

MICROBIOLOGICAL EVIDENCE OF *Helicobacter pylori* FROM PATIENTS SUFFERING FROM GASTRODUODENAL DISEASE : (a)

Reçu le 26/09/2009 – Accepté le 23/10/2011

L. MEDOUAKH¹, A. BENSOLTANE

¹ Laboratoire de microbiologie alimentaire et industrielle, Département de Biologie, Faculté des sciences, Université d'Oran 31100. Algérie.

E.mail: medouakhlinda@yahoo.fr

Résumé

Notre étude a porté sur la recherche de *H.pylori* à partir de biopsies gastriques chez dix patients atteints de maladies gastroduodénales, et ayant subi une endoscopie digestive haute. Cette bactérie a été mise en évidence par le test rapide à l'uréase, l'examen cytologique, l'examen histologique et la culture. Nous avons pu isoler *H.pylori* de sept patients parmi dix patients infectés par *H.pylori*, signalé par le test de l'uréase et l'histologie. Le processus de leur identification est basé sur les caractères morphologiques et biochimiques. Notre recherche est renforcée par l'antibiogramme pouvant guider vers une antibiothérapie visant à éradiquer *H.pylori*.

Mots clés: *H. pylori*, biopsies gastriques, maladies gastroduodénales, antibiogramme, éradication.

Abstract

Studies of gastric biopsies taken from ten hospitalised patients suffering from gastroduodenal diseases and were subjected to a oesogastroduodenal fibroscopy, with a fast urease test, histological, cytological examination and a microbiological culture make the evidence of *H.pylori*. We had isolated *H.pylori* from seven patients among the ten patients having an infection with *H.pylori* announced by histology and urease test. *H.pylori* identification was based on the morphological and biochemical characters. Our research was reinforced by the realization of an antibiogram which can guide us towards an antibiotherapy for the eradication of *H.pylori* infection.

Keywords: *H.pylori*, gastric biopsies, gastroduodenal diseases, antibiogram, eradication.

ملخص

H. pylori.
H. (10) (07) *H. Pylori*
H. pylori *pylori*
H. pylori : _____

Helicobacter pylori is a common bacterium that is present in millions of people world wide [1, 2]. *H.pylori* is a fastidious Gram-negative, curved rod which is associated with active chronic gastritis, a gastroduodenal ulcer disease and in the development of a gastric cancer [3, 4, 5]. It is the principal cause of chronic active (type B) gastritis [6, 7], and a Co-factor of the gastroduodenal ulcer disease [8, 9], and it is a risk factor for gastric cancer [10, 11, 12].

Many works studied the relationship between the antral infection by *Helicobacter pylori* and its implication in the genesis of the majority of the chronic gastritis and ulcers [13]. The role of *Helicobacter pylori* in the gastric inflammatory diseases is very discussed, and since, a strong correlation was shown between the presence of this bacterium in the stomach and the existence either of a gastrite, or of a peptic ulcer [2, 14]. *Helicobacter pylori* is also associated with gastric tumoral pathology because it was associated with certain gastric carcinomas [15, 16]. This bacterium which is not found in the human stomach would be either responsible, or a Co-factor of gastric inflammatory diseases is in any case an excellent marker as of these diseases [17].

H.pylori infection can be diagnosed by invasive techniques requiring endoscopy and biopsy (histological examination, culture and rapid urease test) and by non-invasive techniques (serology, the C13-urea breath test and the stool antigen test) [14, 18,19]. This infection is difficult to cure, requiring combination therapy with antibiotics and a proton pump inhibitor for eradication [20, 21, 22].

The goal of this study is to establish from a population of patients having undergone a high digestive endoscopy, with gastric biopsies, for varied digestive symptoms. Following a diagnosis which was carried out by the gastro- enterologist at the court of the fibroscopy, one can go in the search of this bacterium by bacteriological and histological examinations.

MATERIAL AND METHODS

Study carried out on the gastric biopsies coming from 10 patients, reached gastroduodenal diseases (chronic gastritis, duodenal ulcer or gastric ulcer) and outwards antibiotics taken or anti-secretoiries. These patients presented themselves at jeun and undergone an Oesogastroduodenal fibroscopy. The gastric biopsies were taken up on the level of antrum (2 cm of the pylore) with grips biopsies beforehand disinfected and rinsed [23, 24, 25]. Three antral biopsies intended for histological examination are fixed in formol at 10 % [23,24].Three to four antral biopsies for bacteriological studies are transported in physiological serum maintained at 4°C and are practiced in less than one hour [23, 26].

