis of the  $^1\text{H}$ - $^1\text{H}$  COSY NMR spectrum of fraction LT-3 revealed substructures belonging to the predicted active metabolites. For example, the correlation of the *meta*-coupled aromatic protons at  $\delta_{\text{H}}$  6.23 and 6.28 as well as that of the methyl doublet at  $\delta_{\text{H}}$  1.35 with one oxygenated methine proton at  $\delta_{\text{H}}$  5.15, which further correlated with the aliphatic chain, was characteristic of the desmethyl-lasiodiplodin structure (8) (**Table S1**, **Fig. S1**, Supporting Information). Substructures of palmarumycin CP17 and cladospirone B were elucidated in the same manner.

Fraction LT-7, an outlier in the active group, was also selected based on the OPLS-DA results, which indicated the bioactive metabolite. Among these metabolites, the structure of the compound 6-oxo-de-O-methylasiodiplodin (2) was confirmed by its COSY spectral data, which exhibited correlations similar to desmethyl-lasiodiplodin (8), as shown in **Fig. S2A**, Supporting Information. In the aromatic region, the COSY spectrum revealed correlations as in preussomerin-C (**Fig. S2B**, Supporting Information).

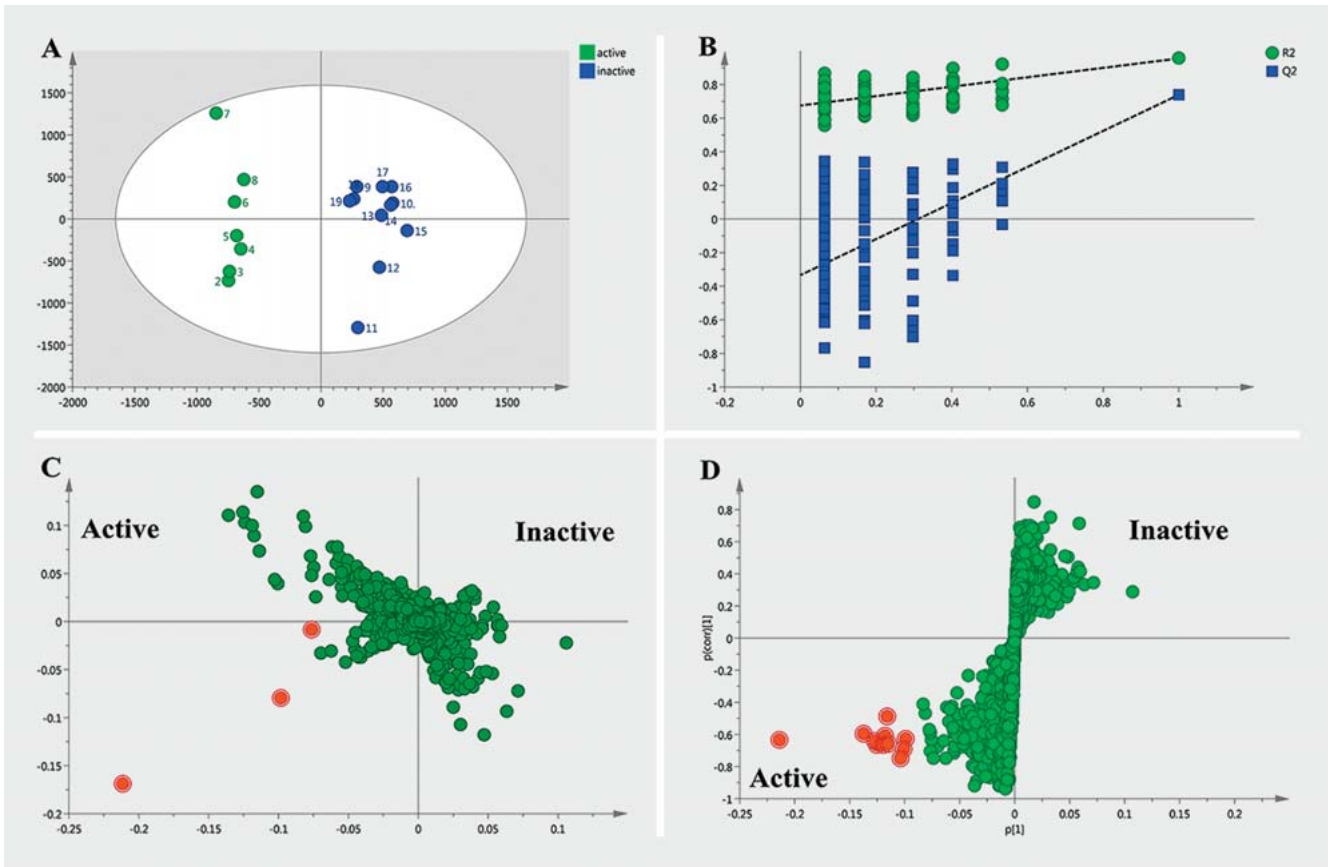
Isolation and purification of the compounds in the active group was performed by high-throughput medium pressure liquid chromatography (MPLC). Three known compounds were isolated and elucidated based on their NMR and MS data as cladospirone B (6) [43], desmethyl-lasiodiplodin (8) [44], and *R*-(-)-mellein (9) [45] (chemical structures shown in ► **Fig. 6**). The isolation of cladospirone B (6) and desmethyl-lasiodiplodin (8) confirmed the putative identification of the bioactive metabolites predicted from the S-plot of the OPLS-DA model. *R*-(-)-mellein (9) is a structurally close analogue of 6,8-dihydroxy-3-methylisocoumarin (1). All isolated compounds were tested against *T. b. brucei*, cladospirone B (6) and desmethyl-lasiodiplodin (8) had MICs of 17.8 and 22.5  $\mu\text{M}$ , respectively. All three metabolites were checked again in the S-plot and *R*-(-)-mellein was located in the middle of the plot, suggesting that the anti-trypanosomal activity would be less (► **Fig. 7**); this is indeed the case as confirmed by the bioassay results (► **Table 3**). Due to low concentration, palmarumycin CP17 was not isolated. Desmethyl-lasiodiplodin (8) has been known to exhibit anticancer activity against MCF-7 via apoptosis with an  $\text{IC}_{50}$  sevenfold more potent than its toxicity on normal cells [46]. On the other hand, cladospirone B (6) has been reported to be inactive in antibacterial and antifungal assays [45].

