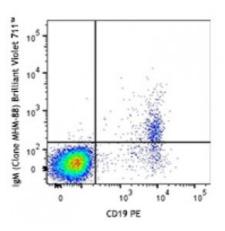
## **Product Data Sheet**

## Brilliant Violet 711® anti-human IgM

Catalog # / Size:	2172700 / 100 tests 2172695 / 25 tests
Clone:	MHM-88
Isotype:	Mouse IgG1, κ
Immunogen:	Human Ig cocktail
<b>Reactivity:</b>	Human
Preparation:	The antibody was purified by affinity chromatography and conjugated with Brilliant Violet 711 <sup>™</sup> under optimal conditions. The solution is free of unconjugated Brilliant Violet 711 <sup>™</sup> and unconjugated antibody.
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and BSA (origin USA).
Concentration:	Lot-specific



Isolated human peripheral blood mononuclear cells were stained with CD19 PE and anti-human IgM (clone MHM-88) Brilliant Violet 711<sup>™</sup> (top) or mouse IgG1, κ Brilliant Violet 711<sup>™</sup> isotype control (bottom).

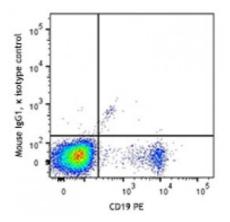
## **Applications:**

Applications:	Flow Cytometry
Recommended Usage:	Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis. For flow cytometric staining, the suggested use of this reagent is ≤5 microL per million cells or 5 microL per 100 microL of whole blood. It is recommended that the reagent be titrated for optimal performance for

each application.

Brilliant Violet 711<sup>™</sup> excites at 405 nm and emits at 711 nm. The bandpass filter 710/50 nm is recommended for detection, although filter optimization may be required depending on other fluorophores used. Be sure to verify that your cytometer configuration and software setup are appropriate for detecting this channel. Refer to your instrument manual or manufacturer for support. Brilliant Violet 711<sup>™</sup> is a trademark of Sirigen Group Ltd.

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U.S. Patent(s), pending patent	
applications and foreign equivalents.	

Application Notes: MHM-88 antibody reacts with both soluble and membrane human immunoglobulin M (IgM). It does not react with other Ig isotypes. Additional reported applications (for the relevant formats) include: use as a primary or secondary reagent for ELISA analysis.

> Due to the presence of excess soluble IgM in whole blood, which competes for antibody binding, staining for IgM on cells in whole blood is not recommended.

**Application** 1. Perez-Shiyama C, *et al.* 2014. *J Immunol.* 192:5192. <u>PubMed</u> **References:** 

**Description:** IgM is the first immunoglobulin made by B cells in the immune response. Surface IgM is expressed on immature and mature B cells, while IgM heavy ( $\mu$ ) chain is expressed intracellularly in pre-B cells.