INTRODUCTION

Soybean is a crop of economic importance, being the most planted oilseed in the world. Brazil is the second largest producer and exporter of soybean in the world, with a planted area of approximately 26.4 to 27.3 million hectares and a production of about 80,08 to 82,99 million tons in 2012/2013 (CONAB, 2013).

The soybean cysts nematode (SCN), *Heterodera glycines*, is one of the most severe plant disease problems of this crop, considered as the most destructive parasite of soybean plants (NOEL, 1992).

In order to control these nematodes the primary common methods used are chemical control (nematicides), the use of resistant varieties and crop rotation. Chemical control by means of nematicide application has proven expensive as well as detrimental to the environment, to the human being, to the wild fauna and to the beneficial organism within the soil. The use of resistant varieties is a natural approach, highly recommended to control plant pests and diseases. However, in the case of this specific nematode, the availability of resistant varieties for the farmer is scarce, due to the great number of races or HG types of SCN occurring in Brazil (FERRAZ; FREITAS, 2004).

In tropical and subtropical countries nematodes find ideal humidity and temperature conditions for reproduction and feeding. Thus, worsening the efficiency of control of these pathogens, that are very difficult to eradicate once they are established within an area (TORRES et al., 2008).

Biological control is an alternative method to control plant parasites, as nematodes. Biological control of nematodes can be achieved by the use of appropriate fungi or bacteria. Predators and egg parasites are certainly the most studied organism and the best fitted as nematode’s biological control agents (JATALA, 1986; NORDBRING-HERTZ et al., 2002).

The first observation of *Pasteuria* in a plant parasite nematode, *Pratylenchus pratensis* de Man, was made by Thorne (1940) whom classified the organism as a Protozoa and named it as *Dubosquia penetrans*. However, studies performed by Mankau (1975), re-classified the organism as a prokaryote of the Genus *Bacillus* Cohn. The re-discovery of *Pasteuria ramose* by Sayre et al. (1977) and its morphological similarities with *Bacillus penetrans* Mankau suggested that these two bacteria had a common generic relation. In 1985, Sayre and Starr named the organism that parasite *Meloidogyne incognita* as *Pasteuria penetrans* (Thorne) Sayre & Starr, as well as *Pasteuria nishizawae* is specific to...
the soybean cyst nematode (*Heterodera glycines*), *Pasteuria thornei* Sayre & Starr is specific to the nematode of the root lesions according to Stirling & Wachtel (1980), while *Pasteuria usgae* is specific to the nematode *Belonolaimus longicaudatus* Rau according to Rau Giblin- Davis et al. (2003).

*Pasteuria* is a gram-positive bacterium that forms an endospore. These bacteria were found as parasites of many economically important nematodes (SAYRE; STARR 1988). The Genus *Pasteuria* spp. was structured from 323 nematode species belonging to 116 Genera, including plant parasitic nematodes, entomopathogenic nematodes, predator nematodes and free living nematodes (CHEN; DICKSON 1998).

The analysis of the 16S rRNA gene sequences is a sensible and well-established tool for the detection and phylogenetic analysis of bacteria (STAHL, 1997). The use of sequences of the gene 16S rRNA may lead to identification of species of *Pasteuria* and to evaluation of the diversity of these microorganisms within a population of nematodes’ samples, thus being a tool of a great utility for the comprehension of the parasite-nematode interactions and the ecology of *Pasteuria* (EBERT et al., 1996; ANDERSON et al., 1999; ATIBALENTJA et al., 2000; BEKAL et al., 2001).

All *Pasteuria* species known are obligate parasites. Until recently the Genus had four species *Pasteuria nishizawai*, *P. penetrans*, *P. ramosa* and *P. thornei* (OOSTENDORP et al., 1990; CHEN; Dickson, 1998).

For many years attempts to obtain an axenic culture of *Pasteuria* spp. resulted in failure (WILLIAMS et al., 1989; BISHOP; ELLAR, 1991). However, an advance towards the success to obtain an *in vitro* culture of *P. penetrans* was announced by Hewlett et al. (2002).

A life cycle study of the bacterium that parasities the nematode *Heterodera glycines* was performed by means of germination of the endospore that infects J2’s for production of a next generation endospores in adult females and cysts. Descriptions were based in microscopic examination of successive juvenile stages of *H. glycines* excised from soybean roots, unlike *P. nishizawai*, that were bused exclusively on the examination of diseased cysts (SAYRE et al., 1991).

