



Instructions for Use

ALA DIFFERENTIATION DISKS

Cat. no. Z7081	ALA Differentiation Disks	50 disks/cartridge
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INTENDED USE

HardyDisk™ ALA (delta-aminolevulinic) Differentiation Disks rapidly detect the presence of porphyrin and cytochrome compounds and are used to differentiate *Haemophilus* species, including *Aggregatibacter aphrophilus*.

SUMMARY

Traditionally *Haemophilus* species have been differentiated by their varying requirements for hemin (X-Factor), NAD (nicotinamide adenine dinucleotide, V-Factor), and a combination of hemin and NAD (XV-Factor). However, erroneous results have been demonstrated when using growth factor requirement tests. These misidentifications are largely attributed to the carryover of X-Factor in the inoculum as well as the presence of trace amounts of X-Factor in the medium. ⁽⁵⁾ The use of HardyDisk™ ALA Differentiation Disk is used as an alternative method to X-Factor requirement testing. The ALA procedure is a more rapid test method as well as a more accurate method for determining hemin requirements by eliminating the erroneous results associated with X-Factor requirement tests. ⁽⁸⁾

HardyDisk™ ALA Differentiation Disk assesses the ability of a *Haemophilus* strain to synthesize hemin from the ALA substrate. The test detects the presence of porphyrin compounds, which are intermediates in the hemin biosynthetic pathway. ^(4,5) Porphyrins are detected by the emission of a red-orange fluorescence under UV light (366nm) and indicate that the organism is capable of synthesizing hemin and is not dependent on hemin (X-Factor) for growth. Conversely, hemin requiring strains lack the enzymes to synthesize hemin and do not produce intermediate porphyrin compounds.

FORMULA

Each HardyDisk™ ALA Differentiation Disk is prepared by impregnating carefully controlled concentrations of delta-aminolevulinic acid onto a high quality 6mm diameter filter paper disk.

STORAGE AND SHELF LIFE

Storage: Upon receipt store at -20 to +8°C. away from direct light. Disks should not be used if there are any signs of discoloration or if the expiration date has passed. Product is light and temperature sensitive; protect from light, excessive heat, moisture, and freezing.

The expiration dating on the product label applies to the product in its intact packaging when stored as directed. The product may be used and tested up to the expiration date on the product label and incubated for the recommended quality control incubation times.

Refer to the document "[Storage](#)" for more information.

PRECAUTIONS

This product may contain components of animal origin. Certified knowledge of the origin and/or sanitary state of the animals does not guarantee the absence of transmissible pathogenic agents. Therefore, it is recommended that these products be treated as potentially infectious, and handle observing the usual universal blood precautions. Do not ingest, inhale, or allow to come into contact with skin.

This product is for *in vitro* diagnostic use only. It is to be used only by adequately trained and qualified laboratory personnel. Observe approved biohazard precautions and aseptic techniques. All laboratory specimens should be considered infectious and handled according to "standard precautions." The "Guidelines for Isolation Precautions" is available from the Centers for Disease Control and Prevention at www.cdc.gov/ncidod/dhqp/gl_isolation.html.

For additional information regarding specific precautions for the prevention of the transmission of all infectious agents from laboratory instruments and materials, and for recommendations for the management of exposure to infectious disease, refer to CLSI document M-29: *Protection of Laboratory Workers from Occupationally Acquired Infections: Approved Guideline*.

Sterilize all biohazard waste before disposal.

Refer to the document "[Precautions When Using Media](#)" for more information.

Refer to the document [SDS Search](#) instructions on the Hardy Diagnostics' website for more information.

PROCEDURE

Specimen Collection: This product is not intended for primary isolation of patient specimens. It should be used only with cultures of isolated organisms. This product is used in conjunction with other biochemical tests to identify cultures of isolated organism.

Method of Use:

1. Perform HardyDisk™ ALA Differentiation Disks only on isolates either growing on Chocolate Agar in 18-24 hours or that satellite around *Staphylococcus aureus* on a blood agar plate. See "Precautions" section above.
2. Prior to use, allow the disks to equilibrate to room temperature.
3. Aseptically place the disk into a sterile petri plate. Wet the disk with a small drop (0.04ml) of sterile saline (Cat. no. R45).
4. Inoculate disk with several well isolated 18-24 hour colonies to yield a visible cell paste on the disk surface. Alternatively, the disk can be touched to a colony and then placed in a petri dish.
5. Moisten a piece of filter paper with water and place it into the lid of the petri plate to ensure that the disk is kept moist during incubation.
6. Incubate the disks aerobically at 35°C. for up to 2 hours.
7. After 2 hours, examine the disk under ultra-violet light (366nm) in a darkened room. Observe the disk for the presence of a red fluorescence while under UV light.

INTERPRETATION OF RESULTS

A positive reaction is recorded when orange/red fluorescence is observed on the HardyDisk™ ALA Differentiation Disk. This color change is a positive result for porphyrin synthesis and indicates that the organism does not require hemin (X-Factor) for growth.

