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Novel applications of mass spectrometry-based proteomics in clinical microbiology

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CHAPTER 5

Proteomic identification of Axc, a novel beta-lactamase with carbapenemase activity in a meropenem-resistant clinical isolate of *Achromobacter xylosoxidans*

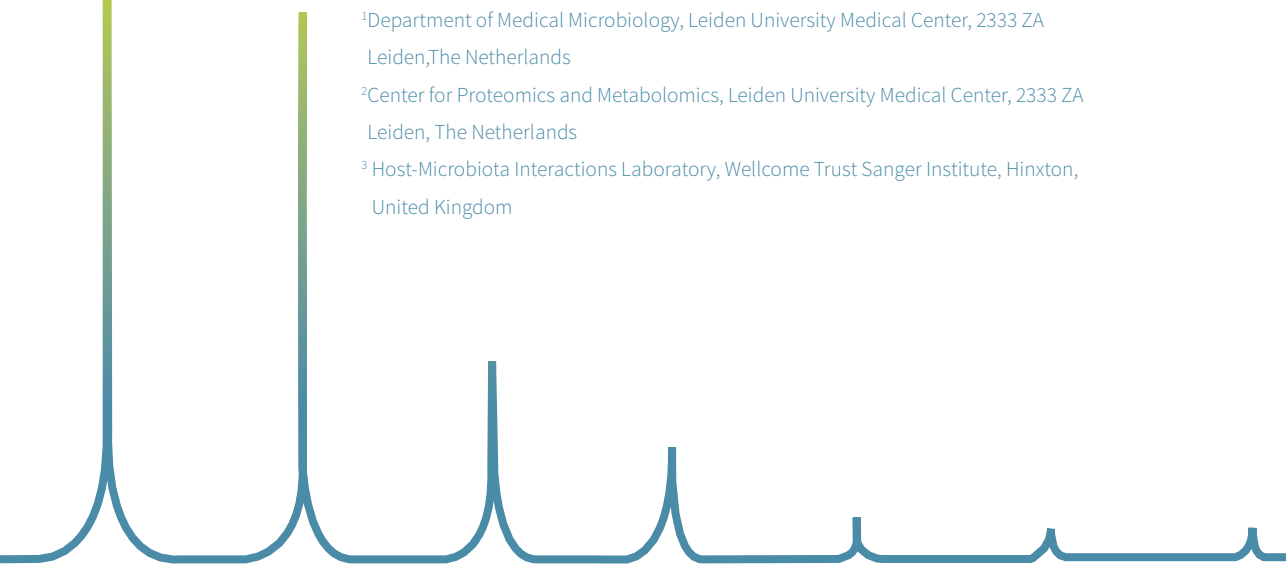
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Abstract

The development of antibiotic resistance during treatment is a threat to patients and their environment. Insight in the mechanisms of resistance development is important for appropriate therapy and infection control. Here, we describe how through the application of mass spectrometry-based proteomics, a novel beta-lactamase *Axc* was identified as an indicator of acquired carbapenem resistance in a clinical isolate of *Achromobacter xylosoxidans*.

Comparative proteomic analysis of consecutively collected susceptible and resistant isolates from the same patient revealed that high *Axc* protein levels were only observed in the resistant isolate. Heterologous expression of *Axc* in *Escherichia coli* significantly increased the resistance towards carbapenems. Importantly, direct *Axc* mediated hydrolysis of imipenem was demonstrated using pH shift assays and $^1\text{H-NMR}$, confirming *Axc* as a legitimate carbapenemase. Whole genome sequencing revealed that the susceptible and resistant isolates were remarkably similar.

Together these findings provide a molecular context for the fast development of meropenem resistance in *A. xylosoxidans* during treatment and demonstrate the use of mass spectrometric techniques in identifying novel resistance determinants.

Introduction

Development and spread of antibiotic resistance by pathogenic microorganisms is an increasing healthcare problem. Moreover, certain resistance determinants spread readily^{1,2}, while the introduction of novel antibiotics is lagging behind. Several clinically important classes of antimicrobials such as the beta-lactams, target the bacterial cell wall³. Resistance to beta-lactams can be mediated by beta-lactamases that are capable of hydrolysing the beta-lactam ring. Following the initial introduction of penicillin, second and third generation beta-lactams have been developed which, in turn, triggered the selection of beta-lactamases with broader specificities. Carbapenem treatment is often used as a last resort, since extended-spectrum beta-lactamases (cephalosporinases) are becoming more prevalent in Gram-negative bacteria. The emergence and spread of carbapenemases such as class A KPC⁴, a number of metallo beta-lactamases^{5,6} (class B: IMP, VIM, NDM) and class D oxacillinases such as OXA-48⁷, in combination with other resistance mechanisms⁸, can jeopardize carbapenem efficacy, leaving little or no treatment options for patients.

Achromobacter xylosoxidans is a rod shaped aerobic non-fermentative Gram-negative bacterium. It is widespread in the environment and generally considered as an opportunistic pathogen. Chronic infections with *A. xylosoxidans* are problematic in cystic fibrosis patients^{9,10} but reported prevalence numbers vary greatly (3-30%)^{11,12}. Moreover, bacteremia as a result of *A. xylosoxidans* can occur in immunocompromised patients¹³. *A. xylosoxidans* is notorious for its intrinsic high level of resistance, especially towards penicillins and cephalosporins¹⁴⁻¹⁶. In general, carbapenem resistance in *A. xylosoxidans* is not widespread and as a result meropenem treatment is routinely applied, even in the case of recurring infections^{17,18}. Though carbapenem resistance is observed, specifically for meropenem¹⁹, there are few reports on the mechanism of carbapenemase resistance in *A. xylosoxidans*. Notable exceptions are the plasmid-encoded carbapenemase IMP^{20,21} and the chromosomally encoded class D beta-lactamase OXA-114¹⁶, that show low level carbapenemase activity. A comparative genomic exploration of two *A. xylosoxidans* isolates revealed many genes that could be involved in drug resistance, such as efflux pumps and β -lactamases. However, most of these genes were conserved between carbapenem susceptible and resistant strains, highlighting the difficulty in translating genomic data to the observed resistant phenotypes²².

In this study, two clinical isolates of *A. xylosoxidans* from an immunocompromised patient with pneumonia were investigated. The initially cultured isolate from the respiratory tract was susceptible to meropenem and treatment was started accordingly. However, a subsequent meropenem resistant isolate was obtained from a blood culture after treatment failure. Since PCR analysis was negative for known carbapenemases, we performed a proteomic analysis which revealed the novel beta-lactamase Axc as highly abundant in the meropenem-resistant, but not in the susceptible isolate. Axc expression led to an increase of minimal inhibitory

concentrations for carbapenems when introduced in a susceptible *Escherichia coli* strain and direct carbapenemase activity of Axc was demonstrated using *in vitro* imipenem conversion assays. Interestingly, the resistant and the susceptible clinical isolates are genetically almost identical, emphasizing the importance of mass spectrometry as a technique to investigate carbapenem resistance in *A. xylosoxidans*.

