

Complete all sections. To avoid delay in processing, completion of fields with (\*) is required.

- ☐ Sign and date the form.
- ☐ Fax the completed form to Philips Remote INR Patient Services at 1-800-779-8560.
- ☐ Send signed and dated clinical documentation of the intent for ordering test materials and/or equipment for home INR monitoring of patient.
- ☐ If available, provide POA documentation.

1	Patient first name*	MI	Last name*	Legal gender _ M _ F	DOB (mm/dd/yyyy)*
	Home address*		City*	State*	Zip/postal code*
	Home phone #* 1-	Cell phone # 1-		Patient email (if available)	
MRN # (Medical record number, if applicable for EHR connectivity)					

### 2 Patient diagnosis code\*

Based on diagnosis of the patient's condition, enter all the applicable ICD-10 diagnosis codes. Below are commonly used ICD-10 diagnosis codes for patients who are monitoring PT/INR at home. This is not a complete list of possible codes. You may also enter separate code(s) in Other. For a full list of ICD-10 codes recognized by CMS, please visit <https://www.cms.gov>. All diagnosis codes must be provided in ICD-10 format.

- |   |  |
|---|--|
| <input type="checkbox"/> 279.01 - Long term (current) use of anticoagulants   | <input type="checkbox"/> Z95.2 - Presence of prosthetic heart valve                    |
| <input type="checkbox"/> I48.11 - Longstanding persistent atrial fibrillation | <input type="checkbox"/> I26.99 - Other pulmonary embolism without acute cor pulmonale |
| <input type="checkbox"/> I48.21 - Permanent atrial fibrillation               | <input type="checkbox"/> D68.59 - Other primary thrombophilia                          |
| <input type="checkbox"/> I48.0 - Paroxysmal atrial fibrillation               | <input type="checkbox"/> Other - _____   |

### 3 Therapeutic range

Low\*: \_\_\_\_\_

High\*: \_\_\_\_\_

### 4 Notification range

Do not call for any INR results.\*

\*This is default option if no option is selected.

Call if INR result is <1.5 and >5.0

Call for customized notification range if result is below \_\_\_\_\_ or above \_\_\_\_\_

Notification below cannot be greater than therapeutic low. Notification above cannot be greater than therapeutic high.

### 5 Prescribed frequency

Tests per month (select one)\*

Frequencies outside of this range are not offered by Philips:

**Weekly**      **2-4 Times per month**  
(flexibility to test weekly or bi-weekly)

### 6 Patient results contact (All results will be faxed and notification calls will be made per your choice in section 4)

Phone (Out of range)* 1-	FAX (All results)* 1-	Medical Records Phone 1-	Medical Records Fax 1-
Practice/Clinic name			
Clinic street address*	Suite #	Clinic city*	Clinic State*    Clinic zip*

### 7 Patient training face-to-face training is required\*

**Already Trained** – Physician certifies patient was face-to-face trained on the Philips Remote INR PT/INR monitoring system

**Philips Train** – Philips will attempt to train patient

**Clinic Train** – Training contract with Philips must be in place

- Notes:**
- If patient has previously been on service with Philips Remote INR, and it has been documented that a training has already taken place, we reserve the right to bypass face to face training.
  - Philips Remote INR Patient Services will attempt to train your patient unless one of the other options above is selected (training is not guaranteed).

### 8 Physician authorization (Signature and date must be hand-written or e-signed)

This form serves as a Physician's Order for the Philips Remote INR PT/INR monitoring system for Patient Self-Testing and related supplies. I certify that this patient has been on oral warfarin therapy for more than 3 months and is a suitable candidate for self-testing. At this time, the patient or his/her caregiver has no condition that makes self-testing unsafe (e.g., cognitive and/or physical disorders). I agree to notify Philips Remote INR if self-testing is no longer prescribed for this patient. I acknowledge that this product is intended for patients 22 years of age and older, and use outside of the intended population is off-label use. Evaluation of risk of use on patients under 22 is determined by the device manufacturer. Use outside of the intended population is not endorsed by Philips.

<b>Sign &amp; date</b>	Prescribing physician signature*	Date (mm/dd/yyyy)*	Physician NPI*
	Prescribing physician printed*		

### 9 Insurance information To expedite patient enrollment please include a copy of front and back of patient's insurance card

<b>Primary health insurance information</b>	Insurance company	Policy ID#	Customer service phone # 1-
<b>Secondary health insurance information</b>	Insurance company	Policy ID#	Customer service phone # 1-

Please fax completed form to the central office **Phone: 1-800-780-0675 Fax: 1-800-779-8560 Alt Fax: 1-877-740-1831**