In our screening program, we observed that the crude extract of *L. theobromae* from agar plates showed anti-trypanosomal activity against *T. b. brucei*. In order to find the optimal conditions to grow *L. theobromae*, HR-LCMS-based metabolomics was ap-





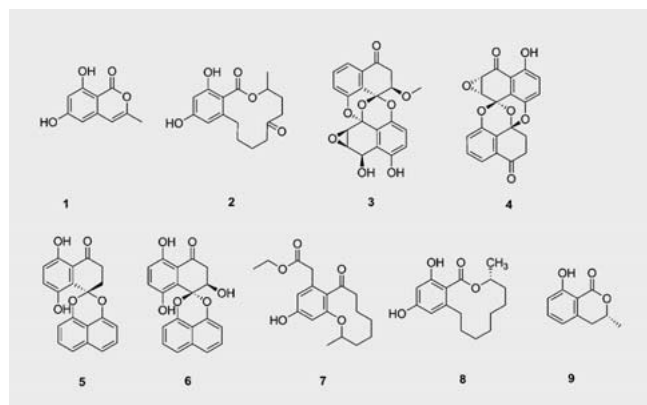
► **Fig. 4** A The  $^1\text{H}$  NMR spectra of the 19 fractions. B The expansion of the  $^1\text{H}$  NMR data of active anti-trypanosome fractions highlighting several unique chemical fingerprints found only in these fractions.



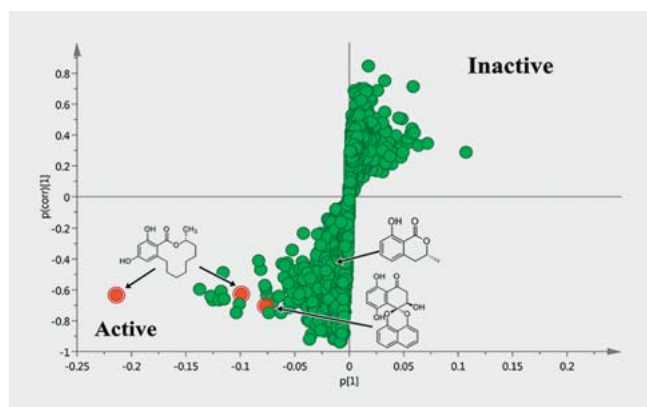
► **Fig. 5** Multivariate analysis of *L. theobromae* fractions and anti-trypanosomal activity data correlation. A The score scatter plot of OPLS-DA shows the samples were grouped based on their bioactivity. B The permutation test result of the OPLS-DA model. (C) The loading scatter plot of OPLS-DA shows the  $m/z$  values of active metabolites. (D) The S-plot generated from the OPLS-DA model shows the end point compounds that are the predicted metabolites responsible for the bioactivity (highlighted in red).