Results

Development of meropenem resistance in *Achromobacter xylosoxidans* during treatment

An antibiogram of a clinical isolate of *Achromobacter xylosoxidans* from a patient treated for pneumonia (see Methods for details about patient history and treatment) revealed a multi-resistant character, but it was susceptible to meropenem (Supplemental Table 1). Hence, meropenem treatment was initiated. During treatment, the patient developed a pneumothorax and died from septic shock 6 days later. An antibiogram of a blood culture from a sample taken one day before the patient's death revealed a meropenem-resistant *A. xylosoxidans* phenotype (Supplemental Table 1). Subsequently, pure cultures from the first and second clinical isolate were prepared (AchroS and AchroR, respectively). In line with the antibiogram analysis described above, Etests showed that both isolates were resistant to imipenem (MIC > 32 mg/L for both) but differed in their susceptibility towards meropenem (MIC of 0.0094 mg/L (first isolate, AchroS) and 2 mg/L (second isolate, AchroR), respectively). An in-house multiplex PCR assay (based on a published PCR³⁴) failed to detect common carbapenemases (KPC, IMP, VIM, NDM and OXA-48, data not shown), suggesting that the change in resistance is not mediated by these enzymes. This finding prompted us to perform a comparative proteomic analysis of the two isolates to attempt to identify the meropenem resistance mechanism.

Comparative proteomic analysis shows differential levels of the beta-lactamase Axc

Protein extracts of the meropenem-susceptible and resistant *A. xylosoxidans* isolates (AchroS and AchroR) were first analysed by SDS-PAGE. Since no major visual differences were observed (Supplemental Figure S1A), all bands were excised and processed for a bottom-up LC-MS/MS proteomic analysis. In total, 2276 unique proteins were identified, of which 1519 proteins were common to both isolates, while 223 and 534 were only found in the susceptible and resistant isolates, respectively. For a semi-quantitative analysis, the spectra assigned to peptides belonging to a certain protein were counted and compared between the two different isolates (Supplemental Figure S1B and S2, Table S2). Of the uniquely observed proteins, most were proteins with low spectral counts (often single peptide identifications), likely resulting from sampling bias of low abundant proteins. One protein was observed with 102 spectra in the resistant isolate (AchroR) but none in the susceptible isolate (AchroS). This

protein, hereafter called Axc (for *Achromobacter xylosoxidans* carbapenemase, GenBank ID: MF767301), is a putative PenP class A beta-lactamase (COG2367/pfam13354).

To confirm that Axc was highly abundant in the resistant isolate in comparison to the susceptible isolate, a second proteomic analysis was performed on whole cell protein extracts that were digested in-solution and analysed without any prior fractionation. Spectral count analysis resulted in a cumulative quantification of 1342 different proteins, of which 981 were found in both isolates, but 196 and 165 were uniquely quantified in the susceptible and resistant isolates, respectively. The spectral count plot reflects the high similarity of the two clinical isolates, with the vast majority of the proteins distributed along the diagonal (Figure 1A, Table S2, Supplemental figure S2). In accordance with the data described above, Axc was the most prominent outlier in the resistant clinical isolate. A number of Axc tryptic peptides (Figure 1B) was clearly visible in the LC-MS/MS analysis of the resistant but not the susceptible isolates (Figure 1C). Of note, cells used for these analyses were grown in the absence of meropenem, so the high level of Axc in the resistant isolate was independent of antibiotic pressure.

Axc is present in meropenem resistant and susceptible *A. xylosoxidans* isolates

To investigate whether *A. xylosoxidans* acquired the *axc* gene in the course of the treatment, we performed a PCR for the *axc* open reading frame on both the resistant and susceptible isolates AchroS and AchroR. This analysis demonstrated that *axc* was present in both clinical isolates (Supplemental Figure S3A). Moreover, Sanger sequence analysis demonstrated that the sequence of *axc* was identical in both isolates (data not shown), indicating that the observed meropenem resistance was not likely caused by an alteration of protein function.

A database search revealed that the *axc* gene is not present in all *A. xylosoxidans* strains (Figure 2A). Like in our clinical isolates, *axc* is present in the NH44784-1996 strain (ENA assembly GCA_000967095.1) 35 but not in the strains NBRC 15126/ATCC 27061 (ENA assembly GCA_000508285.1) and C54 (ENA assembly GCA_000758265.1) for instance. In those strains that contain *axc*, the gene is located next to a putative LysR-type transcriptional regulator (Pfam 03466), hereafter *axcR* (for *axc*-associated regulator). Additional PCR and Sanger sequencing experiments verified that the intergenic regions between *axc* and *axcR* in AchroS and AchroR were identical, However, they differed at two positions with the intergenic region between *axc* and *axcR* in strain NH44784_1996 (Supplemental Figure S3B).

The results showed a high degree of similarity between the meropenem resistant and susceptible strain, raising the possibility that these two strains represented a clonal complex. To further explore the relatedness of both isolates, whole genome sequencing (WGS) analysis was performed. This showed that both patient isolates were highly similar, with only one single-nucleotide polymorphism (SNP) within the gene encoding AxyZ 36. This SNP was

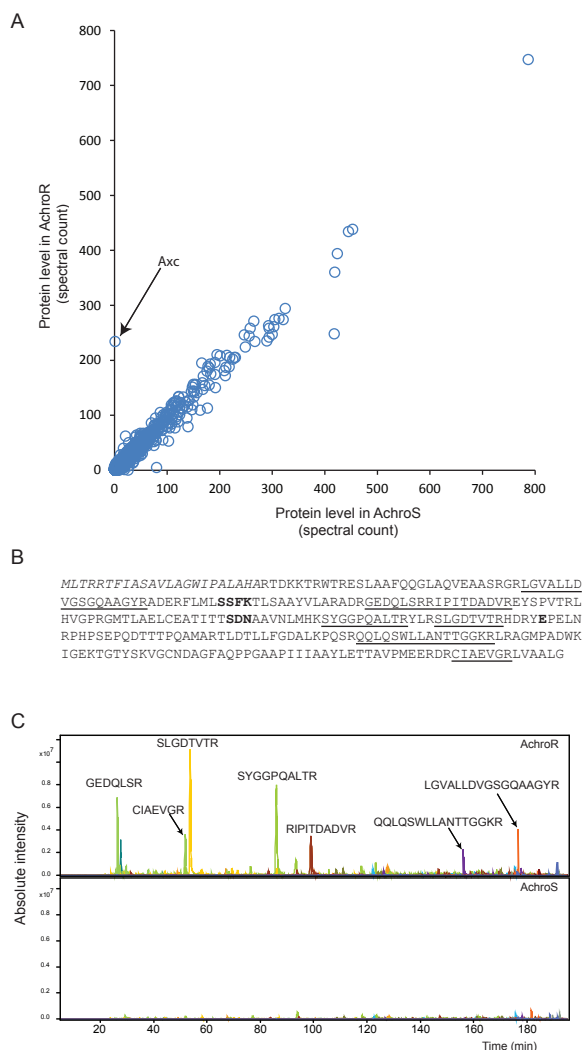


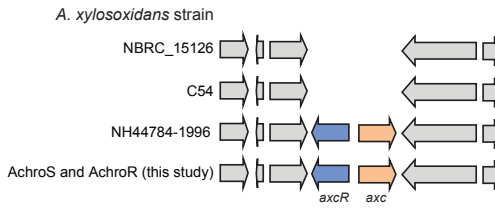
Figure 1: Comparative proteomic analysis of meropenem resistant and susceptible *Achromobacter xylosoxidans* clinical isolates

A: Tryptic digests of protein extracts of the meropenem resistant (AchroR, Table 1) and susceptible (AchroS, Table 1) isolate were analysed by LC-MS/MS. Spectra were assigned to peptides based on database searching. Identified spectra were then assigned to the corresponding proteins and the total number of spectra per protein were counted. Each circle represents one protein with the number of spectra observed in the resistant and the susceptible isolate. Hence, proteins on the diagonal were observed in similar counts in both isolates. Axc (arrow), a classA PenP-family beta-lactamase, is the most prominent outlier. See also Table S2 and Supplemental figure S2.

