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ORIGINAL ARTICLE

A workflow for bacterial metabolic fingerprinting and lipid profiling: application to Ciprofloxacin challenged *Escherichia coli*

**J. William Allwood · Haitham AlRabiah ·
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Abstract The field of lipidomics focuses upon the non-targeted analysis of lipid composition, the process of which follows similar routines to those applied in conventional metabolic profiling, however lipidomics differs with respect to the sample preparation steps and chosen analytical platform applied to the sample analysis. Conventionally, lipidomics has applied analytical techniques such as direct infusion mass spectrometry and more recently reverse phase liquid chromatography–mass spectrometry, for the detection of mono-, di-, and tri-acyl glycerols, phospholipids, and other complex lipophilic species such as sterols. The field is rapidly expanding, especially with respect to the clinical sciences where it is known that changes of lipid composition, especially phospholipids, are commonly associated with many disease processes. As a proof of principle study, a small number of *Escherichia coli* isolates were selected on the basis of their sensitivity to a second generation fluoroquinolone antibiotic, known as Ciprofloxacin (*E. coli* isolates 161 and 171, non-ST131 isolates, which are resistant and sensitive respectively: *E. coli* isolates 160 and 173, ST131 sequence isolates which are resistant and susceptible respectively). It has been proposed that Ciprofloxacin may be a surface active drug that interacts at the surface-water interface of the phospholipid bi-layer where the head groups reside. Further, antibiotic resistance through intracellular exclusion is known to result in remodelling of the phospholipid membrane. Therefore, to study the effects of Ciprofloxacin on both susceptible and resistant bacterial strains, lipid profiling would present an informative approach. Control and antibiotic challenged cultures for each of the isolates were compared for changes in metabolite and lipid composition as detected by FT-IR spectroscopy and RP-UHPLC–MS, and appraised with a variety of chemometric data analysis approaches. The developed bacterial lipidomics workflow was deemed to be highly reproducible (with respect to the

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employed technical and analytical routines) and led to the detection of a large array of lipid classes as well as highlighting a range of significant lipid alterations that differed in regulation between susceptible and resistant *E. coli* isolates.

Keywords Lipidomics · *Escherichia coli* · Ciprofloxacin hydrochloride · Antibiotic-resistance · Fourier transform infrared spectroscopy · Liquid chromatography–mass spectrometry

1 Introduction

The popularity of metabolomics and the closely related discipline of lipidomics has risen greatly over the past decade, across an ever growing range of disciplines and applications, including, gene function analysis (Fiehn et al. 2000; Roessner et al. 2001), environmental perturbation (Kaplan et al. 2004; Allwood et al. 2006, 2010), diseases of clinical significance (Kolak et al. 2007; Mattila et al. 2008; Orešič et al. 2008; Sreekumar et al. 2009; Kenny et al. 2010), and to a large number of organisms including plants (Lisec et al. 2006; De Vos et al. 2007), fruits, vegetables and grains (Allwood et al. 2009; Biais et al. 2009; Tikunov et al. 2010), microbes (Koek et al. 2006; Winder et al. 2006, 2008; Van Der Werf et al. 2008; Lowe et al. 2010), and mammals (Griffin and Kauppinen 2007; Sreekumar et al. 2009; Kenny et al. 2010; Velagapudi et al. 2010; Dunn et al. 2007, 2008, 2011). The metabolome and lipidome perhaps represents the ultimate phenotype of cells, controlled by gene expression and the modulation of protein function, which in turn are controlled by the environment and genome mutation (Fiehn 2002; Saito and Matsuda 2010). The field of lipidomics focuses upon the non-targeted analysis of lipid composition, the process of data collection and statistical analysis follows very similar routines to those applied in conventional metabolic profiling, however lipidomics differs with respect to the sample preparation steps and chosen analytical platform applied to the sample analysis. Conventionally, lipidomics has applied analytical techniques such as direct infusion mass spectrometry (DIMS; Goodacre et al. 2002; Han and Gross 2005; Allwood et al. 2006) and less commonly reverse phase liquid chromatography–mass spectrometry (RP-LC–MS; Kolak et al. 2007; Mattila et al. 2008; Wedge et al. 2011), for the detection of mono-, di-, and tri-acyl glycerols, phospholipids, and other complex lipophilic species such as sterols and ceramides.

Given the influence and importance of microbes upon many areas of biological, clinical and environmental research, they are indeed a very significant target for metabolomics and lipidomics. Many studies have focused

upon the polar central metabolism of microbes (Koek et al. 2006; Winder et al. 2008; MacKenzie et al. 2008), including applications in several systematic studies to increase our level of understanding on the control of central metabolism at the levels of the transcriptome, proteome and metabolome, with such studies typically focusing upon model organisms such as *Escherichia coli* K-12 (Mori 2004) and *Saccharomyces cerevisiae* (Castrillo et al. 2007; Herrgård et al. 2008). Further to *E. coli* being a commonly employed model species (Mori 2004; Riley et al. 2006), it is also of great significance as an enteric pathogen (Kaper et al. 2004), requiring antibiotic treatments in mammals to clear the gastric infection and alleviate symptoms of gastroenteritis. Understanding the actions of antibiotics, which are frequently poorly described, at the levels of the metabolome and lipidome using model bacteria such as *E. coli* is of importance to elucidate the mechanisms of action of antibiotics in current use and those under development. Antibiotics are commonly targeted to specific proteins, for example, fluoroquinolones such as Ciprofloxacin, are targeted to the bacterial DNA topoisomerase II (DNA gyrase) and DNA topoisomerase IV enzymes which function to relax supercoiled DNA and perform important processes in partitioning chromosomal DNA respectively (Greenwood 2000). Therefore, to study the effects of Ciprofloxacin on the metabolism of bacteria, especially those that are sensitive to the antibiotic, it would be logical to employ an analytical technique that measures the central metabolome (Koek et al. 2006; Winder et al. 2008). Such an approach would be capable of detecting changes in nucleotide metabolism and how they impact upon other areas of primary metabolism, potentially revealing direct consequential effects on metabolism due to the antibiotic mode of action. However, with the ever growing problem of bacterial resistance to antibiotics and the emergence of bacteria with resistance to multiple antibiotics (Tomasz 1994; Wiener et al. 1999; Sáenz et al. 2004), for example Methicillin-resistant *Staphylococcus aureus* (MRSA; Diederen and Kluytmans 2006), it is perhaps more important to focus studies on bacterial modes of resistance. Since bacterial resistance to antibiotics is commonly dependent on the ability of the bacteria to exclude the antibiotic from the cell by membrane remodelling via aminoacylation or cell wall modification by peptidoglycan cross-linking (RajBhandary and Söll 2008; Roy and Ibba 2008), or alternatively by removal of the antibiotic from the cytoplasm to the extracellular matrix through energy dependent membrane export (Poole et al. 1993; Okusu et al. 1996), it may be of great relevance to alternatively analyse the lipidome of bacteria.

Much research has been previously undertaken on the optimisation of bacterial growth and metabolic quenching, as well as the development of suitable GC–MS

methodologies for the analysis of central metabolism in *E. coli* as well as other bacterial species and yeasts (Winder et al. 2008; MacKenzie et al. 2008). Therefore, the principle aim of this investigation was to establish and validate a method for FT-IR fingerprinting and RP UHPLC–MS profiling of lipids in antibiotic challenged *E. coli*, taking into consideration sample extraction and analysis steps. FT-IR spectroscopy is a constantly developing analytical technique often used routinely as a method in bacteriology being applied to species level identification and discrimination, as well as a tool to follow phenotypic responses to abiotic and biotic perturbation (Winder et al. 2006; Preisner et al. 2007; Wehlri et al. 2013). RP LC–MS employing C18 column chemistries is the standard approach applied to the analysis of free and membrane lipids in the field of lipidomics. Therefore both FT-IR and RP UHPLC–MS were deemed appropriate analytical tools to apply to the study of changes in lipid quantity and composition in response to antibiotic challenge in bacteria.