Urease test

This test is based on the search for ureasic activity of *H.pylori* [27]. The biopsic fragment deposited in a tube containing the urea-indol, incubated at 37°C. The readings of the results were carried out at the end of 20 and 30 min [25, 28, 29,30].

Histological examination

This examination allows visualizing spiral bacteria [6,23]. It carries out on histological cuts prepared from the gastric biopsies, coloured with modified Giemsa colouring, and then observed in strong enlarging focus [6,31].

Cytological examination

This examination is based on the setting in evidence of spiral bacteria by a Gram stain of impressed biopsy [6, 31]. The smear is prepared by friction of the biopsy mucus on a side blade, coloured with the modified Gram stain (carbofuxine 0,3%), then observed under the optical microscope [32, 33].

Culture

The biopsy is crushed in 1 ml of broth heart-brain so as to release the bacteria. Then the totality of the crush was cultured on two Petri dishes respectively containing the chocolate agar (supplemented with 10% human blood) and the medium based on Wilkings Chalgren enriched by human blood (10%) and antibiotics (vancomycin, cefsulodin and cycloheximid).These plates were incubated for 3 to 5 days at 37°C under microaerophilic conditions [23, 29, 31, 32]. The isolated bacteria were identified as *H. pylori* on the basis of morphological colonies, Gram negative staining, positive urease tests, oxidase and catalase activities [34, 35].

Antibiogramm

This test allows the study of the *H.pylori* sensitivity to antibiotics by the methods of diffusion agar on the Muller-Hinton medium agar additionned with 10% human blood, inoculated with inoculum 3,0 Mc Farland [36, 37]. The incubation carries out at 37°C during 72h in micro aerobic conditions [24, 37].The readings of the results were carried out by the measurements of diameters of inhibition zones [23, 24].

RESULTS AND DISCUSSION

The demonstration the pathogenic role of *H.pylori* in the gastroduodenal diseases returns on the search for this bacterium which is frequently necessary [38]. In our study, we sought *H.pylori* according to invasive methods requiring the realization of the gastric biopsies [18, 25, 29, 39, 40].

The majority of the patients presented a chronic gastritis and a duodenal ulcer. Ten patients having an infection with *H.pylori* documented by the histology, the urease test and sometimes the cytological test were included. We had isolated seven *H.pylori* (Hp 1, Hp2, Hp3, Hp4, Hp5, Hp6, Hp7) from the ten patients infected with *H.pylori* (Table 1).

Table 1: Different invasive methods of *H.pylori* research.

| Number of Patients | Urease test | Cytologica | Histologic al | Microbiolo gical | Diagnostic |
|--------------------|-------------|------------|---------------|------------------|-------------------------------------|
| 1 | + | + | + | + | Chronic Gastritis |
| 2 | + | - | + | + | Chronic Gastritis |
| 3 | + | + | + | + | Duodenal Ulcer |
| 4 | + | - | + | + | Chronic Gastritis |
| 5 | + | + | + | - | Gastric ulcer |
| 6 | + | + | + | - | Chronic Gastritis |
| 7 | + | + | + | - | Duodenal Ulcer |
| 8 | + | + | + | + | Duodenal Ulcer |
| 9 | + | + | + | + | Duodenal Ulcer Chronic Gastritis |
| 10 | + | + | + | + | Duodenal Ulcer |

+: Positive, -: Negative

Urease test

The research, directly on the level of the biopsy is a fast and intense ureasic activity, is a major argument in favour of the presence of *H.pylori* [18, 30, 41, 42]. The urease test gives a positive reaction, represented by the turn of indicator of pH into the pink. This indicates the presence of the bacterium [24,31,39]. Urease produced hydrolysis of the urea normally present in stomach from ammonia and carbon dioxide. Ammonia neutralizes the micro environment of the bacterium by protecting it from gastric acidity [23, 43].

Histological examination

The microscopic observations of the histological cuts showed the presence of particular forms of *H.pylori* curved or spiral in the crypts and on the surface of the gastric mucous. (Fig.1)The presences of these forms were conformed by other scientific works [18, 24, 29, 31].The presence of *H.pylori* on the level of gastric epithelium made in evidence the reports of contact between this bacterium and the epithelial gastric cells, site preferentially of *H. pylori* [44].

