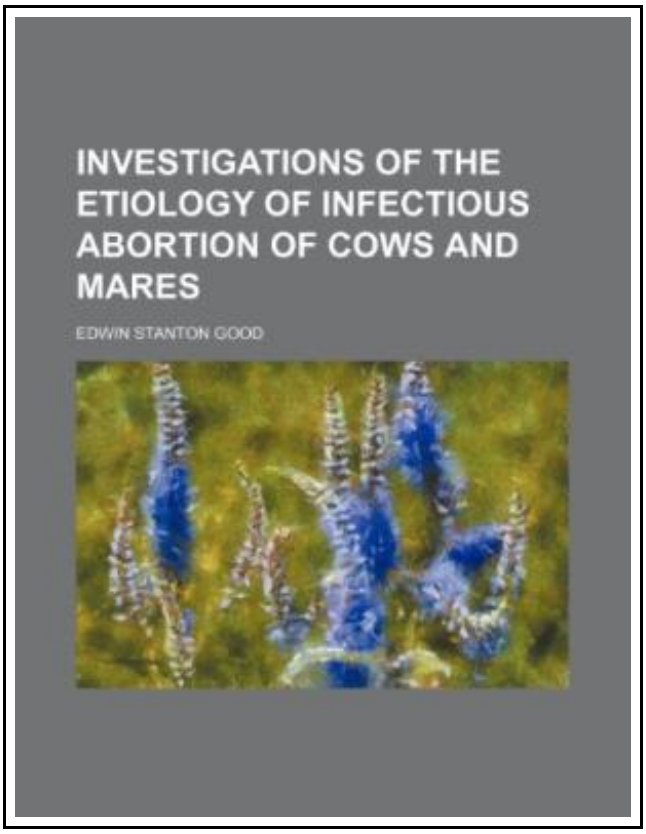
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Rarebooksclub.com, United States, 2012. Paperback. Book Condition: New. 246 x 189 mm. Language: English . Brand New Book ***** Print on Demand *****.This historic book may have numerous typos and missing text. Purchasers can download a free scanned copy of the original book (without typos) from the publisher. Not indexed. Not illustrated. 1912 Excerpt: .at the end and in the center, leaving two unstained portions of the bacillus--even the cell wall of those unstained places can seldom be seen. Under very high magnification, however, a very dim stain may at times be noted in the spaces between the well stained granules. There are some individuals in a preparation that appear wedge shaped. The methods of staining usually employed by us are as follows: Plain fuchsin, two to three minutes, slightly warmed; Lofflers methylene blue, several minutes, slightly warmed; carbolic fuchsin, two to five seconds, unwarmed. (See plates 6, 7, 8.) Gram s Stain.---An actively growing culture four days old and one fourteen days old of the same strain were used in this test. The one fourteen days old had been incubated for four days and then put in the ice chest until used. The germ growing in either tube was Gram negative, and as far as could be seen, exhibited the same characteristics relative to the stain. The two following methods of making Gram s stain were employed in the test: Method 1 is practically that accepted for the standard methods of water analyses as reported in American Public Health Association, Vol. 30, Part 2, page 101. METHOD 1. 1. Cover the preparation with aniline, gentian-violet, for one minute. 2. Wash in water and dry. 3. Cover the preparation with Lugol s solution for 2 minutes. 4. Wash in water and dry. 5. Decolorize in 95 per cent. alcohol with...

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