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Predilection Site for the Meat Lover *Trichinella* spp Larvae and Its Pathogenesis and Potency

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ABSTRACT

Aims: To revisited predilection site of *Trichinella* spp larvae including their pathogenesis and its potency

Discussion: The nematode *Trichinella* spp causes serious zoonosis called trichinellosis, a disease affecting muscles which consider as one of tropical disease. Even though its natural host varied, but infection among popular live stocks, such as pigs and other animals, which are raising community medicine concern. Human infection occurs after consumption of raw or undercooked meat or meat products contain muscle larvae of *T. spiralis*. The tropism of the parasite for individual muscles and/or muscle groups varies significantly. *Trichinella* spp. has a direct life cycle where all three life cycle stages (the infective muscle larvae, adult, and new born larvae) happen serially in one host only. Intestine-dwelling adults of *Trichinella* produce newborn larvae that bypass the enterocyte, enter the bloodstream and colonize skeletal muscle. The muscle larvae assemble excretory-secretory products which play crucial role in establishing and maintaining persistent parasitism and the host's immune modulation and evasion. It turns out that excretory-secretory products from muscle larvae and mature worm also have hidden medical potential that can be used to treat allergic problems, inflammation-based diseases, autoimmunity and even malignancy.

Conclusion: Trichinellosis is a serious and potentially fatal zoonosis which transmitted through consuming raw or uncooked contaminated meat or its comestibles. Its primary tropism is to the host's striated muscle and infection can persist for a long time facilitated by several reciprocities of its product (e.g., excretory-secretory) with the host's cell and immune system. Fortunately, there are several promising potency in the field of therapeutic and prevention medicine which should be explored intensely.

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Keywords: tropical disease, nematode, tropism, excretory, secretory, muscle larvae, carcasses, trichinosis/trichinelosis

1. INTRODUCTION

The term tropical diseases encompass all diseases, communicable and no communicable, that occur principally in the tropical countries or tropics, areas that lie between, and alongside, the Tropic of Cancer and Tropic of Capricorn belts [1]. Among those communicable group of disease, neglected helminth infections including trichinellosis or trichinosis, which is still the major health problem [2]. There are eleven known species within the genus *Trichinella*. These eleven species subdivide into those that invade host muscle cells and encapsulate (surrounded by a collagen capsule) and those that do not encapsulate [3,4]. *Trichinella Spiralis*, the most common species in this genus, belongs to the

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28 encapsulated group and causes most human infections and deaths from trichinosis [3]. This
29 genus specifically causing a disease which affecting the host's muscles [1-5].
30 Even though its natural host varied, but infection among popular live stocks such as pigs [5-
31 8] and other animals, e.g., horses [9], wild game meat (meat from an animal that is typically
32 found in the wild and not raised domestically on a farm for mass consumption; usually free-
33 roaming foragers and hunted for their meat) [10], rats [11], wild birds [12], wild and farmed
34 reptiles [13] etc., which are raising public health concern [14], even though its global burden
35 is much lower than that of other foodborne parasitic diseases (a mean estimated 76 healthy
36 life years lost per billion people per year for human trichinellosis, globally [15].
37 The aim of this study is to revisited predilection site of *Trichinella* spp larvae in its host
38 including their pathogenesis and potency along with its comestible implication and effort
39 conducted to prevent transmission.

40

41 **2. LIFE CYCLE AND PATHOGENESIS**

42 Among many member of helminths which affect human, *Trichinella* spp. are distinctive
43 because it has a direct life cycle [16]; which means that all three life cycle stages of the
44 parasite, namely infective muscle larvae, adult, and new born larvae; Intestine-dwelling
45 adults of *Trichinella* produce newborn larvae that enter the bloodstream and colonize
46 skeletal muscle [17]. Infection is acquired by consumption of infected raw or undercooked
47 meat or meat based comestibles [3-17].

48 Under the biochemically pressure of low pH gastric juice, entrapped larvae which is basically
49 anaerobic are released in the host's stomach, followed by the molting process
50 (approximately four times in 30-40-time span) [19]. Proteases secreted by *Trichinella spiralis*
51 intestinal infective larvae directly damage the surrounding junctions of the intestinal epithelial
52 cell monolayer and also arbitrate larval invasion and develop into the adult stage inside the
53 enterocytes of small intestine [20]. The results of study conducted by Song et al [20]
54 stipulate that the parasite enzyme named serine proteases and cysteine proteases play
55 crucial roles in larvae invasion, growth and survival inside the host and that they may be
56 main candidate target molecules for vaccines against larval invasion and development.

57 After successfully entering enterocyte and become mature, male and female are mating then
58 produce new born larvae [21] are released into circulation and spread throughout the tissues
59 and organs [22] and only those that enter striated muscles mature into muscle larvae [23].
60 During the muscular phase, the larvae invade the skeletal muscle fibers inducing a relevant
61 inflammatory reaction aiming for the elimination of the parasite. However, the larvae
62 eventually succeed to build their own home inside the infected myocytes [23]. Muscle
63 invasion results in formation of a capsule surrounding muscle larvae in the region of infected
64 muscles [24]. Once again, this eccentric meat lover *Trichinella* blessed with the capability to
65 make itself "feel homey like being at home" by way of transforming the infected muscle cell
66 for their own benefit and accomplishing a new type of cell inside the host affected
67 musculature, the so-called nurse cell [25].