According to Atibalenja et al. (2004), bacterium development, germination of the endospore and penetration of the germ tube inside the nematode begin soon after penetration of the J2’s in the radicular system. Otherwise, the endospore does not germinate and consequently no infection by *Pasteuria* will arise. For this reason, observations based only on diseased cysts will result in an incomplete analysis of *Pasteuria*’s life cycle, which would explain why germination of *P. nishizawai* is not observed (SAYRE et al., 1991).

After the endospore’s germination, primary cauliflower-like colonies are formed within the J3. Development of the bacterium was observed only in female adults but not in males. Later, the bacterial sporulation can be observed with the development of a structure similar to a chunk of grapes, which can be observed also in immature females of fourth stage and cysts. Finally, the presence of mature sporangia and endospores can be observed, varying in number (30.000 to 820.000 with mean value and standard deviation of 314.00 and 234.000, respectively) as a function of the size of the cyst or female nematode (ATIBALENTJA et. al., 2004).

Until now, *P. penetrans* is the most studied species due to its potential as a biological control agent. However, technical hitches for mass production *in vitro* had made its commercial production problematic. In order to introduce bacteria in the soil environment for field experiments, dry roots’ powder infected by the bacteria was used, as proposed by Stirling and Wachtel (1980).

Thus, the objective of the present work was to verify the natural occurrence of *Pasteuria nishizawai* Sayre in soils under soybean cultivation.

**MATERIAL AND METHODS**

Soil samples with cysts coming from different established soybean growing areas from Brazil (Table 1), were processed by the centrifugal flotation in sucrose solution technique (Jenkins, 1964).

Part of those ten processed soil samples were added, individually, to ceramic pots and preserved under greenhouse at Syngenta’s experimental station in Uberlândia, MG – Brazil. Those pots were planted with the susceptible soybean cultivar ‘Lee 74E’, in order to promote multiplication of *Heterodera glycines*, as shown in Figure 1.
Table 1. Place of origin of the samples analyzed in the essay for natural occurrence of *Pasteuria nishizawai* in Brazilian soils.

| Identification | County               | State |
|---------------|----------------------|-------|
| H.g 1         | Jataí                | GO    |
| H.g 2         | Sorriso              | MT    |
| H.g 3         | Campo Alegre         | GO    |
| H.g 4         | Lucas do Rio Verde   | MT    |
| H.g 5         | Luís Eduardo Magalhães | BA   |
| H.g 6         | Campo Alegre         | GO    |
| H.g 7         | Nova Mutum           | MT    |
| H.g 8         | Campo Alegre         | GO    |
| H.g 9         | Luís Eduardo Magalhães | BA   |
| H.g 10        | Londrina             | PR    |

Figure 1. Populations of *Heterodera glycines* originated from field soil samples, being multiplied and preserved under greenhouse conditions.

A soil aliquot of 150 cm$^3$ was obtained 28 days after sowing and then processed by the centrifugal flotation in sucrose solution technique, according Jenkins (1964).

From the obtained suspension solution an aliquot of 1 mL was observed in Peters chamber under an inverted light microscope, verifying the presence or absence of endospores of the bacterium attached to the nematode’s cuticle (Figure 2). Subsequently, the percentage of nematodes showing bacteria attached to its body in relation to the total number of nematodes observed in the suspension was calculated.

Figure 2. Counting of nematodes in suspension solution with the aid of a Peters’ chamber under an inverted light microscope to observe bacterial endospores.
RESULTS AND DISCUSSION

Within all ten samples the frequency of occurrence of the bacterium *Pasteuria nishizawae* was of 100%, that is to say, there were observed bacterial endospores attached to the cuticle of all the nematodes *Heterodera glycines* extracted from the soil 28 days after soybean sowing (Table 2), as showed in Figure 3.

Table 2. Natural occurrence and frequency of *Pasteuria nishizawae* in samples originated from different districts in Brazil.

| Sample identification | Total number of *H. glycines* 2nd stage juveniles recovered | Frequency of attached endospores of *P. nishizawae* (%) |
|-----------------------|------------------------------------------------------------|--------------------------------------------------------|
| H.g 1                 | 720                                                        | 100                                                    |
| H.g 2                 | 294                                                        | 100                                                    |
| H.g 3                 | 294                                                        | 100                                                    |
| H.g 4                 | 396                                                        | 100                                                    |
| H.g 5                 | 156                                                        | 100                                                    |
| H.g 6                 | 76                                                         | 100                                                    |
| H.g 7                 | 442                                                        | 100                                                    |
| H.g 8                 | 105                                                        | 100                                                    |
| H.g 9                 | 90                                                         | 100                                                    |
| H.g 10                | 116                                                        | 100                                                    |

Figure 3. Observation of juvenile nematodes with *Pasteuria nishizawae* endospores attached to their cuticle.

