Defining ICR-Mo, an intrinsic colistin resistance determinant from *Moraxella osloensis*

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Abstract

Polymyxin is the last line of defense against severe infections caused by carbapenem-resistant gram-negative pathogens. The emergence of transferable MCR-1/2 polymyxin resistance greatly challenges the renewed interest in colistin (polymyxin E) for clinical treatments. Recent studies have suggested that *Moraxella* species are a putative reservoir for MCR-1/2 genetic determinants. Here, we report the functional definition of ICR-Mo from *M. osloensis*, a chromosomally encoded determinant of colistin resistance, in close relation to current MCR-1/2 family. ICR-Mo transmembrane protein was prepared and purified to homogeneity. Taken along with an *in vitro* enzymatic detection, MALDI-TOF mass spectrometry of bacterial lipid A pools determined that the ICR-Mo enzyme might exploit a possible “ping-pong” mechanism to accept the phosphoethanolamine (PEA) moiety from its donor phosphatidylethanolamine (PE) and then transfer it to the 1(or 4’)-phosphate position of lipid A via an ICR-Mo-bound PEA adduct. Structural decoration of LPS-lipid A by ICR-Mo renders the recipient strain of *E. coli* resistant to polymyxin. Domain swapping assays indicate that the two domains of ICR-Mo cannot be functionally-exchanged with its counterparts in MCR-1/2 and EptA, validating its phylogenetic position in a distinct set of MCR-like genes. Structure-guided functional mapping of ICR-Mo reveals a PE lipid substrate recognizing cavity having a role in enzymatic catalysis and the resultant conference of antibiotic resistance. Expression of *icr-Mo* in *E. coli* significantly prevents the formation of reactive oxygen species (ROS) induced by colistin. Taken together, our results define a member of a group of intrinsic colistin resistance genes phylogenetically close to the MCR-1/2 family, highlighting the evolution of transferable colistin resistance.
Author summary

This work represents the first functional definition of ICR-Mo, a member of a family of MCR-1/2 like genes from Moraxella. As inferred by phylogeny, this is the closest family of chromosomally encoded determinants of colistin resistance. We show evidence that ICR-Mo exploits a possible “ping-pong” mechanism to catalyze the transfer of PEA from its donor PE to the 1(or 4’)-phosphate position of lipid A via an adduct of ICR-Mo-bound PEA. Structure-guided functional studies reveal that ICR-Mo possesses a PE lipid substrate recognizing cavity having a role in enzymatic catalysis and the resulting bestowal of antibiotic resistance against colistin. The expression of icr-Mo in E. coli prevents the generation of ROS in response to treatment with colistin. The fact that the two domains of ICR-Mo cannot be functionally exchanged with their counterparts in MCR-1/2 validates its standing as a distinct phylogenetic entity. In summary, the results define a closer chromosomal relative of MCR-1/2 variants than EptA and highlight its role in the evolution of transferable colistin resistance.

Introduction

Prevalent antibiotic resistance is posing a threat to providing safe and effective health care worldwide. Antimicrobial resistance (AMR) is associated with 700,000 deaths each year [1, 2]. It is estimated by O’Neill and his team that AMR would claim as many as 10,000,000 deaths per year globally by 2050 [3]. Despite this prediction being exaggerated and unreliable [4], we acknowledge that AMR has been posing an ever-growing burden on clinical therapies and public health, thereby highlighting the urgent need for a nationwide response to alleviate this burden [4, 5]. Polymyxins, a class of cationic cyclic polypeptide antibiotics, act as a “last-resort” option against infections by carbapenem-resistant gram-negative pathogens [6–9]. However, the emergence and global spread of plasmid-borne mobilized colistin resistance determinants (mcr-1) has greatly threatened the renewed interest of colistin (polymyxin E) in clinical therapies [10]. To the best of our knowledge, mcr-1-harboring Enterobacteriaceae have been detected in no less than 40 countries worldwide, spanning 5 of 7 continents [11], including the United States of America [12, 13].

MCR-1 is annotated as a transmembrane PEA transferase, which belongs to the “YhjW/ YjdB/YijP” alkaline phosphatase super-family [10, 14]. MCR-1 catalyzes the transfer of phosphoethanolamine (PEA) from phosphatidylethanolamine (PE) to the 1(or 4’)-phosphate position of lipid A glucosamine (GlcN) moieties (S1 Fig) [15–17]. In general, the addition of PEA to lipid A is believed to reduce the net negative charge of the bacterial outer-membrane [6, 18, 19], consequently leading to a phenotypic resistance to polymyxin, a final line of defense against severe infections by pathogens with multi-drug resistance (MDR) [7, 8]. In comparison to the paradigm PEA transferase, the Neisseria EptA protein [16, 20], MCR-1 seems to render the recipient E. coli strains more resistant to colistin [14], implying that it is a more efficient enzyme. In contrast to the more prevalent MCR-1, its cousin MCR-2 with almost 80% amino acid identity, seems to be rare in that it is only detected in Belgium in 2016 [21]. In particular, it seems likely that both MCR-1 and MCR-2 have an evolutionarily-conserved catalytic motif having essential roles in enzymatic activities and resultant antibiotic resistance [14, 22].

Along with other transferable determinants of colistin resistance (MCR-3 [23, 24], MCR-4 [25] and MCR-5 [26]), a growing body of MCR-1 variants with point substitutions have been reported, such as MCR-1.2 [Q3L] [27], MCR-1.3 [I38V] [28] and MCR-1.6 [R536H] [29]. More recently, mcr-1 and mcr-2 variant genes have been discovered on chromosomes of
Moraxella species [30, 31]. In brief, MCR-1.10 with 98.7% amino acid identity to MCR-1 is present in M. porci MSG13-C03 [30], MCR-2.2 with 99% amino acid identity (only 8 amino acid substitutions of 538 residues) to MCR-2 is encoded on the chromosome of M. pluranimarium [31], and MCR-2.3 with 87.9% amino acid identity to MCR-2 is harbored in a M. pluranimalium-like isolate, MSG47-C17. Moreover, it seems that MCR-like enzymes with about 60% amino acid identity are prevalent in Moraxella species [32], suggesting a potential reservoir of mcr-like genes. However, this requires further experimental evidence.

In this study, we have integrated multiple lines of approaches to study the structure and mechanism for one such mcr-like gene, AXE82_07515 in M. osloensis (hereafter designated as icr-Mo [of note: Mo denotes Moraxella osloensis]). This will allow it to serve as a representative member of this family for future studies. Consistent with genetic speculations raised by other groups [30–32], our results formulate a functional definition of icr-Mo and provide an evolutionary relationship between itself, mcr-1/2 variants and non-mcr genes that confer colistin resistance. By probing the biochemical and physiological relevance, our findings highlight the importance of the Moraxella family in understanding the growing body of mcr-1/2-like genetic determinants conferring colistin resistance.

Results

Evolution of mcr-like genes in Moraxella

The gene mcr-1, is a prevalent determinant of plasmid-borne mobilized colistin resistance with a global distribution. In addition to a closely-related but rare mcr-1 homolog, mcr-2 sharing 81% protein identity [21], a number of genetic variants of mcr-1 (like mcr-1.2 [27]) with point substitutions have also been elucidated. Earlier this year, Kieffer and coworkers [32] showed genetic evidence that Moraxella species might be a potential reservoir for mcr-like genes. This prediction is further validated by additional discovery of mcr-1.10 (98.7% amino acid identity) in M. porci MSG13-C03 [30] and mcr-2.1 (99% amino acid identity) in M. pluranimarium [31]. In addition to their similarity, certain members of these chromosomally encoded mcr-1/2-like genes also neighbor almost identical mobilizable elements like the insertion sequence ISApl1, associated with mcr-1/2 dissemination [31, 32]. To probe the evolutionary relationships between the icr-Mo in Moraxella and other extant mcr genes, we performed phylogenetic analyses (Fig 1), using their coding nucleotide sequences. As anticipated, MCR-1 and its variants (namely MCR-1.2, MCR-1.3, . . . MCR-1.9 and MCR-1.10) cluster together to form a tight group (Fig 1A and 1B). In fact, we have indicated 2 additional variants of MCR-1 with single synonymous point mutations. MCR-2 and its two variants (MCR-2.1 and MCR-2.2) also form a distinct group (Fig 1A and 1B) while still being closely-clustered within a single subclade, comprising MCR-1/2 variants (Fig 1A and 1B). Surprisingly, Neisseria EptA, a chromosomal colistin resistance determinant (31.5% amino acid identity to ICR-Mo, S3 Fig) clusters with other putative sulfatases, a large family of enzymes encoding broad functionality that includes PEA transferases. The sulfatase sub-clade along with a cluster comprising MCR-3 and MCR-4 variants are only distantly-related to the MCR-1/2 and MCR-like genes. In fact, Z1140 (539 aa) of E. coli O157:H7, a putative PEA transerase lacking detectable ability to confer phenotypic resistance to polymyxin (S2 Fig) is found within the MCR-3/4 subclade leading us to categorize them as non MCR-like genes (Fig 1A). Given that the identity between ICR-Mo and EptA is only 31.5%, it seems likely that mcr-1/2 and icr-Mo genes have diverged from sulfatases much earlier than the recent colistin usage in agriculture and clinical medicine. A much closer ancestor might be found in bacteria that naturally produce polymyxins as secondary metabolites.

In contrast, a subset of MCR-like members exclusively from the Moraxella species (porci, lincolnii and catarrhalis subspecies, in Fig 1) with about 60% amino acid identities (S3 Fig) are
clustered into a neighboring clade that includes ICR-Mo (Fig 1A and 1B). This also includes the recently crystallized ICR-Mc (Accession No: AIT43666) from *Moraxella catarrhalis* which shares 55.2% amino acid identity with ICR-Mo [33]. We are inclined to believe that plasmid-borne MCR-1/2 variants and the intrinsic ICR-Mo might share a common ancestor. Intriguingly, certain species of *Moraxella* contain chromosomally encoded colistin resistance determinants that are grouped with members of MCR-1/2 family (Fig 1B). Therefore, a systematic evaluation of biochemical and physiological properties of ICR-Mo, is necessary.

