# 生病豬隻分離之產廣效性乙內醯胺酶大腸桿菌 之抗藥性分析

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#### 摘要

廣效性乙內醯胺酶(extended-spectrum  $\beta$ -lactamases; ESBLs) 是由 一些腸內菌科細菌產生,可破壞cephalosporins、penicillin及aztreonam 類抗菌劑之活性。自2000年起,世界各地都有產ESBL腸內菌科細菌 感染的人類病例,而在經濟動物、食物生產鏈及伴侶動物也檢測得到 產ESBL之大腸桿菌(ESBL producing Escherichia coli, ESBL-EC)。 本計畫自2013年至2019年間,從生病豬隻分離得E.coli 239株,其中9.2 % (22/239)確認為ESBL-EC。檢測22株ESBL-EC所帶之bla基因,最多 數為bla<sub>CTX-M</sub> 90.9% (20/22), 其次為bla<sub>TEM</sub>基因31.8% (7/22), 及bla<sub>SHV</sub> 基因為9.1% (2/22),部分菌株帶有兩種bla基因。其中最常見的型別為 bla<sub>CTX-M-55</sub>, 為36.4% (8/22), bla<sub>CTX-M-14</sub> 及bla<sub>TEM-234</sub>則為次要的型別, 各佔27.3% (6/22)。所有的ESBL-EC皆對tetracyclin產生抗藥性,95.5% (21/22) 對於 ceftiofur 及 SXT 具抗藥性,其次對於 gentamycin、 enrofloxacin及amoxicillin之抗藥性別為81.8% (18/22)、54.5% (12/22) 及31.8% (7/22);而本研究之所有菌株對於imipenem皆具感受性。

## **Prevalence and Characteristics of Extended-spectrum**

### $\beta$ -lactamase-producing *Escherichia coli* in Diseased Pigs

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#### **Abstract**

Extended-spectrum β-lactamases (ESBLs) are a rapidly evolving group of β-lactamases that share the ability to hydrolyze third-generation cephalosporins, penicillins, and aztreonam. Since 2000, the incidence of ESBL-producing *Enterobacteriaceae* infections increased globally in human cases. In parallel, ESBL-producing *Escherichia coli* (ESBL-EC) has increasingly been reported in livestock, the food chain and companion animals. In this study, 239 isolates of *E. coli* from diseased pigs samples were included from 2013 to 2019. Twenty-two ESBL-EC out of 239 isolates were confirmed by phenotyping and genotyping, which were mainly presented as *bla* genes, such as *bla*<sub>CTX-M</sub> (90.9%), *bla*<sub>TEM</sub> (31.8%) and *bla*<sub>SHV</sub> (9.1%). The amount in which, the dominant *bla* genes variant were *bla*<sub>CTX-M-55</sub> (36.4%), followed by *bla*<sub>CTX-M-14</sub> and *bla*<sub>TEM-234</sub> (27.3%). All the ESBL-EC were resistant to tetracycline, mostly resistant to ceftiofur and SXT (95.5%), followed by gentamycin (81.8%), enrofloxacin (54.5%) and amoxicillin (31.8%). All isolates remain the sensitivity to imipenem.