As a proof of principle study, a small number of *E. coli* isolates were selected on the basis of their sensitivity to Ciprofloxacin. The isolates chosen are clinically relevant, being recently recovered from cases of urinary tract infection, and included strains from the globally disseminated ST131 clone (Lau et al., 2008). *E. coli* isolates 160 and 173, of the ST131 clone, which are classified as resistant and sensitive respectively, as well as *E. coli* isolates 161 and 171, non-ST131 strains, which are classified as resistant and intermediate resistant respectively, were selected to provide a range of *E. coli* responses to challenge with Ciprofloxacin. It has been proposed that Ciprofloxacin may be a surface active drug that interacts at the surface-water interface of the phospholipid bi-layer where the head groups reside (Merino et al. 2003). Further, since antibiotic resistance through intracellular exclusion is known to result in remodelling of the phospholipid membrane, the developed FT-IR fingerprinting and RP-UHPLC–MS lipid profiling approach was deemed to provide an informative method to study the effects of Ciprofloxacin on both susceptible and resistant bacterial responses.

2 Materials and methods

2.1 Chemicals

All chemicals used were of analytical reagent or a higher purity grade. All reference standards were of 99 % minimum purity. All materials were purchased from Sigma-Aldrich (Gillingham, UK) unless otherwise stated. Ciprofloxacin hydrochloride was obtained from Discovery fine chemicals (Dorset, UK), a stock solution was prepared by dissolving 100 mg in 50 ml of sterile distilled water,

further dilutions were also made with sterile distilled water. HPLC grade methanol, chloroform and water were purchased from Sigma-Aldrich (Gillingham, UK). Formic acid (BDH Aristar 1) was purchased from VWR International (East Grinstead, UK).

2.2 *Escherichia coli* isolates

Four clinical *E. coli* isolates were donated by Dr. Mathew Upton from the Manchester Royal Infirmary. The isolates were selected on the basis of sequence type and sensitivity towards quinolones. A sensitive isolate (isolate 173) and a resistant isolate (isolate 160) of the ST131 clone, as well as two isolates that were resistant and partially-resistant to quinolones, isolates 161 and 171 respectively (non-ST131 strains), were selected for the lipidomic analysis.

2.3 Preparation of *Escherichia coli* cultures, their antibiotic challenge and determination of minimal inhibitory concentration (MIC)

Preparation of *E. coli* inoculates, their culture, challenge with Ciprofloxacin and determination of MIC are described in full detail within the Supplementary methods (Sects. 1.1–1.3) and are also presented within Fig. 1. For both the determination of MIC and also generation of samples for metabolic fingerprinting and profiling, all cultures were generated on a single day, with a single batch of culture media and were incubated in a common orbital shaker. In the case of larger experiments where sample numbers demand the generation of cultures in multiple batches (across days and even weeks), it is recommended that a minimum of one sample class is maintained within all culture batches, therefore providing a measure of variance associated with the different days/batches of culture generation. Further, by generating inoculums and culture media as a single lot for all experimental batches, as well as preparing a single source of both quenching and extraction solvents, biological and technical variation between the experimental batches will be minimised. In total 24 cultures were generated (4 *E. coli* strains × 2 treatments (control and antibiotic challenged) × 3 biological replicates). The experiment was designed to test technical (sample quenching) and analytical reproducibility, with having only a minimal number of biological replicates generated within a single culture batch, the experimental design does not permit a measure of the biological reproducibility of the employed methods.

2.4 *Escherichia coli* sample collection and quenching for metabolic fingerprinting and lipid profiling

The sample collection and quenching of metabolism methods were adapted from the procedures developed by
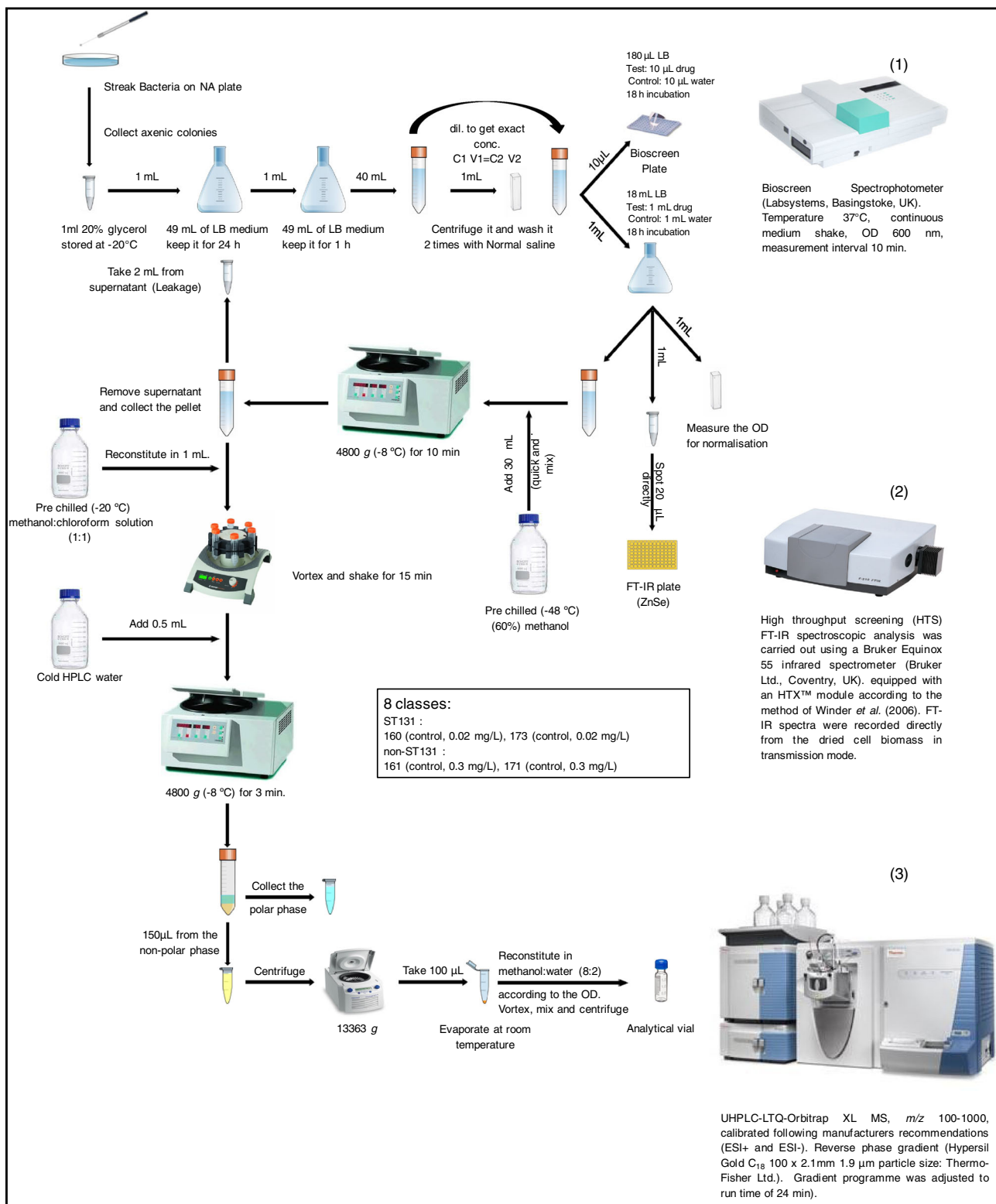
Fig. 1 Schematic of sample generation including (*1*) bioscreen analysis to determine the MIC of ciprofloxacin and produce the growth curves of pathogenic *E. coli*. (*2*) FT-IR analysis of samples. (*3*) LC–MS analysis of samples after quenching with cold (−48 °C) methanol and extraction with (1:1) methanol:chloroform

Winder et al. (2008) (Fig. 1), a full description of which is available in the Supplementary methods Sect. 1.4. All bacterial cultures were collected on a single day. Due to limitations in sample handling capacity, quenching was performed upon randomised batches of six samples at a time and the procedure requires approximately 20 min per a batch, therefore the antibiotic treatment of each randomised batch was staggered across 20 min intervals, thus maintaining that each culture is challenged by the antibiotic for precisely 18 h. By quenching the samples across randomised batches an indication of the procedures technical reproducibility can be obtained.

