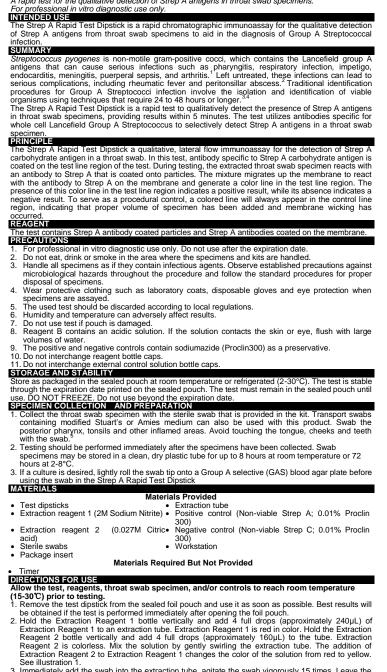
Strep A Rapid Test Dipstick (Throat Swab)

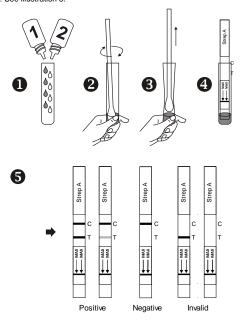
Package Insert

A rapid test for the qualitative detection of Strep A antigens in throat swab specimens.

JusChek



- See illustration 1.
 Immediately add the swab into the extraction tube, agitate the swab vigorously 15 times, Leave the swab in the extraction test tube for 1 minute. See illustration 2
 Press the swab against the side of the tube and squeeze the bottom of the tube while removing the swab so that most of the liquid stays in the tube. Discard the swab. See illustration 3.
 With arrows pointing down, place the dipstick into the tube of solution and then start the timer. If the procedure is followed correctly, the liquid should be at or just below the maximum line (MAX) on the test dipstick. See the illustration 4.
 Weil the procedure lipstopic sector and the source of the sector of
- Wait for the colored line(s) to appear. Read the result at 5 minutes. Do not interpret the result after 10 minutes. See illustration 5.



INTERPRETATION OF RESULTS

(Please refer to the illustration above) **POSITIVE:* Two lines appear.** One colored line should be in the control line region (C) and another apparent colored line should be in the test line region (T). A positive result indicates that Strep A was detected in the specimen.

*NOTE: The interspectrue. Strep A present in the spectrue. NECOTIVE: One of the color in the test line region (T) will vary depending on the concentration of Strep A present in the specimen. Therefore, any shade of color in the test line region (T) should be considered positive.

NEGATIVE: One colored line appears in the control line region (C). No line appears in the test line NEGATIVE: One colored line appears in the control line region (C). No line appears in the test line region (T). A negative result indicates that Strep A antigen is not present in the specimen, or is present below the detectable level of the test. The patient's specimen should be cultured to confirm the absence of Strep A infection. If clinical symptoms are not consistent with results, obtain another specimen for culture. INVALID: Control line fails to appear. Insufficient specimen volume or incorrect procedural techniques are the most likely reasons for control line failure. Review the procedure and repeat the test with a new test. If the problem persists, discontinue using the test kit immediately and contact your local distributor.

QUALITY CONTROL

Internal Quality Control

 Internal Quality Control

 Internal Quality Control

 Internal positive procedural control. It confirms sufficient specimen volume, adequate membrane wicking and correct procedural control. It confirms sufficient specimen volume, adequate membrane wicking and correct procedural technique.

 External Quality Control

 It is recommended that a positive and negative external control be run every 25 tests, and as deemed necessary by internal laboratory procedures. External positive and negative controls are supplied in the kit. Alternatively, other Group A and non-Group A Streptococcus reference strains may be used as external controls are not recommended.

 Procedure for External Quality Control Testing

 1. Add 4 full drops of Extraction Reagent 1 and 4 full drops of Extraction Reagent 1 and 4 full drops of the tube gently to mix the liquid.

 2. Add 1 full drops of positive or negative control solution into the tube, holding the bottle upright.

 3. Place a clean swab in the extraction tube and agitate the swab in the solution by rotating it at least 15 times. Leave the swab in the extraction tube for 1 minute. Then express the liquid from the swab head by rolling the swab agains the inside of the extraction tube and squeezing the extraction tube as the swab is withdrawn. Discard the swab.

 A. Continue with Step 5 of Directions For Use.

 If the controls do not yield the expected results, do not use the test results. Repeat the test or contact your distributor.