► **Table 2** Putatively identified unique metabolites of *L. theobromae* active fractions obtained from S-plot “end point” data as shown on ► **Fig. 5D**. (P = positive mode; N = negative mode).

Ionization mode	MS <i>m/z</i>	Rt (min)	Chemical formula	Name
N	191.035	11.32	C <sub>10</sub> H <sub>8</sub> O <sub>4</sub>	6,8-Dihydroxy-3-methylisocoumarin(1)
N	291.124	13.17	C <sub>16</sub> H <sub>20</sub> O <sub>5</sub>	6-Oxo-de-O-methylasiodiplodin (2)
P	293.139	13.71	C <sub>16</sub> H <sub>20</sub> O <sub>5</sub>	6-Oxo-de-O-methylasiodiplodin (2)
N	395.077	13.60	C <sub>21</sub> H <sub>16</sub> O <sub>8</sub>	Preussomerin-C (3)
N	363.051	14.61	C <sub>20</sub> H <sub>12</sub> O <sub>7</sub>	Preussomerin-H (4)
N	333.077	17.61	C <sub>20</sub> H <sub>14</sub> O <sub>5</sub>	Palmarumycin CP17 (5)
N	349.072	13.71	C <sub>20</sub> H <sub>14</sub> O <sub>6</sub>	Cladospiron B (6)
P	321.170	19.14	C <sub>18</sub> H <sub>24</sub> O <sub>5</sub>	Phomopsin B (7)
N	319.155	19.16	C <sub>18</sub> H <sub>24</sub> O <sub>5</sub>	Phomopsin B (7)
N	277.144	21.79	C <sub>16</sub> H <sub>22</sub> O <sub>4</sub>	Desmethyl-lasiodiplodin (8)
P	279.159	21.80	C <sub>16</sub> H <sub>22</sub> O <sub>4</sub>	Desmethyl-lasiodiplodin (8)
N	555.296	21.79		Complex of 277.144



► **Fig. 6** Chemical structures of unique metabolites (1–8) predicted from the S-plot and further secondary metabolites (6, 8, 9) isolated from an active fraction of *L. theobromae* (Group 1).



► **Fig. 7** Three isolated compounds from *L. theobromae* labelled in the S-plot. *R*-(-)-mellein was in the middle of the plot, suggesting less anti-trypanosomal activity for this compound.

plied, which resulted in the selection of solid rice culture for 30 days as the best conditions for medium-scale fermentation. Fractionation was performed on the crude extract, and based on the 1D <sup>1</sup>H NMR data comparison of 19 fractions, several unique chemical fingerprints in the active fractions were highlighted. Furthermore, by utilizing the HR-LCMS data for multivariate analysis such as OPLS-DA, consequently, a set of the metabolites that were predicted to be active, was generated. All predicted metabolites were easily identified with the aid of AntiBase coupled to MZMine. The application of <sup>1</sup>H NMR and COSY allowed for the detection of the predicted metabolites in the active fractions as well as the confirmation of the dereplication results obtained from the HR-LCMS data. Three known compounds were isolated and identified as cladospiron B (6), desmethyl-lasiodiplodin (8), and *R*-(-)-mellein (9). To the best of our knowledge, this is the first report of isolation of cladospiron B (6) from *L. theobromae*. It is also the first report to indicate the good anti-trypanosomal activity of cladospiron B (6) and desmethyl-lasiodiplodin (8) against *T. b. brucei* in comparison with suramin (MIC value of 0.1 ± 0 μM). Our strategy of emphasizing HR-LCMS and NMR-based metabolomics to search for active anti-trypanosomal compounds has therefore been proven to be effective. In conclusion, this study determined that the combination of HR-LCMS and NMR-based metabolomics is a powerful and advantageous decision-making tool in mining active metabolites of *L. theobromae* against *T. b. brucei* and is also promising for implementation in other drug discovery programs elsewhere.

