USEFULNESS OF SPUTUM GRAM STAIN AND CULTURE FOR DIAGNOSIS OF PNEUMONIA IN A GERIATRIC INSTITUTION

Jean-Jacques Lloveras, Mohamed-Issam Shukr, Claude Pinos, Anissa Lindoulsi, Philippe Grima
Centre de Reeducation Fonctionnelle, Bagnères de Luchon, FRANCE

SUMMARY:
The value of bacteriological assessment of sputum samples is controversial during lower respiratory tract infections. We retrospectively studied sputum Gram stain and culture in a geriatric population during a two-year period. A total of 42 sputum samples were send to the laboratory; among them, 24 (57%) allowed a diagnosis with a predominant morphotype.

Gram positive and Gram negative were equally distributed, but staphylococci and pseudomonas were most frequently found, as it is usually reported in the setting of nosocomial infections. In geriatric units and in long-term care facilities, diagnostic tests for management of lower respiratory tract infections are rarely performed, but the ratio cost-efficacy of this bacteriological testing could be improved, if considered the shortness of evolution of broncho-pulmonary infectious episodes.

Key words: pneumonia, geriatrics, sputum.

INTRODUCTION:
Broncho-pulmonary episodes are the second cause of nosocomial infections; among bacterial complications, they represent in geriatric patients one of the leading cause of morbidity and mortality. Numerous guidelines recommend not to perform bacteriological studies (ie Gram stain sputum and culture) and to treat patients with Amoxicillin – Clavulanate or a third generation cephalosporin without anti-Pseudomonas activity associated with a macrolide or a fluoroquinolone in severe cases. However, in geriatric inpatients, hospital-acquired pneumonia are often caused by different and more resistant pathogens then in the community. In this study, we tried to precisely in this setting the prevalence of positive sputum samples and the causative pathogens.

MATERIAL AND METHODS:
During a two-year period, from January 2007 to December 2008, we retrospectively studied all the sputum samples send to the laboratory in a 26-beds unit. Mean age of the patients was 80,5 years, with a mean hospitalization time of 33 days. The samples were collected, if feasible, when a feature of pneumonia was present. Gram coloration and bacterial culture was performed in every sample in the 24 hours after collection, before antibiotic treatment. Microorganisms were identified and sensitivity to antibiotics precised and samples were classified as negative, positive with bacterial pathogens, or positive with yeasts only.

We considered samples as positive if a microorganism was predominant or if two potential pathogens were present.

RESULTS:
During the study period, 456 patients were admitted, corresponding to 16024 patient-days. Sputum samples have been obtained from 42 patients, corresponding to 9 per 100 admissions and 3 per 1000 patient-days. Among them, 24 (57%) revealed a predominant causative microorganism.

Seven samples (17%) were strictly negative. A total of 30 bacterial pathogens were identified, equally distributed between Gram positive and negative: Staphylococcus aureus: 14, whose 2 MRSA, Diplococcus Pneumoniae: 2 Pseudomonas: 6; Haemophilus: 4; Moraxella: 1; E.Coli: 1; Enterobacter: 1; Serratia: 1 (cf Fig. 1.). In 11 cases, Candida species was isolated as the only causative microorganism.

DISCUSSION:
Lower respiratory tract infections are one of the leading cause of morbidity and mortality in elderly. Usefulness of sputum Gram stain and culture in community-acquired pneumonia or inpatient setting is controversial. In adults, some guidelines do not recommend to perform this bacteriological test and for some authors, it should be restricted to specific situations, mainly suspicion of drug-resistant bacteria.

In geriatric patients, sputum examination is rarely performed: in long-term care facilities, a recent study reported only this test in 3,3% of presumed lower respiratory tract infections. In fact, in older subjects, expectoration is rarely present: its occurrence has been evaluated to 21% of cases in a recent study where mean
The age of patients with nosocomial pneumonia was 77 years (4). However, for some authors, when performed, this technique is able to identify a predominant morphotype in 30 to 40% cases (5). Such values of percentage of positive samples compared to all the examined sputum samples have been found in four others studies (6, 7, 8, 9).

In order to improve reliability of sputum Gram stain and culture, it seems important to process quickly samples to perform examinations before administering antibiotics (10). Failure of these rules, with absence of sending or examination of the samples appears partially responsible for the low sensitivity of the technique.

However, interpretation of sputum samples must also take into account oropharyngeal colonization with Gram negative bacilli, which is frequent in elderly persons (11). But, in this study, examination of sputum was indicated in presence of a clinical suspicion of a lower respiratory tract infection and Gram negative are well recognized as causative pathogens of nosocomial pneumonia. Staphylococcus aureus and pseudomonas are mainly reported in different studies (4). Moreover, laboratory processing was performed in the 24 hours: despite this, the high proportion of bacteria-associated Candida isolates in our study is probably in relation with age and immunodepression of the patients, but a delay in processing is associated with a increased number of yeasts isolates (12).

In this setting of geriatric units or long-term care facilities, cost-efficacy of this test should be evaluated; for that purpose, ratios of number of bacteriological examinations per 100 admissions or per 1000 patient-days could be used.

**CONCLUSION:**

Value of sputum Gram stain and culture is controversial for diagnosis of pneumonia. In a geriatric population, despite a low frequency of feasible examination, sputum examination could be of diagnostic value, particularly in order to isolate a drug-resistant pathogen in a setting of nosocomial pneumonia.

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**Diagram:**

- **Staphylococcus Aureus**: 41%
- **MRSA**: 20%
- **Diplococcus Pneumoniae**: 13%
- **Pseudomonas**: 7%
- **Haemophilus**: 7%
- **Moraxella**: 7%
- **E.Coli**: 7%
- **Enterobacter**: 3%
- **Serratia**: 3%
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Address for correspondence:
Jean-Jacques Lloveras
CRF, 5 Cours des Quinconces, 31110 Bagneres de Luchon, FRANCE
Tel: 33 05 61 79 93 02; Fax: 33 05 61 79 93 13; e-mail: jjlloveras@yahoo.fr