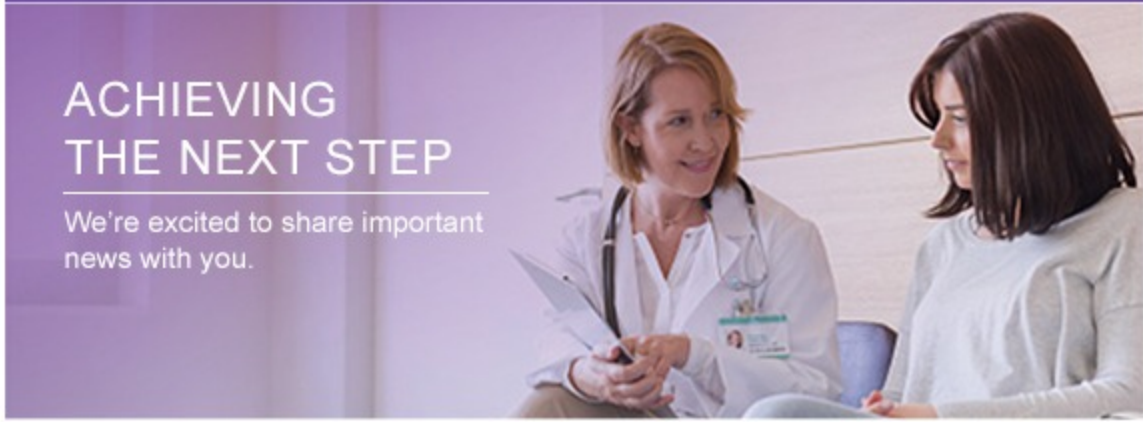


TOUCHPOINTS | Keeping you informed so you can care with confidence.



ACHIEVING THE NEXT STEP

We're excited to share important news with you.

FDA SUBMISSION UPDATE

Patients are now enrolling in our new study

A significant milestone has been reached.

In late spring, we shared that initiating patient enrollment for our new "critically ill" study was a top priority. That's why we are pleased to announce that patient enrollment is now underway. We have incorporated feedback and are implementing new approaches in our study to ensure the highest probability of success. Now, reaching this key objective signifies great progress.

It continues to be a priority to submit the Accu-Chek® Inform II system for FDA clearance for use with all appropriate patients through the hospital, including "critically ill" patients and to meet our customers' clinical policies.

It's important to keep in mind that, due to multiple interdependencies in a clinical trial and development of an FDA submission, it's difficult to predict the exact timing of when a regulatory application will be submitted.

We are working with the greatest sense of urgency for clearance. We'll continue to keep you posted on our progress. [For news of the latest developments, sign up for updates at go.roche.com/informMe](#)



STAY INFORMED

CARING WITH CONFIDENCE

How to manage the "critically ill" limitation

Education is key.

Did you know? Every commercially available strip-based hospital blood glucose monitoring system has a "critically ill" limitation. Therefore, recognizing how these meters are used in your system is key to managing compliance with your device.

To develop the most effective, clinically appropriate procedures, we encourage collaboration with your end users. For regulatory compliant options for managing the "critically ill" limitation, [click here](#).



ACCESS EDUCATION RESOURCES

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