## Materials and Methods

### Fungal sampling

The fungus *L. theobromae* was isolated from fresh healthy leaves of *V. pinnata* collected in April 2011 near Kuala Terengganu, Malaysia. The plant was identified by Dr. Nashriyah Mat from the Faculty of Bioresources and Food Industry, Universiti Sultan Zainal

► **Table 3** Anti-trypanosomal activity of isolated compounds obtained from *L. theobromae* fermented in solid rice culture for 30 days.

Compound	Anti-trypanosomal activity ( <i>T. b. brucei</i> ) MIC average $\pm$ SD (n = 4) ( $\mu$ M)
Cladospirone-B (6)	17.8 $\pm$ 0
Desmethyl-lasiiodiplodin (8)	22.5 $\pm$ 1.50
R-(–)-mellein (9)	> 100 $\pm$ 2.75
Suramin	0.1 $\pm$ 0

Abidin, and a voucher specimen was deposited (collection number VP 01).

Samples were kept in zip lock bags and stored at 4°C until the isolation of endophytic fungi was performed 4 days later upon arrival at the University of Strathclyde in Glasgow. The surfaces of the leaves and stems were sterilized with 70% iso-propanol for 2 min and subsequently rinsed in sterile water. Small tissue samples from inside the leaves and stems were cut aseptically and pressed onto agar plates (composition of isolation medium: 15 g/L malt extract, 15 g/L agar, and 0.2 g/L chloramphenicol in distilled water, pH 7.4–7.8, adjusted with 10% NaOH or 36.5% HCl). Chloramphenicol ( $\geq$  98%; purity Sigma-Aldrich) was added to inhibit bacterial growth. The plates were left for a few days until fungal growth was observed. Reinoculation onto new malt agar plates was repeated several times until pure strains were attained.

### Identification of fungal strains

The fungal strain was identified using DNA amplification and sequencing of the internal transcribed spacer (ITS) region as described previously [47]. The sequence data has been submitted to GenBank with accession number KC960898. The fungal strain was identified as *L. theobromae*. A voucher strain was submitted and is kept at the Natural Product Metabolomics Laboratory, SIPBS.

### Small- and medium-scale fermentation

The fungal strain was cultivated on malt agar plates for 7 days at 30°C. The colonies and agar were cut into small pieces and were placed in liquid or solid media. The liquid medium used was Wickerham medium, consisting of 3 g yeast extract, 3 g malt extract, 5 g peptone, 10 g glucose, and distilled water adding up to 1000 mL in 2 L Erlenmeyer flasks, pH 7.2–7.4, adjusted with 10% NaOH or 36.5% HCl. The solid medium used was rice medium, with 100 g of long grain rice and 100 mL of distilled water autoclaved together in 1 L Erlenmeyer flasks. The strain was grown for 3 different incubation times: 7, 15, and 30 days, under static conditions at room temperature. For medium-scale fermentation, *L. theobromae* was cultivated in 1 L flasks using the optimal conditions as determined by the small-scale cultures. In this case, the growth of the fungus on rice medium for 30 days under static conditions was determined to be the most favorable condition for the production of anti-trypanosomal metabolites.

