



Complete this form:

- At baseline, for the qualifying episode of atrial fibrillation, prior to randomization.
- At baseline, after conversion to normal sinus rhythm, for patients in rhythm control arm.
- At baseline, after completion of drug titration.
- At 4 month follow-up.
- At 1 year follow-up.

1. Date of recording:

		/			/		
Month			Day			Year	

Days03

Affix Patient ID # Here

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Print Acrostic Here

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2. Reason for recording:

- ¹ Baseline, during qualifying episode of atrial fibrillation
- ² Baseline, after conversion to sinus rhythm ⇒ Specify: *(Mark one only.)*
 - ¹ Before randomization
 - ² After randomization
- ³ Baseline, after completion of drug titration
- ⁴ 4 month follow-up
- ⁵ 1 year follow-up

BARand03

Reason03

3. Source of tracing: (Mark one only.)

- Source03* ¹ 12-lead ECG ² Rhythm strip

4. Rhythm(s) at time of recording:

No ⁰ Yes ¹

- Sinus03* Sinus rhythm (normal, tachycardia, or bradycardia)
- AF03* Atrial fibrillation
- Atrial flutter
- Paced03* Paced ⇒ Atrial pacing only
- Ventricular pacing only
- Dual chamber pacing

Rhyth003 Other ⇒ Specify:

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For CTC use only:

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Electrocardiogram

Fax to: (800) 547-0463

Date of recording:

<input type="text"/>	/	<input type="text"/>	/	<input type="text"/>
Month		Day		Year

Affix Patient ID # Here

<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	-	<input type="text"/>	<input type="text"/>	<input type="text"/>	-	<input type="text"/>
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Specify reason for recording:

<input type="radio"/> Baseline, during atrial fibrillation <input type="radio"/> Baseline, after conversion to sinus rhythm <input type="radio"/> Baseline, after completion of drug titration	<input type="radio"/> 4 month follow-up <input type="radio"/> 1 year follow-up
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5. Heart rate: *Rate03* beats/min

6. Intervals:

PR interval *PRInt03* msec PR not measurable

QRS duration *QRS03* msec

QT interval *QT03* msec (Do not record QTc)

7. Drugs taken by the patient at the time of the ECG:

No	Yes	No	Yes			
0	1	0	1			
<i>Amiod03</i>	<input type="radio"/>	<input type="radio"/>	Amiodarone	<input type="radio"/>	<input type="radio"/>	Moricizine
<i>BetaB103</i>	<input type="radio"/>	<input type="radio"/>	Beta blocker	<input type="radio"/>	<input type="radio"/>	Procainamide
<i>Digox03</i>	<input type="radio"/>	<input type="radio"/>	Digoxin	<input type="radio"/>	<input type="radio"/>	Propafenone
<i>Dilt03</i>	<input type="radio"/>	<input type="radio"/>	Diltiazem	<input type="radio"/>	<input type="radio"/>	Quinidine
	<input type="radio"/>	<input type="radio"/>	Disopyramide	<i>Sotal03</i>	<input type="radio"/>	Sotalol
	<input type="radio"/>	<input type="radio"/>	Flecainide	<i>Verap03</i>	<input type="radio"/>	Verapamil
	<input type="radio"/>	<input type="radio"/>	Ibutilide	<i>CalcB103</i>	<input type="radio"/>	Calcium channel blocker (other than diltiazem and verapamil)

Class I - Class I Drug (Disopyramide or Flecainide or Moricizine or Procainamide or Propafenone or Quinidine)

Name of person completing this form _____

Date _____

Please print

mm/dd/yy

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