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ANNEX

Testing scheme necessary to verify the achievement of the Community target for the reduction of Salmonella enteritidis. Salmonella hadar. Salmonella infantis. Salmonella typhimurium and Salmonella virchow in adult breeding flocks of Gallus gallus

- 3. Examination of the samples
- 3.1. Preparation of the samples
- 3.1.1. Hatcher basket liners
- place in 1 litre Buffered Peptone Water (BPW) which has been been prewarmed at (a) room temperature and mix gently;
- (b) continue culture of the sample by using the detection method in 3.2.
- 3.1.2. Boot swabs samples
- carefully unpack the pair of boot swabs (or 'socks') to avoid dislodging adherent faecal (a) material and place in 225 ml BPW which has been prewarmed to room temperature;
- (b) where five pairs of boot swabs are pooled into two samples, place five individual samples into a minimum of 225 ml BPW and ensure that all the samples are totally immersed in the BPW;
- swirl to fully saturate the sample and continue culture by using the detection method (c) in 3.2.
- 3.1.3. Other faecal material samples
- (a) at the laboratory place each sample (or pooled sample as appropriate) into an equal weight of Buffered Peptone Water and mix gently;
- (b) allow the sample to soften for 10-15 minutes then mix gently;
- immediately after mixing remove 50 g of the mixture and add to 200 ml of Buffered (c) Peptone Water which has been pre-warmed to room temperature;
- (d) continue culture of the sample by using the detection method in 3.2.

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