B: The full amino acid sequence of Axc, with the peptides identified by LC-MS/MS analysis underlined. Conserved residues from serine beta-lactamases, Ser-X-X-Lys, Ser-Asp-Asn and the active site Glu, are in bold (37).

C: Extracted ion chromatograms of m/z values corresponding to tryptic peptides of Axc in the meropenem resistant isolate (AchroR, upper panel) and susceptible isolate (AchroS, lower panel). The corresponding tryptic peptides are indicated above the corresponding peaks.

A



B

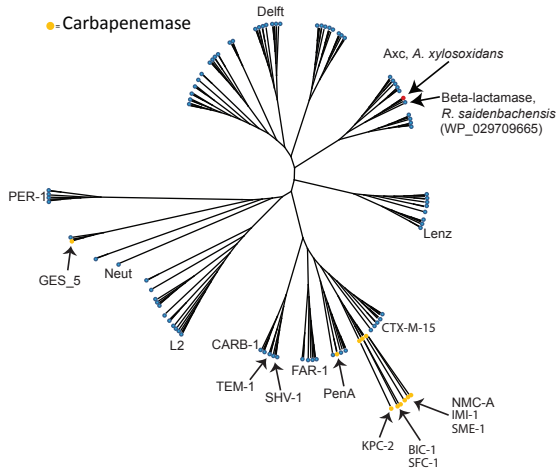


Figure 2: Genomic context of *axc* in *Achromobacter xylosoxidans* strains and comparison of *Axc* with other class A beta-lactamases

A: *Axc* (a putative PenP class A beta-lactamase, 1.17 e⁻⁵⁴) and the gene encoding its putative transcriptional repressor (*axcR*), were found in both clinical isolates (AchroS and AchroR, Table 1). Three other fully sequenced genomes of *Achromobacter xylosoxidans* were examined for the presence of *axc*; NH44784_1996 (ENA assembly GCA_000967095.1), C54 (ENA assembly GCA_000758265.1) and NBRC_15126/ATCC27061 (ENA assembly GCA_000508285.1). Only within the strain NH44784_1996 (used as a reference to search our proteomics data), *axc* and the putative regulator are also present.

B: Unrooted cladogram obtained for 176 class A beta-lactamases including *Axc*. The class A β-lactamase protein sequences of Gram negative bacteria were obtained by querying the refseq_protein database using Blastp (<1e-10, <http://blast.ncbi.nlm.nih.gov/blast/Blast.cgi>) and a consensus β-lactamase-alignment (37). Duplicate sequences and sequences causing a strong overrepresentation of branches produced in the tree were removed. Names of known carbapenemases (orange) and names of identifier of branches (37) are indicated. *Axc* is indicated with a red dot. The closest homologue to *Axc*, a β-lactamase from *R. saidenbachensis* (WP_029709665), is also indicated.

confirmed by PCR and subsequent Sanger sequencing, ruling out the possibility that this was an artefact of the assembly procedure. Moreover, *axc* and its putative regulator *axcR* were found to be located in the same genomic region as in NH44784_1996 strain (Figure 2A). Overall, the genome sequences suggest that both isolates are clonal, and that the meropenem resistance evolved within the same strain during the course of treatment.

To demonstrate how *Axc* is related to other class A beta-lactamases, we compared the *Axc* amino acid sequence with the sequence of another 176 representatives of this family using an alignment consensus based on a report by Walther-Rasmussen and Hoiby 37. The resulting unrooted cladogram shows that *Axc* is most closely related to a class A beta-lactamase of *Rhodofera* *saidenbachensis* (WP_029709665, Figure 2B). Only a limited number of class A beta-lactamases have activity towards carbapenems, but none of these cluster with the *Axc* sequence (Figure 2B).

Functional characterisation of *Axc*

To establish whether *Axc* indeed has activity towards carbapenems, *Axc* was expressed in a heterologous host and hydrolysis of carbapenems was measured indirectly and directly. The *E. coli* C43 strain, suitable for the expression of toxic proteins³⁰, is susceptible to carbapenems. We generated a derivative of C43 that allows for IPTG-dependent expression of a plasmid-based copy of *axc* (*E. coli_Axc* (JC107)). Susceptibility testing for imipenem and meropenem showed that the MICs for these carbapenems increased 8-fold, following induction of *Axc* expression (Table 2). Though these levels were lower than for the positive control (KPC) for our assay, they were specific for *Axc* as the expression of an unrelated protein (PPEP-1³¹) did not lead to an increase in MIC values (Table 2). Thus, expression of *Axc* is correlated to resistance towards carbapenems.

To directly demonstrate carbapenemase activity, hydrolysis of imipenem was monitored *in vitro* through colorimetric assays (Figure 3A) and ¹H-NMR (Figure 3B). Imipenem hydrolysis results in the formation of a carboxylic acid, and monitoring the accompanying pH drop colorimetrically is a well-established method for the detection of carbapenemase activity^{32,38}. Indeed, hydrolysis of imipenem was readily observed using KPC cells (without IPTG). Consistent with our previous results, imipenem hydrolysis was observed for *E. coli* cells harbouring the *Axc* expression plasmid (*E. coli_Axc*) grown in the presence, but not in the absence, of IPTG. As before, these results were specific for *Axc*, as induction of PPEP-1 expression (*E.coli_control*) did not result in imipenem hydrolysis (Figure 3A). In a parallel assay, ¹H-NMR was used to directly observe the opening (hydrolysis) of the lactam ring in imipenem (Figure 3B). The chemical shifts of the peaks change as a result of this hydrolysis, with the protons closest to the ring opening undergoing the largest change. The H-6 proton of imipenem generates an adequately resolved multiplet at 3.42 ppm that decreases in intensity upon hydrolysis. Concomitantly, the doublet generated by the protons of the