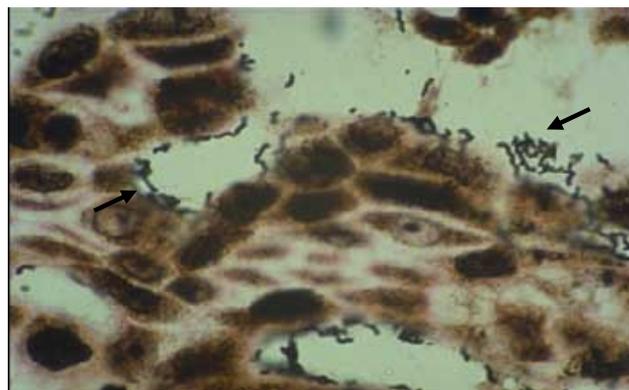


Figure 1: Examination anatomopathology of biopsic gastric by the technique of colouring of "Warthin Starry ", indicating the presence of *H. pylori*

Cytological examination

The cytological examination of gastric biopsy coloured with Gram stain revealed in the majority of the cases the presence of the shapes of curved Gram negative bacilli or slightly spiral (Fig.2).They are tendency to be many in certain zones of the smear. The presence of these forms in biopsic imprint was already notified by many authors [18, 23, 24, 29, 31]. These authors confirmed the importance of this test which informed us for the presence or the absence of the bacterium.

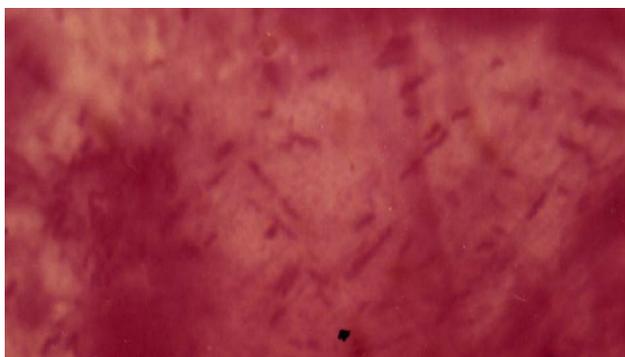


Figure 2: Microscopic observation of a print biopsy after Gram stain

Culture

The morphological methods leave sometimes a doubt contrary to the culture which only allows an unquestionable identification of the bacterium. [38, 45]. The process of identification is founded on the morphological and biochemical characters [18, 24, 29, 31].

Macroscopic aspect

The culture of the gastric biopsy on the enriched medium with blood revealed appearance of small colonies, of 1 to 2mm of diameter. They are greys or transparent, gleaming, discreetly convex, round and of regular contour (Fig.3). This aspect corresponded to a *H.pylori* and correlate with different scientists toward the world [6, 18, 28, 29, 31, 33].



Figure 3: Aspect of the colonies of *H.pylori* developed on chocolate agar after 48h of incubation with 37°C in microaerobiose.

Fresh state

The examined bacteria with fresh state appear very mobile by spiral movements and of rotation. This mobility is due a share to its spiral form and different leaves to the presence of polar flagella. [17, 23].

Microscopic aspect

Colonies coloured with Gram stain showed that this bacterium which is negative gram bacilli, curved, in comma or the shape of C, V or S, with regular contour (Fig.4). This morphology is in particular of *H.pylori*. These results correlate with the findings of *H.pylori*. [6, 17, 18, 32].



Figure 4: Microscopic observation of *H.pylori* after Gram stain from a culture of 48 h on chocolate agar.

Biochemical study

Identification is supplemented by the research of the biochemical characters (catalase, oxydase, and urease) [28, 29, 31]. We noted that the presence of oxydase, catalase and a very powerful urease in all seven isolated bacteria. Based on these biochemical characters we confirmed that was a *H.pylori* [6, 17, 23, 46].

In our study, the microbiological culture was negative among 3 patients from the 10 hospitalised patients having an infection with *H. pylori* announced by the histology, the urease test and sometimes the cytological test. Failure of the culture even in the most tested centers can be explained by the sensitivity of the technique which is extremely depended on the transport conditions, storage and culture provided in the laboratory [25, 29, 46].