2.5 Fourier transform infrared (FT-IR) spectroscopy

High throughput screening (HTS) FT-IR spectroscopic analysis was carried out using a Bruker Equinox 55 infrared spectrometer (Bruker Ltd., Coventry, UK) equipped with an HTX^{TM} module according to the method of Winder et al. (2006). Preparation of samples, FT-IR sample plates and acquisition of spectra are fully described in the Supplementary methods Sect. 1.5. Each biological sample was repeat spotted on the FT-IR plate and analysed in triplicate to provide an indication of analytical variance.

2.6 Sample extraction for UHPLC–MS lipid profiling

The bacterial samples were extracted applying methanol:chloroform (1:1), followed by addition of water so that polar and non-polar metabolites could be isolated via phase separation. The method is fully described in the Supplementary methods Sect. 1.6 and was adapted from the methods described in Winder et al. (2008), which were originally conceived by Bligh and Dyer (1959). The extraction method leads to the capture of both free and membrane lipids within the chloroform phase of the extract, all of which are amenable to separation and detection by RP-UHPLC–MS approaches.

2.7 UHPLC–MS analysis

All samples were analysed on the Accela UHPLC system (Thermo-Fisher Ltd. Hemel Hempsted, UK) coupled to an electrospray LTQ-Orbitrap XL hybrid mass spectrometry system (ThermoFisher, Bremen, Germany) adapting the methods previously described by Wedge et al. (2011) and provided in full detail in the Supplementary methods Sect. 1.7.

2.8 Processing of raw UHPLC–MS profiles and lipid identification

The UHPLC–MS raw data profiles were deconvolved using the freely available XCMS software (http://masspec.

scripps.edu/xcms/xcms.php), the data were quality assured as described in (Dunn et al. 2008; Wedge et al. 2011) and normalised by peak sum for each sample (i.e. TIC normalisation). The putative identification of lipid features was performed applying the PUTMEDID-LCMS set of workflows (Brown et al. 2011), and a table of the putative lipid assignments is available in the supplementary information (Table S2). Detailed descriptions of deconvolution, quality assurance and putative lipid identification are provided in Supplementary methods Sect. 1.8.

2.9 Statistical analyses

Statistical analyses were performed applying the multivariate method of principal component-discriminant function analysis (PC-DFA). PC-DFA scores plots were employed for data visualisation and loadings plots to indicate the most significant metabolic features. Further feature selection was performed on the UHPLC–MS data for each individual bacterial isolate, comparing the response between the antibiotic treated and non-treated controls. First, a multivariate method known as partial least squares (PLS) regression was applied, this resulted in a filtered list of the most significant features ordered by the magnitude of their PLS regression coefficients from largest to smallest. Secondly, a univariate approach, known as ANOVA ($p < 0.05$), was applied to the significant PLS regression coefficient lists. Finally, a list of the 50 most significant features, ordered according to the magnitude of their PLS regression coefficients (from high to low), which also had an ANOVA p value of <0.05, was produced for each bacterial strain. Full descriptions of the statistical methods are provided in Supplementary methods Sects. 1.9 and 1.10.

3 Results and discussion

3.1 Determination of the Ciprofloxacin minimal inhibitory concentration (MIC)

So that the lipidomics approach under development did not lead to the detection of lipid changes associated with the death of bacterial cells as opposed to detecting lipid alterations that may contribute to the ability of the bacteria to exclude and remove the antibiotic, thus mediating antibiotic resistance, it was first necessary to establish the MIC of each bacterial isolate when cultured in Lysogeny Broth (LB) and challenged by Ciprofloxacin. Each of the four *E. coli* clinical isolates were cultured in LB media and challenged with Ciprofloxacin solutions ranging from 0.0025 to 100 mg/l in concentration. Bacterial growth curves were generated by measuring the OD 600 nm across an 18 h growth period

Fig. 2 OD 600 growth curves from 0 to 18 h post antibiotic challenge to indicate minimal inhibitory concentrations (MIC) of Ciprofloxacin hydrochloride against *Escherichia coli* ST131 strains **a** 160 (resistant) and **b** 173 (sensitive) and non-ST131 strains **c** 161 (resistant) and **d** 171 (intermediate resistance), **e** table of predicted MIC (mg/l Ciprofloxacin)

(Fig. 2). It was determined that the *E. coli* isolates 161 (non-ST131) and 160 (ST131), regarded as fully resistant, were not inhibited in growth even at Ciprofloxacin concentrations as high as 100 mg/l. For the intermediate-resistant *E. coli* isolate 171 (non-ST131) the Ciprofloxacin MIC was determined to fall between 0.25 and 0.5 mg/l. Whereas for the sensitive *E. coli* isolate 173 (ST131) the Ciprofloxacin MIC was determined to fall between 0.02 and 0.03 mg/l. Based upon these results, the concentrations of antibiotic were selected for the challenge of bacteria for sample generation for metabolic fingerprinting and lipid profiling. For the

