NUTRITIONAL REQUIREMENT AMONG PSEUDOMONAS AERUGINOSA ISOLATES RECOVERED FROM RESPIRATORY CLINICAL SPECIMENS AT A TERTIARY HOSPITAL FROM SOUTH OF BRAZIL

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ABSTRACT

We screened 349 isolates of P. aeruginosa from cystic fibrosis (CF+) and non-cystic fibrosis (CF-) patients for the auxotrophy. Fourteen (4.0%) were auxotrophic and among them only one was recovered from CF-patient showing that this characteristic is strongly associated with cystic fibrosis. In total, a requirement for 5 different compounds (or combination) was verified and, of these, methionine was the most common single amino acid required. Only one auxotrophic isolate was no able to produce biofilm in vitro.

Key words: Pseudomonas aeruginosa, auxotrophy, cystic fibrosis, biofilm

Pseudomonas aeruginosa is the most common organism isolated from the respiratory tract of patients with cystic fibrosis (CF). Once established in the airways of patients with CF, P. aeruginosa undergoes some modifications to adapt and persist in this environment. Some isolates of P. aeruginosa from patients with CF also have specific nutritional requirements and they are called auxotrophic because of their inability to grow in a basic medium containing mineral salts and glucose as the sole carbon source (3).

Generally, P. aeruginosa from clinical and environmental sources are prototrophic, but approximately half of CF patients colonized with P. aeruginosa are harboring isolates that require the provision of specific factors for growth (3, 4). The most common nutritional requirement of these auxotrophic isolates is methionine, but other amino acids such as leucine, arginine, or ornithine may also be required by a minority of isolates. Sometimes, auxotrophic isolates are also found in patients with bronchiectasis, but they have not been isolated from patients with other conditions (1). Difficulty for eradication and antimicrobial resistance are characteristics presented by auxotrophic isolates that have been related to its easy adaptation inside lung with CF and the ability to produce biofilm (1).

The aim of the study presented here was to evaluate the nutritional requirement for P. aeruginosa isolates recovered from sputa and endotracheal aspirate of CF+ and CF- patients. Furthermore, we evaluated the ability of biofilm production for the auxotrophic isolates.

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A total of 349 *P. aeruginosa* isolates were included in the study, 165 (47.3%) and 184 (52.7%) isolates from CF+ and CF- patients, respectively. Non-CF hospitalized patients had no record of bronchiectasis documented. All isolates were identified by using a NC32 panel and WalkAway® 96SI automated system (Siemens, Newark, DE, USA) for reading. Also, Gram stain, oxidase test and colony morphology were concomitantly inspected. All isolates were recovered from respiratory clinical specimens (endotracheal aspirate and sputum). More than one isolate from the same culture could be included in the study once that occurrence of auxotrophic and prototrophic isolates has already been concomitantly described.

A 0.5 McFarland suspension of each isolate was diluted 1:10 in saline. An aliquot of 0.3µL of the suspension was applied onto Mueller-Hinton agar (complete medium) and onto minimal agar medium (MAM) just containing mineral salts and glucose (minimal medium) using a Steers multipoint inoculator. The plates were incubated at 37ºC and were screened after 48h. Prototrophic isolates were able to grow in both media while auxotrophic isolates were unable to grow on MAM. Two methods were used to verify the amino acid nutritional requirement. Firstly, MAM plates individually containing 20µg/mL one of the 23 amino acids (L-forms of alanine, arginine, asparagine, aspartic acid, cysteine, cystine, glutamic acid, glutamine, glycine, histidine, hydroxy-L-proline, isoleucine, leucine, lysine, methionine, ornithine, phenylalanine, proline, serine, threonine, tryptophan, tyrosine, and valine; Sigma Chemical Co., St. Louis, Mo.) were inoculated as described above. Growth in the presence of a specific amino acid indicates that the bacteria need this component, while the failure to grow in this condition suggests that one or more amino acids in combination or a distinct growth factor is needed. Secondly, all amino acids less one were added into MAM plates. The specific requirement of one or more amino acid is evident when the isolate is unable to grow on medium without this amino acid.

The ability to produce biofilm for the auxotrophic isolates was determined by microtiter plate assay according to the protocol previously described (5).

