Before the

Federal Communications Commission

Washington, DC 20554

In the Matter of Experimental Radio License Grant to Biotronik, Inc.	File No. 0223-EX-PL-2002

EXHIBIT A

TO

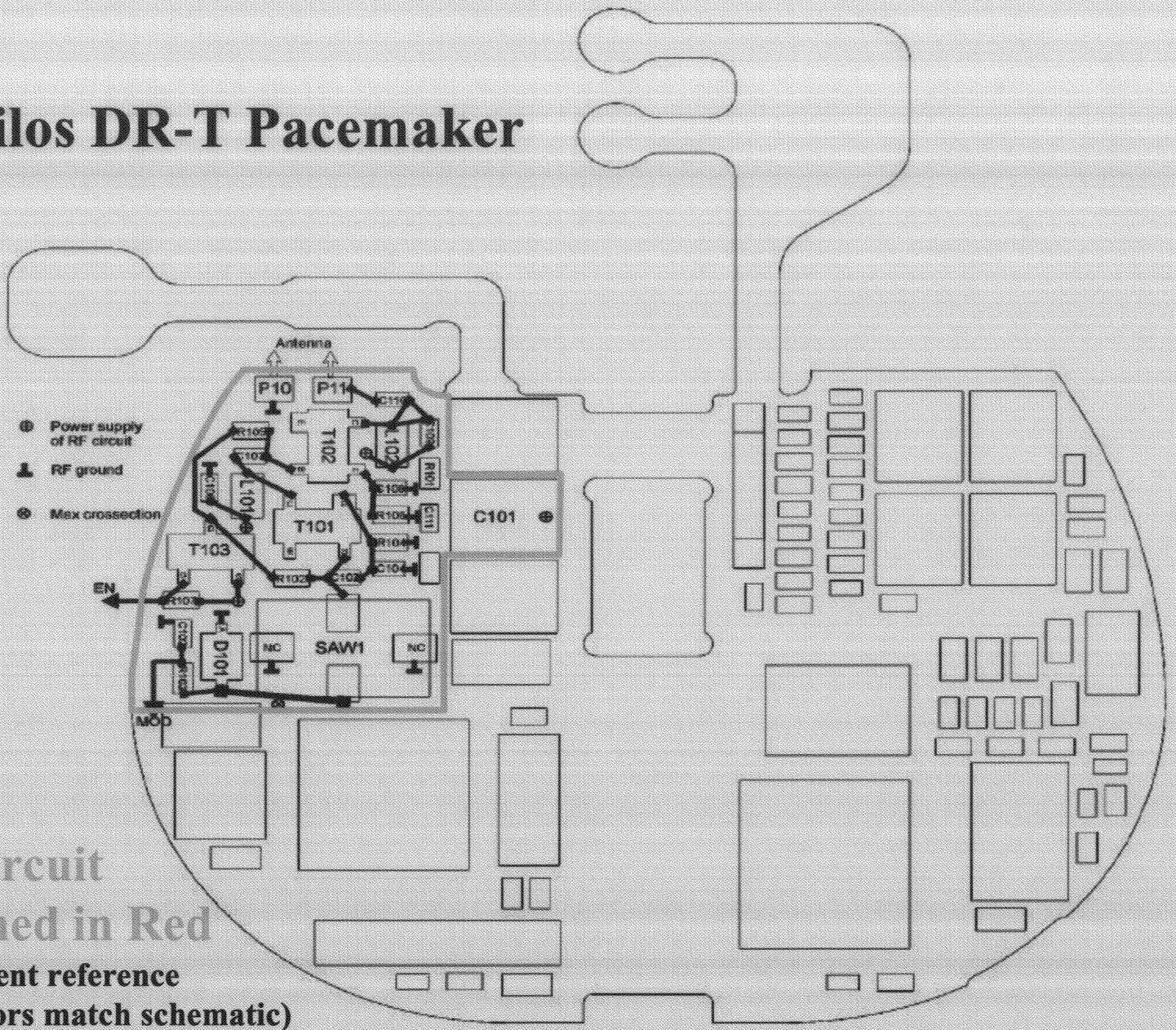
MEDTRONIC'S OPPOSITION TO MOTION FOR LEAVE