E. coli ST131 isolates 160 and 173, in order to permit direct comparison of metabolic responses, a common antibiotic concentration of 0.02 mg/l was selected based upon it being the MIC of the sensitive isolate 173. For the non-ST131 isolates 161 and 171, again in order to permit direct comparison of metabolic responses, a common antibiotic concentration of 0.3 mg/l was selected based upon it being intermediate to the MIC of 0.25–0.5 mg/l of isolate 171, and as being of a great enough concentration to lead to an inhibitory effect upon growth without resulting in total bacterial death (Fig. 2).
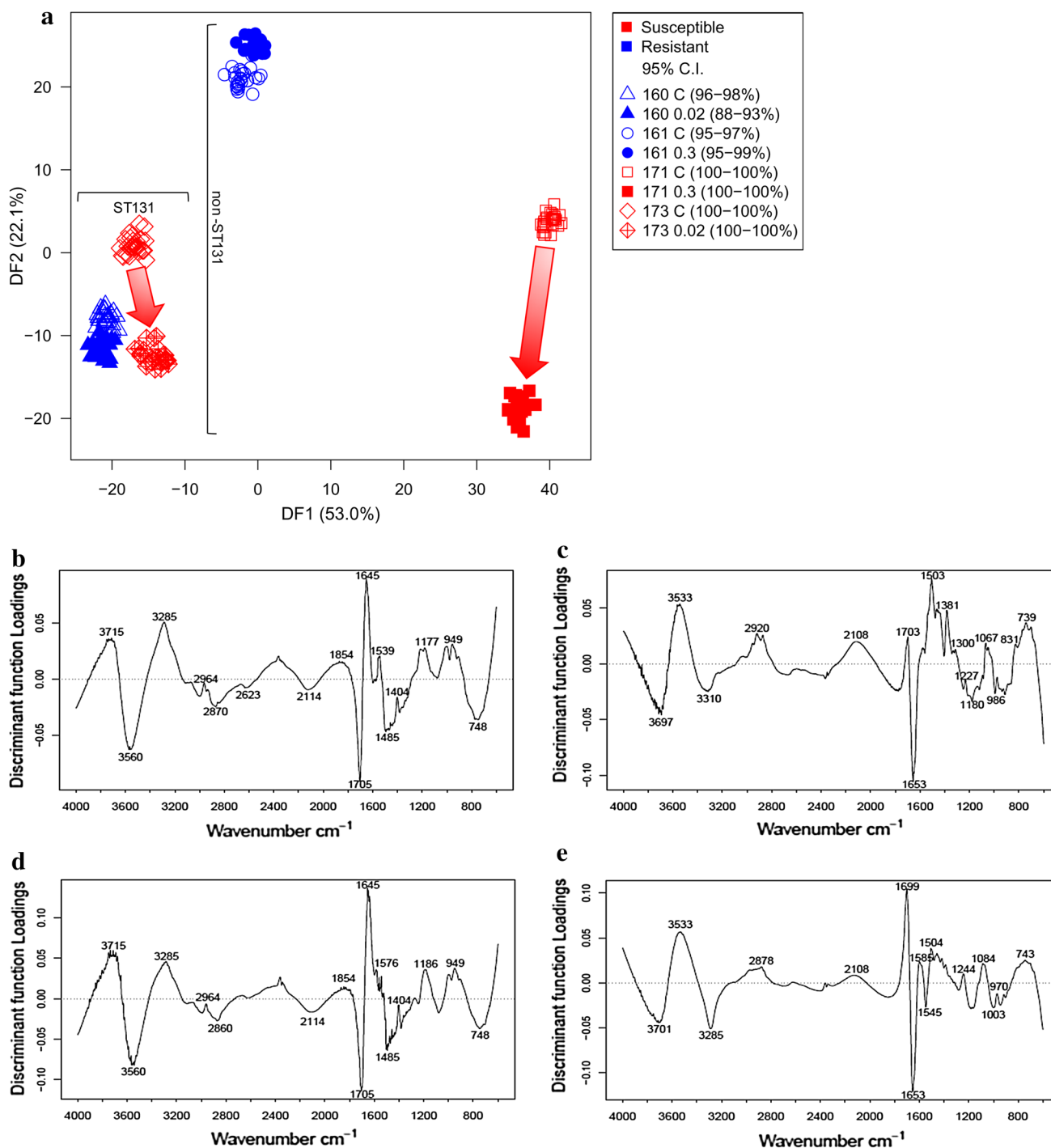
Fig. 3 FT-IR spectroscopy principle component-discriminant function analysis (PC-DFA) of 18 h Ciprofloxacin challenged and control *Escherichia coli*. **a** PC-DFA scores plot (DF1 vs. DF2). PC-DFA loadings plots derived by comparisons of Ciprofloxacin challenged and control samples for each respective isolate **b** ST131 strain 160

(resistant), **c** ST131 strain 173 (sensitive), **d** non-ST131 strain 161 (resistant), **e** non-ST131 strain 171 (intermediate resistance). Within the figures key, *C* refers to control samples, 0.02 and 0.3 refer to the antibiotic dose (mg/l), the 95 % confidence Interval (CI) is given in *brackets*

3.2 Fourier transform infrared (FT-IR) spectroscopy based metabolic fingerprinting

FT-IR spectroscopy is an established and constantly developing analytical technique, which enables the

extremely rapid (seconds per sample), high-throughput (1,000 s of samples per day) and non-destructive analysis of a wide-range of sample types (Ellis and Goodacre 2006; Allwood et al. 2008). FT-IR is also relatively inexpensive and therefore readily lends itself as a rapid first-round

screening method (Allwood et al. 2008) prior to more expensive and labour intensive approaches. Whilst FT-IR does not provide high levels of functional information, it does enable the collection of a holistic fingerprint that can indicate quantitative changes in peptide, carbohydrate and lipid levels. Therefore, FT-IR could be employed as a first round step where comparisons of non-treated and antibiotic exposed samples are made for numerous bacterial strains and antibiotic combinations. The FT-IR data would then be applied to classify the bacterial strains response as resistant, intermediate resistant, or fully susceptible. An informed selection may then be made to identify which bacterial strains and antibiotic combinations present the most interesting metabolic phenotypes to study in more depth with the lower throughput and financially more expensive chromatography-MS based approaches. The FT-IR data may also be consulted in order to select an appropriate method of chromatography with respect to analysing changes in lipid or polar metabolite composition.

To visualise the distribution of samples on the basis of their IR metabolic fingerprints, a PC-DFA scores plot was generated (Fig. 3a). It was observed that *E. coli* isolates 160 and 173 (ST131), cluster in close proximity to the negative side of the DF1 axis. *E. coli* isolate 161 (non ST131) was observed to separate from the ST131 isolates largely along the DF2 axis, which agrees with the results of a previous FT-IR study (AlRabiah et al. 2013, 2014). Interestingly, *E. coli* isolate 171 (non ST131), which is classed as intermediate resistant, was distantly clustered from all of the other isolates on the positive side of the DF1 axis. The *E. coli* isolates classed as fully resistant (isolate 160 and 161), revealed very little variance between control and Ciprofloxacin challenged samples, suggesting that on the level of the metabolic fingerprint, very little metabolic change occurs as a result of antibiotic challenge. The sensitive *E. coli* isolate 173 and the intermediate resistant isolate 171 both showed a clear separation between the control and Ciprofloxacin challenged sample clusters, with the Ciprofloxacin challenged samples clustering to the negative side of the DF2 axis.

As a next step, the PC-DFA loadings were derived on the basis of the wavelengths responsible for the separation of the control samples from the Ciprofloxacin challenged samples for each respective *E. coli* isolate. The fully resistant *E. coli* isolates, 160 and 161, despite being of different sequence type, ST131 and non-ST131 (resulting in their control samples being clearly clustered from each other in the PC-DFA scores plot), revealed very similar spectral changes associated with Ciprofloxacin challenge, suggesting a similar metabolic shift in the response of the two isolates (Fig. 3b, d). The most pronounced spectral changes due to Ciprofloxacin challenge in the fully resistant isolates included peaks at 1,645 and 1,705 cm^{-1} corresponding to the amide spectral region indicating alterations in protein content in

response to Ciprofloxacin challenge. The sensitive *E. coli* isolate 173 and the intermediate resistant isolate 171 also revealed similar spectral changes associated with Ciprofloxacin challenge (Fig. 3c, e). The most pronounced spectral changes due to Ciprofloxacin challenge in these isolates included peaks at 1,653 and 1,700 cm^{-1} again corresponding to the amide spectral region. Although the preliminary sample screening approach revealed significant protein alterations, this is perhaps unsurprising given that Ciprofloxacin and other fluoroquinolones are predicted to target enzymatic proteins such as DNA topoisomerase II and IV, therefore it may be expected that alterations in protein content would dominate the alterations observed in the lipid, polysaccharide and mixed spectral regions of the IR spectra. Given the tight clustering of samples of the same experimental class within the PC-DFA scores plot (Fig. 3a), it was clear that the within experiment technical reproducibility of sample growth and collection was high and therefore likely suited to UHPLC–MS lipid profiling.

3.3 Reverse phase ultra high performance liquid chromatography–mass spectrometry (UHPLC–MS) based lipid profiling

Following UHPLC–MS analysis in both ESI positive and negative ionisation modes, peak deconvolution was performed using the freely available XCMS software (http://masspec.scripps.edu/xcms/xcms.php) (Dunn et al. 2008; Wedge et al. 2011), producing a MS Excel based XY matrix of mass spectral features (with related accurate *m/z* and retention time variable pairs) × sample, with peak area inputted where the mass spectral feature was detected in each sample. In the UHPLC–MS ESI positive mode and negative mode datasets a total of 2,933 and 1,505 mass spectral features were deconvolved respectively, features showing greater than 20 % relative standard deviation within quality control (QC) samples were removed as were features detected in the first 0.5 min of LC (void volume), resulting in a total of 2,642 and 1,317 features in ESI positive and negative modes respectively.