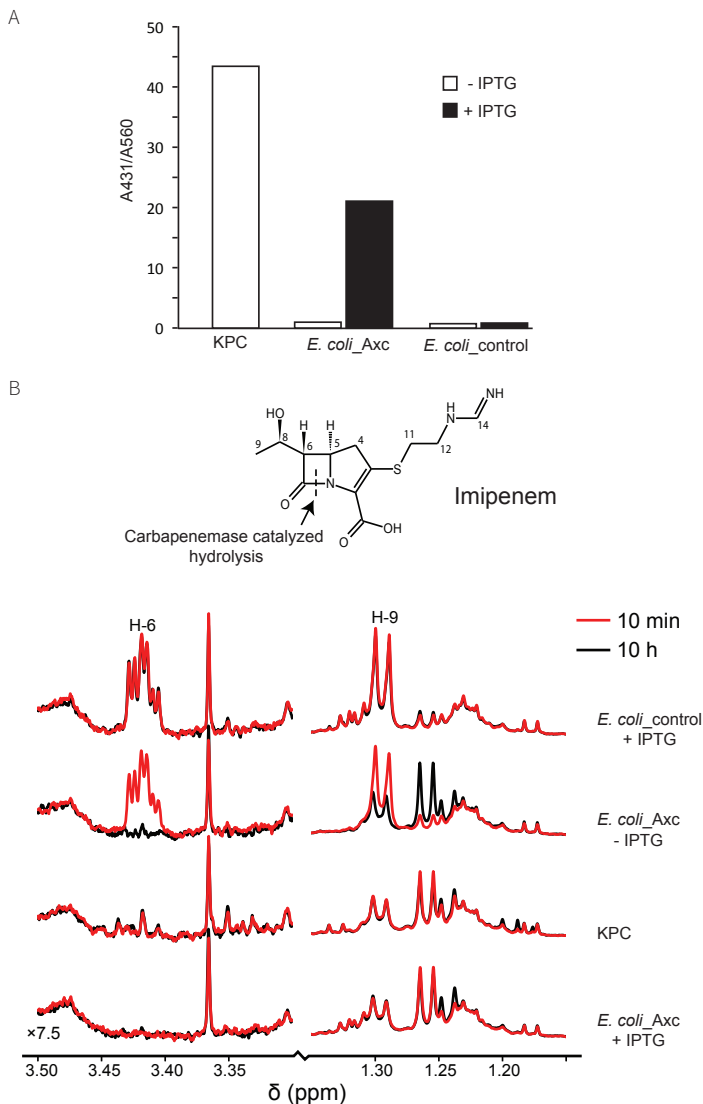


Figure 3: Axc has carbapenemase activity.

A: Axc was expressed in *E. coli* and cell extracts were tested for the ability to hydrolyse imipenem. Bars show the A431/A560 ratio, which is a measure for the shift in pH due to imipenem hydrolysis.

B: NMR-based identification of imipenem hydrolysis by Axc. The structure of imipenem is shown, with the proton numbering used in the spectrum. The line color indicates the incubation time (red, 10 min at room temperature or black, 10 hrs at 6 °C). Imipenem hydrolysis is accompanied by the loss of the H-6 multiplet at 3.4 ppm, and a shift in the H-9 doublet, resulting in a decrease of the doublet at 1.3 ppm.

Strains used (see also Table 1): KPC (JC113): Carbapenem resistant *Klebsiella pneumoniae*. *E. coli* Axc (JC107): *E. coli* strain C43(DE3), containing plasmid pET21-Axc; *E. coli_control* (JC108): *E. coli* strain C43, containing plasmid pET21-PPEP-1 (Pro-Pro endopeptidase 1). IPTG: isopropyl β -D-1-thiogalactopyranoside

°methyl group (H-9) move upfield, resulting in a decrease of the doublet at 1.3 ppm. After 10 min incubation of bacterial cells with imipenem, hydrolysis of imipenem was observed with *E. coli*_Axc grown in the presence, but not in the absence, of IPTG. Hydrolysis was also apparent for KPC, but not for *E. coli* cells expressing PPEP-1 (*E. coli*_control), even after long incubation times (10h). Under these conditions, imipenem hydrolysis was also observed for samples with *E. coli* cells harbouring the Axc expression plasmid grown in absence of IPTG, due to leaky expression from the inducible promoter.

Taken together, our data establish that Axc has carbapenemase activity.

Discussion

In this paper we identified a new resistance mechanism that explains the difference in meropenem susceptibility of two clinical isolates of *A. xylosoxidans* that were collected during treatment. Using a combination of comparative proteomic analyses and functional assays, we have shown that the class A beta-lactamase Axc is highly abundant in the meropenem resistant isolate in comparison to the susceptible isolate and that Axc has carbapenemase activity.

Detection of carbapenemases from sequence data is challenging. First, carbapenemases belong to different subgroups of beta-lactamases which have probably evolved by convergent evolution. Although sequence identities are moderate, most of the class A carbapenemases have a disulphide bridge between Cys-69 and Cys-238, but this is dispensable for activity against carbapenems³⁹ and our data show that Axc does not contain these residues. Next, some studies also revealed a mechanism where beta-lactam trapping, without actual degradation, can be involved in resistance towards carbapenems when levels are sufficiently high. In such cases, concomitant loss of porins is often observed^{40,41}. Thus, it is crucial to determine whether a beta-lactamase actually induces hydrolysis of carbapenems. Our NMR and pH-shift analyses data clearly demonstrate Axc-mediated opening of the beta-lactam ring in imipenem. Finally, many unexplained mechanisms of carbapenem resistance remain. For instance, a recent study demonstrated plasmid derived carbapenem resistance in *Klebsiella pneumoniae* strains which could not be explained by the most commonly found carbapenemases (KPC-type). Even though none of the plasmid-encoded genes were obvious candidates for the observed resistance towards carbapenems, several TEM-homologs were detected⁴². Though the prediction of the activity of a certain beta-lactamase against carbapenems is not straightforward studies such as the present one highlight that mass spectrometry approaches can be used to gain insight in the mechanism of action and role of specific proteins in the observed phenotypes.

We do not know whether the high level of Axc is the only mechanism conferring meropenem resistance to our isolate of *A. xylosoxidans*. When expressed from an inducible promoter, Axc confers moderate resistance to carbapenems to *E. coli*; MIC values compared to the KPC strain suggest that Axc has a lower efficiency than KPC, but may also indicate lower overall levels of expression. Differing efficiencies in carbapenemases are well documented, to the point where the activity of a specific class, such as OXA-48, is difficult to detect but of great clinical importance⁴³. Full biochemical characterization of Axc, including kinetic experiments, is subject to further study. Such experiments, in combination with crystallography analysis, will provide more insight in the activity of Axc against different beta-lactams and could resolve the structural characteristics of the binding pocket which facilitates its activity towards carbapenems. We note, however, that two other changes in the antibiogram between the meropenem susceptible and resistant isolates involve beta-lactam antibiotics. Augmentin (amoxicillin/clavulanate) resistance changed from intermediate (8 mg/L) to resistant (>32 mg/L), and piperacilline/tazobactam from susceptible (<=4 mg/L) to intermediate resistance (8 mg/L). This suggests that Axc may have a broad substrate specificity and is insensitive to inhibition by clavulanate and tazobactam. Strikingly, both the meropenem-susceptible and meropenem-resistant isolates were resistant to imipenem (MIC values higher than the maximum concentration tested (32 mg/L)). This indicates that, notwithstanding the activity of Axc towards imipenem as presented here, imipenem resistance in the clinical isolates is not dependent on Axc. Differences in the sensitivities towards different carbapenems results from the chemical differences between the individual drugs⁴⁴ and are often linked to the differential permeability of the outer cell membrane^{45,46}.