### Extraction and isolation of pure compounds

The medium-scale rice cultures were extracted with ethyl acetate and homogenized as finely as possible using a T18 Basic Ultra-Turrax (IKA) at maximum speed. These were then kept overnight. The extract was subsequently dried under *vacuo* using a rotary evaporator (Buchi Labortechnik AG). The crude extract (5.0 g) was then partitioned with 10% *n*-hexane and 90% methanol to remove fatty acids. The extract containing methanol-soluble compounds (2.0 g) was collected for further isolation work. The fractionation of the methanol extract was accomplished using MPLC. Linear gradient elution was employed with hexane (A) and ethyl acetate (B) as the mobile phase at a flow rate of 20 mL/min. A pre-packed silica column (20–45  $\mu$ m, 23  $\times$  110 mm, Silica VersaPak cartridge) was used. It was connected to a Buchi Pump Manager C-615 coupled to binary pumps (Buchi Modules C-601). 100% A was run for 5 min, followed by 100% A to 100% B for 20 min, and finished with 100% B for the last 5 min. The total run time was 30 min. Fractions were collected in collection tubes automatically every 2 mL using a fraction collector Frac 920 (GE Healthcare Bio-Sciences AB). Fractions with similar TLC profiles were pooled together, yielding a total of 19 fractions. Fraction LT-2 (435 mg) was further subjected to MPLC on a prepacked silica column (20–45  $\mu$ m, 23  $\times$  53 mm, Silica VersaPak cartridge) utilizing an isocratic gradient system (80% *n*-hexane: 20% ethyl acetate) for 30 min at a flow rate of 20 mL/min. In total, 300 collection tubes of fractions (2 mL in each tube) were collected automatically using their TLC profiles. Similar fractions were pooled, resulting in 17 fractions and giving 3 known compounds, **6** (3 mg), **8** (73 mg), and **9** (11 mg) with a purity > 95% (determined by HPLC).

### *In vitro* anti-trypanosomal assay

The samples were prepared to a final concentration of 10 mg/mL (stock solution) by being dissolved in the appropriate amount of DMSO. To screen for *in vitro* activity, a concentration of 200  $\mu$ g/mL was used. This was achieved by diluting the stock solution 1 in 10 with HMI-9 (drug solution). Four  $\mu$ L of drug solution were transferred to 96  $\mu$ L of HMI-9 in the 96-well plate. One hundred  $\mu$ L of trypanosome suspension, consisting of *T. b. brucei* S427 blood stream form at  $3 \times 10^4$  trypanosomes/mL, were then added to the 96-well plate to make the final concentration of the compounds range from 100  $\mu$ g/mL to 0.17  $\mu$ g/mL. DMSO was used as the negative control (concentration of 1 to 0.002%) and suramin (Calbiochem-Novabiochem Co., purity > 98% by HPLC) was selected as the positive control (concentration of 1 to 0.008  $\mu$ M). The plate was incubated for 48 h at 37°C, 5% CO<sub>2</sub> with a humidified atmosphere, after which 20  $\mu$ L of Alamar blue were added. The plate was again incubated for another 24 h under the same conditions. The fluorescence was measured using a Wallac Victor microplate reader (PerkinElmer) with excitation at 530 nm and emission at 590 nm. The results were calculated as percentages of control values. All samples that exhibited > 90% inhibition were selected for the MIC assay to determine the MIC value.

### Nuclear magnetic resonance instrumentation

One- and two-dimensional <sup>1</sup>H and <sup>13</sup>C NMR spectra were recorded at 400 MHz on a JEOL-LA400 FT-NMR spectrometer system with a 40<sup>TH</sup>5AT/FG probe (JEOL LTD.). Compounds **6** and **8** were reconsti-

tuted in deuterated chloroform (CDCl<sub>3</sub>) while compound **9** was reconstituted in deuterated DMSO (DMSO-*d*<sub>6</sub>).

### High-resolution-liquid chromatography mass spectrometry procedure

HR-LCMS was measured using an Accela 600 HPLC pump with Accela autosampler and UV/Vis detector (Thermo Scientific) and an Orbitrap Exactive mass spectrometer (Thermo Fisher Scientific, Inc.). Analysis of samples was done using similar protocols described previously [23, 47, 48].

### High-resolution-liquid chromatography mass spectrometry data processing

Initially the raw HR-LCMS data were sliced into two data sets according to ionization mode using the MassConvert tool from ProteoWizard (<http://proteowizard.sourceforge.net/>). The sliced data were imported to MZMine 2.10 (<http://sourceforge.net/projects/mzmine/>), software developed for the differential analysis of mass spectrometry data. The data processing step was performed in the same manner as explained previously [23], albeit with slightly modified parameters. In this analysis, the data set was crop filtered from 0.1 to 35 min and the retention time normalizer was not applied because only one batch of data was used.

### Statistical analysis

MS spectral data were converted to an ASCII text file and imported to MS Excel. The data was sorted to exclude background peaks that belonged to the MeOH blank. The sorted data were then exported to the SIMCA-P software 14.0 version (Umetrics). Pareto scaling was employed on the MS data set. Finally, PCA, OPLS-DA, and S-plot were performed.