In the same way, the irregular distribution of the bacterium in the gastric mucous can contribute to wrongfully negative results [46, 47, 48]. This also explains the existence of the negative results in certain cases of the cytological test (cytology negative whereas infection is present). For that, certain authors recommended the realization of several biopsies at different sites from the gastric mucous, thus improving the output of culture [46,48, 31].

Antibiogramm

Study of the *H.pylori* sensitivity to antibiotics, sometimes it is useful in therapeutic choice which showed the excellent activity of the majority of the antibiotics tested.

All strains isolated were sensitive for the majority of antibiotics tested in particular the B-lactamines (amoxicillin, penicillin) where the diameter of inhibition was superior to 21 mm and 29 mm respectively. Also, strains of *H.pylori* present sensitivity against erythromycin, chloramphenicol, tetracycline, gentamycin, metronidazole, tobramycine and furane. This sensitivity was already signalled by many authors [37, 49, 50, 51].

On the other hand, strains of *H.pylori* showed a remarkable resistance to antibiotics with nalidixic acid, cefsulodin where the diameter of inhibition was inferior to 20 mm and 22 mm respectively, and trimethoprim and sulfamide with the diameters of inhibition were below to 16mm. This was also quoted by several authors [31, 50, 52]. The resistance of *H.pylori* to antibiotics is one of the major determinants of failure of the treatments for the eradication of the *H.pylori* [5, 49].

Acknowledgments: The author's wish to thank the University of Mostaganem for their help for the practical works especially the microbiological laboratory. Also, we wish to thank the personal of the military medical laboratory and the gastro-enterology service-Oran.

REFERENCES

- [1]- Myllyluoma E., Kajander k., Mikkola H., Kyronpalo S., Rasmussen M., Kankuri E., Sipponen P., Vapaatalo H. and Korpela R., "Probiotic intervention decreases serum gastrin-17 in *Helicobacter pylori* infection". *Dig. Liv. Dis.* 39 (2007), pp. 516-523.
- [2]- Carolyn J., Hildreth M.D., Cassio Lynn M.A., Richard M. and Glass M.D., " *Helicobacter pylori*". *JAMA.* 300 (11) (2008).
- [3]- Lee A., Fox J. and Hazell S., "Pathogenicity of *Helicobacter pylori*: a perspective". *infect. Immun.* 61 (1993), pp.1601-1610.
- [4]- Marshall, B.J., "Helicobacter pylori". *Gastroenterol.* 89 (1994), pp. 116-27.
- [5]- de Korwin J.D., "Recommandations d'éradication de *Helicobacter pylori* en 2008". *Hépat. Gastro.* 15 (5) (2008), pp. 363-70.
- [6]- Sobhani I., Vallot T. and Mignon M., "*Helicobacter pylori*, une bactérie Redécouverte . Son implication dans les maladies gastroduodénales ". *Press. Med.* 24 (1995), pp. 67-79.
- [7]- Megraud, F. and Wyatt J.L., "Maladie ulcéreuse et gastrite à l'heure d'*H. pylori*". Conclusion et recommandation du jury. *Gastroenterol. Clin. Biol.* 20 (2000), pp. 155-162.
