Effect of Temperature on the Growth of *Mycoplasma gallisepticum* and on Its Inhibition by Antiserum

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In the adult chicken, *Mycoplasma gallisepticum* does not extend readily beyond the nasal cavity, although lesions are caused even by small numbers of organisms following their inoculation directly into the abdominal air sacs (Grumbles, Boney & Delaplane, 1952; Nelson, 1953; McMartin & Adler, 1961). This may be explained by the difference in temperature of several degrees between the nasal cavity and abdominal air sacs, the latter being slightly below the body temperature of 40·7 °C (Herin, Booth & Johnson, 1960), as Gill (1962) reported that growth was inhibited at 42 °C. To investigate the influence of temperature on the organism, studies were made on the rate of growth, inhibition by antiserum and the role of heat labile components of normal serum on this inhibition, at incubation temperatures of 37 and 40·5 °C, using methods previously described (Woode & McMartin, 1973).

The critical temperature for growth of *M. gallisepticum* was 41·5 °C, determined on four separate occasions by incubating replicate cultures at 41, 41·5 and 42 °C. Growth occurred at 41 °C, at 41·5 °C on two of four occasions, and was inhibited at 42 °C.

The metabolic inhibition (m.i.) test was performed in tubes held in water baths at 37 and 40·5 °C. Growth at 40·5 °C was delayed by 48 h as compared with cultures at 37 °C, but the early m.i. titre was the same at both temperatures. As the higher temperature had not influenced the inhibitory effect of high dilutions of antiserum, the effect on loss of viability by low dilution of antiserum was studied. Replicate cultures were incubated in 10 % (v/v) horse serum (HS) broth or 10 % (v/v) freshly prepared chicken serum (CS) broth, in the presence of a 1:20 dilution of antiserum. Control HS broth cultures without antiserum were included (Fig. 1a). Growth at 40·5 °C was appreciably less than at 37 °C, and the rate of loss of viability by antiserum was greater with CS in the broth and at 40·5 °C.

We did not determine whether lysis occurred during inactivation of the organism by low dilutions of antiserum, as at the concentration used visualization of the organism was not possible. However, lysis appears unlikely as the main mechanism of loss of viability when the length of time required for complete loss is considered. Taylor-Robinson & Berry (1969) reported that heat labile components of guinea-pig serum had little effect on the m.i. test, but it is known that avian complement is required for the complement-associated antigen–antibody reaction (Benson, Brumfield & Pomeroy, 1961). The role of heat labile com-

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Fig. 1. The effect of temperature on (a) growth and loss of viability of *M. gallisepticum* in antiserum with HS or CS in the broth, and (b) on loss of viability in antiserum in the presence of heated or unheated CS in the broth. Incubation at 37 °C (full line) or 40·5 °C (broken line), with HS (▲), HS and antiserum (Δ), CS and antiserum (○) or heated CS and antiserum (●) present.

Components was studied by performing the m.i. test in broth containing normal or heated (56 °C for 30 min) HS and CS. Heat treatment of HS did not influence the early or late m.i. titre, nor did heating of the CS influence the early m.i. titre. However, late m.i. titres with CS broth were, on average, twofold lower after heat treatment. The rate of loss of viability of the organism in a 1:20 dilution of antiserum in broth with heated CS was similarly reduced at 37 °C, but was unaffected at 40·5 °C (Fig. 1b).

These results show that the effects of temperature and antiserum on the organism are complex. The rate of growth is reduced, the inhibitory effect of low dilutions of antiserum are enhanced, and the heat labile components only affect the rate of loss of viability at 37 °C.
Short communication

These observations support the suggestion by Davies (1969) that the bacteriostatic and bactericidal activities of antiserum are dependent on a number of factors including the stage of antigenic stimulation and response to complement.

REFERENCES


