

# Hydroxystilbamidine \*CAS 223769-64-0\*

PRODUCT INFORMATION SHEET

Catalog number: 17514 Unit size: 10 mg

Component	Storage	Amount
Hydroxystilbamidine *CAS 223769-64-0*	Freeze (< -15 °C), Minimize light exposure	1 vial (10 mg)

# OVERVIEW

Hydroxystilbamidine (also called Fluoro Gold) is used for staining DNA and RNA. It exhibits distinctively different fluorescence emission profiles when bound to DNA compared to RNA. This cationic dye is also frequently used as a retrograde neuronal tracer. Fluoro-Gold is a trademark of Fluorochrome, Inc.

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### **Biological Applications**

Hydroxystilbamidine (also called Fluoro Gold) is used for staining DNA and RNA. It exhibits distinctively different fluorescence emission profiles when bound to DNA compared to RNA. This cationic dye is also frequently used as a retrograde neuronal tracer.

#### **Spectral Properties**

Excitation = 360 nm; Emission = 625 nm

# PREPARATION OF WORKING SOLUTION

#### Hydroxystilbamidine working solution

1 to 10% Hydroxystilbamidine in water has been successfully used. Initially, a 4% concentration is advised. If undesirable necrosis occurs at the injection site, or labeling is too intense, reduce the concentration to a 2% solution.

## SAMPLE EXPERIMENTAL PROTOCOL

The use of Hydroxystilbamidine is essentially the same as other fluorescent tracers. The main difference is that Fluoro-Gold is more flexible in terms of post-injection survival times, concentration range, tissue treatment and compatibility with other histochemical techniques. It is also more resistant to fading, brighter and more permanent than most other fluorescent tracers.

#### **Dye Administration**

- 1. Pressure injection This is probably the most frequently used mode of application. Volumes injected range from 0.05-1  $\mu$ L, typically 0.1-0.2  $\mu$ L.
- 2. Crystal A crystal of the tracer can be administered from the tip of a micro-pipette.

#### Fixation

Although any fixative, or no fixative, can be used, PBS containing 4% formaldehyde is most frequently employed. Fixatives containing high concentrations of heavy metals (eg. osmium, mercury) will quench the fluorescence, while high concentrations (over 1%) of glutaraldehyde may increase background fluorescence

#### Histochemical Processing

Tissue containing Hydroxystilbamidine may be processed according to virtually any common histological technique. Frozen sections of fixed tissue are most frequently used.

## **Combined Methods**

At this point of processing, sections may be further processed for a second marker such as autoradiography, HRP histochemistry, immunocytochemistry, a second fluorescent tracer, fluorescent counterstain, etc.

#### Mounting, Clearing and Coverslipping

Sections are typically mounted on gelatin-coated slides, air-dried, immersed in xylene, and coverslipped with nonfluorescent DPX plastic mounting media. Sections may be dehydrated with graded alcohols, unless this is not compatible with a second tracer. If Hydroxystilbamidine is to be combined with fluorescence immunocytochemistsry, then sections are air-dried and directly coverslipped with DPX.

#### **Examination and Photography**

Hydroxystilbamidine can be visualized with a fluorescence microscope using a wide band ultraviolet excitation filter (excitation - 323 nm, emission - 620 nm at neutral pH). A gold color is emitted when tissue has been processed with neutral pH buffer whereas a blue color is emitted when tissue is processed with acidic (eg. PH 3.3) pH buffer. It can be photographed digitally or with film (use Ektachrome 200-400 ASA film for color prints and comparable speed film for black and white prints). Most exposure times range from 10-60 second exposures, depending on the objective magnification and the intensity of the label. Thirty (30) second exposures are about average. Multiple exposures may be exploited to simultaneously visualize Hydroxystilbamidine and another tracer. Thus, UV would be combined with bright field illumination to simultaneously locate Fluoro-Gold with HRP or silver grains in autoradiography. Similarly, blue light excitation can be combined to also visualize the green emission color of FITC, while green excitation light may be used to simultaneously observe the red emission color of propidium iodide, or ethidium bromide (a fluorescent counterstain).

# EXAMPLE DATA ANALYSIS AND FIGURES

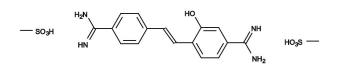


Figure 1. Chemical structure for Hydroxystilbamidine \*CAS 223769-64-0\*

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