Following deconvolution, the putative identification of lipid features was performed applying the PUTMEDID-LCMS set of workflows (Brown et al., 2011) (Table S2). An initial appraisal of the putatively identified lipid species suggested that the phospholipids of *E. coli* are predominantly phosphatidyl cholines (PCs) and/or phosphatidyl ethanolamines (PEs). Given previous reports that state the major lipid classes of *E. coli* are represented by PEs, phosphatidyl glycerols (PGs) and Cardiolipins, with minor contributions made by phosphatidyl serines (PSs) and phosphatidic acids (PAs) (Ames 1968), it is perhaps tempting to speculate that the major lipid species detected in this study are more likely to be PEs than PCs. The PC
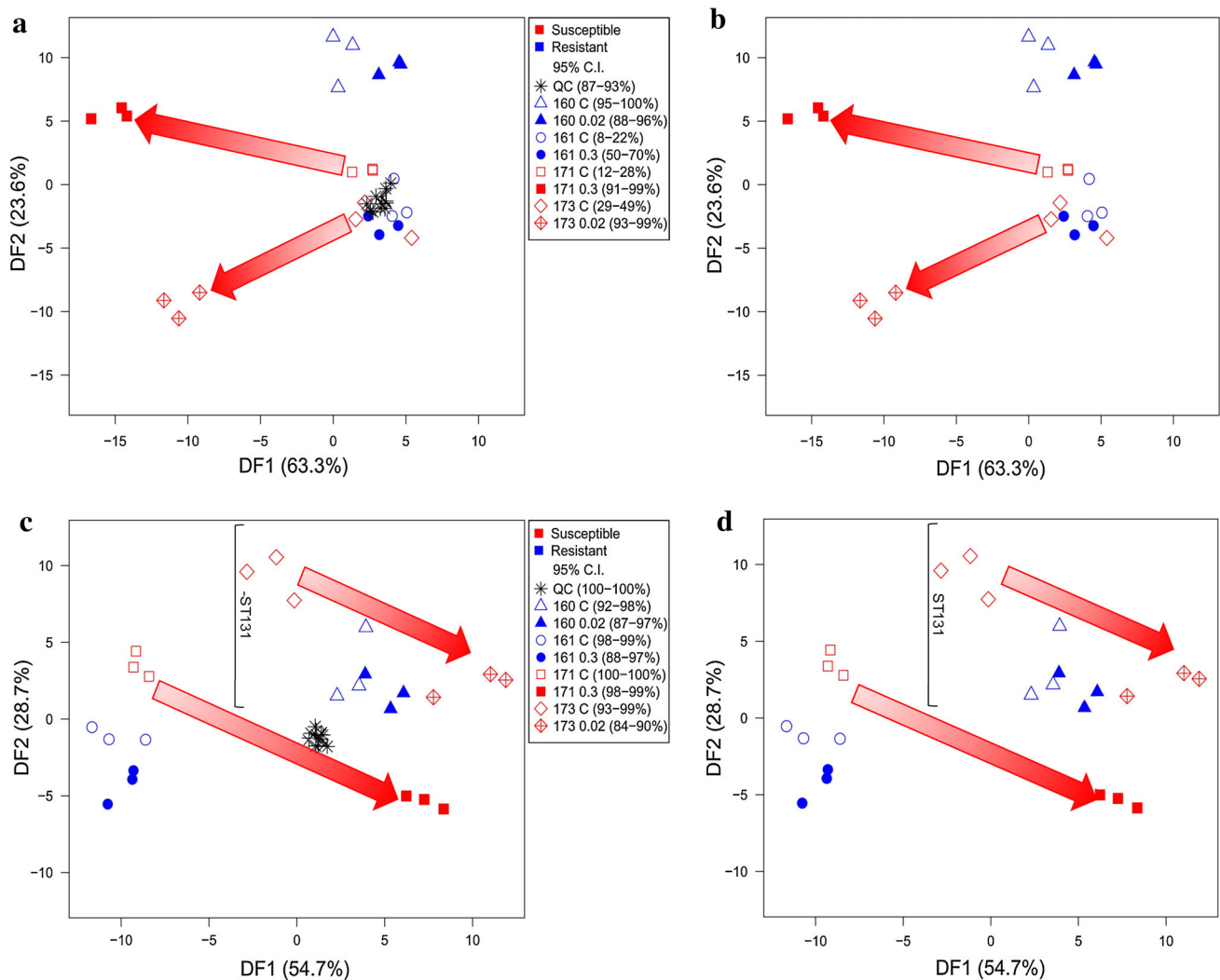
Fig. 4 UHPLC–MS principle component-discriminant function analysis (PC-DFA) of 18 h Ciprofloxacin challenged and control Escherichia coli. PC-DFA scores plot (DF1 vs. DF2) of UHPLC–MS ESI + mode data **a** with QCs included and **b** with QCs removed, PC-DFA scores plot (DF1 vs. DF2) of UHPLC–MS ESI- mode data **c** with QCs included and **d** with QCs removed. Within the figures key, *C* refers to control samples, 0.02 and 0.3 refer to the antibiotic dose (mg/l), the 95 % confidence interval (CI) is given in *brackets*

and/or PE phospholipids were largely detected in ESI positive ionisation mode although a smaller number were also observed in ESI negative mode. Further, a small number of PAs were observed in ESI positive and negative modes and a number of PGs were observed predominantly in ESI negative mode. Further to phospholipids, a large number of acylglycerols and free fatty acids were detected, as well as further diverse lipid species such as sterols, ceramides and sphingolipids. The putative level of identification provided by this approach can be categorised as identification levels 2 and 3 according to the Metabolomics Standards Initiative guidelines (Sumner et al. 2007).

As a first step towards the appraisal of the UHPLC–MS datasets, PC-DFA was applied and score plots were generated (Fig. 4) in order to visualise the distribution of the various samples and sample groups in relation to each other.

The central positioning of the QC samples through the 0 axes of both DF1 and DF2 (Fig. 4a, c), indicate that after deconvolution and quality control steps, the QC samples still represent an average data point, and that the levels of analytical and technical reproducibility were high. Similar to observations within the FT-IR dataset when subjected to PC-DFA (Fig. 3), the UHPLC–MS datasets also indicated that alterations due to Ciprofloxacin challenge within the resistant bacterial isolates (isolates 160 and 161), resulted in only a small level of sample separation within the PC-DFA score plots for both ESI positive (Fig. 4a, b) and negative (Fig. 4c, d) ionisation modes. Perhaps of the greatest significance was the clear differentiation between control and Ciprofloxacin challenged samples of the sensitive *E. coli* isolate 173 and the intermediate resistant *E. coli* isolate 171, with lipid changes due to Ciprofloxacin challenge being associated

a

Phospholipids

PC 27:1 / PE 30:1 ($C_{35}H_{68}NO_8P$)

ESI+ mode Peak no. 2332 RT 14.52 min.
m/z 662.4763

PC 29:1 / PE 32:1 ($C_{37}H_{72}NO_8P$) / PA 34:2
($C_{37}H_{69}O_8P$) /
PE-NMe (O -28:0) ($C_{34}H_{72}NO_6P$)

ESI+ mode Peak no. 2398 RT 14.97 min.
m/z 690.5076

PE 34:1 / PC 31:1 ($C_{39}H_{76}NO_8P$) / PA 36:2
($C_{39}H_{73}O_8P$) / PC (O-28:0) ($C_{36}H_{76}NO_6P$)