Of the 349 isolates evaluated, 14 (4%) were auxotrophic. Among these auxotrophic isolates, only one isolate were recovered from a CF- patient. Table shows that among the 14 auxotrophic strains identified, the most common single amino acid required was methionine (8 isolates), followed by arginine and asparagine (2 isolates each). One isolate grew only on a combination of alanine and proline, and other required a combination of methionine, cysteine and threonine (Table 1). Because the presence of some amino acids are required in order to allow the growth of auxotrophic isolate, it is possible that these amino acids are present in the airways of CF patients. *P. aeruginosa* wild-type are nutritionally versatile. Thus, we speculate that this would be the original mode of the respiratory tract colonization and that wild-type isolate would be substituted by its auxotrophic variants during the infection process (3). The selection process of auxotrophic isolates in the lung is not clear; however it is known to be a bacterial strategy to prevent the waste of energy through downregulation of a metabolic way, in view to reduce the production of some substances freely present in the environment. Synthesis of amino acid is energetically expensive and bacteria with loss of ability to produce some amino acid would have metabolic and energetic advantages (7).

Some studies have shown that auxotrophic *P. aeruginosa* isolates are highly associated with acute exacerbations of respiratory infection in CF patients (2, 3, 4), and that these isolates are often more resistant to antimicrobial agents than prototrophic wild-type isolates (2). This antimicrobial resistance can be explained, in part, due to the biofilm production, once that there is a microenvironment in the CF lung that permit the ideal condition for the biofilm formation. Indeed, in our study, thirteen of the 14 auxotrophic strains presented weak- (10 isolates) and moderate- (3 isolates) biofilm production in vitro (Table 1). Resistance rates for the auxotrophic isolates were of 57.1% to ciprofloxacin; 35.7% to
gentamicin; 28.6% to amikacin, aztreonam, ceftazidine; 21.4% to imipenem; and 14.3% to meropenem and piperacillin/tazobactam. One isolate, auxotrophic to arginine, showed resistance to all antibiotics tested except polymixin (data not shown).

In this study, the most required amino acids were methionine and arginine, which were previously reported as the most common ones for the auxotrophic isolates (1, 4). High level serum of the key-precursor of methionine, called sulphadenosylmethionone, is verified in CF patients (6). Thus, methionine synthetase inhibitors and antimicrobial therapy together could be useful for the treatment of these patients (6).

To our knowledge, this is the first description of auxotrophy phenomenon among Brazilian isolates of \textit{P. aeruginosa} from CF patients. On the other hand, a potential limitation of this study lies in the fact that the genetic background of the isolates was not evaluated and, thus, the origin of auxotrophic from prototrophic mutants can not be clearly defined.

Finally, we conclude that methionine was the main nutritional requirement among auxotrophic isolates and that this phenomenon was more frequently observed in \textit{P. aeruginosa} recovered from sputa of CF patients indicating that this is a characteristic of this bacteria in this group of patients.

### Table 1. Nutritional requirement and biofilm production among 14 auxotrophic \textit{Pseudomonas aeruginosa} isolates.

| Isolate number | Cystic fibrosis | Nutritional requirement | Biofilm production\textsuperscript{a} |
|----------------|-----------------|-------------------------|----------------------------------------|
| 86             | yes             | methionine              | no-producing                           |
| 101            | yes             | methionine              | weak                                   |
| 144            | no              | methionine              | weak                                   |
| 146            | yes             | methionine              | weak                                   |
| 165            | yes             | arginine                | weak                                   |
| 178            | yes             | arginine                | weak                                   |
| 197            | yes             | alanine and proline     | weak                                   |
| 214            | yes             | asparagine              | moderate                               |
| 233            | yes             | asparagine              | moderate                               |
| 236            | yes             | methionine              | weak                                   |
| 338            | yes             | methionine              | moderate                               |
| 340            | yes             | methionine              | weak                                   |
| 342            | yes             | methionine              | weak                                   |
| 345            | yes             | methionine, cysteine and threonine | weak                                   |

\textsuperscript{a}Classification of biofilm production determined by a microtiter plate assay according to Stepanovic et al (6).

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