**Characterization of ICR-Mo**

Prediction with TMHMM server v2.0 (http://www.cbs.dtu.dk/services/TMHMM) suggests that ICR-Mo is an integral membrane protein possessing five N-terminal helices (S3A Fig), which is almost identical to those of MCR-1 [14] and MCR-2 [22]. As recently described for MCR-1/2 [14, 22], the transmembrane protein ICR-Mo was overexpressed and purified to homogeneity with nickel affinity column in the presence of 1% detergent dodecyl-β-D-
maltoside (DDM). Following visualization by 12% SDS-PAGE (S4A and S8A Figs), identity of the recombinant ICR-Mo protein was determined with MALDI-TOF MS (Matrix-Assisted Laser Desorption/Ionization Time of Flight Mass Spectrometry) with a 84.67% fragment sequence (S10C Fig). The structure of full-length EptA (PDB: 5FGN) was used as a template [34] for the structural modeling of ICR-Mo via Swiss-Model (https://swissmodel.expasy.org/interactive) [35]. Despite possessing only 31.5% amino acid identity to EptA (S2 Fig), the modeled structure of ICR-Mo shows an appreciable coverage of 93% (24–558), indicating a satisfied prediction (S4 Fig). ICR-Mo also displays a similar topology to EptA (S4B Fig) [34]. The TM domain spanning the inner-membrane involves five α-helices (S4B Fig), and the catalytic domain has a hydrolase-fold (S4C Fig) comprising 10 α-helices and 7 β-sheets (S4C Fig). The two domains are linked by four short periplasm loops (PH2, PH2', PH3 and PH4), a bridge helix (BH) and a long-coiled loop (S4B and S4C Fig).

**Enzymology of ICR-Mo**

As described by Anandan et al. [34] for EptA activity, we established an in vitro system for ICR-Mo enzymatic reaction. In this reaction system, the PEA donor substrate used as a fluorescent label, 1-acyl-2-{12-[7-nitro-2-1,3-benzoxadiazol-4-yl) amino] dodecanoyl}-sn-glycerol-3-phosphoethanolamine (abbreviated as NBD-glycerol-3-PEA, Fig 2A). It is hypothesized that ICR-Mo removes the PEA moiety from the alternative substrate NBD-glycerol-3-PEA, and gives rise to a NBD-glycerol product (Fig 2A). Given that an adduct of PEA-Thr280-EptA has been observed by Vrielink and coworkers [20], we anticipate that an intermediate of ICR-Mo-bound PEA (i.e., PEA-Thr287-ICR-Mo) is produced in the hydrolytic reaction (Fig 2A). Thereafter, we utilized thin layer chromatography (TLC) to separate the mixture of the ICR-Mo reaction (Fig 2B and 2C and S5 Fig). As expected, the product NBD-glycerol is released in the presence of ICR-Mo, which is consistent with observations of EptA and MCR-1/2 (S5A and S5B Fig) [36, 37]. Under blue light (455–485 nm), TLC-based assay illustrates that the NBD-glycerol product (Fig 2C) migrates faster than the substrate NBD-glycerol-3-PEA does (S5 Fig, Fig 2B and 2C). Subsequently, liquid chromatography mass spectrometry (LC/MS) confirms the identities of NBD-glycerol-3-PEA (Fig 2B) and its resultant product, NBD-glycerol (Fig 2C). This demonstrates clearly that ICR-Mo displays an enzymatic activity of removing the PEA moiety from the lipid substrate in vitro, which is similar to the well-studied EptA [34], as well as MCR-1/2 (S5 Fig).

To further the in vivo transfer of PEA from PE lipids (S1A Fig and Fig 2D–2G), we engineered *E. coli* MG1655 strains to carry arabinose inducible plasmid pBAD24-borne icr-Mo (and eptA/mcr-1). We prepared and purified the LPS-lipid A from an array of *E. coli* strains. Unlike the negative control MG1655 exhibiting a single peak (m/z of 1796.370–1797.843) for lipid A alone (Fig 2E–2G), MALDI-TOF mass spectrometry unveiled an additional peak associated with PPEA-1(or 4')-lipid A, the modified form of lipid A and this peak appeared in the *E. coli* strain harboring mcr-like genes. We believe that lipid A accepts PEA moiety from the intermediate of ICR-Mo-bound PEA, giving PPEA-4'-lipid A (Fig 2D). In fact, the molecular mass of PPEA-lipid A in different strains exhibits slight variation, namely m/z 1920.014 for EptA (Fig 2F), m/z 1920.334 for MCR-1 (Fig 2F), and m/z 1919.577 for ICR-Mo (Fig 2G). The in vitro and in vivo evidence leads us to believe that ICR-Mo catalysis proceeds by PEA transfer from the donor PE lipid substrate to the receiver lipid A, giving the two final products, PPEA-4'-lipid A and diacyl glycerol (S1A Fig). Taking into account the similarity between the adduct of Thr280-PEA in EptA [20, 34] and its counterparts in the reactions involving alkaline phosphatase-type phosphate transferase [38], it can be hypothesized that i) the ICR-Mo-bound PEA intermediate is released from the PE lipid molecule in the first-half reaction of ICR-Mo.
catalysis (S1B Fig); ii) In the second half-reaction, PEA is transferred from the ICR-Mo-bound PEA adduct to the 1(4')-phosphate position of lipid A GlcN moieties, generating PPEA-4'-lipid A (S1C Fig). It seems likely that ICR-Mo exploits a possible “ping-pong” mechanism for enzymatic catalysis, similar to those proposed for EptA [34] and MCR-1/2 [36, 37, 39].

A cavity in ICR-Mo for the entry of PE substrate
On the basis of the newly-determined complex structure of EptA with the detergent DDM, an analogue of its physiological PE substrate [34], we applied molecular docking to reanalyze the binding to PE molecule (Fig 3A). As a result, it allows us to propose a 12 residue-containing cavity for PE entry/binding (Fig 3). Superposition of the modeled structures of MCR-1 (ICR-Mo) to EptA also gives a similar PE-recognizable cavity (Fig 3C and 3D). Amongst them, five amino acids (E248, T287, H397, D472 and H473) are centering with the zinc ion (Fig 3E–3G and S7A and S7B Fig), and remaining 7 residues (N110, T114, E118, S332, K335, H402 and H485) are anticipated to be involved in the recognition of its physiological lipid PE substrate (Fig 3H–3J and S7C and S7D Fig). Similar observations have been made earlier for E. coli (Fig 4). Hence, this hypothesis needs further experimental validation.

We thereafter conducted structure-guided site-directed mutagenesis, generating 12 point-mutants of ICR-Mo (S2 Fig and Figs 3E–3J and 4A). Prior to functional assays in vivo, western blotting demonstrates that all the mutated versions of ICR-Mo express well in E. coli (Fig 4A). Assays of colistin resistance elucidate that i) ICR-Mo permits the recipient E. coli to grow on LBA plates with up to 8 μg/ml colistin, [which is equivalent to that of EptA (S6 Fig), but significantly lower than MCR-1 does (16 μg/ml)] (Fig 4 and S6 Fig); (ii) all the 5 point-mutants of ICR-Mo with defects in zinc-binding residues fail to support the growth of the recipient E. coli strains under the condition with up to 0.5 μg/ml colistin (Fig 4B); (iii) Among the 7 ICR-Mo mutants with the alanine substitution in putative PE substrate-interacting residues, 3 of them seem to retain partial activity (namely N110A [2 μg/ml], T114A [1 μg/ml] and S332A [1 μg/ml], Fig 4C) and the remaining 4 mutants are nonfunctional [0.5 μg/ml] (Fig 4C). These results are in agreement with the observations in the colistin MIC measurements (Fig 4D).

Biochemical analyses of PE-interactive cavity in ICR-Mo
To gain a mechanistic glimpse of the PE substrate-interactive cavity, we purified the 12 point-mutants of ICR-Mo protein to homogeneity (S8A Fig). First, catalytic activities of the ICR-Mo mutants were examined using the in vitro enzymatic reaction with the fluorescence-labelled substrate, NDB-glycerol-3-PEA and LBD-glycerol-3-PEA lipid substrate by the ICR-Mo (AXE82_07515) into NBD-glycerol. D. Transfer of PEA from ICR-Mo-bound PEA to lipid A, generating the PPEA-lipid A product. Position of PPEA depicted is only suggestive. MALDI-TOF-MS evidence for the structural alteration of the lipid A moieties of lipopolysaccharide (LPS) in E. coli expressing EptA (E), MCR-1 (F) and ICR-Mo (AXE82_07515) (G). The peak of the bis-phosphorylated hexa-acylated lipid A varies at m/z of 1796.063 – 1797.426, whereas resultant derivative with PEA modification (PPEA-1(4')-lipid A) exhibits at m/z varying from 1919.409 to 1920.087.
Fig 3. Paralleled PE-recognizing cavities amongst ICR-Mo, EptA and MCR-1. A. Chemical structure of the PE lipid substrate molecule. B. EptA has a cavity for the entry of the PE lipid substrate. C. A PE-recognizable cavity is present in MCR-1 enzyme. D. A conservative PE-binding cavity is also shared by ICR-Mo (AXE82_07515). A Zn$^{2+}$-bound five-residues forming motif is conserved in PE lipid substrate-interactive cavities of three enzymes EptA (E), MCR-1 (F) and ICR-Mo (G). Comparative
glycerol-3-PEA is not converted into NDB-glycerol, almost identical to the negative controls] (S8B Fig).

In general, the variation in enzymatic activities of ICR-Mo mutants was further verified by structural analyses of LPS-lipid A moieties in a physiological context (Fig 5). As expected, MALDI-TOF MS analysis reveals that a unique peak of PPEA-1(or 4')-lipid A (m/z, 1920.140) is present in the E. coli expressing the wild-type ICR-Mo (Fig 5C), relative to the recipient strain MG1655 with a single peak of lipid A [m/z, 1796.782~1796.946] (Fig 5A and 5B). Consistent with the observations of the TLC experiments, 9 of 12 ICR-Mo point-mutants cannot modify lipid A species in vivo since only a single peak of lipid A is detected (Fig 5F–5H and 5J–5O). In contrast, the remaining 3 mutants seem to retain partial abilities of PEA transfer, in that an extra-peak of PPEA-4'-lipid A [m/z, 1919.865~1920.457] is produced in addition to the lipid A [1796.709~1797.230] (Fig 5D, 5E and 5I). Of note, as for the ICR-Mo(T114A) mutant retaining partial activity in vivo (Figs 4C and 5E), no enzymatic activity in vitro is detected (S8B Fig). This could be partially attributed to the lack of sufficient sensitivity of TLC assays in this case. Evidently, this highlights the importance of cavity-forming residues in ICR-Mo biochemistry.