ESI+ mode Peak no. 2479 RT 15.46 min. m/z
718.5383

PE 40:1 ($C_{45}H_{88}NO_7P$)

ESI+ mode Peak no. 2730 RT 18.03 min. m/z
786.6402

PC (29:1) / PE (32:1) ($C_{37}H_{72}NO_8P$)

ESI-mode Peak no. 1050 RT 19.71 min. m/z
688.4900

PE-NMe(O -28:0) ($C_{34}H_{72}NO_6P$):

ESI+ mode Peak no. 2458 RT 12.67 min. m/z
696.3792

Diacylglycerols

DG 35:4 ($C_{38}H_{66}O_5$)

ESI+ mode Peak no. 2303 RT 16.72 min.
m/z 647.4589

Non-ST131   ST131

b

Phospholipids

PE (38:2) / PC (35:2) ($C_{43}H_{82}NO_8P$) /
PE-NMe2 (36:2) ($C_{42}H_{80}NO_8P$)

ESI- mode Peak no. 1220 RT 21.17 min. m/z
770.5670

PA(38:4) (C41H73O8P) / PS (32:3) ($C_{38}H_{68}NO_{10}P$) /
2-octadecanoyl -1-hexadecyl-sn-glycero-3-phosphate ($C_{37}H_{75}O_7P$)

ESI- mode Peak no. 1168 RT 19.24 min. m/z
745.4808

PC 32:2 / PC O-32:3 ($C_{40}H_{76}NO_7P$) /
1-(6-[5]-ladderane -hexanyl)-2-(8-[3]-ladderane-octanyl)-sn-
glycerophosphoethanolamine ($C_{43}H_{72}NO_6P$)

ESI+ mode Peak no. 2737 RT 15.25 min. m/z
788.4753

1-16:1-lysoPC ($C_{24}H_{48}NO_7P$)

ESI+ mode Peak no. 2004 RT 12.98 min.
m/z 516.3063

Cholesteryl-beta-D-glucoside ($C_{33}H_{56}O_6$)

ESI-mode Peak no. 899 RT 20.32 min. m/z
593.4069

17-Hydroxy-3-oxo-17alpha-pregna-1,4-diene-21-carboxylic acid,
gamma-lactone / Canrenone / Norethindrone acetate / Norethisterone
acetate ($C_{22}H_{28}O_3$): Steroid Derivatives

ESI-mode Peak no. 369 RT 16.87 min. m/z
339.1980

Others

c

**Phospholipids /
Diacylglycerols**

PA (32:1) (C35H67O8P) / DG (37:7) ($C_{40}H_{64}O_5$) / DG
(35:4) ($C_{38}H_{66}O_5$)

ESI-mode Peak no. 973 RT 20.23 min. m/z
645.4474

PA (34:1) ($C_{37}H_{71}O_8P$)

ESI-mode Peak no. 1021 RT 21.40 min. m/z
673.4791

◄**Fig. 5** UHPLC–MS trend plots of lipids significantly altered in response to Ciprofloxacin: Lipids differentially regulated between susceptible and resistant isolates. **a** Significantly up-regulated in susceptible isolates. **b** Significantly down-regulated in susceptible isolates. **c** Significantly down-regulated in resistant isolates. *Error bars* represent the standard error within the non averaged data for each experimental class

with a trajectory across the DF1 axis for both ESI positive (Fig. 4a, b) and negative (Fig. 4c, d) ionisation modes. The PC-DFA scores plots of the UHPLC–MS datasets overall indicate that whilst there are alterations in lipid profiles of the resistant bacterial isolates (isolates 160 and 161) when challenged by Ciprofloxacin, these are overall minimal in nature compared to the vast alterations in lipid profiles resulting from Ciprofloxacin challenge within the sensitive and intermediate resistant isolates (isolates 173 and 171 respectively).

Following PC-DFA, the next objective was to select which lipids were significantly altered between control and Ciprofloxacin challenged samples for each bacterial isolate independently, by applying significance testing. Two significance tests were applied and a consensus drawn between them. Firstly, a PLS algorithm was applied to the data and the mass spectral features were sorted by the magnitude of their corresponding PLS coefficient values. Secondly, the ANOVA statistical test was applied to all mass spectral features. Finally, starting with the mass spectral feature with the highest PLS coefficient, the top 50 were selected whose p-value computed by ANOVA was less than 0.05 (Table S3 and S4). For each bacterial isolate, the significant features were sorted according to retention time, since certain lipid species were represented by multiple mass spectral features, the top 50 list was next reduced to just the major ion (most intense) observed for each putatively identified lipid species. The top 50 mass spectral features lists for each isolate and ESI polarity, after having been reduced in this manner, were next compiled and a total list of significant lipid mass spectral features across all bacterial isolates was obtained. As a next step, trend plots were generated for each of the significant lipid features, which were then grouped according to their response to Ciprofloxacin: lipid features that were commonly up- or down-regulated across both resistant and sensitive *E. coli* isolates (Fig. S1); lipid features that were up- or down-regulated specifically in susceptible or in resistant isolates (Fig. 5); lipid features that were differentially regulated between resistant and sensitive isolates (Fig. 6).
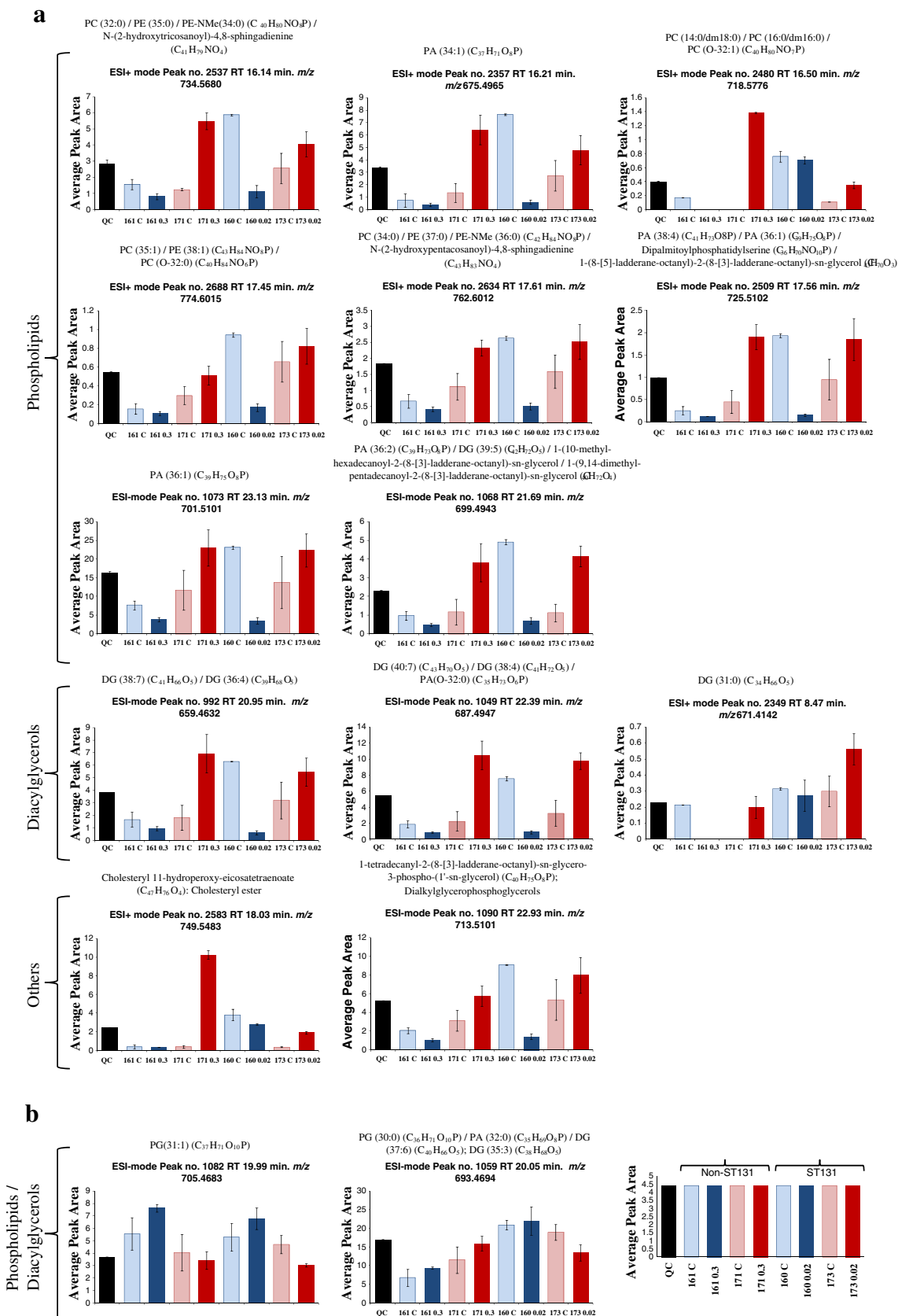
3.4 A wide selection of *E. coli* lipid species are altered by challenge with Ciprofloxacin, with many showing differential regulation between sensitive and resistant *E. coli* isolates