- [8]- Megraud F. and Lamouliatte H., "*H.pylori* and duodenal ulcer". Evidence suggesting causation *digdis.* 38 (1997), pp. 632-644.
- [9]- Megraud F. and Lamouliatte H. "*H. pylori* and gastric ulcer" .Evidence suggesting causation *digdis.* 37 (1999), pp.769-72.
- [10]- Brenner H., Bode G. and Boeing H., "*Helicobacter pylori* among offspring of patients with stomach cancer". *Gastroenterol.*118 (2000), pp. 31-5.
- [11]- El Omar E. M., Oien K. Murray L. S. *et al.*, "Increased prevalence of precancerous changes in relatives of gastric cancer patients: critical role of *H. pylori*". *Gastroenterol.* 118 (2000), pp. 22-30.
- [12]- Uemura N., Okamoto S., Yamamoto S. *et al.*, "*Helicobacter pylori* infection and the development of gastric cancer". *N. Engl. J. Med.* 345 (2001), pp. 784-788.
- [13]- Colin R., "Heliobacter pylori en route vers l'an 2000. Quels sont les résultats de la conférence de consensus et pourquoi ? *Actual. Innovat. Med.* 2 (1996), pp. 1-5.
- [14]- Rehcinski T., Chmiela M., Malecka-Panas E., Planeta-Malecka I. and Rudnicka W., "Serological indicators of *Helicobacter pylori* infection in adult dyspeptic patients and healthy blood donors". *Microbiol. Immunol.* 41 (1997), pp. 337-393.
- [15]- Suganuma M., Kurusu M., Okabe S. *et al.*, "*Helicobacter pylori* membrane protein 1: a new carcinogenic factor of *Helicobacter pylori*". *Cancer Res.* 61 (2001), pp. 6356-9.
- [16]- Peek R.M. and Blaser M.J., "*Helicobacter pylori* and gastrointestinal tract Adenocarcinomas ". *Nat Rev Cancer.* 2 (2002), pp. 28-37.
- [17]- Fauchère J.L., "Infections gastriques à *Helicobacter pylori*". *Gastroenterolo. Clin. Biol.* 18 (1994), pp. 212-216.
- [18]- Megraud F., "Méthode diagnostic directe et indirecte d'*Helicobacter pylori*". Centre National de référence des Campylobacters et Helicobacters; Bordeaux (2004).
- [19]- Ricci C., Holton J. and Vaira D., "Diagnosis of *Helicobacter pylori*: Invasive and non-invasive tests. Best". *Pract. Res. Clin. Gastroenterol.* 21(2) (2007), pp. 299-313.
- [20]- Takahashi M., Taguchi H., Yamagushi H., Osaki T. and Kamiya S., "Studies of the effect of *Clostridium butyricum* on *Helicobacter pylori* in several test models including gnotobiotic mice". *J. Med. Microbiol.* 49 (2000), pp. 635-642.
- [21]- Glupezynskiy N., Cantagrel A., Andersen L.P., Alarcon T., Lopez-Bren M. and Megraud F., "Comparaison of the E Test and agar dilution method for antimicrobial susceptibility testing of *Helicobacter pylori*". *Eur. J. Clin. Microbiol. Infect. Dis.* 21 (2002), pp. 549-552.
- [22]- Zhao W., Wu W. and Xu X., "Oral vaccination with liposome-encapsulated recombinant fusion peptide of urease B epitope and cholera toxin B subunit affords prophylactic and therapeutic effects against *H. pylori* infection in BALB/c mice". *Vaccine.* 25 (2007), pp. 7664-7673
- [23]- Lamouliatte H., Mégrand F. and Cayla R., "*Helicobacter pylori* et pathologie Gastroduodénales". Encyclopédie Médico-chirurgicale. Editions techniques. EMC. (1992).