### Domain-swapping analyses of ICR-Mo

Phylogenetically, ICR-Mo is distinct from EptA and MCR-1/2. This prompted us to probe the possible association of the two modules of MCR-like enzymes [transmembrane region (TM) and PEA transferase domain] with respect to their functional evolution (Fig 1). As described with MCR-2 [22], we adopted the method of domain swapping to engineer a collection of hybrid versions of the icr-Mo gene (Fig 6A). In addition to three parental proteins (EptA, MCR-1 and ICR-Mo (AXE82_07515)), four hybrid proteins engineered here include i) TM (EptA)-AXE82, a modified ICR-Mo carrying TM region of EptA; ii) TM(AXE82)-EptA, a derivative of EptA whose TM domain is replaced with its ICR-Mo counterpart; iii) TM(MCR-1)-AXE82, a modified version of ICR-Mo containing the TM domain of MCR-1; and iv) TM (AXE82)-MCR-1, a hybrid derivative of MCR-1 carrying the TM region of ICR-Mo (Fig 6A). Western blotting analyses prove that all the MCR-like enzymes (ICR-Mo and its domain-swapped derivatives) express pretty well in E. coli (Fig 6B). Subsequently, they were subjected to functional evaluation under both in vitro and in vivo conditions (S9–S11 Figs). Following an MS-based confirmation of their polypeptide sequence identity (S9 and S10 Figs), circular dichroism (CD) experiments reveal that all the hybrid derivatives of ICR-Mo consistently display typical CD spectra of being rich in α-helices (S11D–S11G Fig and S3 Table), almost identical to that of MCR-1 (S11A Fig), EptA (S11B Fig) and ICR-Mo (S11C Fig). As observed with soluble domains of both MCR-1 [40–43, 45] and MCR-2 [46], the result of ICP/MS (inductively coupled plasma mass spectrometry) reveals that zinc is specifically occupied with ICR-Mo and its derivatives (S12A Fig), and the relative ratio of zinc ion to protein is calculated to be around 1:1 (S12B Fig).

Assays of antibiotic resistance on LBA plates show that i) 3 of 4 chimeric versions are inactive; and the only one variant, TM(MCR-1)-AXE82, has partial activity to allowing the recipient E. coli to grow on the condition with up to 4 μg/ml colistin (Fig 6C), less than those of its parental versions MCR-1 (16 μg/ml colistin) and ICR-Mo (8 μg/ml colistin) (Fig 6C). Similar scenarios were seen in the measurement of colistin MIC (Fig 6D). In addition, MALDI-TOF
MS analyses of lipid A species demonstrated that like the wild-type MCR-like proteins (Fig 7C–7E), TM(MCR-1)-AXE82 is the only one hybrid derivative of ICR-Mo having a role in the transfer of the PEA moiety from PE to the acceptor LPS-lipid A (Fig 7H). In contrast, the
Fig 5. MALDI-TOF MS identification of LPS-lipid A structure suggests different roles of putative cavity-forming sites in catalysis mechanism of ICR-Mo (AXE82_07515). MALDI-TOF MS profile of the LPS-lipid A pool isolated from the two negative controls, *E. coli* strain MG1655 alone (A) and with the empty vector pBAD24 (B).
C. Appearance of a unique peak of the mono-modified lipid A, PPEA-1(or 4')-lipid A, in the positive control, MG1655 strain carrying the wild-type of ICR-Mo (AXE82_07515). The two mutations of N110A (D) and T114A (E) failed to fully inactivate the enzymatic activity of ICR-Mo in that the modified peak, PPEA-lipid A is present. MALDI-TOF-MS analyses suggests that the three mutations of E118A (AxE82_07515). The two mutations of N110A (C). Appearance of a unique peak of the mono-modified lipid A, PPEA-1(o r 4')-lipid A, in the positive control, MG1655 strain carrying the wild-type of ICR-Mo still possesses partial activity of catalyzing the transfer of PPEA to the 1(4')-phosphate group of lipid A moieties. The six point-mutants of ICR-Mo are consistently inactive in the enzymatic activity of PEA transferase in vivo, including K335A (J), H397A (K), H402A (L), D472A (M), H473 (N) and H485A (O), respectively. The MS peak of lipid A species in E. coli is shown at m/z of 1796.744~1797.843, whereas its modified form appears at m/z of 1919.865~1920.457, when functional (even partial active) versions of ICR-Mo (AXE82_07515) are present in E. coli.

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other 3 derivatives (Fig 7G-7I), have no detectable activities similar to those of negative controls (Fig 7A and 7B). Evidently, MS-based visualization of altered structures of lipid A moieties agrees with scenarios observed in the both enzymatic tests and colistin resistance assays (Fig 6). In summary, these data suggest that protein evolution has created a gradient with respect to the acquisition/differentiation of antibiotic resistance amongst EptA, ICR-Mo and MCR-1/2 to some extent.

### ICR-Mo quenches hydroxyl radical death pathway

Successful killing and eradication of the Gram-negative bacteria by polymyxins is determined by two critical factors: i) Efficient binding-entry to the initial target, bacterial LPS-lipid A [9]; ii) Activation of a downstream hydroxyl radical death pathway [47, 48]. The aforementioned results have proved that ICR-Mo can modify the lipid A moieties, whereas it is unclear as to whether or not it influences the downstream event. Therefore, we integrated two different methods [confocal microscopy (Figs 8 and 9) and chemical rescue assay (Fig 10)] to address this question. First, an oxidant-sensitive dye DCFH2_DA (2',7'- dichlorodihydrofluorescein diacetate) was applied in monitoring intra-cellular H2O2 species in E. coli with/without ICR-Mo (or EptA/MCR-1) (Fig 8). As expected, the treatment with colistin boosts ROS production in the E. coli MG1655 (Fig 8A and 8B). However, expression of icr-Mo alleviates colistin-triggered ROS formation regardless of colistin treatment (Figs 8G and 9). This is in agreement with observations in E. coli strains expressing either EptA (Fig 8C and 8D) or MCR-1 (Fig 8E and 8F).

In chemical rescue trials, the presence of a ferric chelator, bipyridine, significantly bypasses cell death of the icr-Mo-negative E. coli with that are subjected to treatment with colistin (Fig 10A and 10C). It seems likely that a Fenton reaction is involved in the formation of free hydroxyl radicals (Fig 10A). Similarly, the presence of the ROS scavenger, L-cysteine alone [or accomplished with bipyridine] also greatly improves bacterial survival of E. coli under conditions of colistin treatment (Fig 10A and 10C). Intriguingly, the presence of icr-Mo (or eptA/ mcr-1) prevents the recipient E. coli from entering into the hydroxy radical death pathway in E. coli (Fig 10B and 10C). Of note, this impact is independent of presence of pyridine (and/or L-cysteine). Thus, we believe that ICR-Mo (EptA/MCR-1) modifies the bacterial LPS-covered membrane, prevents efficient entry of polymyxin into cells, quenches/altibiotics ROS production in vivo, and consequently bypasses antibiotic killing by colistin (Fig 10).

### Discussion

Polymyxin is a paradigm for cationic antimicrobial polypeptides (CAMP). Physical basis for CAMP-type resistance is due to structural alteration of bacterial LPS, leading to a reduction of the net negative charge of the cell surface. In total, three types of chemical modifications of lipid A species are involved in surface remodeling of Gram-negative bacterium. Among them, the most prevalent form refers to the modification of the LPS-lipid A with an addition of the cationic sugar 4-amino-4-deoxy-l-arabinose [49–51]; Second, the transfer of PEA to the 1(or 4')-phosphate position of lipid A moieties [16, 17, 51–54]; Third, the glycine attachment to
### A. Domain-swapping analyses of ICR-Mo.

| Enzyme | TM Domain | Extra-cellular Domain | Activity |
|--------|-----------|-----------------------|----------|
| EptA [544 aa] | TM (1-179 aa) | PEA (180-544 aa) | ++ |
| MCR-1 [541 aa] | TM1 (1-181 aa) | PEA (182-541 aa) | +++ |
| ICR-Mo (AXE82_07515) [548 aa] | TM (1-183 aa) | PEA (184-548 aa) | ++ |
| TM(EptA)-AXE82 [544 aa] | TM (1-179 aa) | PEA (184-548 aa) | - |
| TM(AXE82)-EptA [548 aa] | TM1 (1-183 aa) | PEA (180-544 aa) | - |
| TM(MCR-1)-AXE82 [546 aa] | TM (1-181 aa) | PEA (184-548 aa) | + |
| TM(AXE82)-MCR-1 [543 aa] | TM2 (1-183 aa) | PEA (182-541 aa) | - |

### B. Use of Western blotting to detect the expression of icr-Mo and its mosaic versions.

![Western Blot](image)

### C. Functional evaluation of ICR-Mo derivatives in ability of conferring appreciable growth of *E. coli* on LBA plates with varied levels of colistin.

![Bar Graph](image)

### D. Measurement of colistin MIC of *E. coli* strains expressing ICR-Mo and its hybrid derivatives.

| Strain | MG1655 | MG1655+Vec | EptA/MCR-1/AXE82_07515 |
|--------|---------|------------|------------------------|
| MIC    | 0.25    | 0.25       | 2.0 2.0 2.0 2.0 2.0 0.25 0.25 0.25 1.0 |

**Fig 6.** Domain-swapping analyses of ICR-Mo. **A.** Schematic illustration for domain-swapping designing amongst the three transmembrane enzymes EptA, MCR-1 and ICR-Mo (AXE82_07515). **B.** Use of Western blotting to detect the expression of icr-Mo and its mosaic versions. **C.** Functional evaluation of ICR-Mo derivatives in ability of conferring appreciable growth of *E. coli* on LBA plates with varied levels of colistin. **D.** Measurement of colistin MIC of *E. coli* strains expressing ICR-Mo and its hybrid derivatives.