TO FILE A RESPONSE TO REPLY

EXHIBIT H – Internal Photos

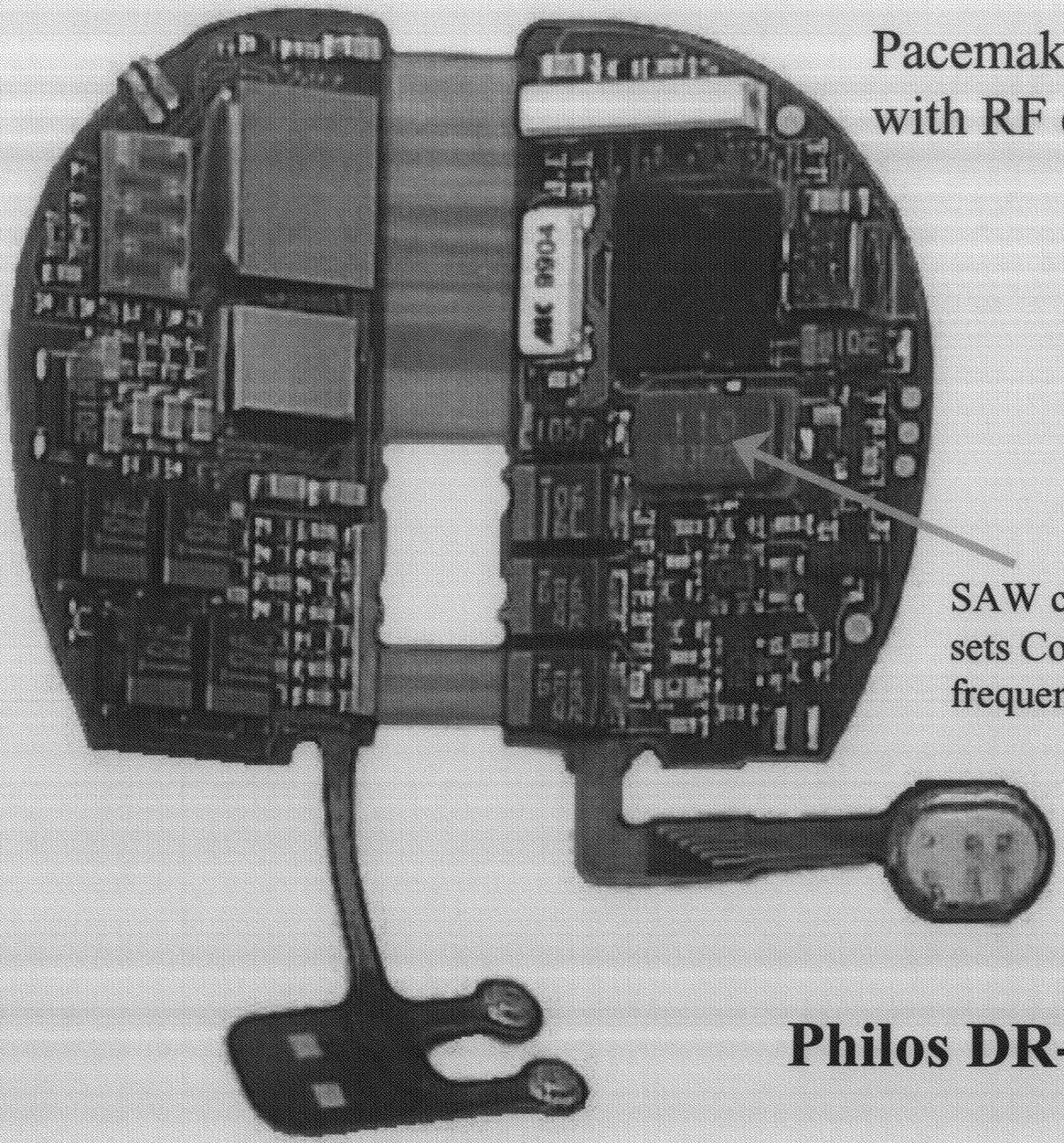
FCC ID# PG6BA0T

Philos DR-T Pacemaker



RF Circuit Outlined in Red

(component reference designators match schematic)



Pacemaker electronics
with RF circuitry

SAW component
sets Colpitts oscillator
frequency at 403.55MHz

Philos DR-T

Folded Pacemaker Circuit with
flex arms for attachment to battery
and feedthrough

Orange coil is part of
magnetically coupled
communications system
(not part of RF circuit).

