

TIMERTAG GO INSTRUCTIONS FOR USE

English (EN) · IFU-TTG-001 Rev B · Issued 15AUG26

Read this document before first use. Retain for reference. Free template · Single-patient use

SUPERSESION NOTICE

This document supersedes IFU-TTG-001 Rev A (issued 26JUL26) in full. Rev A specifies a 20 mm × 40 mm label applied flat to dressings and drainage bags, and refers to an Annex A that does not exist. Do not rely on, distribute, or reproduce Rev A.

1 PRODUCT DESCRIPTION AND INTENDED PURPOSE

Timertag Go is a free label template that gives patients and families plain-language guidance about a medical consumable in place, such as a peripheral IV catheter, an indwelling urinary catheter, or a dressing.

Timertag Go labels must be professionally printed from the template as 20 mm × 90 mm rounded-corner labels supplied on rolls. In use, the label is folded on itself around tubing to form a double-sided flag. The artwork carries the same QR code at each end, so that both faces of the folded flag can be scanned. Scanning with a smartphone camera opens a medical consumable-specific guidance page in the phone's web browser, built on language-agnostic graphics and a simple checklist of signs to report to nursing staff.

No app, no account, no login, and no patient data. Timertag Go labels are static: they have no timing or monitoring function. For dwell-time tracking, see Timertag Pro (IFU-TTP-003).

2 CLASSIFICATION NOTICE

Timertag Go labels are informational and educational aids only. They do not monitor the medical consumable or the patient, do not alert staff, do not process patient data, and do not replace clinical observation or judgment.

3 INTENDED USERS

For ordering by the deploying organisation and application by clinical staff. For scanning by patients, families, and caregivers using any internet-enabled smartphone with a camera. Not intended for application by patients.

4 WHAT YOU NEED

- The current Timertag Go label template for the relevant medical consumable type, downloaded from www.timertag.com.
- A commercial label printing company that can supply 20 mm × 90 mm rounded-corner rectangular labels on rolls.
- An internet connection at the point of care for the phone opening the guidance page.

5 ORDERING AND PRINTING

Download the current template for the relevant medical consumable type from www.timertag.com and supply it to your label printer unaltered. Labels must be produced as 20 mm × 90 mm rounded-corner rectangles on rolls.

The artwork places a QR code at each end of the label. Both must print at full size and full contrast, because the label is folded in half in use and each code becomes one face of the flag. Do not resize, crop, redraw, or reposition the artwork: distortion of a QR code prevents scanning.

Specify non-metallic, non-foil label stock. Commercial printers apply their own cutting tolerances and colour profiles, so review and approve the printer's proof before the production run. Fold and test-scan labels from each delivered batch, checking both faces, before clinical use. Reorder from the current template rather than reusing stored files, so that guidance reflects the latest version.

1 ORDER → 2 FOLD AND APPLY → 3 SCAN

6 APPLICATION

Timertag Go is applied as a folded flag wrapped around tubing. It is not applied to the patient's skin, to dressings, to securement devices, or to fluid containers.

Complete insertion, dressing, and securement of the medical consumable in accordance with hospital protocol before applying a label.

1. Peel one label from the roll.
2. Lay the tubing across the centre of the adhesive face, at the fold line.
3. Fold the label back on itself so that the two adhesive halves meet around the tubing and bond to each other.
4. Press firmly along the join. A QR code now faces outward on each side of the flag.
5. Confirm that both faces scan before leaving the bedside.

TIMERTAG GO INSTRUCTIONS FOR USE

English (EN) · IFU-TTG-001 Rev B · Issued 15AUG26

Read this document before first use. Retain for reference. Free template · Single-patient use

Apply the flag at the following points:

- **Peripheral IV catheter** — the extension set or connector tubing, clear of the dressing. See 6.1.
- **Urinary catheter** — the drainage tubing, above the sampling port. Not the drainage bag.
- **Dressing, central venous catheter, or PICC** — the catheter limb or extension leg, distal to the securement device. Never between the securement device and the insertion site.

