

Maximize Detection of Light Chains

Reduce dubiously negative, dual-positive, or difficult-to-interpret results by using Agilent Dako rabbit polyclonal anti-kappa or lambda light chain reagents.



Key characteristics of Agilent Dako rabbit polyclonal reagents

Agilent
Dako

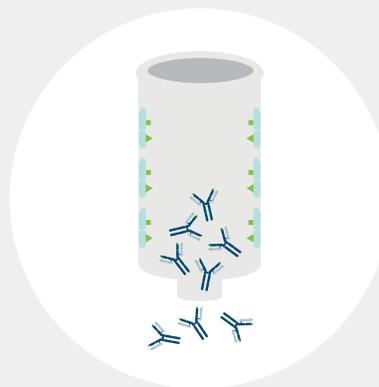
Robust immunization

The strain and pool of rabbits used in producing our polyclonal antibodies ensures steady response and yields of optimal reproducibility. Robust production standards result in reliable lot-to-lot output.



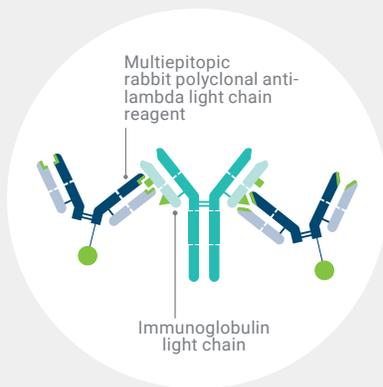
High-grade purity

Several purification steps select solely for light reactivity removing unspecific antibodies and other serum proteins. The process results in highly purified and specific reagents with minimal background noise or cross-reactivity.



Inherent consistency

By targeting multiple epitopes of the light chain, our rabbit polyclonal reagents have an inherent capacity to accommodate antigen polymorphisms, ensuring consistent staining in difficult cases.



Enhanced specificity

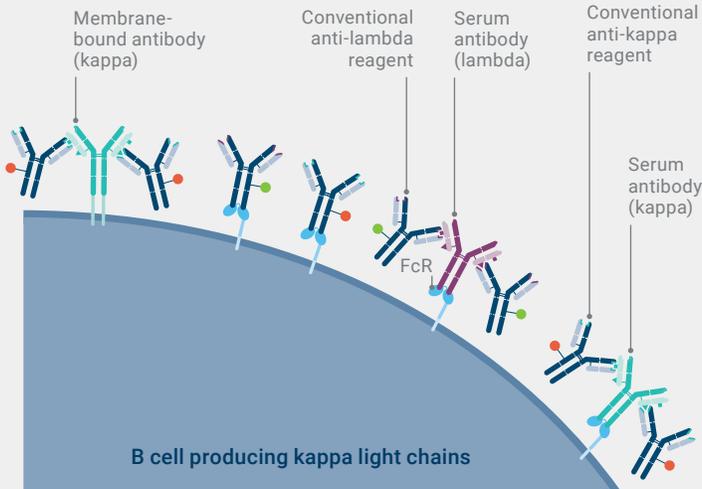
F(ab)₂ fragments do not contain the Fc region and cannot bind to Fc receptors. The elimination of off-target binding is expected to reduce noise caused by FcR interference.



Eliminate interference of the reagent with cellular Fc receptors and ensure meaningful separation of kappa vs lambda light chain populations despite residual cross-staining from serum antibodies

Using conventional anti-light chain reagents containing Fc region

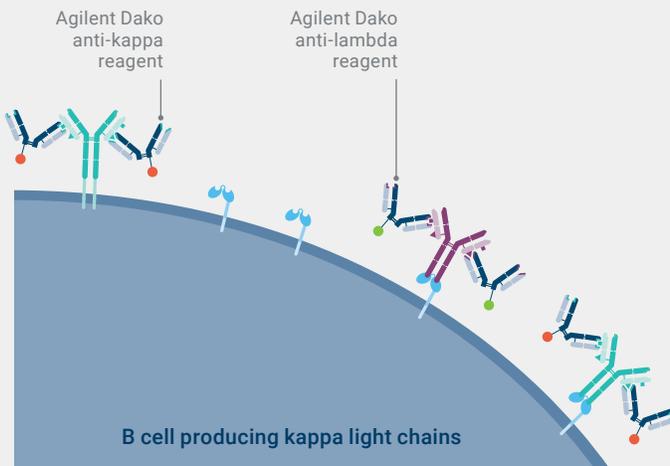
Target labeling | High FcR interference



Conventional reagents with high FcR interference are difficult to interpret

Using Agilent Dako anti-light chain reagents F(ab')₂ fragments of high purity and specificity

Target labeling | Residual FcR interference



Results with Agilent Dako polyclonal reagents are easier to interpret due to clearer separation of cell populations

For more information, please visit:
www.agilent.com

Please contact us for more details:
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This information is subject to change without notice. Always refer to the regulatory labeling of the products in the country of interest. For US users, in accordance with 21 CFR Part 809.30 (G), the current material is intended only for IVD manufacturers or organizations that use the reagents to make tests for purposes other than providing diagnostic information to patients and practitioners.

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