Backside of Electronics Circuit Board

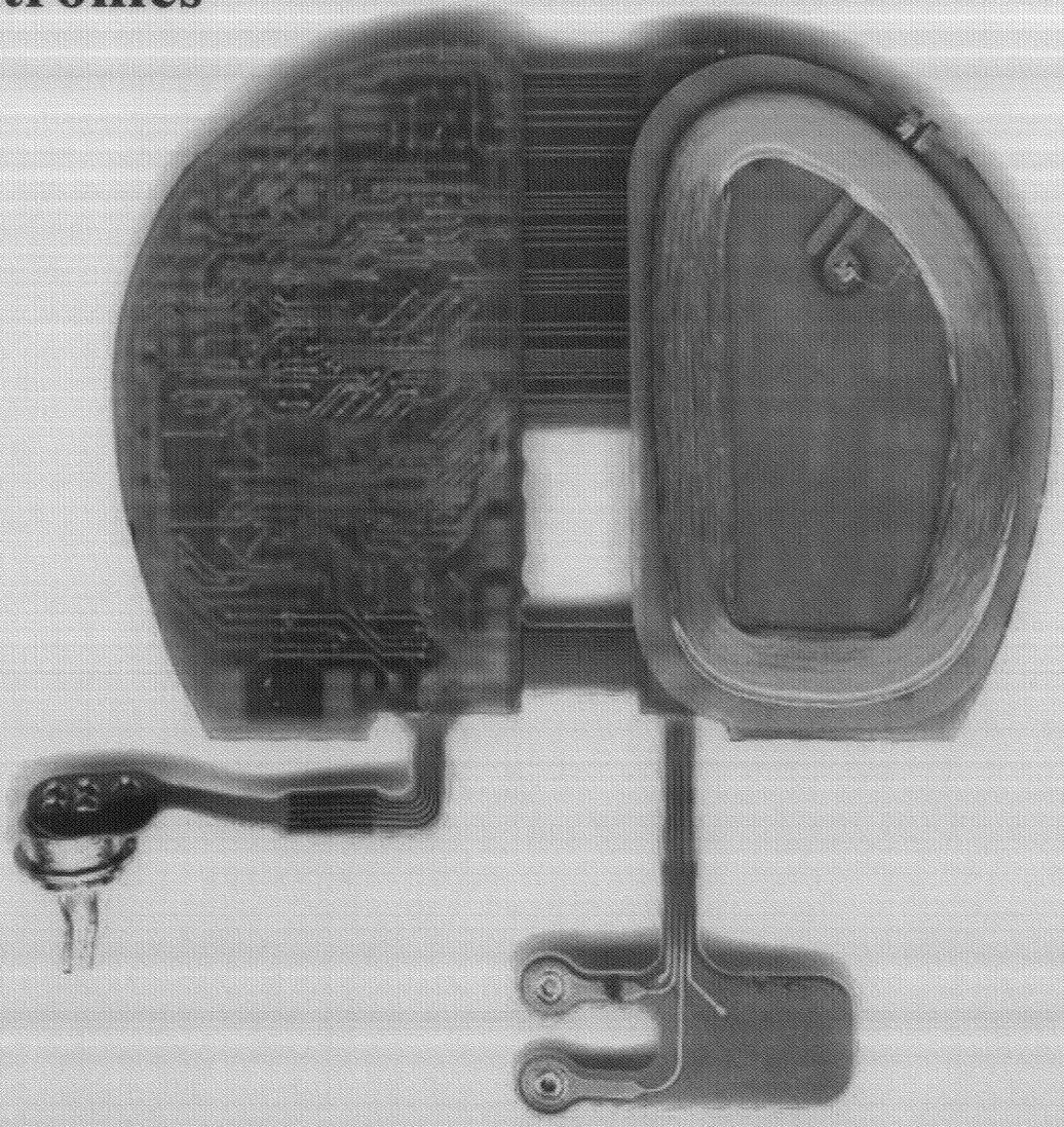


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EXHIBIT B

TO

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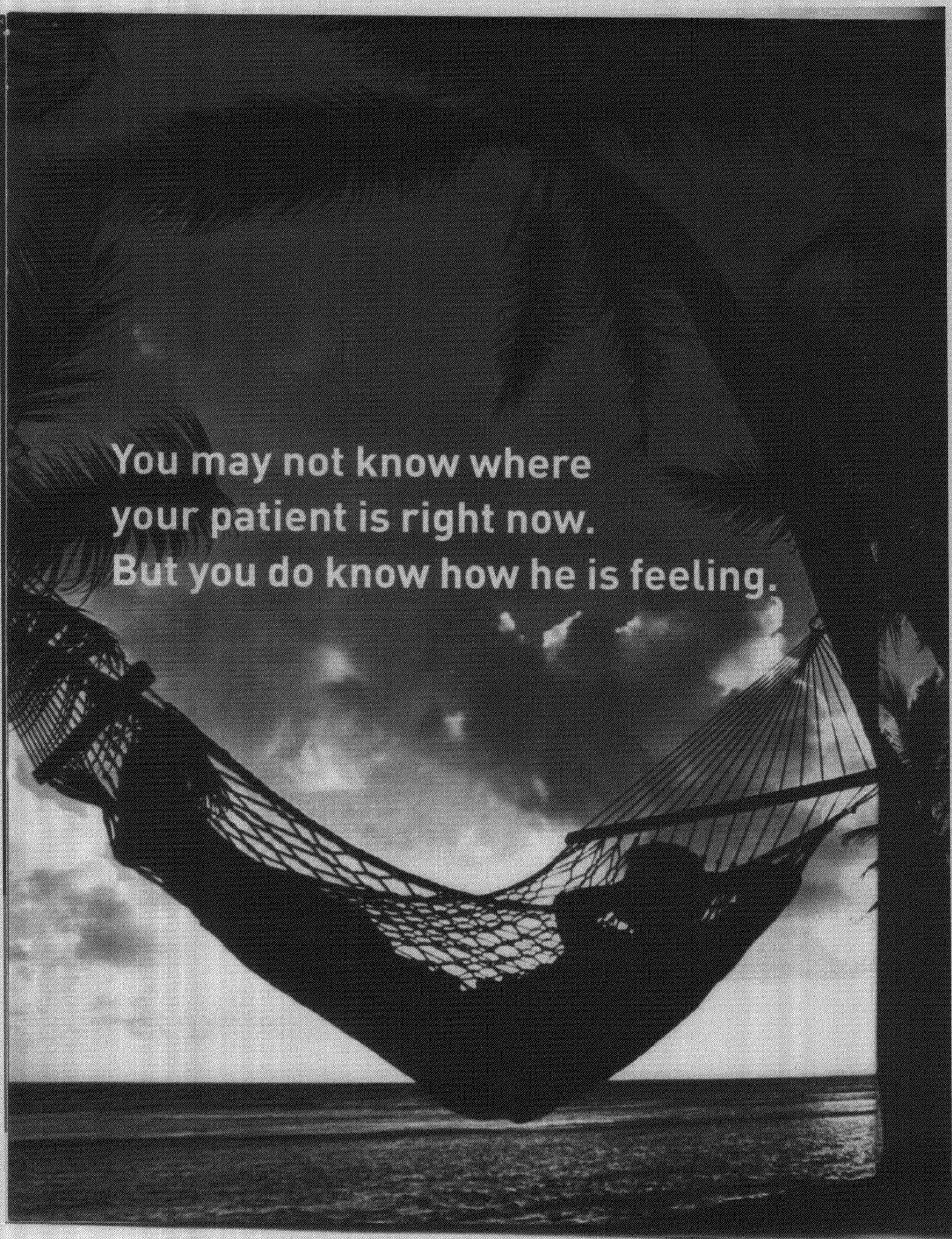
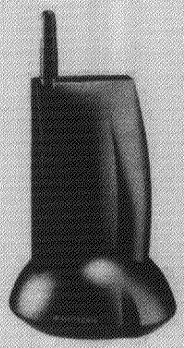

You may not know where
your patient is right now.
But you do know how he is feeling.

BIOTRONIK implants send messages straight from the heart. Conveniently. Automatically. Safely. With no patient intervention whatsoever – day and night. And your patient maintains complete mobility. A Cardio Report with all current, comprehensive data comes to you following an event or at any specific time or frequency that you decide.



BIOTRONIK's innovative Home Monitoring System:
A revolutionary approach to patient management.

www.biotronik.com

BIOTRONIK

Indications and Contraindications

BIOTRONIK's Home Monitoring system is currently available in the United States for both pacemakers and Implantable Cardioverter Defibrillators (ICDs). The approved indications and contraindications for the pacemakers and ICDs are identical regardless if the Home Monitoring functionality is available.

Warnings and Precautions

Certain therapeutic and diagnostic procedures may cause undetected damage to pacemakers or ICDs, resulting in malfunction or failure at a later time. The approved warnings and precautions for the pacemakers and ICDs are identical regardless if the Home Monitoring functionality is available. However there are specific precautions that are applicable to the Home Monitoring system. Please note the following Home Monitoring specific warnings and precautions:

- **Patient's Ability** – Use of the Home Monitoring system requires the patient and/or caregiver to follow the system instructions and cooperate fully when transmitting data. If the patient cannot understand or follow the instructions because of physical or mental challenges, another adult who can follow the instructions will be necessary for proper transmission.
- **Cellular Phone Availability** – The Home Monitoring system is not practical for patients who live in areas where cellular telephone networks, utilizing the GSM standard, are not available or are not likely to become available in the near future.
- **Use in Cellular Phone Restricted Areas** – The mobile patient device (transmitter/receiver) should not be utilized in areas where cellular phones are restricted or prohibited (i.e., commercial aircraft).
- **Event Triggered Report** – A timely receipt of the event report cannot be guaranteed. The receipt is also dependent on whether the patient was physically situated in the required coverage range of the patient device at the time the event information was sent.
- **Patient-Activated Report** – The magnet effect must be programmed "synchronous" if the [Patient Report] function is activated.
- **Not for Diagnosis** – The data transmitted by Home Monitoring are not suitable for diagnosis, because not all information available in the implant is being transmitted.
- **Follow-Ups** – When using Home Monitoring, the time period between follow-up visits may not be extended.

Potential Adverse Events

The approved labeling with potential adverse events for the pacemakers and ICDs are identical regardless if the Home Monitoring functionality is available. Home Monitoring does not introduce additional potential adverse events per approved labeling.

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