The regulatory mechanism leading to higher levels of Axc expression are unclear. Sequence analyses showed that the meropenem susceptible and resistant *A. xylosoxidans* clinical isolates are highly similar, with no differences in the *axc* promoter and coding sequence, nor in its putative regulator AxcR and the *axc-axcR* intergenic region. The only SNP we identified is located in the gene encoding AxyZ, the TetR-type repressor of the *axyXY-oprZ* operon⁴⁷. This leads to an amino acid substitution (V29G) in a region of AxyZ that is involved in DNA binding in other members of TetR family⁴⁸. AxyX, AxyY and OprZ form an efflux pump of the resistance-nodulation division (RND) family and are predominantly found in aminoglycoside resistant *Achromobacter* species⁴⁹⁻⁵¹. A recent paper showed higher expression levels of *axyY* in *A. xylosoxidans* strains containing AxyZ_Gly29, suggesting that this mutation leads to reduced repression of AxyZ⁵². Closer inspection of our proteomics data indicates that also AxyX and AxyY are more abundant in the resistant isolate (spectral counts of 62 vs 21 for AxyX and 10 vs 2 for AxyY in the results presented in Figure 1 (see Table S2)). However, the difference is not as pronounced as found for Axc and more accurate quantitative proteomics experiments have to be performed to validate these data.

Mutations in TetR-like repressors have been linked to differences in carbapenem resistance⁵³. For example, a 162 bp deletion in *axyZ* has been identified in certain carbapenem resistant strains⁵⁴. However, even though *AxyZ_Gly29* leads to higher expression of *axyY*, resulting in higher MIC values for aminoglycosides, fluoroquinolones and tetracyclines, no correlation between *axyY* expression and meropenem resistance was observed⁵². In line with this observation, a deletion of *axyY* in several *A. xylosoxidans* strains is reported to result in only a modest increase in the susceptibility towards carbapenems⁴⁷. Taken together, it is likely that *AxyXY-OprZ* per se contributes little if any to the meropenem resistance phenotype of our clinical isolate. Instead, our data show a critical role for *Axc* and suggest that *axc* expression is regulated by *AxyZ*. If this is indeed the case, we postulate that the increase in the meropenem MIC for the ACH-CF-911_{v29G} strain⁵² is accompanied by increased *Axc* levels.

From the clinical perspective, the development of resistance to meropenem within days following meropenem treatment is remarkable. Previous longitudinal analyses of different *A. xylosoxidans* isolates from one patient have revealed large phenotypic and genetic differences, for example in the resistance towards different classes of antibiotics but they were generally performed over longer time periods^{35,53,54}. Such changes are believed to be the result of adaptive evolution of the initial strain which infected the patient, but there is also evidence that genetically different strains of *A. xylosoxidans* can co-exist within the same chronically infected individual⁵⁵. Though we cannot exclude a co-infection, it is likely that the mutation in *axyZ* in our case occurred during treatment.

Finally, from a diagnostic point of view, the presence of *Axc* in both the sensitive and resistant *A. xylosoxidans* isolates complicates straightforward detection by molecular methods in the future and warrants detection based on protein abundance levels. Even though more elaborate mass spectrometry-based platforms clearly have potential for the detection of resistance and virulence proteins^{23,56}, as also exemplified in our study, it will take more time and effort before they could find their way into the medical microbiology laboratory to complement the now well-established application of MALDI-ToF-MS for bacterial species identification.

Methods

Patient history

All procedures performed in studies involving human participants were in accordance with the ethical standards of LUMC Medical Ethical Committee and with the 1964 Helsinki Declaration and its later amendments or comparable ethical standards.

A 65-year old patient, diagnosed with chronic lymphocytic leukemia in 1989, underwent a non-myeloablative stem cell transplantation in July 2014. The patient was discharged on July 25th but readmitted on August 1st with a mucositis-associated neutropenic fever and blood cultures positive for *Gemella haemolysans*. The patient was treated with cefuroxim intravenously but developed pneumonia one week later. Analysis of a sputum sample showed a microscopically and culture confirmed fungal infection and abundant growth of a Gram-negative bacterium. This bacterium was identified as *A. xylosoxidans* by matrix-assisted laser desorption/ionization time-of-flight mass spectrometry. The patient was treated with liposomal amphotericin B for the fungal infection, but this did not result in improvement of the patient's status. Due to clinical deterioration the patient received ceftazidim (3 dd 1 gram intravenously) between September 3rd and 9th to treat the ceftazidim sensitive *A. xylosoxidans* isolate. On September 10th, the antibiogram (Supplemental Table 1), revealed a multi-resistant character for the *A. xylosoxidans* isolate, but it was susceptible to meropenem. Hence, meropenem treatment was initiated (4 dd 1 gram intravenously) on September 10th. During treatment the patient developed a pneumothorax and died from septic shock 6 days later. Because of this, informed consent of the patient was not possible to obtain. We therefore anonymized all patient data and presented the data in such a form that it is not traceable to the patient.

Materials

MilliQ water was obtained from a Q-Gard 2 system (Merck Millipore, Amsterdam the Netherlands). Acetonitrile of LC-MS grade was obtained from Biosolve (Valkenswaard, the Netherlands). Porcine trypsin was purchased from Promega (Madison, WI). If not indicated otherwise, chemicals were from Sigma Aldrich (St Louis, MN, USA).

Susceptibility profiling of clinical isolates

The minimum inhibitory concentrations (MICs) for different antibiotic compounds on the clinical isolates (Clinical IDs: M 14073954-7 (first isolate), M 14076260-2 (second isolate)) were initially determined using a Vitek-2 system (bioMérieux, Marcy-l'Étoile, France). Several colonies of plate grown cultures were inoculated and suspended in 0.45% sterile physiological saline solution. Suspensions for testing had densities between of approx. 0.5 McFarland standards. The testing procedure was performed according to the manufacturer's instructions.

Pure cultures of the susceptible and resistant isolates (AchroS and AchroR, respectively (Table 2)) were subjected to further susceptibility testing and used for the proteomic and genomic analyses. Etests (bioMérieux) were performed according to the recommendations from EUCAST (http://www.eucast.org/clinical_breakpoints/). EUCAST does not provide recommendations for the interpretation of MIC values or clinical breakpoints for *Achromobacter xylosoxidans*. Therefore, scoring was performed using locally developed

protocols that are based on clinical breakpoints for other non-fermentative Gram-negative rods.

Mass spectrometry-based proteomics

Cells were grown in BHI (Oxoid, Basingstoke, UK). Cells were collected by centrifugation (4000xg, 5 minutes) from 1 mL cell culture and washed with phosphate-buffered saline (PBS, pH 7.4). Pellets were stored at -80 °C until further use.

