# A Note On Survival Of Infective Stage Larva Of Gnathostoma Spp. In BME Culture Medium

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#### **Abstract**

Dear editor, human gnathostomiasis in an important tropical parasitic infection (1). The major manifestation as nodular migratory eosinophilic panniculitis is suggested for this disease. More severe presentations, resulting from visceral larva migran, such as ocular and CNS gnathostomiasis are also reported. The diagnosis is confirmed by identification of the parasite from the pathological specimen. However, it is difficult to get the histological diagnosis, therefore, the immunological diagnosis for this infection is necessary.

At present, Gnathostoma spinigerum third stage larvae (L3) antigen is necessary for Western blot analysis in the diagnosis of Gnathostomiasis (2,3). Acid pepsin solution is required for digestion of eel's liver (Fluta alba) to yield the larvae for antigen preparation. However, the specific antigen is excretion secretory (ES) antigen, which can be derived from viable larva. Since the larva start to degenerate within a short period, therefore, the larva derived from digestion must be used before its death. Here, we reported our result in using the BME culture medium for cultivation of the derived larva to prolong the period of usage of the harvested larva in antigen preparation.

We used 48 L3 harvesting from standard acid digestion of

eel's viscera as described in our recent report ( $_4$ ) for this experiment. These larva were culture in BME culture medium (Gibco, Grand Island, N.Y., U.S.A.): preparation of this medium was done by using one pack of BME powder (9.2 gm) dissolved in 1 litre of distilled water and adjusted to pH 7.2-7.4 by NaHCO3. Sterilization was done by 0.45  $\mu$ m millipore membrane filtration. Of interest, after a week of cultivation, we could detect 41 survived L3 giving the survival rate of 85.4 %. Therefore, using the BME culture media for cultivated of the derived L3 from pepsin digestion can help prolong the period in using the L3 for antigen preparation.

#### References

- 1. Daengsvang S: Gnathostomiasis in Southeast Asia. Southeast Asian J Trop Med Public Health. 1981; 12: 319-32.
- 2. Tapchaisri P, Nopparatana C, Chaicumpa W, Setasuban P, : Specific antigen of Gnathostoma spinigerum for immunodiagnosis of human gnathostomiasis. Int J Parasitol. 1991; 21: 315-9.
- 3. Saksirisampant W, Chawengkiattikul R, Kraivichain K, Nuchprayoon S. Specific IgE antibody responses to somatic and excretory-secretory antigens of third stage G. spinigerum larvae in human gnathostomiasis. J Med Assoc Thai. 2001;84 Suppl 1:S173-81.
- 4. Sugaroon S, Wiwanitkit V. Gnathostoma infective stage larvae in swamp eels (Fluta alba) at a metropolitan market in Bangkok, Thailand. Ann Clin Lab Sci. 2003;33:94-6.

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