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LPS-lipid A GlcN moieties by a tripartite system [Vc1577 (AlmG), Vc1578 (AlmF), and Vc1579 (AlmE)] in *Vibrio cholerae* [55]. It is known that rare cases of an intrinsic colistin resistance are generated by spontaneous point-mutations of chromosomal genes, especially two-component systems of PhoB/Q [53, 56] and PmrA/B [17, 52–54]. In addition, the EptA enzyme of the *Neisseria* species, which is a representative member of PEA lipid A transferase family, also contributes to the generation of an intrinsic resistance to colistin [16]. In contrast, the discovery of MCR-like determinants belonging to PEA lipid A transferases constitutes a new mechanism for the transferability of polymyxin resistance [10, 14, 21, 22, 24–26]. It seems

https://doi.org/10.1371/journal.pgen.1007389.g007

![Graphs and images showing MALDI-TOF mass spectrometry results](https://doi.org/10.1371/journal.pgen.1007389.g007)
Fig 8. Colistin-stimulated ROS production is significantly impaired by the presence of colistin resistance-conferring proteins (EptA/MCR-1/ICR-Mo). A & B Accumulation of hydrogen peroxide is boosted by colistin stress in *E. coli*. C & D The presence of plasmid-borne EptA inhibits colistin-induced production of hydrogen peroxide in *E. coli*. E & F Colistin-triggered production of hydrogen peroxide is decreased upon the expression of MCR-1 in *E. coli*. G & H The expression of *icr-Mo*
more worrisome that global transmission of \textit{mcr-1} genetic determinant on diversified plasmids raises significant challenge to clinical therapy and public health \cite{11, 57, 58}. Therefore, it is essential to understand the evolution of \textit{mcr-1}-like variants.

Very recently, \textit{Moraxella} species have been proposed as a reservoir for \textit{mcr-1/2} genetic determinants \cite{30–32}. Our phylogenetic analyses showed that the MCR-1/2 family is much more closely related to the \textit{Moraxella} family of MCR-like genes than either Sulfatases or other non MCR-like genes. In fact, the \textit{pap2} gene originally found next to \textit{mcr-1} on plasmids, has also been detected on the chromosome next to \textit{mcr-1} (and its genetic variants) in certain other species of \textit{Moraxella} \cite{30}. Given the fact that i) both \textit{M. pluranimimalium} \cite{59} and \textit{M. porci} \cite{60} colonize in pigs; and ii) pigs are major host reservoirs for \textit{mcr-1}-harboring Enterobacteriaceae \cite{10}, it is ecologically reasonable that genetic exchange of \textit{mcr-1/2}-like determinants between \textit{Moraxella} species and Enterobacteriaceae like \textit{E. coli} occur. This leads us to believe that evaluating the chromosomally-encoded genetic determinants of colistin resistance in \textit{Moraxella} might further our understanding of evolution in the MCR1/2 family. The data we report here presents a biochemical and physiological understanding of ICR-Mo, found on the chromosome of \textit{Moraxella osloensis}. Relative to MCR-1, ICR-Mo confers a lower level of resistance to colistin to a recipient \textit{E. coli} strain (S6 Fig), which argues the possibility that \textit{mcr}-like genes are in either a developing or degenerating state \cite{61}. Of note, any direct evidence of ICR-Mc \cite{33} or ICR-Mo conferring resistance to colistin in native \textit{Moraxella} strains is also lacking. The two domains (TM region and extra-domain of PEA transferase) of MCR-1 and MCR-2 are functionally-exchangeable \cite{22}, while those in ICR-Mo and MCR-1 cannot be fully switched (Fig 6). This indicates a fundamental difference between these two proteins that further validates their phylogenetic placement. Structural and functional characterization of ICR-Mo facilitates...
Fig 10. A working model that ICR-Mo stops colistin-induced hydroxyl radical killing in E. coli. A. Scheme for ROS production triggered by colistin in E. coli. B. Impairment of colistin-induced ROS formation in icr-Mo-bearing E. coli. C. Chemical rescue experiments reveal that a Fenton reaction is involved in the colistin-activated hydroxyl radical killing pathway in E. coli. The LPS-lipid A moiety refers to an initial target for colistin treatment. Bipyridine is a well-studied ferric chelator, and L-cysteine is the ROS scavenger.

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the identification of an important PE substrate cavity, which is essential for its enzymatic activity and for the conferring a phenotypic resistance to colistin (Figs 3 and 4). In light of its similarity to those of EptA and MCR-1 (Figs 3 and 4), this hints an evolutionarily-conserved mechanism for intrinsic and transferable colistin resistance. In fact, this is in close agreement with observations made from a very recent X-ray crystal structure of an intrinsic colistin resistance gene product, ICR-Mc, in *Moraxella catarrhalis* [33]. Moreover, ICR-Mo displays similar abilities in both modifying lipid A structure (Fig 2) and quenching the production of ROS *in vivo* (Figs 8 and 9), when compared with that of MCR-1. Intriguingly, it seems likely that a putative “ping-pong” mechanism is shared amongst these MCR-like enzymes (S1 Fig and Fig 2). It was thought that the ancestral source of colistin resistance might be from proteins containing sulfatase/hydrolase domains. These domains possess the same catalytic architecture as PEA transferases. Further, polymyxin is naturally produced by (and originally isolated in) certain members of the *Paenibacillus* family as a secondary metabolite. However, based on our biochemical and phylogenetic data it is tough to conclude whether the members of the non-MCR-like family (comprising MCR-3/4), which cluster closely with the Sulfatase family, are in a state of development or degeneration of colistin resistance. Thus, we simply propose that ICR-Mo and the other members of the *Moraxella* family have a same ancestor as the MCR-1/2 family does, providing a current source of genetic variation.

In response to the emergence of MCR-1 colistin resistance in the very late of 2015 [10], Chinese agricultural government has taken on a more active role in preventing the further spread of MCR-1 colistin resistance by formally banning the use of colistin as a growth booster in China in early 2017. Moreover, it is also important to reconsider the clinical use of colistin as a final line of refuge against lethal infections with carbapenem-resistant superbugs. Our findings represent a full mechanistic understanding of ICR-Mo, a representative member of a family of chromosomal relatives to transferable MCR-1 colistin resistance. It might provide molecular basis for the rational development of small molecules targeting the reversal of MCR-like resistance to colistin, a last-resort antibiotic.

**Materials and methods**

**Strains, plasmids and growth conditions**

All the strains used here are derivatives of *E. coli* MG1655 (S1 Table). As described with *mcr-2* [22], the full-length *icr-Mo* (AXE82_07515) was synthesized *in vitro*. To generate hybrid versions of *icr-Mo* and *mcr-1* (and/or *eptA*), overlapping PCR experiments were conducted [22]. The point-mutants of *icr-Mo* were produced using the Mut Express II Fast Mutagenesis Kit V2 (Vazyme Biotech Co., Ltd) with suitable primers (S1 Table). The pET21a expression vector was used for protein production, and arabinose-inducible plasmid pBAD24 was utilized for functional complementation (S1 Table). All the constructs were confirmed with direct DNA sequencing. Luria-Bertani (LB) broth (either liquid culture or solid agar plates) was applied and appropriate antibiotics such as ampicillin and colistin were supplied.

**Assays for colistin susceptibility**

The minimum inhibitory concentration (MIC) of colistin was determined using a liquid broth dilution test as recommended by EUCAST with Cation-adjusted Mueller-Hinton Broth (CAMHB) [22]. When necessary, 0.2% arabinose was added into CAMHB media to induce the expression of pBAD24-borne *icr-Mo* and its derivatives in *E. coli*. Also, the viabilities of *E. coli* carrying *icr-Mo* and its derivatives were judged with solid LBA broth dilution test [14, 62, 63].
Expression, purification and identification of transmembrane enzyme ICR-Mo

As described with MCR-2 [22] with little change, ICR-Mo and its chimeric versions were overexpressed. The membrane fraction containing the protein of interest was solubilized in buffer B (20 mM Tris-HCl [pH 8.0], 100 mM NaCl, 5% glycerol, 1% DDM (M/V)) and then centrifuged at 38000 r.p.m for 1.5 h at 4˚C. The resultant supernatant was subjected to affinity purification of protein with pre-equilibrated Ni-NTA agarose beads. The expected integral membrane proteins of ICR-Mo and its derivatives were eluted using an elution buffer (20 mM Tris-HCl [pH 8.0], 100 mM NaCl, 100 mM imidazole, 5% glycerol, 0.03% DDM (M/V)) [14, 22]. Following determination of the purified protein with 12% SDS-PAGE, the protein band with expected size was cut from gels and subjected to MS-based identification. The data of acquired polypeptides was confirmed by BLAST against NCBI NR database.

Experiments of circular dichroism and inductively coupled plasma mass spectrometry

Circular dichroism (CD) tests were conducted to characterize secondary structures of ICR-Mo and its chimeric variants in the Tris buffer [20 mM Tris-HCl [pH 8.0], 100 mM NaCl, 5% glycerol, 0.03% DDM]. The CD spectra were recorded on a Jasco Model J-1500 spectrometer (Jasco Corp., Tokyo, Japan) through continuous wavelength scanning (in triplicate) from 190 to 240 nm at a scan rate of 50 nm/min [64] and smoothed with a Savitsky-Golay filter [65]. CD spectra were analyzed using SELCON3 program (protein basis set 10) in the CDPro software package (http://lamar.colostate.edu/~sreeram/CDPro/), developed at the Department of Biochemistry and molecular Biology of Colorado State University [66]. The different percentages were measured, which corresponded to four major types of secondary structure motifs (α-helices, β-sheet, η-turn and coils) [67].

To further probe whether or not zinc ions are occupied in ICR-Mo and their derivatives, inductively coupled plasma mass spectrometry (ICP-MS) was performed. The protein samples (~0.2 mg/ml) were subjected to the NexION 300X ICP-MS instrument (PerkinElmer, USA) with helium as carrier gas [68]. The mass-to-charge ratio (m/z) was measured in the mode of kinetic energy discrimination (KED) mode. In total, seven proteins were examined, which include 3 parental enzymes (ICR-Mo [AXE82_07515], EptA and MCR-1) and 4 hybrid versions (namely TM(AXE82)-EptA, TM(EptA)-AXE82, TM(AXE82)-MCR-1 and TM(MCR-1)-AXE82).