In Brazil, studies regarding *Pasteuria* spp. are still developing, especially when concerning to its employment to control species of *Pratylenchus* and *Heterodera*. The vast majority research about the subject is developed in the country focusing the root-knot nematode (*Meloidogyne* spp.).

According to Tzortzakakis and Gowen (1994), a great number of endospores attached to the J2 nematodes, does not ensure reduction in the number of eggs produced, if a high variance of attachment and infectivity of endospores occurs. These remain viable in the soil by many years, are resistant to desiccation (STIRLING, 1984) and relatively resistant to high temperatures.

Lordello (1966) reported for the first time in Brazil, the occurrence of *Pasteuria* spp. in females of *Meloidogyne javanica*, infecting plants of tomato coming from Vargina in Minas Gerais.

Santos (1981), observed the bacterium in juveniles of the second stage of *M. javanica* extracted from soil collected from the rhizosphere of bean plants intensively infected by the nematode in Petrolina, Pernambuco. Subsequently, several works were performed on the parasitism of *Pasteuria* spp. in nematodes, verifying in the average, activity of the bacterium over *Meloidogyne* species (PIMENTA; CARNEIRO, 2005).

According to Souza and Campos (1996), *Pasteuria* spp. was observed in 29,69% of the samples collected in 128 locations and 28 vegetal species in Minas Gerais, similar results were also verified in other countries: South Africa (Spaull, 1984), Australia (STIRLING; WHITE, 1982; BIRD;
Natural occurrence of *Pasteuria nishizawae*… VICENTE, C. B.; SANTOS, M. A.

BRISBANE, 1988), USA (WALTER; KAPLAN, 1990; HEWLETT et al., 1994) and Spain (VERDEJO; LUCAS, 1992). These authors observed infection percentages of 14.61%, 8.66%, 7.30% and 4.21% respectively, on populations of *Helicotylenchus dihystera* (Cobb) Sher, *Meloidogyne javanica* Goeldi, *Pratylenchus brachyurus* Godfrey and *Tylenchus* sp. Bastian extracted from the soil.

There is no record of the occurrence of *Pasteuria* spp. in regions with annual mean temperature under 10ºC, with the higher occurrence being observed under annual mean temperatures above 21ºC (CHEN; DICKSON, 1998).

**CONCLUSIONS**

*Pasteuria nishizawae* has natural occurrence in Brazilian soils under soybean cultivation, specifically in the States of Bahia, Goiás, Mato Grosso and Paraná.

The frequency of *Pasteuria nishizawae* occurrence within the analyzed soils was of 100%.

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**RESUMO:** A soja (*Glycine max* (L.) Merrill) é uma cultura de grande importância econômica. O Brasil é o segundo maior produtor e exportador mundial. Um dos mais sérios problemas fitossanitários desta cultura é o nematoide do cisto, *Heterodera glycines* Ichinohe, considerado o parasito mais destrutivo. A primeira ocorrência desta doença no Brasil foi relatada na safra 1991/1992. O controle de nematoides é mais difícil quando comparado com outras doenças. Torna-se cada vez mais importante a busca de controle alternativo, que não ofereça risco ao meio ambiente e nem ao aplicador. Sendo assim, a bactéria *Pasteuria* spp. Metchnikoff com alta especificidade, representa um promissor agente de controle biológico dos nematoides. O controle biológico do nematoide do cisto da soja pela bactéria *Pasteuria nishizawae* Sayre vem sendo estudado por diversos pesquisadores. O objetivo deste trabalho foi determinar a ocorrência natural de *Pasteuria nishizawae* em solos brasileiros. O ensaio foi conduzido em casa-de-vegetação na Unidade de Pesquisa da Syngenta em Uberlândia – MG, com amostras de solo provenientes de áreas de cultivo de soja dos estados brasileiros Bahia, Goiás, Mato Grosso e Paraná. Foram retiradas uma alíquota de 150 cm$^3$ de solo e então a mesma foi processada pela técnica da flotação centrífuga em solução de sacarose. Da suspensão obtida foi observada uma alíquota de 1 mL em câmara de Peters, com o auxílio do microscópio invertido, verificando a presença ou ausência de endósporos da bactéria aderidos na cutícula dos nematoides extraídos. A frequência de ocorrência da *Pasteuria nishizawae* foi de 100% nas amostras analisadas.

**PALAVRAS-CHAVE:** Bactéria. Controle biológico, Nematoide de cisto, *Glycine max*. 

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