# Rev A/2014-04-09/SF-2923DSCA\VK

# TRPM2 Hu-Cy5 SmartFlare<sup>TM</sup> RNA Detection Probe

Cat. # SF-2923

pack size: 50µL (250 rxns)

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

Species: Hu

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

Validated Accession #: NM 003307.3



# **Product Data Sheet**

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Gene Aliases: KNP3; EREG1; TRPC7; LTRPC2; NUDT9H; NUDT9L1

# Confirmation of TRPM2 SmartFlare Performance:

TRPM2 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.

TRPM2 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	TRPM2			
5.7400000000000000	37.810000000000002	171.919999999999			

**Figure 1:** TRPM2 Mean Fluorescence Intensity (green) measured by flow cytometry in living Jurkat 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 200µL 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.