In Vitro enzymatic assays

As described with EptA by Anandan et al. [34], we tested the enzymatic activity of ICR-Mo in vitro. In this enzymatic reaction system, the fluorescent substrate, is abbreviated as NBD-PEA from 1-acyl-2-[12-[(7-nitro-2-1,3-benzoxadiazol-4-yl) amino] dodecanoyl]-sn-glycero-3-phosphoethanolamine (Avanti Lipids, USA). The reaction system (50 μl in total) consists of 50 mM HEPES (pH 7.50), 100 mM NaCl, 0.03% of DDM, 0.2 mM NBD-PEA and 40 μM ICR-Mo [MCR-1/MCR-2 and derivatives]. The reaction proceeded at 25˚C for around 24 hrs [36, 37]. Thin layer chromatography (TLC) was performed to detect the presence of the NBD-glycerol-3-PEA. Of note, fluorescence signal of NBD-glycerol-3-PEA (and/or NBD-glycerol) separated with TLC plates was visualized by Epi blue light (455–485 nm) and a corresponding filter of a ChemiDoc MP imaging system (Biorad, CA, USA) [34].

The identity of both NBD-glycerol-3-PEA and NBD-glycerol was determined by Liquid Chromatography Mass Spectrometry(LC/MS) system (Agilent technologies 6460 Triple Quad...
LC/MS) [69]. Mass spectrometry was coupled with electrospray ionization (ESI) source, in which neutral loss ion (m/z 141) mode was set for scanning of the positive ion. The samples were eluted with the solution of methanol/0.1% methanoic acid (95:5) at 0.3 ml/min and separated with an analytical chromatographic column of Zorbax SB C18 (2.1*50 mm, 3.5 μm).

**Extraction, purification and identification of LPS-lipid A**

LPS-lipid A was extracted as described by Liu et al. [70]. Following separation with SDS-PAGE (10%), the purity of LPS specimens was judged with silver staining [36, 37]. The lipid A species that satisfied the purity criteria were then subjected to structural identification with MALDI-TOF-MS (Bruker, ultrafleXtreme) in negative ion mode with the linear detector [70].

**Confocal microscopy**

Mid-log phase cultures of *E. coli* strains were used in the challenge by colistin (4 μg/ml, 0.5 h). The oxidant sensor dye DCFH2-DA (i.e., 2',7'-dichlorodihydrofluorescein diacetate, Sigma) was utilized to detect intra-cellular accumulation of reactive oxygen species (ROS). The fluorescent dichlorofluorescein (DCF), the oxidation product of DCFH2-DA, was visualized using a Zeiss LSM 510 Meta confocal laser scanning microscope [71]. Totally, four types of *E. coli* strains used here correspond to FYJ796 [MG1655 with empty vector] which served as the negative control, FYJ832 [MG1655 carrying pBAD24::eptA], FYJ795 [MG1655 carrying pBAD24::mcr-1] and FYJ968 [MG1655 with pBAD24::axe82_07515], respectively (S1 Table).

**Chemical rescue tests**

Using two ROS inhibitors: bipyridine (a ferric chelator) and L-cysteine (a ROS scavenger), chemical rescue experiments were carried out as described by Collins and coauthors [72, 73] with minor modifications. Following different challenges, bacterial viability of *E. coli* strains (with or without icr-Mo (or mcr-1/2)) was recorded. The four different kinds of challenges employed here were i) colistin alone; ii) colistin combined with 2,2'-dipyridine [74]; iii) colistin mixed with L-cysteine [75] and iv) the mixture consisting of colistin, 2,2'-dipyridine [74] and L-cysteine [75]. In this assay, compounds were supplemented as follows: 20 μg/ml for colistin (Sigma), 500 μM for 2,2’-dipyridine (Sangon Biotech) and 10 mM for L-cysteine (Sangon Biotech). After 30 mins of incubation at 37˚C, cultures were serially diluted. The selected dilutions (10⁻⁴~10⁻⁷) were dropped onto LB agar plates (5 μl each). Colony forming units (CFU) were enumerated [76].

**Bioinformatics, structural modelling and molecular docking**

Sequence alignment of ICR-Mo (AXE82_07515) with its paralogues was conducted with Clustal Omega (http://www.ebi.ac.uk/Tools/msa/clustalo/) and processed via the program ESPript 3.0 (http://esprima.ibcp.fr/ESPript/cgi-bin/ESPript.cgi)[77]. The trans-membrane region of ICR-Mo was predicted using TMHMM server v2.0 (http://www.cbs.dtu.dk/services/TMHMM/). Using the *N. meningitis* EptA (PDB accession number, 5FGN) as structural template [34], architecture of ICR-Mo was modeled via Swiss-Model [35]. The value of GMQE (Global Model Quality Estimation) is 0.68 and the score of QMEAN (which provides a global and local absolute quality estimate on the modeled structure [78]) is -3.62. Thus, it suggests a reliable qualified structural prediction. The ready-to-dock chemical structure of PE (ID: ZINC32837871) and head group of PE (ID: ZINC02798545) was sampled from ZINC database [79].

The UCSF DOCK 6 software (version 6.7) was applied to predict binding patterns of PE molecule vs EptA and head group of PE to ICR-Mo [80]. LigPlot+ was used to illustrate the
diagrams for possible ligand-protein interaction [81]. Concretely, protein structure was processed for molecular docking using UCSF Chimera software [82]. Solvent molecules were removed. Hydrogens were added and charges were assigned using chimera tool Dock Prep. Preferred orientation of PE in EptA was searched in 20 Å space around complexed ligand dodecyl-β-D-maltoside (DDM) in 5FGN. A set of spheres located in 20 Å space around DDM was calculated by the use of the program Sphere_select. The accessory program GRID in DOCK6 software package was used to compute van der Waals potential grid and electrostatic potential grid for energy scoring. Given that PE molecule contains many rotatable bonds and is thus more flexible, anchor-and-grow algorithm was set for conformational search in docking studies. This type of flexible ligand docking allows the ligand to structurally rearrange in response to the receptor.

**Phylogenetic analysis**

The nucleotide sequence of AXE82_07515 (designated as icr-Mo) from *M. osloensis* was used as a query to perform a nucleotide BLAST with options enabled to exclude models and uncultured environmental samples. The blastn algorithm parameters were modified to display 1000 target sequences with the query word size set to 7, allowing it to return ‘somewhat similar sequences’ to the query. All sequences with a minimum of 30% identity and greater than 70% query coverage were selected and exported. In addition, members of the distantly related Sulfa-tase family, if not identified in the blast search, were included. Unique nucleotide sequences were identified using Uniqueseq (https://www.ncbi.nlm.nih.gov/CBBresearch/Spouge/html_ncbi/html/fasta/uniqueseq.cgi) and aligned using MUSCLE (https://www.ebi.ac.uk/Tools/msa/muscle/). A total of 54 unique nucleotide sequences were utilized for the subsequent phylogenetic analysis. The 5’ end of the aligned sequences was trimmed to improve conservancy and then utilized for phylogenetic analysis. jModeltest (via MEGA 7 [83]) was used to identify the best-fit nucleotide substitution model. Using MEGA 7 [83], the best model obtained from before was used to generate a maximum-likelihood tree with 1000 bootstrap replicates.

Initial tree(s) for the heuristic search were obtained automatically in MEGA 7 by applying Neighbor-Joining and BioNJ algorithms [84] to a matrix of pairwise distances estimated using the Maximum Composite Likelihood (MCL) approach, and then selecting the topology with superior log likelihood value. A GTR model with discrete Gamma distribution and Invariant sites [85] was used to model evolutionary rate differences among sites (5 categories (+G, parameter = 1.1625)).

**Supporting information**

S1 Table. Bacteria and plasmids used in this study.

(DOCX)

S2 Table. Primers used in this study.

(DOCX)

S3 Table. Comparative analyses of secondary structure motifs among the EptA, MCR-1, ICR-Mo and their derivatives.

(DOC)

S1 Fig. A scheme for the enzymatic reactions catalyzed by ICR-Mo (AXE82_07515). A. Chemical reaction proposed for the transfer of PEA to lipid A by ICR-Mo. ICR-Mo catalyzes the addition of PEA to the 1 or 4’-position of lipid A moiety anchored on lipopolysaccharides, in which final products referred to PEA-4’-lipid A and diacylglycerol. The chemical structures of molecules are depicted with ChemDraw software. B. First-half reaction proposed for the
ICR-Mo catalysis is illustrated with the removal of PEA from PE, giving the final product DG and an adduct of ICR-Mo-bound PEA. C. Second-half reaction proposed for ICR-Mo catalysis, which involves the generation of the final product Kdo2-lipid A-1 (or 4’)-PEA through the transfer of the intermediate product ICR-Mo-bound PEA to the recipient Kdo2-lipid A.

S2 Fig. The Z1140 of the *E. coli* O157: H7 strain EDL933 (a putative member of the lipid A PEA transferase family) is not functional in the trials of LBA-based colistin susceptibility. The *E. coli* MG1655 cultures in mid-log phase were spotted on LBA plates with different levels of colistin (0, 0.5, 1.0, 2.0, 4.0, 8.0, 16.0, and 32.0 μg/ml) and maintained for 20 hrs at 37˚C. Prior to bacterial spotting, 0.2% arabinose was mixed into the melted media of LBA to trigger expression of pBAD24-based MCR-like genes as we described [86, 87]. Designations: Vec, the arabinose-inducible pBAD24 vector; TM-MCR-1, a hybrid version of MCR-1 in which the TM region is replaced with the counterpart in Z1140 of the *E. coli* O157:H7 strain EDL933; TM1-Z1140, a mosaic derivative of Z1140 carrying the TM domain of MCR-1. Here, it seems likely that Z1140 and its two domain-swapped versions (TM-MCR-1 & TM1-Z1140) have no roles in conferring the colistin-susceptible MG1655 resistance to polymyxin. A representative photograph of three independent assays is given.

S3 Fig. Bioinformatic analyses of ICR-Mo (AXE82_07515). A. Transmembrane prediction of ICR-Mo. B. Sequence comparison of ICR-Mo with EptA and MCR-1/2 homologs. The polypeptide sequences of four enzymes (EptA, ICR-Mo, MCR-1 and MCR-2) were subjected to Clustal Omega (http://www.ebi.ac.uk/Tools/msa/clustalo/) for sequence alignments. The final output is generated with the program ESPript 3.0 (http://escript.ibcp.fr/ESPr ipt/cgi-bin/ESPript.cgi) [22]. The trans-membrane (TM) region was underlined in blue following the prediction with TMHMM server v2.0 (http://www.cbs.dtu.dk/services/TMHMM) and underlined in blue. The five Zn$^{2+}$-interacting residues (E248, T287, H397, D472 and H473) are indicated with green arrows, and the seven PE substrate-binding residues highlighted with red arrows included N110, T114, E118, S332, K335, H402, and H485, respectively. Identical residues are in white letters with red background, similar residues are in red letters with white background, and the varied residues are in black letters. The protein secondary structure was shown in cartoon (on top). Designations: α: α-helix; β: β-sheet; T: Turn; η: coil; PH: Periplasmic-facing helix.

