

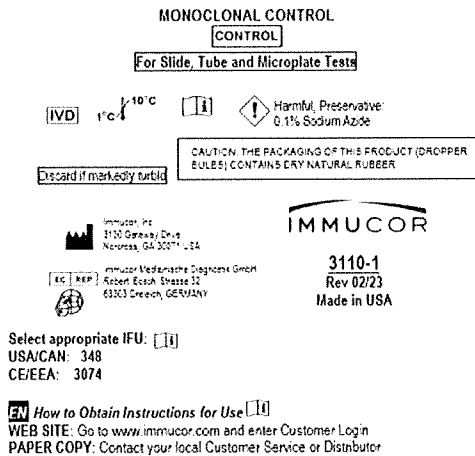
Dear Valued Immucor Customer,

As a result of ongoing due diligence to ensure compliance with global regulatory requirements as well as our commitment to protecting our environment through our sustainability initiatives, Immucor will begin the transition to electronic Instructions For Use (eIFU) for our Transfusion Serology products legally manufactured by Immucor, Inc.

We are also taking this opportunity to update Instructions For Use to meet global in-vitro diagnostic regulations. We will continue to underline significant changes made to Instructions for Use to easily identify modifications to the current revision.

Immucor is planning to begin implementation of eIFUs in **mid-April**. Beginning April 17, 2023, you may see products that are no longer shipped with a printed IFU, since we will be implementing this change in coordination with manufacturing cycles. Therefore, due to the shelf life of our products we anticipate that the implementation of eIFUs for our entire product line will take **approximately 18 months**.

As product is transitioned to an eIFU, it will be packaged with an insert instructing you to login into the Immucor Customer Center and will provide information to select the appropriate IFU. This insert will include instructions in multiple languages. An example is found below:



If you have any questions, please don't hesitate to contact your local Immucor Representative.

Best Regards,  
Amanda Gallagher  
Sr. Director, Global Transfusion Marketing  
Immucor, Inc.



## Electronic Instructions For Use COMING SOON!

Immucor <marketinginformation@immucor.com>

Wed 2023-04-05 10:42

To: Slayten, Jayanna K <jslayten@IUHealth.org>

\*\*\*\* EXTERNAL Message From 355734626000020fb4f02-b23095-

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