Two different proteomics experiments were performed. For the first, cell extracts from *A. xylosoxidans* strains (AchoS and AchoR, Table 2) were prepared in LDS (Lithium dodecyl sulphate) sample buffer (Novex, Thermo Scientific) and put at 95 °C for 5 minutes for cell lysis and protein extraction. Proteins were separated on Novex precast 4–12% Bis-Tris gels (Thermo Scientific, Waltham, MA, USA) with MOPS (3-(N-morpholino)propanesulfonic acid) running buffer (Thermo Scientific). After overnight staining using a Colloidal Blue Staining Kit (Thermo Scientific), destained gel lanes were processed into 31 slices per lane. Gel pieces were sequentially washed with 25 mM ammonium bicarbonate and acetonitrile. Reduction and alkylation were performed with dithiothreitol (DTT, 10 mM, 30 minutes at 56 °C) and iodoacetamide (IAA, 55 mM, 20 minutes at room temperature) respectively. Following several washes with 25 mM ammonium bicarbonate and acetonitrile, bands were overnight digested with trypsin (12.5 ng/mL in 25 mM NH₄HCO₃). Digest solutions were lyophilized and reconstituted in 0.5% trifluoroacetic acid (TFA). Nano-LC separation was carried out using a Ultimate 3000 RSLC nano System equipped with an Acclaim PepMap RSLC column (C18, 75 µm x 15 cm with 2 µm particles, Thermo Scientific) preceded by a 2 cm Acclaim PepMap100 guard column (Thermo Scientific). Peptide elution was performed by applying a mixture of solvents A and B with solvent A being 0.1% formic acid (FA) in water and solvent B 0.1% FA in 80 % acetonitrile (ACN). Peptides were eluted from the column with a multi-step gradient from 5% to 55% solvent B in 55 minutes (0-5 minutes 5%B, then in 15 minutes to 20%B and finally in 25 minutes till 55%B), at a constant flow rate of 300 nl min⁻¹. MS analysis was performed employing a maXis Impact UHR-TOF-MS (Bruker Daltonics, Bremen, Germany) in data dependent MS/MS mode in the *m/z* 150-2200 range. The source settings were as follows: voltage 1500 V, dry gas 3.0 l.min⁻¹, temperature 150 °C. Ten ions were selected at a time, based on relative abundance and subjected to collision-induced-dissociation with helium as collision gas. A 1 minute dynamic exclusion window was applied for precursor selection. The precursor isolation window and collision energy were *m/z* dependent.

For the second proteomics approach, in-solution digests were prepared as described previously²³. In short, following cell disruption proteins were solubilized in 50% trifluoroethanol (TFE). Subsequent reduction with DTT and alkylation with IAA were performed prior to overnight tryptic digestion. Samples were lyophilized and reconstituted in 0.5% TFA for injection. The nano-LC system and solvents were the same as in the above experiment, but

using an Acclaim PepMap RSLC column (C18, 75 μm x 50cm with 2 μm particles, Thermo Scientific) with a 2 cm Acclaim PepMap100 guard column (Thermo Scientific). Peptides were eluted from the column with a multi-step gradient from 5% to 55% of solvent B in 180 minutes (0-5 minutes 5%B, then in 65 minutes till 13%B, then in 90 minutes till 35%B and finally in 20 minutes till 55%B), at a constant flow rate of 300 nL/min⁻¹. MS analysis was carried out as described above.

Mass spectrometry data export and spectral count analysis

Conversion of Bruker Impact files into mzXML format using CompassXport version 3.0.9, led to an initial total of 1,207,329 MS/MS and 444,565 MS/MS spectra, for the gel and total lysate based comparisons, respectively (in-solution total lysate digests were analysed twice and the data were merged). These spectra were searched using a concatenated forward and decoy strategy. The forward database was constructed from the 6386 unreviewed sequences from Uniprot for the organism *A. xylosoxidans* NH44784-1996 (November 2015) together with the cRAP contaminant sequences, as downloaded in January 2015. An in-house developed program, Decoy version V1.0.1-2-gfddc, that preserves homology, amino-acid frequency and peptide length distribution, was used with default flags to construct the decoy search space, which was concatenated to the forward sequences. The search against the resulting database was performed using Comet version 2014.02 rev.2, with precursor mass tolerance equal to 50 ppm and a fragment bin width of 0.05 Da, considering only fully tryptic digests with at most 2 missed cleavages. All cysteines were assumed carbamidomethylated, while methionine oxidation and N-terminal acetylation were regarded as variable modifications. The confidence of these results was assessed by means of Xinteract from the Trans Proteomics Pipeline suite version 4.8.0, retaining peptides longer than 6 amino acids and running in semi-parametric mode. A second in-house developed program, Pepxmltool version 2.5.1, was used to construct a protein quantification table based on spectral counting of only non-degenerate peptides (peptides mapping to a single protein) with corresponding q-value of at most 1%. Plots of the resistant versus susceptible quantifications of all proteins revealed for both methods a predominantly linear relationship, suggesting the applicability of an in-house developed program, Qntdiff version 0.1.1, to assign p-values to deviations from linear behaviour and provide lists of the most significantly differential protein expression levels between the two isolates. The programs Pepxmltool and Qntdiff are available from the authors on request.

Bacterial culture and genomic DNA preparation

AchroS and AchroR (Table 2) were cultured on tryptic soy agar plates (BioMérieux, Marcy l'Etoile, France), inoculated into liquid medium brain-heart-infusion (BHI) broth (Oxoid, Basingstoke, UK) and grown overnight (~16-hours) at 37 °C. Cells were harvested, washed with phosphate-buffered saline (PBS, pH 7.4), and genomic DNA extraction was performed using a phenol-chloroform extraction as previously described²⁴.

Whole genome sequencing and SNP calling

Paired-end multiplex libraries were created as previously described²⁵. Sequencing was performed on an Illumina HiSeq 2000 platform (Illumina, San Diego, Ca, USA), with a read-length of 100 basepairs. High-throughput *de novo* assembly of sequenced genome was performed as previously described^{26,27}. The assemblies are then automatically annotated using PROKKA²⁸ with genus-specific databases from RefSeq²⁹. To identify single nucleotide polymorphisms (SNPs), the Illumina sequence data of the meropenem-susceptible *A. xylosoxidans* isolate (AchroS) was mapped on the assembled genome of the meropenem-resistant isolate (AchroR) using SMALT software (<http://smalt.sourceforge.net/>), after which SNPs were determined as previously described²⁵.

PCR and heterologous expression of Axc

To corroborate the proteomics results and confirm results from the whole genome sequence analysis, the *axc* ORF, the *axc-axcR* intergenic region and part of the *axyZ* ORF were Sanger sequenced at a commercial provider (Macrogen, Amsterdam, the Netherlands). PCR products (for primers see Table 3) were sequenced using the same primers as those used for generating the product from genomic DNA isolated from the *A. xylosoxidans* isolates. We have submitted the Axc sequence to Genbank, ID MF767301.