S4 Fig. Structural characterization of ICR-Mo. A. SDS-PAGE (12%) profile of the purified ICR-Mo protein. B. Scheme for the ICR-Mo topology. C. Overall architecture of the full-length ICR-Mo integral membrane protein. The overall structure of ICR-Mo (AXE82_07515) in ribbon was modeled using the *Neisseria meningitis* EptA (PDB: 5FGN) as structural template. The catalytic domain is illustrated in blue, the TM region is highlighted in magenta and the two helices (PH2 and PH2’) are in light grey. The rectangle with light peach background refers to the layer of inner-membrane. The red sphere denotes zinc ion. Designations: TM, Transmembrane; PH, Periplasmic-facing helices; H, α-helix; S, β-sheet; N, N-terminus; C, C-terminus.

S5 Fig. ICR-Mo, a homolog of MCR-1/2 can hydrolyze NBD-glycerol-3-PEA to give NBD-glycerol. A. Scheme for ICR-Mo-based hydrolysis reaction of NBD-glycerol-3-PEA into NBD-glycerol. B. TLC analyses of the alternative substrate NBD-glycerol-3-PEA and its resultant product NBD-glycerol. Thin layer chromatography (TLC) was performed as Anandan
et al. [34] described with minor changes. The identities of both NBD-glycerol-3-PEA and NBD-glycerol were validated with LC/MS (seen in Fig 1). Minus denotes no addition of the enzyme of EptA/MCR-1/MCR-2/ICR-Mo.

(TIF)

S6 Fig. Comparison of bacterial growth of *E. coli* strains expressing EptA, MCR-1 or ICR-Mo on the LBA plates supplied with varied levels of colistin. The serial diluted cultures (in mid-log phase) were spotted on LBA plates with various concentrations of colistin (0, 0.5, 1.0, 2.0, 4.0, 8.0, 16.0, and 32.0 μg/ml) and maintained at 37°C for ~20 hrs. Plasmid-borne expression of icr-Mo (axe82_07515)/eptA/mcr-1 is dependent on the arabinose-inducible expression vector pBAD24 in *E. coli* MG1655. 0.2% arabinose is supplied here as an inducer. A representative result of three independent tests is given. (TIF)

S7 Fig. Functional dissection of the PE lipid substrate-recognizable cavity of ICR-Mo (AXE82_07515). A. Surface structure of ICR-Mo with the cavity required for entry and binding of PE substrate. B. An enlarged illustration for a five residues-containing, Zn$^{2+}$-binding motif. The five residues in Zn$^{2+}$-binding motif of ICR-Mo refers to E248, T287, H397, D472 and H473, respectively. C. Surface structure of ICR-Mo in counter-clockwise rotation (30°). D. Fine structural illustration of the seven residues-containing motif that is involved in binding of ICR-Mo to PE lipid substrate. The seven residues corresponded to N110, T114, E118, S332, K335, H402 and H485, respectively. (TIF)

S8 Fig. Use of *in vitro* enzymatic assays to evaluate the impact of PE-binding cavity on ICR-Mo catalysis. A. SDS-PAGE (12%) profile of the purified ICR-Mo transmembrane enzyme and its 12 derivatives. The estimated mass of ICR-Mo (and its derivatives) is ~60 kDa, indicated with an arrow. B. TLC-based analyses of *in vitro* enzymatic activities of ICR-Mo and its 12 point-mutants. The identities of both NBD-glycerol-3-PEA and NBD-glycerol were validated by LC/MS (seen in Fig 1). This suggests that only three point-mutants of ICR-Mo (N110A, T114A & S332A) retain partial enzymatic activities to hydrolyze the alternative substrate NBD-glycerol-3-PEA into NBD-glycerol. TLC experiments were conducted as described by Anandan et al. [34] with minor modifications. A representative result from three independent trials is given, and this photograph was generated through combining three different TLC plates with limited wells. (TIF)

S9 Fig. MALDI-TOF measurement of molecular mass of ICR-Mo (AXE82_07515) and its derivatives. A. MALDI-TOF determination of molecular weight of ICR-Mo protein. B. Use of MALDI-TOF to measure molecular mass of an intact hybrid version of ICR-Mo protein, TM(AXE82)-EptA. C. Molecular mass of TM(EptA)-AXE82 (a mosaic derivative of AXE82_07515 protein) revealed by MALDI-TOF. D. Molecular mass of a hybrid version of ICR-Mo (AXE82_07515) protein, TM(AXE82)-MCR-1. E. Molecular weight of TM(MCR-1)-AXE82, a hybrid version of AXE82_07515 (TIF)

S10 Fig. MS identification of ICR-Mo (AXE82_07515) and its derivatives. A. Protein sequence of *Neisseria gonorrhoea* EptA. B. Protein sequence of MCR-1. C. MS-based identification of ICR-Mo. D. MS verification of the chimeric version of ICR-Mo, TM(AXE82)-EptA. E. MS determination of a hybrid version of ICR-Mo, TM(AXE82)-MCR-1. F. MS-based elucidation of the mosaic version of ICR-Mo, TM1(MCR-1)-AXE82. G. MS identification of the
hybrid derivative of ICR-Mo, TM(EptA)-AXE82. The bold letters with yellow background denote the TM regions, and the other letters refer to catalytic domains. The underlined letters correspond to the polypeptides identified by mass spectrometry. Designations: TM(AXE82)-EptA, a derivative of AXE82_07515 whose extracellular region is replaced with its counterpart in EptA; TM(AXE82)-MCR-1, a derivative of AXE82_07515 whose extracellular region is replaced with its counterpart in MCR-1; TM1(MCR-1)-AXE82, a derivative of AXE82_07515 whose TM region is replaced with its counterpart in MCR-1; TM(EptA)-AXE82, a derivative of AXE82_07515 in which TM region is replaced with that of EptA.

S11 Fig. Circular dichroism (CD) analyses of ICR-Mo (AXE82_07515) and its chimeric versions. A. CD-based visualization of secondary structure of MCR-1 protein. B. CD spectrum of EptA protein. C. CD assays for ICR-Mo. D. CD spectrum of the ICR-Mo derivative, TM(AXE82)-EptA. E. CD spectrum of the hybrid version of AXE82_07515, TM(EptA)-AXE82. F. CD spectrum of the mosaic version of AXE82_07515, TM(AXE82)-MCR-1. G. CD profile of the mosaic version of AXE82_07515, TM(MCR-1)-AXE82. Here, the CD results of EptA/MCR-1/ICR-Mo and their domain-swapped versions indicate they share a similar conformation in protein secondary structures. Abbreviations: CD, Circular dichroism; TM(AXE82)-EptA, a derivative of AXE82_07515 whose extracellular region is replaced with its counterpart in EptA (Panel B); TM(EptA)-AXE82, a derivative of AXE82_07515 whose TM region is replaced with its counterpart in EptA (Panel B); M(AXE82)-MCR-1, a derivative of AXE82_07515 whose extracellular region is replaced with its counterpart in MCR-1 (Panel A); TM1(MCR-1)-AXE82, a derivative of AXE82_07515 whose TM region is replaced with its counterpart in MCR-1 (Panel A).

S12 Fig. Physical evidence for the presence of zinc ions in the ICR-Mo protein and its derivatives. A. Inductively coupled plasma mass spectrometry (ICP/MS)-based detection of Zn$^{2+}$ in ICR-Mo/MCR-1/EptA and the related chimeric derivatives. B. Use of ICP/MS to calculate the relative ratio of protein-bound Zn$^{2+}$ to protein. Similar to the observations we recently reported [36, 37], ICP/MS data of ICR-Mo confirms that it might possess zinc ions at the relative ratio of 1:1 (zinc: protein). Indeed, it is also consistent with scenarios seen with the crystal structure of the neissertial EptA [20, 34]. The designations of ICR-Mo derivatives are identical to those of S11 Fig.

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References
1. Centers for Disease Control and Prevention, Office of Infectious Disease. Antibiotic resistance threats in the United States, 2013. April 2013. Available at: http://www.cdc.gov/drugresistance/threat-report-2013. Accessed September 5, 2016.
2. Laxminarayan R, Amabile-Cuevas CF, Cars O, Evans T, Heymann DL, Hoffman S, et al. UN High-Level Meeting on antimicrobials—what do we need? Lancet. 2016; 388(10041):218–20. https://doi.org/10.1016/S0140-6736(16)31079-0 PMID: 27479554.
3. O’Neill LJ. Antimicrobial resistance: Tackling a crisis for the health and wealth of nations. London: Review on Antimicrobial Resistance. 2014.
4. de Kraker ME, Stewardson AJ, Harbarth S. Will 10 million people die a year due to antimicrobial resistance by 2050? PLoS Med. 2016; 13(11):e1002184. https://doi.org/10.1371/journal.pmed.1002184 PMID: 27898664; PubMed Central PMCID: PMC5127510 and AJS participated in an investigator-initiated study on the burden of antimicrobial resistance (TIMBER) supported by an unrestricted research grant provided by Pfizer Europe and bioMerieux. SH is also a member of the Editorial Board of PLOS Medicine. AJS is an employee of Hand Hygiene Australia, a national quality improvement program, and is a member of the Healthcare Associated Infection Advisory Committee of the Australian Commission for Safety and Quality in Healthcare.
5. de Kraker ME, Davey PG, Grundmann H, group Bs. Mortality and hospital stay associated with resistant Staphylococcus aureus and Escherichia coli bacteraemia: estimating the burden of antibiotic resistance in Europe. PLoS Med. 2011; 8(10):e1001104. https://doi.org/10.1371/journal.pmed.1001104 PMID: 22022233; PubMed Central PMCID: PMC3191157.
6. Caniaux I, van Belkum A, Zambardi G, Poirel L, Gros MF. MCR: modern colistin resistance. Eur J Clin Microbiol Infect Dis. 2017; 36(3):415–20. https://doi.org/10.1007/s10096-016-2846-y PMID: 27873028.
7. Li J, Nation RL, Turnidge JD, Milne RW, Coulthard K, Rayner CR, et al. Colistin: the re-emerging antibiotic for multidrug-resistant gram-negative bacterial infections. Lancet Infect Dis. 2006; 6(9):589–601. https://doi.org/10.1016/S1473-3099(06)70580-1 PMID: 16931410.
8. Falagas ME, Kasiakou SK. Collistin: the revival of polymyxins for the management of multidrug-resistant gram-negative bacterial infections. Clin Infect Dis. 2005; 40(9):1333–41. https://doi.org/10.1086/429323 PMID: 15825037.
9. Dixon RA, Chopra I. Polymyxin B and polymyxin B nonapeptide alter cytoplasmic membrane permeability in Escherichia coli. J Antimicrob Chemother. 1986; 18(5):557–63. PMID: 3027012.
10. Liu YY, Wang Y, Walsh TR, Yi LX, Zhang R, Spencer J, et al. Emergence of plasmid-mediated colistin resistance mechanism MCR-1 in animals and human beings in China: a microbiological and molecular biological study. Lancet Infect Dis. 2016; 16(2):161–8. https://doi.org/10.1016/S1473-3099(15)00424-7 PMID: 26603172.
11. Wang X, Zhang H, Sun J, Liu YH, Feng Y. The MCR-1 colistin resistance: A new challenge to global public health. Chinese Science Bulletin. 2017; 62(10):1018–29. Epub 01-22-2017. https://doi.org/10.1360/N972016-01084
27. Di Pilato V, Arena F, Tascini A, Cannatelli A, Henrici De Angelis L, Fortunato S, et al. MCR-1.2: a new carbapenemase in Salmonella enterica subsp. enterica serovar Paratyphi B. J Antimicrob Chemother. 2017; 72(12):3317–24. https://doi.org/10.1093/jac/dkx327 PMID: 28907927; PubMed Central PMCID: PMC54914657.