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- UNITATIONS
 The Strep A Rapid Test Dipstick is for in vitro diagnostic use only. The test should be used for the detection of Strep A antigen in throat swab specimens only. Neither the quantitative value nor the rate of increase in Strep A antigen concentration can be determined by this qualitative test.
- This test will only indicate the presence of Strep A antigen in the specimen from both viable and non-viable Group A Streptococcus bacteria.
 A negative result should be confirmed by culture. A negative result may be obtained if the concentration of the Strep A antigen present in the throat swab is not adequate or is below the
- detectable level of the test. 4. Excess blood or mucus on the swab specimen may interfere with test performance and may yield a false positive result. Avoid touching the tongue, cheeks, and teeth³ and any bleeding areas of the mouth with the swab when collecting specimens.
 5. As with all diagnostic tests, all results must be interpreted together with other clinical information
- , physician. vailable to the

As with all diagnostic tests, all results must be interpreted together with other clinical information available to the physician.
 EXPECTED VALUES
 Approximately 15% of pharyngitis in children ages 3 months to 5 years is caused by Group A beta hemolytic Streptococcus.⁶ In school-aged children and adults, the incidence of Strep throat infection is about 40%. This disease usually occurs in the winter and early spring in temperate climates.⁹
 PEREORMANCECHARACTERISTICS
 Sensitivity and Specificity
 Using three medical centers for evaluation, a total of 526 throat swabs were collected from patients exhibiting symptoms of pharyngitis. Each swab was rolled onto a sheep blood agar plate, and then tested by the Strep A Rapid Test Dipstick (Throat Swab). The plates were further streaked for isolation, and then incubated at 37°C with 5-10% CO2 and a Bacitracin disk for 18-24 hours. The negative culture plates were incubated for an additional 18-24 hours. Possible GAS colonies were subcultured and confirmed with a commercially available latex agglutination grouping kit. Of the 526 total specimens, 404 were confirmed to be negative and 122 were confirmed to be positive by culture. During this study, one Strep F specimens yielded positive results with the Test. One of these specimens was re-cultured and tested for cross-reactivity and also yielded negative results.
 Method
 Results Positive Negative

| inctriou | Method | | uio | Total Results | |
|-----------------------------|----------|----------|----------|---------------|--|
| | Results | Positive | Negative | Total Results | |
| Strep A Rapid Test Dipstick | Positive | 116 | 9 | 125 | |
| | Negative | 6 | 395 | 401 | |
| Total Results | | 122 | 404 | 526 | |

Relative Sensitivity: 95.1% (95%CI*: 89.6%-98.2%) Relative Specificity: 97.8% (95%CI*: 95.8%-99%)

| Accuracy: | 97.1% | (95%CI*: 95.3%-98.4%) | |
|-----------|-------|-----------------------|--|

| Positive Culture Classification | Strep A Rapid Test/Culture | % Agreement |
|------------------------------------|----------------------------|-------------|
| Rare | 8/12 | 80.0% |
| 1+ | 18/22 | 90.0% |
| 2+ | 19/20 | 95.0% |
| 3+ | 33/34 | 97.1% |
| 4+ | 38/38 | 100.0% |

Cross Reactivity The following organisms were tested at 1.0 x 10' organisms per test and were all found to be negative when tested with the Strep A Rapid Test Dipstick. No mucoid-producing strains were tested.

| Group B Streptococcus | Neisseria meningitidis | Serratia marcescens |
|----------------------------|---------------------------------------|------------------------|
| Group F Streptococcus | Neisseria sicca | Klebsiella pneumoniae |
| Streptococcus pneumoniae | Branhamella catarrhalis | Bordetella pertussis |
| Streptococcus mutans | Group C Streptococcus | Neisseria gonorrhea |
| Staphylococcus aureus | Group G Streptococcus | Neisseria subflava |
| Corynebacterium diphtheria | Streptococcus sanguis | Hemophilus influenza |
| Candida albicans | Staphylococcus epidermidis | Pseudomonas aeruginosa |
| Entorococcus faocalis | , , , , , , , , , , , , , , , , , , , | 3 |

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| Adults 30 to 65 years of age. Southern Medical Journal (May 1999), 491-492. | | | | | | | |
|---|--|--|----------|---------------|--|--------|------------------------------|
| Index of Symbols | | | | | | | |
| \triangle | Attention, see instructions for use | | Σ | Tests per kit | | EC REP | Authorized Representative |
| IVD | For in vitro diagnostic use only | | | Use by | | R | Do not reuse |
| - 30°C | Store between 2-30°C | | LOT | Lot Number | | REF | Catalog # |

Store between 2-30°C LOT Lot Number

Do not use if package is damaged Hangzhou AllTest Biotech Co., Ltd.

#550, Yinhai Street Hangzhou Economic & Technological Development Area Hangzhou - 310018, P. R. China

EC REP MedNet GmbH Borkstrasse 10 48163 Muenster Germany

Confidence Interval