To construct an *E. coli* strain expressing Axc with a C-terminal 6xHis-tag from an IPTG (isopropyl β -D-1-thiogalactopyranoside) inducible promoter, the *axc* open reading frame was amplified using primers Axfor2 and Axrev3 (Table 3), using Accuzyme polymerase (GC Biotech, Alphen aan den Rijn, The Netherlands) and genomic DNA from *A. xylosoxidans* (AchroR) as a template. The amplified PCR product was digested using NdeI (Bioké, Leiden, The Netherlands) and XhoI (Roche, Almere, The Netherlands), and cloned into similarly digested pET-21b(+), yielding plasmid pET21B/Axc. The *axc* expression region was confirmed using Sanger sequencing. Expression of Axc-his6 was carried out in *E. coli* C43(DE3)³⁰ in Luria-Bertani broth (Affymetrix, Cleveland, OH, USA) with ampicillin (50 μ g/mL) and 1mM IPTG (GC Biotech, Alphen aan de Rijn, the Netherlands) for 3 hours at 37 °C and verified by immunoblotting using anti-His antibody (Agilent Technologies, Santa Clara, CA, USA).

Susceptibility testing of *E. coli* expressing Axc

Minimal inhibitory concentration (MIC) values for the carbapenems imipenem (Etest, BioMérieux) and meropenem (microbroth dilution) were established for *E. coli* C43/pET21B-Axc (strain JC107, Table 2) in the presence and absence of 1mM IPTG. For imipenem, cells were grown overnight in LB broth at 37 °C in the presence of ampicillin (50 μ g/mL). The overnight cultures were diluted 1:100 in LB broth with ampicillin and grown to mid logarithmic phase ($OD_{600nm} \sim 0.5$). Two hundred μ l of bacterial culture was spread on LB-ampicillin (50 μ g/mL) plates (with or without 1 mM IPTG) and an Imipenem Etest (BioMérieux) was applied. MIC values were determined after 24 hours incubation at 37°C. The meropenem MIC values were

established by microbroth dilution. Bacterial cultures in logarithmic phase ($OD_{600nm} \sim 0.5$) were diluted into LB-ampicillin medium to an OD_{600nm} of 0.05 in the presence or absence of 1mM IPTG and subsequently seeded in a 96-well plate. A two-fold serial dilution of meropenem (starting at 12.5 $\mu\text{g}/\text{mL}$) was made by adding equal amounts of meropenem (25 $\mu\text{g}/\text{mL}$) to the first row, from which a two-fold dilution series was made in the rest of the plate. Samples were investigated for growth by measuring the OD_{600nm} after 24 hours incubation at 37 °C, while shaking. The MIC was the lowest concentration of meropenem at which no growth was observed. As controls for our assays, a carbapenem resistant *Klebsiella pneumoniae* clinical isolate (KPC; JC113) and an unrelated expression construct (JC108; which expresses PPEP-1³¹ in an IPTG-dependent manner in the same *E. coli* C43 background, Table 2) were included.

Colorimetric imipenem hydrolysis assay

Overnight bacterial cultures were diluted 100 fold in LB-ampicillin (50 $\mu\text{g}/\text{mL}$) medium and grown to exponential growth phase at 37 °C while shaking. At the time of induction, the OD_{600nm} was determined, the cultures were split in two and 1mM IPTG was added to one of the cultures, followed by a further incubation for 3 hrs at 37 °C while shaking. At T=3h, the OD_{600nm} was determined and cells were harvested by centrifugation (4000 g, 5 min) and stored at -20 °C overnight. Cells were resuspended in water to yield equal densities based on measured OD_{600nm} values. Then, 7.5 μL of bacterial suspension was mixed with 25 μL of imipenem/phenol red/ ZnSO_4 solution (3 mg/mL imipenem, 0.35% (wt/vol) phenol red, pH 7.8, 70 μM ZnSO_4) and incubated at 37 °C for 1 hr. Conversion of imipenem leads to a pH drop that can be visualized by the color change of the buffer from red to yellow³². To quantify this effect, the UV-Vis spectrum was determined with a Nanodrop ND1000 spectrophotometer (Thermo Scientific) and the ratio between the absorption peaks at 431 and 560 nm was taken as a measure of imipenem hydrolysis.

NMR spectroscopy of imipenem conversion

All proton nuclear magnetic resonance (¹H-NMR) experiments were performed on a 600 MHz Bruker Avance II spectrometer (Bruker BioSpin, Karlsruhe, Germany) equipped with a 5-mm triple resonance inverse (TCl) cryogenic probe head with a Z-gradient system and automatic tuning and matching. All experiments were recorded at 310 K. Temperature calibration was done before each batch of measurements³³. The duration of the $\pi/2$ pulses was automatically calibrated for each individual sample using a homonuclear-gated nutation experiment on the locked and shimmed samples after automatic tuning and matching of the probe head. The samples were prepared by adding 70 μL imipenem aqueous solution (5 mg/mL) to 280 μL milliQ water. This solution was mixed with 350 μL 75 mM phosphate buffer (pH 7.4) in water/deuterium oxide (80/20) containing 4.6 mM sodium 3-[trimethylsilyl] d4-propionate. Twenty μL of bacterial cell suspension were added and the sample was mixed. Samples were manually transferred into 5-mm SampleJet NMR tubes. The cell suspension samples were kept at 6 °C on a SampleJet sample changer while queued for acquisition. For water

suppression, presaturation of the water resonance with an effective field of $\gamma B_1 = 25$ Hz was applied during the relaxation delay. A 1D-version of the NOESY (Nuclear Overhauser effect spectroscopy) experiment was performed with a relaxation delay of 4 seconds. A NOESY mixing time of 10 ms was used during which the water resonance was irradiated with the presaturation RF field. After applying 4 dummy scans, a total of 98,304 data points covering a spectral width of 18,029 Hz were collected using 16 scans. The Free Induction Decay was zero-filled to 131,072 complex data points, and an exponential window function was applied with a line broadening factor of 0.3 Hz before Fourier transformation. The spectra were automatically phased and baseline corrected.

Bioinformatic analysis

Comparison of Axc with other class A beta lactamases was performed by multiple alignment using the Geneious 9.0 (Biomatters Ltd, Auckland, New Zealand)) software algorithm for Global alignment with free end gaps, cost Matrix Blosum62. The tree was then built using Jukes-Cantor genetic distance model with the Neighbor Joining tree build method.

Data availability

Illumina raw reads were deposited at the European Nucleotide Archive (ENA). Study ID: PRJEB19781. Sample IDs: ERS1575148 (AchoR) and ERS1575149 (AchoS).