Do not apply the label:

- to the patient's skin, or over broken skin;
- to a dressing, a dressing edge, a dressing inspection window, or a securement device;
- to a urinary drainage bag, an infusion bag, or any other fluid-filled container;
- over an insertion site;
- where it obstructs a clamp, roller, injection port, connector, or printed graduation.

Position the flag so that it hangs free without pulling on the catheter, the securement device, or a connector, and so that it does not snag on bed rails or bedding. Replace the label if it becomes soiled, damaged, or unreadable.

6.1 Peripheral IV limitations

Timertag Go is applied to tubing. A peripheral IV catheter dressed with a needle-free connector or bung only, with no extension set or accessible tubing, provides no suitable application point. In that configuration Timertag Go cannot be used. Do not apply the label to the dressing, to the cannula hub, or to the skin as an alternative.

Where an extension set or connector tubing is present, apply the flag to that tubing, and discard the label whenever the extension set or the cannula is replaced.

7 PATIENT USE

Open the phone's camera, point it at the QR code on either face of the flag, and tap the link that appears. The guidance page explains key warning signs to report to nursing staff immediately: looseness, leakage, swelling, redness, pain, or fever.

8 WARNINGS AND PRECAUTIONS

! READ BEFORE USE

- Guidance only. Timertag Go does not observe the medical consumable or the patient, does not measure time, and does not notify staff. Clinical observation schedules remain unchanged.
- Single-patient use. Do not move a label between patients.
- Do not write on a Timertag Go label. No patient name, identifier, room number, or other patient information may be recorded on a label.
- Do not apply the label to skin, over broken skin, to dressings, to securement devices, or to fluid containers. Apply only as directed in Section 6.
- Apply the flag so that it cannot transmit traction to a catheter, a securement device, or an insertion site, and so that it cannot obstruct a clamp, port, or connector.
- Confirm that both faces of the folded flag scan before leaving the bedside.
- An internet connection is required at the point of care to open the guidance page.
- Specify non-metallic, non-foil label stock when ordering.

9 MATERIALS AND MRI NOTE

As supplied by Timertag, a Go label consists of printed artwork only and contains no electronic, metallic, or magnetic components.

Where printed on the non-metallic, non-foil label stock that the deploying organisation is required to specify (Sections 5 and 8), the label presents no known hazard in the MR environment. Responsibility for specifying compliant stock rests with the deploying organisation.

10 RELATIONSHIP TO TIMERTAG PRO

Timertag Go and Timertag Pro are different products and are not interchangeable.

TIMERTAG GO INSTRUCTIONS FOR USE

English (EN) · IFU-TTG-001 Rev B · Issued 15AUG26

Read this document before first use. Retain for reference. Free template · Single-patient use

- **Timertag Go** is a printed label only. It carries patient guidance and has no timing function. It is printed by the deploying organisation at 20 mm × 90 mm and applied folded on itself.
- **Timertag Pro** is supplied by the manufacturer, carries a passive NFC inlay, and provides a timing function. It is supplied at 20 mm × 40 mm and applied on its release liner.

Where both products are in use in the same facility, store them separately and confirm which product is in hand before application.

The Timertag Go guidance page is also reachable from a Timertag Pro status page, via the information button below the guidance images. No session data passes between the two.

11 DATA, PRIVACY, AND TERMS OF USE

No accounts, logins, or patient identifiers, and no facility for entering patient information. As with any web page, standard technical information such as IP address and browser type is processed to deliver the page.

Downloading and printing Timertag Go is subject to the Timertag Download and Usage Agreement (Version 2026.2), published at www.timertag.com. Free for use in patient care. Not for resale.

12 DISPOSAL

Dispose of used labels as general waste, or in accordance with facility clinical waste procedures where contaminated with blood or body fluids.

13 MANUFACTURER

Senver Group Pty Ltd (ABN 90 622 761 380), trading as Timertag

53/61–65 Glencoe Street, Sutherland NSW 2232, Australia

www.timertag.com

TIMERTAG®, Timertag Pro, and Timertag Go are trademarks of Senver Holdings Pty Ltd, used under license by Senver Group Pty Ltd. © 2026 Senver Group Pty Ltd. All rights reserved. IFU-TTG-001 Rev B, issued 2026-08. English (EN).