After the statistically significant lipids had been grouped according to their response under Ciprofloxacin challenge,

the patterns of response were considered taking into account the putative assignment of the lipid species and the levels of saturation within the fatty acid components of the lipid species, in order to identify potentially significant trends within the results. Figure S1 reveals a series of lipid alterations that were grouped as showing common responses between both resistant and sensitive isolates. Typically, higher levels of lipid up-regulation were observed within the sensitive isolate 173 and intermediate resistant isolate 171, with the lipids being putatively assigned as PAs, PGs, PAs or PGs, and PGs or DGs (Diacylglycerides). Again, for the lipid species that were down-regulated, typically higher levels of down-regulation were observed in the sensitive isolate 173 and intermediate resistant isolate 171, with the lipids being putatively assigned as PC or PE or PE-NmE (n-Methyl PE) and one lipid putatively assigned as a highly unsaturated DG with total chain composition of 40:10 or 38:7 (dependent on the potential ESI adduct that is formed). The phospholipids that were generally up-regulated, according to their putative assignments, had more saturated fatty acids (typically one or two unsaturated bonds) than the phospholipids that were down regulated which typically had two-to-four unsaturated bonds within the fatty acids.

The lipid features that were up- or down-regulated specifically in susceptible or in resistant isolates are presented in Fig. 5. The lipids that were significantly up-regulated in the sensitive isolates generally showed a higher level of up-regulation within the fully sensitive isolate 173 than the intermediate resistant isolate 171. A total of five phospholipids assigned as PEs or PCs were up-regulated, the putative assignments suggest high levels of saturation within the phospholipid fatty acids, with only one or two unsaturated bonds being present. The lipids that were significantly down-regulated specifically in the sensitive isolates generally showed similar levels of down-regulation between the fully sensitive isolate 173 and the intermediate resistant isolate 171. A small number of phospholipids fell within this response grouping and were putatively assigned as PCs or PEs and PAs or PSs. A lyso-PC and Cholesteryl-beta-D-glucoside were also observed to be down-regulated. Only a small number of lipids were altered specifically within the resistant isolates 160 and 161, all of which were highly down-regulated upon challenge with Ciprofloxacin, these ions were putatively assigned as DGs and PAs.

Perhaps the most interesting series of lipid alterations observed within the investigation, were those that were differentially regulated between the sensitive and resistant isolates (Fig. 6). A large number of phospholipids were found to be down-regulated in the resistant isolates (isolates 160 and 161) and up-regulated in the sensitive and intermediate resistant isolates 173 and 171. Most commonly the significant lipids were down-regulated in the

a

Phospholipids

PC (32:0) / PE (35:0) / PE-NMe(34:0) (C$_{40}$H$_{80}$NO$_8$P) / N-(2-hydroxytricosanoyl)-4,8-sphingadienine (C$_{41}$H$_{79}$NO$_4$)

ESI+ mode Peak no. 2537 RT 16.14 min. *m/z* 734.5680

PA (34:1) (C$_{37}$H$_{71}$O$_8$P)

ESI+ mode Peak no. 2357 RT 16.21 min. *m/z* 675.4965

PC (14:0/dm18:0) / PC (16:0/dm16:0) / PC (O-32:1) (C$_{40}$H$_{80}$NO$_7$P)

ESI+ mode Peak no. 2480 RT 16.50 min. *m/z* 718.5776

PC (35:1) / PE (38:1) (C$_{43}$H$_{84}$NO$_8$P) / PC (O-32:0) (C$_{40}$H$_{84}$NO$_6$P)

ESI+ mode Peak no. 2688 RT 17.45 min. *m/z* 774.6015

PC (34:0) / PE (37:0) / PE-NMe (36:0) (C$_{42}$H$_{84}$NO$_8$P) / N-(2-hydroxypentacosanoyl)-4,8-sphingadienine (C$_{43}$H$_{83}$NO$_4$)

ESI+ mode Peak no. 2634 RT 17.61 min. *m/z* 762.6012

PA (38:4) (C$_{41}$H$_{73}$O8P) / PA (36:1) (C$_{39}$H$_{75}$O$_8$P) / Dipalmitoylphosphatidylserine (C$_{38}$H$_{70}$NO$_{10}$P) / 1-(8-[5]-ladderane-octanyl)-2-(8-[3]-ladderane-octanyl)-sn-glycerol (C$_{41}$H$_{70}$O$_3$)

ESI+ mode Peak no. 2509 RT 17.56 min. *m/z* 725.5102

PA (36:1) (C$_{39}$H$_{75}$O$_8$P)

ESI-mode Peak no. 1073 RT 23.13 min. *m/z* 701.5101

PA (36:2) (C$_{39}$H$_{73}$O$_8$P) / DG (39:5) (C$_{42}$H$_{72}$O$_5$) / 1-(10-methyl-hexadecanoyl-2-(8-[3]-ladderane-octanyl)-sn-glycerol / 1-(9,14-dimethyl-pentadecanoyl-2-(8-[3]-ladderane-octanyl)-sn-glycerol (C$_{41}$H$_{72}$O$_5$)

ESI-mode Peak no. 1068 RT 21.69 min. *m/z* 699.4943

Diacylglycerols

DG (38:7) (C$_{41}$H$_{66}$O$_5$) / DG (36:4) (C$_{39}$H$_{68}$O$_5$)

ESI-mode Peak no. 992 RT 20.95 min. *m/z* 659.4632

DG (40:7) (C$_{43}$H$_{70}$O$_5$) / DG (38:4) (C$_{41}$H$_{72}$O$_5$) / PA(O-32:0) (C$_{35}$H$_{73}$O$_6$P)

ESI-mode Peak no. 1049 RT 22.39 min. *m/z* 687.4947

DG (31:0) (C$_{34}$H$_{66}$O$_5$)

ESI+ mode Peak no. 2349 RT 8.47 min. *m/z* 671.4142

Others

Cholesteryl 11-hydroperoxy-eicosatetraenoate (C$_{47}$H$_{76}$O$_4$): Cholesteryl ester

ESI+ mode Peak no. 2583 RT 18.03 min. *m/z* 749.5483

1-tetradecanyl-2-(8-[3]-ladderane-octanyl)-sn-glycero-3-phospho-(1'-sn-glycerol) (C$_{40}$H$_{75}$O$_8$P); Dialkylglycerophosphoglycerols