28. Yang YQ, Li YX, Song T, Yang YX, Jiang W, Zhang AY, et al. Colistin resistance gene mcr-1 and its variant in Escherichia coli isolates from chickens in China. Antimicrobial agents and chemotherapy. 2017; 61(5):pii: e01204-16. https://doi.org/10.1128/AAC.01204-16 PMID: 28242671.
Wei P, Song G, Shi M, Zhou Y, Liu Y, Lei J, et al. Substrate analog interaction with MCR-1 offers insight.

29. Lu X, Hu Y, Luo M, Zhou H, Wang X, Du Y, et al. MCR-1.6, a new MCR variant carried by an IncP plasmid in a colistin-resistant Salmonella enterica Serovar Typhimurium isolate from a healthy individual. Antimicrobial agents and chemotherapy. 2017; 61(5). https://doi.org/10.1128/AAC.02632-16 PMID: 28264851.

30. AbuOun M, Stubberfield EJ, Duggett NA, Kirchner M, Dormer L, Nunez-Garcia J, et al. mcr-1 and mcr-2 variant genes identified in Moraxella species isolated from pigs in Great Britain from 2014 to 2015. J Antimicrob Chemother. 2017; 72(10):2745–9. https://doi.org/10.1093/jac/dkx286 PMID: 29091227.

31. Poirel L, Kieffer N, Fernandez-Garayzabal JF, Vela AI, Larpin Y, Nordmann P. MCR-2-mediated plasmid-borne polymyxin resistance most likely originates from Moraxella pluranimalium. J Antimicrob Chemother. 2017. https://doi.org/10.1093/jac/dkx225 PMID: 29091193.

32. Kieffer N, Nordmann P, Poirel L. Moraxella species as potential sources of MCR-like polymyxin resistance determinants. Antimicrobial agents and chemotherapy. 2017; 61(6). https://doi.org/10.1128/AAC.00129-17 PMID: 28320720; PubMed Central PMCID: PMC5444148.

33. Stogios P, Cox G, Zubyk H, Evdokimova E, Wawrzak Z, Wright G, et al. Substrate recognition by a colistin resistance enzyme from Moraxella catarrhalis. ACS Chem Bio. 2018;Epub ahead of print.

34. Anandan A, Evans GL, Condic-Jurkic K, O’Mara ML, John CM, Phillips NJ, et al. Structure of a lipid A phosphoethanolamine transferase suggests how conformational changes govern substrate binding. Proceedings of the National Academy of Sciences of the United States of America. 2017; 114(9):2218–23. https://doi.org/10.1073/pnas.1612927114 PMID: 28193899; PubMed Central PMCID: PMC5338521.

35. Biasini M, Bienert S, Waterhouse A, Arnold K, Studer G, Schmidt T, et al. SWISS-MODEL: modelling protein tertiary and quaternary structure using evolutionary information. Nucleic acids research. 2014; 42(Web Server issue):W252–8. https://doi.org/10.1093/nar/gku340 PMID: 24782522; PubMed Central PMCID: PMC4086889.

36. Xu Y, Lin J, Cui T, Srinivas S, Feng Y. Mechanistic insights into transferable polymyxin resistance among gut bacteria. The Journal of biological chemistry. 2018; 293(12):4350–65. https://doi.org/10.1074/jbc.RA117.000924 PMID: 29462787; PubMed Central PMCID: PMC5668246.

37. Xu Y, Wei W, Lei S, Lin J, Srinivas S, Feng Y. An evolutionarily conserved mechanism for intrinsic and transferable polymyxin resistance. mBio. 2018; 9(2):e02317–17. https://doi.org/10.1128/mBio.02317-17 PMID: 29936432.

38. Stec B, Holtz KM, Kantrowitz ER. A revised mechanism for the alkaline phosphatase reaction involving three metal ions. J Mol Biol. 2000; 299(5):1303–11. https://doi.org/10.1006/jmbi.2000.3799 PMID: 10873454.

39. Sun J, Zhang H, Liu YH, Feng Y. Towards understanding MCR-like colistin resistance. Trends in microbiology. 2018;Epub ahead of print. https://doi.org/10.1016/j.tim.2018.02.006 PMID: 29525421.

40. Hu M, Guo J, Cheng Q, Yang Z, Chan EW, Chen S, et al. Crystal structure of Escherichia coli originated MCR-1, a phosphoethanolamine transferase for colistin resistance. Scientific reports. 2016;6:38793.

41. Ma G, Zhu Y, Yu Z, Ahmad A, Zhang H. High resolution crystal structure of the catalytic domain of MCR-1. Scientific reports. 2016;6:39540.

42. Stojanoski V, Sankaran B, Prasad BV, Poirel L, Nordmann P, Palzkill T. Structure of the catalytic domain of the colistin resistance enzyme MCR-1. BMC Biol. 2016; 14(1):81. https://doi.org/10.1186/s12915-016-0303-0 PMID: 27655155; PubMed Central PMCID: PMC5031297.

43. Biasini M, Bienert S, Waterhouse A, Arnold K, Studer G, Schmidt T, et al. SWISS-MODEL: modelling protein tertiary and quaternary structure using evolutionary information. Nucleic acids research. 2014; 42(Web Server issue):W252–8. https://doi.org/10.1093/nar/gku340 PMID: 24782522; PubMed Central PMCID: PMC4086889.

44. Serovar Typhimurium isolate from a healthy individual. Antimicrob Chemother. 2017; 72(10):2745–9. https://doi.org/10.1093/jac/dkx286 PMID: 29091227.

45. Poirel L, Kieffer N, Fernandez-Garayzabal JF, Vela AI, Larpin Y, Nordmann P. MCR-2-mediated plasmid-borne polymyxin resistance most likely originates from Moraxella pluranimalium. J Antimicrob Chemother. 2017. https://doi.org/10.1093/jac/dkx225 PMID: 29091193.

46. Stogios P, Cox G, Zubyk H, Evdokimova E, Wawrzak Z, Wright G, et al. Substrate recognition by a colistin resistance enzyme from Moraxella catarrhalis. ACS Chem Bio. 2018;Epub ahead of print.