### Supporting information

The <sup>1</sup>H, <sup>13</sup>C NMR and HMBC as well as ESI-MS data of the isolated compounds are available as Supporting Information.

## Acknowledgements

We thank Ministry of Education Malaysia for a scholarship for Nurkhalida Kamal.

### Conflict of Interest

The authors declare no conflict of interest.

### References

- [1] Newman DJ, Cragg GM. Natural products as sources of new drugs over the 30 years from 1981 to 2010. *J Nat Prod* 2012; 75: 311–335
- [2] Challis GL. Mining microbial genomes for new natural products and biosynthetic pathways. *Microbiology* 2008; 154: 1555–1569
- [3] Gongora-Castillo E, Buell CR. Bioinformatics challenges in *de novo* transcriptome assembly using short read sequences in the absence of a reference genome sequence. *Nat Prod Rep* 2013; 30: 490–500
- [4] Yang JY, Karr JR, Watrous JD, Dorrestein PC. Integrating ‘-omics’ and natural product discovery platforms to investigate metabolic exchange in microbiomes. *Curr Opin Chem Biol* 2011; 15: 79–87
- [5] Verpoorte R, Choi Y, Kim H. NMR-based metabolomics at work in phytochemistry. *Phytochem Rev* 2007; 6: 3–14
- [6] Moco S, Vervoort J, Bino RJ, De Vos RCH, Bino R. Metabolomics technologies and metabolite identification. *Trends Analyt Chem* 2007; 26: 855–866
- [7] Sumner LW, Lei Z, Nikolau BJ, Saito K. Modern plant metabolomics: advanced natural product gene discoveries, improved technologies, and future prospects. *Nat Prod Rep* 2015; 32: 212–229
- [8] Lang G, Mayhudin NA, Mitova MI, Sun L, van der Sar S, Blunt JW, Cole ALJ, Ellis G, Laatsch H, Munro MHG. Evolving trends in the dereplication of natural product extracts: new methodology for rapid, small-scale investigation of natural product extracts. *J Nat Prod* 2008; 71: 1595–1599
- [9] Vasilev N, Ebel R, Edrada RA, Fuss E, Alfermann AW, Ionkova I, Petrova A, Replinger M, Schmidt TJ. Metabolic profiling of lignan variability in *Linum* species of section *Syllinum* native to Bulgaria. *Planta Med* 2008; 74: 273–280
- [10] Frisvad JC, Rank C, Nielsen KF, Larsen TO. Metabolomics of *Aspergillus fumigatus*. *Med Mycol* 2008; 47: S53–S71
- [11] Wolfender JL, Marti G, Ferreira Queiroz E. Advances in techniques for profiling crude extracts and for the rapid identification of natural products: dereplication, quality control and metabolomics. *Curr Org Chem* 2010; 14: 1808–1832
- [12] Funari CS, Eugster PJ, Martel S, Carrupt PA, Wolfender JL, Silva DHS. High resolution ultra high pressure liquid chromatography – time-of-flight mass spectrometry dereplication strategy for the metabolite profiling of Brazilian Lippia species. *J Chromatogr A* 2012; 1259: 167–178
- [13] Geiger M, Desanglois G, Hogeveen K, Fessard V, Leprière T, Mondeguer F, Guittou Y, Hervé F, Séchet V, Grovel O, Pouchou YF, Hess P. Cytotoxicity, fractionation and dereplication of extracts of the dinoflagellate *Vulcanodinium rugosum*, a producer of pinnatoxin G. *Mar Drugs* 2013; 11: 3350–3371
- [14] Zhang T, Omar R, Siheri W, Al Mutairi S, Clements C, Fearnley J, Edrada-Ebel R, Watson D. Chromatographic analysis with different detectors in the chemical characterisation and dereplication of African propolis. *Talanta* 2014; 120: 181–190
- [15] Hou Y, Braun DR, Michel CR, Klassen JL, Adnani N, Wyche TP, Bugni TS. Microbial strain prioritization using metabolomics tools for the discovery of natural products. *Anal Chem* 2012; 84: 4277–4283
- [16] Ellis GA, Hou Y, Braun DR, Wyche TP, Adnani N, Vazquez-Rivera E, Bugni TS. LC/MS Untargeted metabolomics for prioritizing marine invertebrate-associated bacteria for discovery of natural products. *Planta Med* 2013; 79: PK7
- [17] Samat N, Tan PJ, Shaari K, Abas F, Lee HB. Prioritization of natural extracts by LC–MS–PCA for the identification of new photosensitizers for photodynamic therapy. *Anal Chem* 2014; 86: 1324–1331
- [18] Macintyre L, Zhang T, Viegelmann C, Martinez IJ, Cheng C, Dowdells C, Abdelmohsen UR, Gernert C, Hentschel U, Edrada-Ebel R. Metabolomic tools for secondary metabolite discovery from marine microbial symbionts. *Mar Drugs* 2014; 12: 3416–3448
- [19] Kim HK, Wilson EG, Choi YH, Verpoorte R. Metabolomics: a tool for anti-cancer lead-finding from natural products. *Planta Med* 2010; 76: 1094–1102
- [20] Ali K, Iqbal M, Yuliana N, Lee YJ, Park S, Han S, Lee JW, Lee HS, Verpoorte R, Choi Y. Identification of bioactive metabolites against adenosine A1 receptor using NMR-based metabolomics. *Metabolomics* 2013; 9: 778–785
- [21] Bohni N, Cordero-Maldonado ML, Maes J, Siverio-Mota D, Marcourt L, Munck S, Kamuhabwa AR, Moshi MJ, Esguerra CV, de Witte PA. Integration of microfractionation, qNMR and zebrafish screening for the *in vivo*