ESI-mode Peak no. 1090 RT 22.93 min. *m/z* 713.5101

b

Phospholipids / Diacylglycerols

PG(31:1) (C$_{37}$H$_{71}$O$_{10}$P)

ESI-mode Peak no. 1082 RT 19.99 min. *m/z* 705.4683

PG (30:0) (C$_{36}$H$_{71}$O$_{10}$P) / PA (32:0) (C$_{35}$H$_{69}$O$_8$P) / DG (37:6) (C$_{40}$H$_{66}$O$_5$); DG (35:3) (C$_{38}$H$_{68}$O$_5$)

ESI-mode Peak no. 1059 RT 20.05 min. *m/z* 693.4694

Non-ST131   ST131

QC 161 C 161 0.3 171 C 171 0.3 160 C 160 0.02 173 C 173 0.02

◄**Fig. 6** UHPLC–MS trend plots of lipids significantly altered in response to Ciprofloxacin: Lipids specifically regulated in susceptible or resistant isolates. **a** Up-regulated in susceptible isolates and down regulated in resistant isolates. **b** Up-regulated in resistant isolates and down regulated in susceptible isolates. *Error bars* represent the standard error within the non averaged data for each experimental class

resistant isolates and up-regulated in the sensitive isolate, with the majority of the significant lipids being putatively assigned as PCs or PEs and PAs. A small number of DGs also fell within this response class, with putative assignments suggesting potentially high levels of un-saturation within the fatty acid components. Interestingly, an ion putatively assigned as Cholesteryl 11-hydroxy-eicosatetraenoate was extremely highly up-regulated, but only within the intermediate resistant isolate 171, perhaps indicating an alternative mode of antibiotic exclusion to that observed within the resistant isolates 160 and 161. In comparison, only a very small number of lipids were observed to be up-regulated within the resistant isolates and down-regulated within the sensitive isolate. One such ion was putatively assigned as a PG with highly saturated fatty acids and the other as being either a PA or PG again with highly saturated fatty acids.

Several trends have emerged within the LC–MS lipidomics dataset with the most significant perhaps being that all phospholipids were assigned as PCs or PEs and PAs or PGs in class, with their putative assignments suggesting high levels of saturation within the fatty acids. Alternatively, a large number of ions putatively assigned as DGs were shown to be significant, many of the putatively assigned DGs possessed more unsaturated fatty acids than those observed for the phospholipids. Typically the levels of up- or down-regulation observed for the sensitive isolate 173 and intermediate resistant isolate 171 were much higher than the levels of regulation observed within the resistant isolates, as indicated also by PC-DFA (Fig. 4). Perhaps of further interest, is that the majority of statistically significant lipids, were up-regulated in the sensitive isolate whilst being down-regulated in the resistant isolates, with only small numbers of significant lipids revealing the converse trend. It appears that the resistant isolates are largely characterised by down-regulation of PCs or PEs in response to Ciprofloxacin challenge, whereas the sensitive isolates show high levels of up-regulation of PCs or PEs in response to Ciprofloxacin challenge, perhaps suggesting that high levels of lipid membrane modulation are taking place as the antibiotic enters and takes effect upon the bacterial cell. The developed experimental design has accounted that the only difference between control and Ciprofloxacin challenged samples for each respective strain is the presence of the antibiotic. Therefore it is tempting to make the assumption that all lipid changes are in direct response to the effects exerted on the bacterial cell by the antibiotic. However, previous research has shown that fatty acid alterations can occur as a response to environmental changes including temperature, pH, and osmotic pressure, a phenomena known as phenotypic shift (Crompton et al. 2014). Therefore, with respect to the current study, it must be considered that whilst some of the observed changes in lipid composition in response to Ciprofloxacin may be the result of the antibiotics interaction with the phospholipid membrane, or in the case of resistant strains, the antibiotics exclusion via membrane transport, other lipid alterations may be indirect responses due to the antibiotic altering environmental conditions such as pH within the bacterial culture resulting in phenotypic shift.

Given that previous research has suggested that the major phospholipid species detected in *E. coli* are PEs (Arnes 1968), it is perhaps unsurprising that the majority of lipids that were significantly altered by Ciprofloxacin challenge within this study were putatively assigned as being PEs or PCs. This may suggest that Ciprofloxacin treatment is specifically affecting PEs or PCs, or alternatively that these are the major lipid species of *E. coli* and thus are more likely to have changes in their levels detected due to their higher concentrations and that the alternate species of phospholipids may be close to or under the detection limit of the employed LC–MS instrumentation. However, if the latter was the case, then perhaps one would expect to also observe alterations in the other major lipid classes of *E. coli*, namely PGs and Cardiolipins (Arnes 1968). Several previous investigations, although based upon extracted liposomes (Bensikaddour et al. 2008) or extracted membrane lipids (Leying et al. 1986; Merino et al. 2003) of *E. coli*, rather than on the bacterial cells within culture as performed in this study, also suggested that Ciprofloxacin interacts at the water-phospholipid head group interface of the lipid membrane as revealed by atomic force microscopy (Merino et al. 2003) and also by combination of quasi-elastic light scattering and steady-state fluorescence anisotropy, combined with attenuated total reflectance (ATR)-FT-IR and ^{31}P NMR spectroscopy (Bensikaddour et al. 2008), with the studies suggesting that re-modulation of the phospholipid fatty acids takes place rather than a conversion of headgroup thus forming a different class of phospholipid. Taking these observations into account, it is clearly of much importance that for the approach applied in this study utilising the high sensitivity of LC-LTQ Orbitrap MS to investigate lipid alterations within the "true" bacterial cell to be of greater value, it will be of massive significance to develop online LC–MS/MS methods and offline MS[n] (van der Hooft et al. 2011, 2012; Rojas-Cherto et al. 2012; Roux et al. 2012) analyses of pre-fractionated samples, in order to not just unambiguously assign a lipid identification, but also to identify the levels of saturation and positions of unsaturated

bonds within fatty acids. Once developed, such methods and their unrivalled abilities to unambiguously confirm the identifications of the lipids and elucidate unsaturated fatty acid bond positions will provide the highest levels of information currently possible and will aid greatly our understanding of the effects of Ciprofloxacin modulation of fatty acid structure within the putatively assigned PE and PC phospholipids.

4 Concluding remarks

The LC–MS workflow developed for the analysis of lipid species within *E. coli* when under the challenge of an antibiotic such as Ciprofloxacin, has been shown to be a technically and analytically reproducible method that is useful for discovery phase studies on bacterial modes of antibiotic resistance. Further, by combining a lipid profiling approach with alternative metabolomics platforms suited to profiling primary metabolism (e.g. GC–MS: Winder et al. 2008), the approach may be highly successful for studying the mechanisms underlying antibiotic modes of action within sensitive isolates, as well as mechanisms of resistance. Further, within this study, the developed workflow has been shown to identify a range of interesting and potentially clinically significant lipid alterations in *E. coli* upon challenge with Ciprofloxacin. However, to obtain a more satisfactory level of information, the next step is the development of online RP-UHPLC–MS/MS and offline direct infusion MS^n methods with the LTQ-Orbitrap XL MS system for identification of lipid species across all classes. Such an approach will allow identification of phospholipid classes and fatty acid constituents by employing orthogonal Retention Time information and MS/MS spectral fragmentation compared to analytical standards. By further employing online fraction collection and performing $MS^{3>5}$ level experiments employing collision induced dissociation and/or higher energy collision dissociation methods then the positions of the unsaturated bond positions within the fatty acid constituents of phospholipids may potentially be identified. The current study has suggested the great potential of LC–MS lipidomics for the study of antibiotic modes of action or bacterial modes of resistance, although greater abilities to unambiguously identify the lipid species are required to maximise the biological information content obtained by such a non-targeted lipidomics approach.

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