Rev A/2014-02-12/SF-2464DSCA\VK

EGR1 Hu-Cy5 SmartFlare™ RNA Detection Probe

Cat. # SF-2464

pack size: 50µL (250 rxns)

Store at 2-8°C, after reconstitution store at 23-27°C DO NOT FREEZE

FOR RESEARCH USE ONLY.
NOT FOR USE IN DIAGNOSTIC PROCEDURES
NOT FOR HUMAN OR ANIMAL CONSUMPTION



Product Data Sheet

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Validated Accession #: NM_001964.2 Species: Hu Gene Aliases: TIS8; AT225; G0S30; NGFI-A; ZNF225; KROX-24; ZIF-268

Confirmation of EGR1 SmartFlare Performance:

EGR1 SmartFlare probe has been tested in a buffer system to detect the release of the fluorophore in the presence of a complementary base pair sequence for each lot to confirm target specificity.

EGR1 SmartFlare probe has also been tested in a cell model system and demonstrated increased fluorescence in cells expressing the target compared to a scrambled negative control SmartFlare probe (Figure 1).

| Mean Fluorescence Intensity (MFI) Values | | | | | |
|--|--------------------|------------------|--|--|--|
| Unflared | Scrambled | EGR1 | | | |
| 6.90299999999999 | 16.527999999999999 | 67.4920000000000 | | | |

Figure 1: EGR1 Mean Fluorescence Intensity (green) measured by flow cytometry in living HEK293 cells demonstrated a significant increase over unflared cells (blue) as well as scramble control (red). Data shown in graph is representative.

Storage and Handling:

Material has been 0.22µm filtered. Stable for 5 years at 2-8°C degrees in lyophilized format ONLY. Room temperature is required for reconstituted product.

Warning-after reconstitution product is sensitive to cold and hot temperatures, a stable room temperature of 23-27°C is required.

Handling Recommendations:

Reconstitute with sterile nuclease free water in a drop wise fashion and tap tube repeatedly to fully dissolve lyophilized material. Vortex for 5-10 sec.

Upon reconstitution, store at room temperature for up to 1 year protected from light. Product must be handled with gloves as product can be absorbed through the skin.

Recommended Cell Testing Protocol:

(example: 30,000 cells in a $200\mu L$ media volume within each well of a 96 well plate)

- Reconstitute reagent in 50µL of sterile nuclease free water.
- Create a working solution based on your experiment by diluting 1:20 in sterile PBS.
- Add 4µL directly to cells (at approx 80% confluency).
- Allow to incubate overnight for 16 hrs.
- Detect using fluorescence detection platform of choice.