47. Xu Y, Wei W, Lei S, Lin J, Srinivas S, Feng Y. An evolutionarily conserved mechanism for intrinsic and transferable polymyxin resistance. mBio. 2018; 9(2):e02317–17. https://doi.org/10.1128/mBio.02317-17 PMID: 29936432.
52. Beceiro A, Llobet E, Aranda J, Bengoechea JA, Doumith M, Hornsey M, et al. Phosphoethanolamine
51. Zhou Z, Ribeiro AA, Lin S, Cotter RJ, Miller SI, Raetz CR. Lipid A modifications in polymyxin-resistant
50. Trent MS, Ribeiro AA, Lin S, Cotter RJ, Raetz CR. An inner membrane enzyme in *Salmonella* and
*Escherichia coli* that transfers 4-amino-4-deoxy-L-arabinose to lipid A: induction on polymyxin-resistant
mutants and role of a novel lipid-linked donor. J Biol Chem. 2001; 276(46):43122–31. https://doi.org/10.1074/jbc.M106961200 PMID: 11535604.
53. Zhou Z, Ribeiro AA, Lin S, Cotter RJ, Miller SI, Raetz CR. Lipid A modifications in polymyxin-resistant
*Salmonella typhimurium*: PMRA-dependent 4-amino-4-deoxy-L-arabinose, and phosphoethanolamine incorporation. J Biol Chem. 2001; 276(46):43111–21. https://doi.org/10.1074/jbc.M106960200 PMID: 11535603.
54. Beceiro A, Llobet E, Aranda J, Bengoechea JA, Doumith M, Hornsey M, et al. Phosphoethanolamine modification of lipid A in colistin-resistant variants of *Acinetobacter baumannii* mediated by the pmrAB two-component regulatory system. Antimicrobial agents and chemotherapy. 2011; 55(7):3370–9. https://doi.org/10.1128/AAC.00079-11 PMID: 21576434; PubMed Central PMCID: PMC3122444.
55. McPhee JB, Lewenza S, Hancock RE. Cationic antimicrobial peptides activate a two-component regulatory system, PmRA-PmRB, that regulates resistance to polymyxin B and cationic antimicrobial peptides in *Pseudomonas aeruginosa*. Mol Microbiol. 2003; 50(1):205–17. PMID: 14507375.
56. Tamayo R, Ryan SS, McCoy AJ, Gunn JS. Identification and genetic characterization of PmRA-regulated genes and genes involved in polymyxin B resistance in *Salmonella enterica* serovar typhimurium. Infect Immun. 2002; 70(12):6770–8. https://doi.org/10.1128/IAI.70.12.6770-6778.2002 PMID: 12438352; PubMed Central PMCID: PMC133008.
57. Hankins JV, Madsen JA, Giles DK, Brodbelt JS, Trent MS. Amino acid addition to *Moraxella porci* sp. nov., isolated from pigs. International journal of systematic and evolutionary microbiology. 2010; 60(Pt 12):4246–50. https://doi.org/10.1099/ijs.0.016626-0 PMID: 2095-8137.2010.003 PMID: 28409502; PubMed Central PMCID: PMC5996029.
58. Wang Q, Sun J, Li J, Ding Y, Li XP, Lin J, et al. Expanding landscapes of the diversified *mcr-1*-bearing plasmid reservoirs. Microbiome. 2017; 5(1):70. https://doi.org/10.1186/s40168-017-0288-0 PMID: 28683827.
59. Velas AI, Arroyo E, Aragon V, Sanchez-Porco C, Latre MV, Cerda-Cuello M, et al. *Moraxella pluranimium* sp. nov., isolated from animal specimens. International journal of systematic and evolutionary microbiology. 2009; 59(Pt 4):671–4. https://doi.org/10.1099/ijs.0.006205-0 PMID: 19329585.
60. Velas AI, Sanchez-Porco C, Aragon V, Olivera A, Dominguez L, Ventosa A, et al. *Moraxella porci* sp. nov., isolated from pigs. International journal of systematic and evolutionary microbiology. 2010; 60(Pt 10):2446–50. https://doi.org/10.1099/ijs.0.016626-0 PMID: 19946050.
61. Feng Y. Transferability of MCR-1-2 polymyxin resistance: Complex dissemination and genetic mechanism. ACS infectious diseases. 2018. https://doi.org/10.1021/acsinfecdis.7b00201 PMID: 29397687.
62. Ye H, Li Y, Li Z, Gao R, Zhang H, Wen R, et al. Diversified *mcr-1*-harbouring plasmid reservoirs confer resistance to colistin in human gut microbiota. mBio. 2016; 7(2):e00177. https://doi.org/10.1128/mBio.00177-16 PMID: 27048797.
63. Lv J, Mohsin M, Lei S, Srinivas S, Wigaw RT, Lin J, et al. Discovery of a *mcr-1*-bearing plasmid in commensal colistin-resistant *Escherichia coli* from healthy broilers in Faisalabad, Pakistan. Virulence. 2018; 9(1):1–13. https://doi.org/10.1080/21505594.2018.1462060 PMID: 29667494.
64. Alves A, Marques A, Martins E, Silva T, Reis R. Cosmetic Potential of Marine Fish Skin Collagen. Cosmetics. 2017; 4(4):39. https://doi.org/10.3390/cosmetics4040039
65. Powl AM, O’Reilly AO, Miles AJ, Wallace BA. Synchrotron radiation circular dichroism spectroscopy-defined structure of the C-terminal domain of NaChBac and its role in channel assembly. Proceedings
of the National Academy of Sciences of the United States of America. 2010; 107(32):14064–9. https://doi.org/10.1073/pnas.1001793107 PMID: 20663949; PubMed Central PMCID: PMC2922593.

66. Sreerama N, Venyaminov SY, Woody RW. Estimation of protein secondary structure from circular dichroism spectra: inclusion of denatured proteins with native proteins in the analysis. Analytical biochemistry. 2000; 287(2):243–51. https://doi.org/10.1006/abio.2000.4879 PMID: 11112270.

67. Sreerama N, Woody RW. On the analysis of membrane protein circular dichroism spectra. Protein Sci. 2004; 13(1):100–12. https://doi.org/10.1110/ps.03258404 PMID: 14691226; PubMed Central PMCID: PMC2286510.

68. Loeschner K, Harrington CF, Kearney JL, Langton DJ, Larsen EH. Feasibility of asymmetric flow field-flow fractionation coupled to ICP-MS for the characterization of wear metal particles and metalloproteins in biofluids from hip replacement patients. Anal Bioanal Chem. 2015; 450(16):4541–54. https://doi.org/10.1007/s00216-015-8631-4 PMID: 25874977.

69. Gros M, Rodriguez-Mozaz S, Barcelo D. Rapid analysis of multiclass antibiotic residues and some of their metabolites in hospital, urban wastewater and river water by ultra-high-performance liquid chromatography coupled to quadrupole-linear ion trap tandem mass spectrometry. J Chromatogr A. 2013; 1292:173–88. https://doi.org/10.1016/j.chroma.2012.12.072 PMID: 23347979.

70. Liu YY, Chandler CE, Leung LM, McElheny CL, Mettus RT, Shanks RM, et al. Structural modification of lipopolysaccharide conferred by mcr-1 in Gram-negative ESKAPE pathogens. Antimicrobial agents and chemotherapy. 2017; 61(6):e00580-17. https://doi.org/10.1128/AAC.00580-17 PMID: 28373195.

71. Chen Z, Zhou Q, Zou D, Tian Y, Liu B, Zhang Y, et al. Chloro-benzoquinones cause oxidative DNA damage through iron-mediated ROS production in Escherichia coli. Chemosphere. 2015; 135:379–86. https://doi.org/10.1016/j.chemosphere.2015.04.076 PMID: 25996850.

72. Dwyer DJ, Belenky PA, Yang JH, MacDonald IC, Martell JD, Takahashi N, et al. Antibiotics induce redox-related physiological alterations as part of their lethality. Proceedings of the National Academy of Sciences of the United States of America. 2014; 111(20):E2100–9. https://doi.org/10.1073/pnas.1401876111 PMID: 24803433; PubMed Central PMCID: PMC4031491.

73. Lobritz MA, Belenky P, Porter CB, Gutierrez A, Yang JH, Schwarz EG, et al. Antibiotic efficacy is linked to bacterial cellular respiration. Proc Natl Acad Sci U S A. 2015; 112(27):8173–80. https://doi.org/10.1073/pnas.1509743112 PMID: 26100896; PubMed Central PMCID: PMC4500273.

74. Kohanski MA, Dwyer DJ, Hayete B, Lawrence CA, Collins JJ. A common mechanism of cellular death induced by bactericidal antibiotics. Cell. 2007; 130(5):797–810. https://doi.org/10.1016/j.cell.2007.06.049 PMID: 17803904.

75. Spagnuolo G, D’Anto V, Cosentino C, Schmalz G, Schweikl H, Rengo S. Effect of N-acetyl-L-cysteine on ROS production and cell death caused by HEMA in human primary gingival fibroblasts. Biomaterials. 2006; 27(9):1803–9. https://doi.org/10.1016/j.biomaterials.2005.02.022 PMID: 16289659.

76. Chen CY, Nace GW, Irwin PL. A 6 x 6 drop plate method for simultaneous colony counting and MPN enumeration of Campylobacter jejuni, Listeria monocytogenes, and Escherichia coli. J Microbiol Methods. 2003; 55(2):475–9. PMID: 14529971.

77. Feng Y, Cronan JE. The Vibrio cholerae fatty acid regulatory protein, FadR, represses transcription of pslB, the gene encoding the first enzyme of membrane phospholipid biosynthesis. Mol Microbiol. 2011; 81(4):1020–33. https://doi.org/10.1111/j.1365-2958.2011.07749.x PMID: 21771112.

78. Benkert P, Kunzli M, Schwede T. QMEAN server for protein model quality estimation. Nucleic Acids Res. 2009; 37(Web Server issue):W510–4. https://doi.org/10.1093/nar/gkp322 PMID: 19426885; PubMed Central PMCID: PMC2703985.

79. Irwin JJ, Sterling T, Mysinger MM, Bolstad ES, Coleman RG. ZINC: a free tool to discover chemistry for biology. J Chem Inf Model. 2012; 52(7):1757–68. https://doi.org/10.1021/ci3001277 PMID: 22587354; PubMed Central PMCID: PMC4020220.

80. Allen WD, Balliu TE, Mukherjee S, Brozell SR, Moustakas DT, Lang PT, et al. DOCK 6: Impact of new features and current operating performance. J Comput Chem. 2015; 36(15):1132–56. https://doi.org/10.1002/jcc.23905 PMID: 25914306; PubMed Central PMCID: PMC4469538.

81. Laskowski RA, Swindells MB. LigPlot+: multiple ligand-protein interaction diagrams for drug discovery. J Chem Inf Model. 2011; 51(1):2778–86. https://doi.org/10.1021/ci200227u PMID: 21915903.

82. Pettersen EF, Goddard TD, Huang CC, Couch GS, Greenblatt DM, Meng EC, et al. UCSF Chimera—a visualization system for exploratory research and analysis. J Comput Chem. 2004; 25(13):1605–12. https://doi.org/10.1002/jcc.20084 PMID: 15264254.

83. Kumar S, Stecher G, Tamura K. MEGA7: Molecular evolutionary genetics analysis version 7.0 for bigger datasets. Molecular biology and evolution. 2016; 33(7):1870–4. https://doi.org/10.1093/molbev/msw054 PMID: 27004904.
84. Criscuolo A, Gascuel O. Fast NJ-like algorithms to deal with incomplete distance matrices. BMC bioinformatics. 2008; 9:166. https://doi.org/10.1186/1471-2105-9-166 PMID: 18366787; PubMed Central PMCID: PMC2335114.

85. Gatto L, Catanzaro D, Milinkovitch MC. Assessing the applicability of the GTR nucleotide substitution model through simulations. Evolutionary bioinformatics online. 2007; 2:145–55. PMID: 19455208; PubMed Central PMCID: PMC2674669.

86. Gao R, Li Y, Lin J, Tan C, Feng Y. Unexpected complexity of multidrug resistance in the mcr-1-harbouring Escherichia coli. Sci China Life Sci. 2016; 59(7):732–4. https://doi.org/10.1007/s11427-016-5070-1 PMID: 27178474.

87. Li Z, Tan C, Lin J, Feng Y. Diversified variants of the mcr-1-carrying plasmid reservoir in the swine lung microbiota. Sci China Life Sci. 2016; 59(9):971–3. https://doi.org/10.1007/s11427-016-5111-9 PMID: 27520829.