

**Clinical Innovations LLC**  
**URGENT MEDICAL DEVICE RECALL**  
***Kiwi Complete Vacuum Delivery System (VAC-6000MT)***

Re: Customer Notification regarding Recall of the Kiwi Complete Vacuum Delivery System (with Traction Force Indicator) (VAC-6000MT)

March 12, 2026

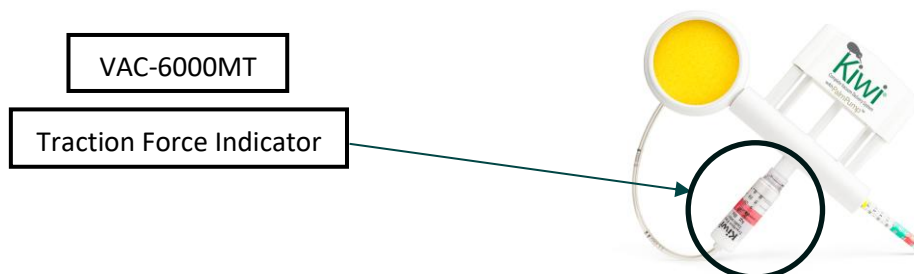
Dear Valued Laborie Customer:

This is to inform you of a voluntary recall of Clinical Innovations' **Kiwi** Complete Vacuum Delivery System (VAC-6000MT) involving the lots in Table 1 below. The purpose of the recall is to address the potential failure of handle break at the Traction Force Indicator handle joint when angular force is applied to the handle. This recall is being conducted due to a manufacturing issue which can compromise the integrity of the bond between the Traction Force Indicator (TFI) and handle joint when certain use techniques are utilized.

It is important to note that this recall **does not affect the following**:

- 1) Kiwi VAC-6000MT manufactured before August 31, 2025
- 2) Kiwi VAC-6000M, Kiwi VAC-6000S or VAC-6000C

For clarity, a picture of the **affected** model is shown below\*.



The company has received 47 complaints believed to involve this issue out of 94,000 devices sold globally. Four complaints indicated that there may have been an undefined patient injury. Although

these events have not been confirmed, the company is currently working with these accounts to obtain additional information. Based on the company's investigation into this, the issue may occur on units from affected lots when the device is pulled at an angle and/or torque is applied to the joint. We remind users that the Instructions for Use includes a warning "not to twist, torque or use excessive force."

The company's investigation has determined that the breakage is not likely to occur when pulling force is applied in line with the TFI.

## **Risk to Health**

Breakage of the device results in complete loss of vacuum which can lead to a delay in the procedure while another mode of delivery is employed or the possible conversion to cesarean section delivery. Delaying a procedure may pose an increased risk of harm when the indication for vacuum assisted delivery is fetal distress. In cases of use in cesarean sections, it is also theoretically possible that fragments could fall into the patient.

## **Actions to be Taken by Customers and Distributors**

Clinical Innovations' (Laborie) records indicate you have received a product that is affected by this action and we are notifying all customers of this recall to confirm that no product (Impacted Lots listed in Table 1 below) remains in distribution channels. Please take the following actions immediately:

1. **Identify and quarantine** all unused inventory of the affected product.
2. **Discontinue distribution** of the affected product.
3. **Return all unused, affected product** to Clinical Innovations (Laborie), regardless of the labeled expiration date.
4. **Complete and return the attached *Recall Acknowledgement and Receipt Form*** within 15 **business days**, even if no inventory remains. Please return the forms via email to [recalls@Laborie.com](mailto:recalls@Laborie.com).
5. **If you have product to return**, a company representative will contact you once we receive your *Recall Acknowledgement and Receipt Form* to issue you an RMA per company policy and arrange for a product return.

## **Product Return Instructions**

- Once we receive your Recall Acknowledgement and Receipt Form, a company representative will contact you and arrange for the return of the product.
  - If you have any questions, call Weiron on (60) 167536258 or email: [singmin.tan@weiron-my.com](mailto:singmin.tan@weiron-my.com)
- Returned product will be **credited to your account**

### **Communication to Others**

Please provide this information to your hospitals. If you have further distributed impacted lots of this product listed in Table 1, please identify your customers and notify them at once of this communication and/or contact Clinical Innovations (Laborie) with contact information so that we can follow up with the owner of the device.

### **Product and Distribution Information**

See Table 1 for a list of affected Kiwi VAC-6000MT lots.

### **Action Being Taken by Clinical Innovations (Laborie)**

We are working diligently to resolve this issue and to support our customers in minimizing any disruption. Once replacement product of Kiwi VAC-6000MT becomes available, we will contact you directly.

Clinical Innovations (Laborie) is notifying and working with the appropriate regulatory authorities in each country in accordance with applicable laws.

***\*Please note, Kiwi VAC-6000M is NOT impacted and remains available for use.***

We regret any inconvenience this may cause and appreciate your understanding and partnership. Our priority remains ensuring that every Laborie product meets the high-quality standards you expect and rely on to deliver safe maternal and neonatal care.

If you have any questions regarding this recall, please contact us using the information above.

Sincerely,

Nicole Boser

Executive Vice President, Quality and Regulatory

[nboser@laborie.com](mailto:nboser@laborie.com)

**Table 1: Lots of *Kiwi Complete Vacuum Delivery System (VAC-6000MT)* UDI :00814247020642 Impacted by this Notice**

<b>Lot</b>	<b>Product Code:</b>
251188	VAC-6000MT
251189	VAC-6000MT
251190	VAC-6000MT
251230	VAC-6000MT
251276	VAC-6000MT
251308	VAC-6000MT
251324	VAC-6000MT
251327	VAC-6000MT
251328	VAC-6000MT
251330	VAC-6000MT
251333	VAC-6000MT
251370	VAC-6000MT
251388	VAC-6000MT
251408	VAC-6000MT
251409	VAC-6000MT
251545	VAC-6000MT
251546	VAC-6000MT
251582	VAC-6000MT
251583	VAC-6000MT
251680	VAC-6000MT
251681	VAC-6000MT
251723	VAC-6000MT
260219	VAC-6000MT
260240	VAC-6000MT

## MEDICAL DEVICE RECALL RETURN RESPONSE

### Acknowledgement and Receipt Form

Response is Required

**Customer Information:**

Customer Name:

Street Address:

Town, State, Zip Code:

### Kiwi Complete Vacuum Delivery System (VAC-6000MT with Traction Force Indicator)

**Lot/Serial Numbers:**

I have read and understand the recall instructions provided in the March 9, 2026 letter.

Yes \_ No\_

Any adverse events associated with recalled product? Yes \_\_ No \_\_

If yes, please explain:

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**Affected Product Information:** Include information that is applicable for affected product.

Affected Product Information Table		
Product/Brand Names	Lot/Serial Number/UDI of product in inventory	Quantity in Inventory

**Return this form to:** [Recalls@laborie.com](mailto:Recalls@laborie.com)