- [24]- Megraud, F., "Méthodes diagnostiques pour *Helicobacter pylori*". *GEFH*. (1992), pp. 15-19.
- [25]- Lozniewski A., "Méthodes diagnostiques de l'infection de *Helicobacter pylori*". *Gastroenterol. Clin. Biol.* 20 (1996), pp. 111-118.
- [26]- Megraud F., "*Helicobacter pylori*. Diagnostic. Comment transporter les biopsies pour un examen bactériologique ?". *Gastrographie*. 19 (1994), pp. 35 – 41.
- [27]- Delchier J.C., "*Helicobacter pylori*. Diagnostic. Que penser de l'uréase test (CLO- test) ?". *Gastrographie*. 19 (1994), pp. 35 – 41.
- [28]- Megraud F., "Méthodes bactériologiques pour le diagnostic de *Campylobacter pylori*". *Gastroenterol. Clin. Biol.* 13 (1989), pp. 31-36.
- [29]-Megraud F., "Comment rechercher *Helicobacter pylori*". *Gastroenterol. Clin. Biol.* 20 (1996), pp. 38-43.
- [30]-Megraud F., Francis G.J. and Labigne., "Rapid urease test in the management of *H. pylori* associated gastritis". *Am. J. Gastrienterol.* 82 (1998), pp. 200-210.
- [31]-Mégraud F., "Méthodes diagnostiques directes et indirectes de *Helicobacter pylori*". *Gastroenterol. Clin. Biol.* 18 (1994), pp. 217-222.
- [32]- Megraud F., "Mise en évidence de *Helicobacter pylori* au niveau de biopsies gastriques par les méthodes bactériologiques". *Revue française des laboratoires.* (1994).
- [33]- Cellini L., Allocati N., Piatteli A., Petrelli L., Fanci P. and Dianelli B., Microbiological evidence of *Helicobacter pylori* from dental plaque in dyspeptic Patients. *Microbiol.* 18 (1995), pp. 187-192.
- [34]- Cellini L., Allocati N., Di Campli E., Masulli M., Di Bortoloneo S. and Dainelli B., "*Helicobacter pylori* isolated from stomach corpus and antrum: comparison of DNA patterns". *J. INFEC.* 32 (1996), pp. 1-3.
- [35]- Minnis J.A., Taylor T.E., Knesek J.E., Peterson W.L. and Mcintire S. A., "Characterization of a 3,5-kbp plasmid from *Helicobacter pylori*". *Plasmid.* 34 (1995), pp. 22-36.
- [36]- Riachi G. and colin R., "*Helicobacter pylori*. Méthodes de recherche". *Les dossiers du praticien. Impact. Med.* 303 (1995), pp. 1-22.
- [37]- Franzin L., Pennazio M., Cabadi D., Paolo Rossini F. and Gioannini P., "Clarithromycin and amoxicillin susceptibility of *Helicobacter pylori* strains isolated from adult patients with gastric or duodenal ulcer in Italy". *Curr. Microbiol.* 40 (2000), pp. 96 – 100.
- [38]- Bruley Des Varannes S., "*Helicobacter.pylori*. Les méthodes diagnostiques". *Quot. Med.* 321 (1996), pp. 14-48.
- [39]- Dorval E. D., "Gastrite chronique et ulcère duodéal: des maladies infectieuses? " *Rev. Prat. Med. Gén.* 162 (1992), pp. 82-83.
- [40]- Fagniez P. L., "*Helicobacter* bactérie de la décennie". *Rev. Prat. Med.* 313 (1995), pp. 10-14.
- [41]- Megraud F., "Ulcère et gastrite : un nouveau domaine d'exploitation au laboratoire de biologie". *Feuillets de biologie.* 15 (1989), pp. 25-28.
- [42]- Megraud F., "*Helicobacter pylori*. Historique et épidémiologie". *Quot. Med.* (1996), pp. 4 –83.
- [43]-Bretagne J.F., *Helicobacter pylori*. De la bactérie à la pathologie gastroduodénale . *Quot. Med.* (1996), pp.9-13.
- [44]- Sobhani I., Flourié B., Lavergne A., Colimon R., Mignon M., Modigliani R. and Rambaud J.C., "*Helicobacter pylori* et pathologie gastroduodénale. Première partie : physiopathologie et méthodes de diagnostic". *Gastroenterol. Clin. Biol.* 15 (1991), pp. 405-411.
- [45]- Souquet J.C., "Diagnostic de l'infection de *Helicobacter pylori*". *Hepato. Gastro.* 2 (1995), pp. 17-22.
- [46]- Yousfi M.M., eddy R., Osato M.S. and Graham D., "Antre ou corps gastrique : quel est le meilleur site biopsique pour mettre en culture *Helicobacter pylori*". *Helicobacter.* 1 (1997), pp. 25-26.
- [47]- Altman C.W., Lordouch A., Briantais M.J., Martin E., Jaques L. and Buffet C., Gastrite antrale au cours de l'alcoolisme chronique: rôle de la cirrhose et d'*Helicobacter pylori*. *Pres. Med.* 24 (1995), pp. 708-710.
- [48]- Megraud F., "Méthodes de diagnostic de l'infection à *Helicobacter pylori*. Le point de vue du biologiste". *Lettre. Infec.* 5 (1996), pp. 17-22.

- [49]- Megraud F., “Difficultés pratiques de l'éradication d'*Helicobacter pylori* dans la maladie ulcéreuse”. *Quot. Med.* (1994), pp. 9 – 13.
- [50]- Cayla R., “ Comment éradiquer *Helicobacter pylori* ?” *Gastroenterol. Clin. Biol.* 20 (1996), pp. 119-130.
- [51]- Frasier B. A., Russel D.G., Flak P., Olsen A.N., Hammar M., Westbon T.U., Frazin L. *et al.*, “Susceptibility of *Helicobacter pylori* to 20 antimicrobiol agence”. *Bio future.* 55 (2000), pp. 10-11.
- [52]- Tankovic J., Lamarque D., Lascols C., Soussy C.J. and Delchier J.C., “Impact of *Helicobacter pylori* resistance to clarithromycin on the efficacy of the omeprazole-amoxicillin-clarithromycin therapy”. *Aliment. Pharmacol. Ther.* 15 (2001), pp. 707-13.