bioassay-guided isolation and quantitative bioactivity analysis of natural products. *PLoS One* 2013; 8: e64006

- [22] Chagas-Paula DA, Zhang T, Oliveira TB, Edrada-Ebel R, Da Costa FB. Discovery of plant anti-inflammatory biomarkers by machine learning algorithms and metabolomic studies. *Planta Med* 2013; 79: SL27
- [23] Abdelmohsen U, Cheng C, Viegelmann C, Zhang T, Grkovic T, Ahmed S, Quinn R, Hentschel U, Edrada-Ebel R. Dereplication strategies for targeted isolation of new antitrypanosomal actinosporins a and b from a marine sponge associated-*Actinokineospora* sp. EG49. *Mar Drugs* 2014; 12: 1220–1244
- [24] Harvey AL, Edrada-Ebel R, Quinn RJ. The re-emergence of natural products for drug discovery in the genomics era. *Nat Rev Drug Discov* 2015; 14: 111–129
- [25] Bacon CW, White J. *Microbial Endophytes*. Basel: CRC Press; 2000
- [26] Dreyfuss M, Chapela IH. Potential of fungi in the discovery of novel, low-molecular weight pharmaceuticals. In: Gullo VP, ed. *The Discovery of natural Products with therapeutic Potential*. London, United Kingdom: Butterworth-Heinemann; 1994: 49–80
- [27] Hawksworth DL. The fungal dimension of biodiversity: magnitude, significance, and conservation. *Mycol Res* 1991; 95: 641–655
- [28] Hawksworth DL. The magnitude of fungal diversity: the 1.5 million species estimate revisited. *Mycol Res* 2001; 105: 1422–1432
- [29] Tsuchinari M, Shimanuki K, Hiramatsu F, Murayama T, Koseki T, Shiono Y. Fusapyridons A and B, novel pyridone alkaloids from an endophytic fungus, *Fusarium* sp. YG-45. *Z Naturforsch B J Chem Sci* 2007; 62: 1203–1207
- [30] Zilla MK, Qadri M, Pathania AS, Strobel GA, Nalli Y, Kumar S, Guru SK, Bhushan S, Singh SK, Vishwakarma RA, Riyaz-Ul-Hassan S, Ali A. Bioactive metabolites from an endophytic *Cryptosporiopsis* sp. inhabiting *Clidemia hirta*. *Phytochemistry* 2013; 95: 291–297
- [31] Isaka M, Berkaew P, Intereya K, Komwijit S, Sathitkunanon T. Antiplasmodial and antiviral cyclohexadepsipeptides from the endophytic fungus *Pullularia* sp. BCC 8613. *Tetrahedron* 2007; 63: 6855–6860
- [32] Singh SB, Ondeyka JG, Tsipouras N, Ruby C, Sardana V, Schulman M, Sanchez M, Pelaez F, Stahlhut MW, Munshi S, Olsen DB, Lingham RB. Hinnuliquinone, a C2-symmetric dimeric non-peptide fungal metabolite inhibitor of HIV-1 protease. *Biochem Biophys Res Commun* 2004; 324: 108–113
- [33] Chen X, Shi Q, Lin G, Guo S, Yang J. Spiroisnaphthalene analogues from the endophytic fungus *Preussia* sp. *J Nat Prod* 2009; 72: 1712–1715
- [34] Ge HM, Yu ZG, Zhang J, Wu JH, Tan RX. Bioactive alkaloids from endophytic *Aspergillus fumigatus*. *J Nat Prod* 2009; 72: 753–755