68 The lowest infectious dose of *Trichinella* larvae is remains unrevealed, but the clinical
69 manifestations of trichinellosis starts to displayed as the number of parasite entering the host
70 increases [26]. Asymptomatic infection could remain silent in human if it is only involving a
71 minimum amount of larvae; gastrointestinal symptoms manifested as a specific syndrome
72 consist of nausea, diarrhea, vomiting, fatigue, fever, and abdominal discomfort [27], starts
73 very early to develop in case of unintentionally ingestion of hundreds of larvae, perhaps
74 manifest itself clinically within the first 48 hours after consuming contaminated meat. The
75 condition that followed by development of a series of condition which are serious, but
76 scarcely fatal illness [5]. Clinical signs of the disease usually last 4–6 months, rarely longer
77 (up to 2 years).

78 Chronic form of trichinellosis rarely reported, once in 1983 revealed by two German doctor in
79 their case report regarding biopsies conducted on muscles of five patients with clinical
80 diagnosis chronic neuromuscular disorder, mostly manifested as spinal muscle atrophy. All

81 of them had previous history of acute trichinellosis, the interval between acute parasitic
82 infection and the appearance of the slowly progressive neuromuscular syndrome being of 13
83 to 26 years respectively. Analysis conducted on the biopsy specimens showed
84 morphological and enzyme-histochemical alteration which indicative the presence of
85 progressive neurogenic muscular atrophy. From the Parasite perspective, distinctive
86 encapsulated but still living, enzyme-positive parasites were clearly identified with definite
87 signs of focal myositis in the muscle portion surrounding the larva. The possibility
88 pathogenesis correlations between the "chronic" trichinellosis and the "degenerative"
89 neuromuscular disorder cannot yet be excluded and this still remains to be an uncharted
90 sea of exploration.

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3. THE POTENCY

93 Infective larvae remain alive in striated muscles of the vulnerable host for years [20]; an
94 evidence supported by the study of Sofronic-Milosavljevic et al which revealed the chronic
95 existence of specific antibody responses that still could be recognized even 30 years post
96 primary infection [29]. In case of invasion by *Trichinella* larvae against the host's immune
97 system, it actually arouses a complex immune response; in human host is better designated
98 by humoral immune response [30] rather than the cellular responses; and this emphasize
99 future prospect of the human host's dynamic humoral response [30] for diagnostic [29] or
100 even vaccine development purposes [31,32] such as reported by Bi et al [32] that revealed
101 the newly identified rTs-ES-1 is potent immunodominant protein secreted by *Trichinella*
102 stichocytes during natural infection and permits the arousal of fractional protective immunity
103 in vaccinated mice inimical to intentional *Trichinella* infection. Therefore, findings of this rTs-
104 ES-1 specific protein with the better understanding of its antigenic shift-dynamicity [31] is a
105 potential candidate for vaccine development against trichinellosis. In contrast to what
106 happened inside their vulnerable human host, in animals *T. spiralis* can outstretch a high
107 worm burden without causing prominent clinical symptoms [33].

108 The initiation of infection depends on first by the annexation of prone intestinal epithelium by
109 infective muscle larvae (ML) and followed secondly with the preservation of parasitism which
110 is marked by the presence of ML in affected muscle cells. The parasite regulatory protein
111 accountable for enzymatic process of these two steps are very important for future
112 investigation.

113 Excretory-secretory products of invading larvae believed to be originate from stichocyte
114 granules in the stichosome, the secretory organelle of the *Trichinella*'s mature muscle larvae
115 [35]. These excretory-secretory products play a pivotal role in parasite's immune evasion
116 and regulation inimical to the host's innate immune system by way of (1) suppressing NET
117 (neutrophil extracellular traps which primary function as a trap for pathogens and facilitating
118 phagocytosis and cytokine production) production and (2) negatively didacte cytokine
119 secretion. The understanding of this excretory-secretory products function for the larvae or
120 worm provides an encouraging area for manufacturing new intervention strategies in other
121 areas of medicine, e.g., in tackling sepsis induced acute lung injury [36] or allergic plethora
122 [29] or autoimmune condition/diseases such as colitis [37] and even malignancies [29].

123 These important excretory-secretory products engage mainly in the reciprocity with various
124 host cells: firstly, the immune cells, secondly the enterocytes and thirdly the muscle cells,
125 and, through those interaction establishing their role in parasitism and immune response
126 induction, modulation and even evasion [29-32,34-37]. Through these approaches, this
127 nematode generates a perfect milieu for its own suitability and survival in two ways either by
128 modulation of host immune response or affecting host cell gene expression. Extensive
129 exploration of these molecules is important in order to build better understanding regarding
130 (1) the establishment of triumphant parasitism, (2) the development of novel therapies and
131 (3) preventive treatments for inflammatory based disorder.

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134 **4. CONCLUSION**

135 Trichinellosis with its related clinical syndrome must always be considered as serious and
136 potentially fatal zoonosis. Transmission occurs through consuming raw or uncooked meat or
137 its comestibles which contaminated with its muscle larvae. Its primary tropism is to the host's
138 striated muscle and can affect the muscle strength and composition in long term. Infection
139 can persist for a long time facilitated by several reciprocities of its product (e.g., excretory-
140 secretory) with the host's organ specific cells (e.g., enterocytes, myocytes) and immune
141 system. Fortunately, there are several promising potency in the field of therapeutic and
142 prevention medicine which should be explored intensely.

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147

148 None to declare

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151 **COMPETING INTERESTS**

152

153 None to declare

154

155 **AUTHORS' CONTRIBUTIONS**

156

157 The sole author designed, analyzed, interpreted and prepared the manuscript

158

159 **CONSENT (WHERE EVER APPLICABLE)**

160

161 Not needed

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163 **ETHICAL APPROVAL (WHERE EVER APPLICABLE)**

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165 Not needed

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