A negative result is recorded when no fluorescence is observed on the disk and indicates that porphyrin was not

synthesized. Consequently, the organism requires hemin for growth.

<i>Haemophilus</i> species	Growth Factor Requirements		Porphyrin Synthesis
	X	V	ALA
<i>Haemophilus ducreyi</i>	+	-	-
<i>Haemophilus haemolyticus</i>	+	+	-
<i>Haemophilus influenzae</i>	+	+	-
<i>Haemophilus parahaemolyticus</i>	-	+	+
<i>Haemophilus parainfluenzae</i>	-	+	+
<i>Aggregatibacter aphrophilus</i> (formerly <i>H. aphrophilus</i> and <i>H. paraphrophilus</i>)	-	V	+

LIMITATIONS

It is recommended that biochemical, immunological, molecular, or mass spectrometry testing be performed on colonies from pure culture for complete identification.

In order to avoid inaccurate results it is recommended that organisms be less than 24 hours old, a heavy inoculum is used to inoculate the disk, and that the disk is kept moist during incubation.⁽⁴⁾

H. ducreyi may not be identified using this procedure as some strains do not grow as satellite colonies on blood agar and grow more slowly, on the magnitude of several days, on enriched media.

Because similarities exist in growth factor requirements of *Haemophilus* species, it is not recommended that this procedure be the sole criterion for species identification.^(4,5) Consult listed references for additional information regarding the recommended tests for complete identification of *Haemophilus* species.⁽²⁻⁵⁾

The ALA test, even in conjunction with the satellite test, does not differentiate *H. influenzae* and *H. haemolyticus*. *H. haemolyticus* is not considered pathogenic and is separated from *H. influenzae* by a positive hemolysis reaction on rabbit or horse blood agar.

Francisella spp., including *Francisella tularensis*, is ALA negative and grows only on chocolate agar. It can be differentiated from *H. influenzae* as it grows more slowly, does not satellite on blood agar, and is not V-Factor dependent.

Refer to the document "[Limitations of Procedures and Warranty](#)" for more information.

MATERIALS REQUIRED BUT NOT PROVIDED

Standard microbiological supplies and equipment such as UV light, loops, other culture media, swabs, applicator sticks, incinerators, and incubators, etc., as well as serological and biochemical reagents, are not provided.

QUALITY CONTROL

Hardy Diagnostics tests each lot of commercially manufactured media using appropriate quality control microorganisms and quality specifications as outlined on the Certificates of Analysis (CofA). The following organisms are routinely used for testing at Hardy Diagnostics:

		Incubation	
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Test Organisms	Inoculation Method*				Results
		Time	Temperature	Atmosphere	
<i>Aggregatibacter aphrophilus</i> (formerly <i>H. parainfluenzae</i>) ATCC® 7901	*	2hr	35°C	Aerobic	Positive for porphyrin synthesis; orange/red fluorescence in the presence of UV light
<i>Haemophilus influenzae</i> ATCC® 10211	*	2hr	35°C	Aerobic	Negative for porphyrin synthesis; no red fluorescence in the presence of UV light

* Refer to the document "[Inoculation Procedures for Media QC](#)" for more information.

USER QUALITY CONTROL

End users of commercially prepared culture media should perform QC testing in accordance with applicable government regulatory agencies, and in compliance with accreditation requirements. Hardy Diagnostics recommends end users check for signs of contamination and deterioration and, if dictated by laboratory quality control procedures or regulation, perform quality control testing to demonstrate growth or a positive reaction and to demonstrate inhibition or a negative reaction, if applicable. Hardy Diagnostics quality control testing is documented on the certificates of analysis (CofA) available from Hardy Diagnostics [Certificates of Analysis](#) website. In addition, refer to the following document "[Finished Product Quality Control Procedures](#)," for more information on QC or see reference(s) for more specific information.

PHYSICAL APPEARANCE

HardyDisk™ ALA Differentiation Disks are 6mm (in diameter) filter paper disks with the letters ALA printed on both sides, and should appear white in color.



Showing positive (left disk) and negative (right disk) HardyDisk™ ALA Differentiation Disks (Cat. no. Z7081) under UV light.

Disks were moistened with a drop of sterile saline and growth from 24 hour cultures was applied to the disks with a sterile loop. The disks were incubated aerobically in a petri dish for two hours. A sterile piece of filter paper soaked with deionized water was placed in the dish to maintain proper humidity.
 LEFT DISK: *Haemophilus parainfluenzae* (ATCC® 7901) growth applied to ALA Disk. The orange/red color under UV light was indicative as **positive**.
 RIGHT DISK: *Haemophilus influenzae* (ATCC® 10211) growth applied to ALA Disk. No orange/red color under UV light was indicative as **negative**.

REFERENCES

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[Ordering Information](#)

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