- [35] Campos FF, Rosa LH, Cota BB, Caligiorne RB, Rabello ALT, Alves TMA, Rosa CA, Zani CL. Leishmanicidal metabolites from *Cochliobolus* sp., an endophytic fungus isolated from *Piptadenia adiantoides* (fabaceae). *PLoS Negl Trop Dis* 2008; 2: e348
- [36] Moreno E, Varughese T, Spadafora C, Arnold AE, Coley PD, Kursar TA, Gerwick WH, Cubilla-Rios L. Chemical constituents of the new endophytic fungus *Mycosphaerella* sp. nov. and their anti-parasitic activity. *Nat Prod Commun* 2011; 6: 835
- [37] Simarro PP, Diarra A, Postigo JAR, Franco JR, Jannin JG. The human African trypanosomiasis control and surveillance programme of the World Health Organization 2000–2009: the way forward. *PLoS Negl Trop Dis* 2011; 5: e1007
- [38] Jacobs RT, Nare B, Phillips MA. State of the art in African trypanosome drug discovery. *Curr Top Med Chem* 2011; 11: 1255
- [39] Webster J, Weber R. *Introduction to Fungi*. Cambridge: Cambridge University Press; 2007
- [40] Pluskal T, Castillo S, Villar-Briones A, Orešič M. MZmine 2: modular framework for processing, visualizing, and analyzing mass spectrometry-based molecular profile data. *BMC Bioinformatics* 2010; 11: 395
- [41] Robotti E, Marengo E. Chemometric multivariate tools for candidate biomarker identification: LDA, PLS-DA, SIMCA, ranking-PCA. *Methods Mol Biol* 2016; 1384: 237–267
- [42] Haoula Z, Ravipati S, Stekel DJ, Ortori CA, Hodgman C, Daykin C, Raine-Fenning N, Barrett DA, Atiomo W. Lipidomic analysis of plasma samples from women with polycystic ovary syndrome. *Metabolomics* 2015; 11: 657–666
- [43] Schulz B, Sucker J, Aust H, Krohn K, Ludewig K, Jones P, Döring D. Biologically active secondary metabolites of endophytic *Pezizula* species. *Mycol Res* 1995; 99: 1007–1015
- [44] Aldridge D, Galt S, Giles D, Turner W. Metabolites of *Lasiodiplodia theobromae*. *J Chem Soc C* 1971: 1623–1627
- [45] Bode HB, Walker M, Zeeck A. Cladosporones B to I from *Sphaeropsidales* sp. F-24'707 by variation of culture conditions. *European J Org Chem* 2000; 2000: 3185–3193
- [46] Hazalin NAM, Lim SM, Cole AL, Majeed ABA, Ramasamy K. Apoptosis induced by desmethyl-lasiodiplodin is associated with upregulation of apoptotic genes and downregulation of monocyte chemotactic protein-3. *Anticancer Drugs* 2013; 24: 852–861
- [47] Tawfike A, Viegelmann C, Edrada-Ebel R. *Metabolomics and Dereplication Strategies in natural Products*. In: Roessner U, Dias DA, eds. *Metabolomics Tools for natural Product Discovery*. New Jersey: Humana Press; 2013: 227–244
- [48] Bafor EE, Lim CV, Rowan EG, Edrada-Ebel R. The leaves of *Ficus exasperata* Vahl (Moraceae) generates uterine active chemical constituents. *J Ethnopharmacol* 2012; 145: 803–812