



**GV-SOLAS**

Gesellschaft für Versuchstierkunde  
Society for Laboratory Animal Science

# **Expert Information**

**from the Working Group on Hygiene**

**Implication of infectious agents on  
results of animal experiments**

***Spironucleus muris***

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## Contents

Background .....	3
Prevalence .....	3
Host species.....	3
Properties.....	3
Susceptibility .....	3
Organotropism.....	3
Clinical disease .....	3
Pathology .....	4
Morbidity and mortality .....	4
Zoonotic potential .....	4
Interference with research .....	4
<i>Oncology</i> .....	4
<i>Teratology</i> .....	4
<i>Infectiology / Interactions with other infectious agents</i> .....	4
<i>Immunology</i> .....	5
<i>Toxicology</i> .....	5
<i>Physiology</i> .....	5
<i>Cell biology</i> .....	5
<i>Assisted reproductive technology</i> .....	5
Special considerations.....	5
References.....	6

## ***Spironucleus muris***

### **Background**

- *Hexamita muris* is used as synonym for *Spironucleus (S.) muris*.

### **Prevalence**

- Rare infections in Europe mice
- Present in 7.14% of laboratory mice in Brasil.<sup>1</sup>

### **Host species**

- Mouse, rat<sup>2,3,4</sup>, European hamster<sup>5,6</sup>, Syrian hamster<sup>7,8,9</sup>, *Mastomys coucha*<sup>10</sup>, Macaques (*Macaca mulatta*)<sup>11</sup>
- There is evidence for a certain degree of host specificity: a mouse isolate can obviously infect golden hamster and vice versa. A rat isolate, however, can infect only another rat.<sup>8,12</sup>

### **Properties**

- Transmission is oro-fecal with cysts.
- Minimum infective dose is one cyst.<sup>13</sup>
- Dirty bedding transmission to mice with non-functional T and B cells seems to be insufficient.<sup>14</sup>
- The cysts resist to high and low temperatures, low pH, most usual disinfectants, high osmotic pressure and centrifugation.<sup>15</sup>

### **Susceptibility**

- Inbred mouse strains differ in susceptibility.<sup>5,6</sup>
- Classified as opportunistic pathogen, trophozoites can damage the microvilli and penetrate into the epithelium.<sup>16</sup>
- Previously infected mice may show resistance to reinfection after recovery.<sup>17</sup>
- It is suggested that the major histocompatibility complex haplotype may influence susceptibility to *S. muris*.<sup>18</sup>

### **Organotropism**

- Intestine (trophozoites, i.e., active stage of the parasite); in the caecum and colon there are mainly cysts.<sup>19</sup>

### **Clinical disease**

- Enlarged abdominal cavity (due to chronic enteritis), sometimes meteorism, diarrhoea and retarded growth in younger animals<sup>19</sup>
- Roughened hair coat, hunched position, sticky stool<sup>5</sup>
- Enhanced mortality, shortened life span in athymic mice<sup>20,21</sup>
- In athymic mice, severe enteritis and weight loss, sporadic deaths by 4 weeks of age<sup>22</sup>
- Liquid, yellow / green and often foamy contents of the small and large intestines<sup>22</sup>

- Some additional weakening/stressing factor(s), for instance athymic status, are necessary to elicit clinical disease.<sup>20,23</sup>
- Shortened life span in athymic mice<sup>18,19</sup>

### **Pathology**

- Enteritis, sometimes subepithelial edema, mononuclear inflammatory infiltrations in the submucosa, desquamation of the epithelia, proliferation and thickening of the intestinal wall<sup>7</sup>
- Accumulation of catarrhal fluid in the small intestine, sometimes hyperplasia of the epithelium<sup>5</sup>
- Damages of microvilli, reduction of their height, increase in crypt depth<sup>17</sup>
- Marked crypt hyperplasia, occasional crypt abscesses and variable degree of villous atrophy<sup>22</sup>
- Degeneration of enterocytes and necrosis; in such areas, penetration of the intestinal barrier by individual trophozoites, exceptionally: invasion of plasma cells<sup>8</sup>

### **Morbidity and mortality**

- Young animals are more sensitive<sup>3</sup>, in older non-immunocompromised animals a spontaneous recovery from the infection occurs.<sup>24</sup>
- Increased mortality in cadmium exposed mice<sup>25</sup>
- Increased sensitivity to X-irradiation<sup>26</sup>

### **Zoonotic potential**

- No data

### **Interference with research**

#### **Oncology**

- Nonspecific activation of macrophages and, hence, enhanced elimination of tumour cells<sup>27</sup>

#### **Teratology**

- No data

### **Infectiology / Interactions with other infectious agents**

- Sometimes enhanced or impaired resistance to experimental infection with other agents<sup>10</sup>
- Concomitant infections with *Babesia microti*, *Plasmodium berghei* and *Plasmodium yoelii* decrease the output of trophozoites and cysts of *Spiroucleus muris*.<sup>28</sup>
- There is a temporary decrease of flagellate cyst output coincident with the peak of the blood parasite infections, followed by a rapid return to normal levels in mice infected with the intestinal flagellates *Giardia muris* or *Spiroucleus muris*, together with the blood parasites *Babesia microti* or *Plasmodium yoelii*. This decrease in cyst output is correlated with decreased numbers of trophozoites in the small intestine. The effect on *Spiroucleus muris* is more marked than that on *Giardia muris*.<sup>28</sup>
- Enhanced resistance to experimental infection with *Listeria monocytogenes*<sup>29,30</sup>

### ***Immunology***

- Infected mice are unsuitable for immunologic studies.<sup>24</sup>
- Sometimes weakened immune response to some agents<sup>29</sup>, depression to mount an immune response to a thymus dependent antigen<sup>30</sup>
- Decreased pneumococci antigen in infected mice but not in infected rats<sup>31</sup>

### ***Toxicology***

- No data

### ***Physiology***

- Microscopically, acute disease is associated with distension of crypts and intervillous spaces by pear-shaped trophozoites and inflammatory edema of the lamina propria.<sup>32</sup>
- During overt infestation, organisms are seen extracellularly in crypts and intervillous spaces associated with blunting of intestinal villi, epithelial degeneration and mucin depletion, reactive epithelial hyperplasia, edema and leukocyte infiltration.<sup>33</sup>

### ***Cell biology***

- Impairment of the RNA-synthesis and of enzyme synthesis of macrophages<sup>34</sup>

### ***Assisted reproductive technology***

- No data

### ***Special considerations***

- No data

Ivo Kunstyr, actualized by Brunhilde Illgen-Wilcke, Liestal Suisse, September 2022

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