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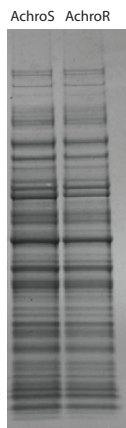
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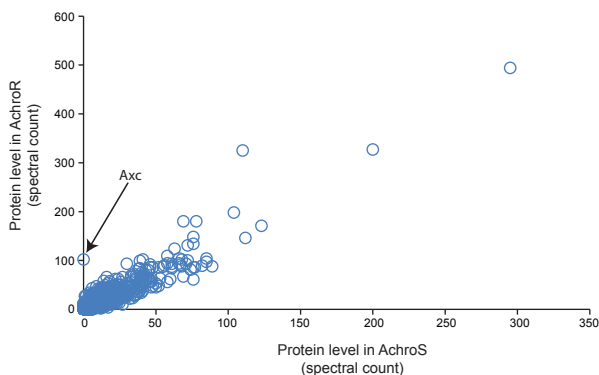
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Supplemental data

A



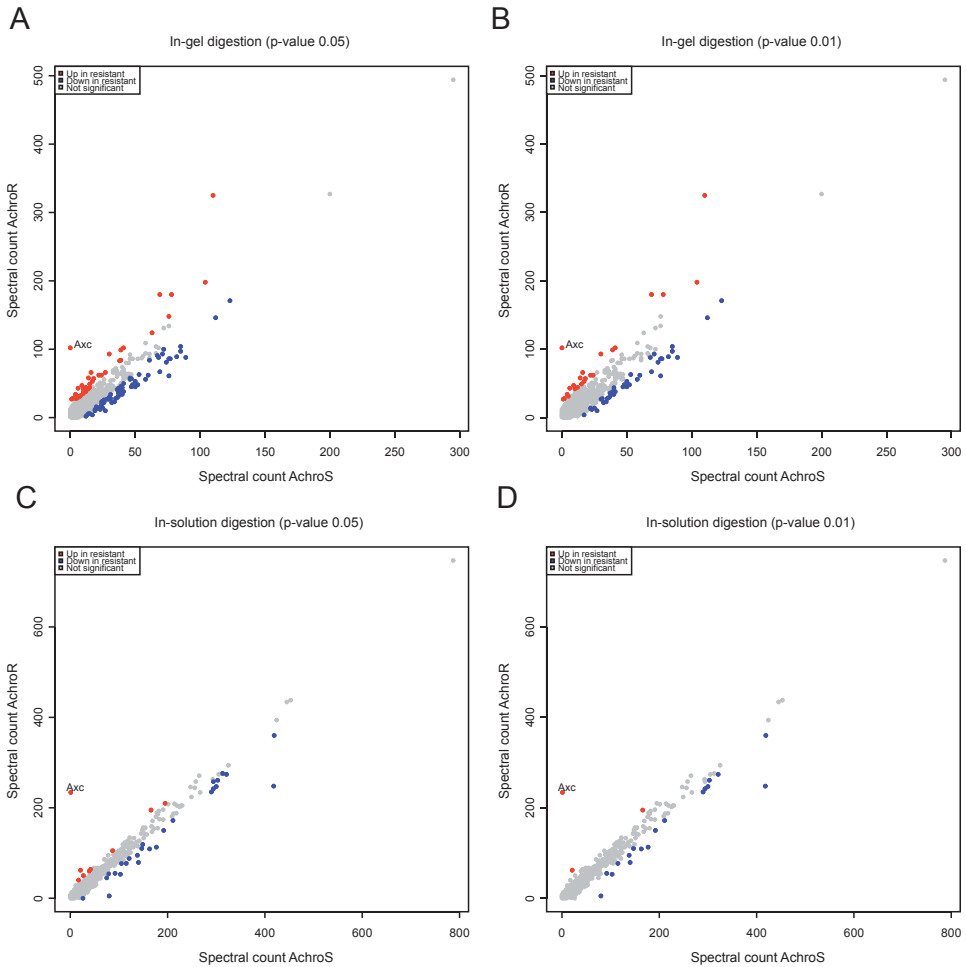
B



Supplemental figure S1: Quantitative proteomic analysis of the meropenem susceptible and resistant *Achromobacter xylosoxidans* clinical isolates.

A: Whole cell lysates from the two isolates (AchroS and AchroR, Table 1) were analyzed by SDS-PAGE and visualized by Coomassie staining. See supplemental figure S4 for a full gel picture, including MW markers.

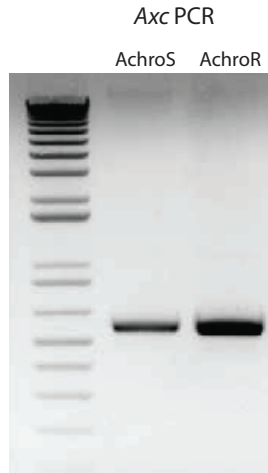
B: Lanes from panel A were sliced into small bands and proteins were subsequently digested using trypsin. Following LC-MS/MS analyses and protein identification, spectral count analysis was used for relative protein quantification. Each dot represents the number of spectral counts observed per unique protein. Axc is the protein with the biggest difference in protein abundance between the resistant and susceptible isolate. In Table S2 and figure S2, the spectral counts and Qntdiff output can be found, respectively.



Supplemental figure S2: Analysis of differential proteins based on spectral counting using Qntdiff

The scatterplots display all quantified proteins using each protein's AchroS spectral count as x-coordinate and the corresponding ArchroR spectral count as y-coordinate, revealing a linear nature in the correlation between the spectral count values for each protein in both isolates. We devised a simple differential detection model that lumps all technical and biological variation into a null distribution for the perpendicular distances of proteins to a linear regression line through the data. If samples are similar enough, this method results in a p-value for the deviation from linearity of each protein. This method is implemented in our program Qntdiff. In figures A and B this method is applied to the in-gel digested data-set and shows differential proteins for p-value thresholds of 0.05 and 0.01, respectively. Figures C and D show the differential proteins selection in the in-solution digestion data-set for p-value thresholds of 0.05 and 0.01, respectively.

A



B

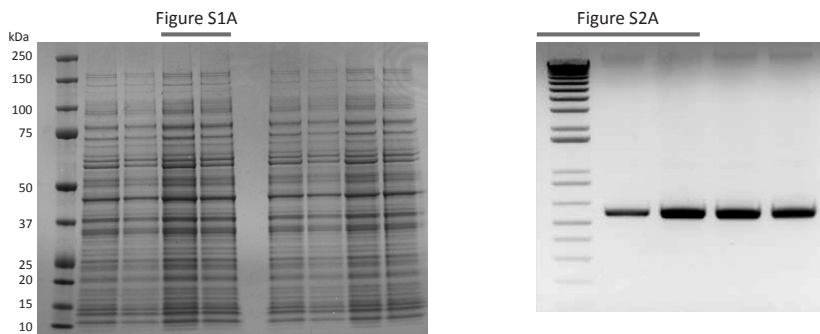


Supplemental figure S3: Presence of *axc* and sequence of its regulatory region in the meropenem resistant and susceptible clinical isolates

A: PCR using primers AxFor and AxRev (Table 3) showed that *axc* is present in both the meropenem resistant (AchroR) and meropenem susceptible (AchroS) clinical isolate. See Figure S4 for a full gel picture from which the cropped version shown is derived.

B: Sequence analysis showed that the intergenic region between *axc* and the gene encoding its putative repressor (*axcR*), were identical in AchroS and AchroR. At two positions (in red) this sequence differed from *A. xylosoxidans* strain NH44784-1996).

Full size gel pictures belonging to fig. S1A and S3A



Supplemental figure S4: Full size gel pictures belonging to Fig. S1A and S3A

