



Urgent:

Medical device field action

HeartSine® samaritan® PAD SAM 350P/SAM 360P/SAM 450P/SAM 500P

Customer name: <<merge file>>

Customer #: <<merge file>>

Attn: Safety Manager

Field action number:

PR3977961 – FA318

PR4280335 – FA319

PR4257283 – FA334

<Month DD, YYYY/ DD Month YYYY>



Model	Serial Numbers	UDI-DI
350P	A subset of serial numbers has been affected by this notice. Please check if your AED serial number is affected by visiting the HeartSine PFA Global Landing Page below: https://heartsinerecall.com/HeartSineSamaritanPAD/LookUp If your device is listed, please follow the instructions provided in this letter.	08858251002132S4
360P		
500P		

Stryker has initiated a serial-specific Urgent Recall and a Safety Notice for the devices listed in the table above. This device recall notification is being issued to alert customers with HeartSine® samaritan® PAD 350P/360P/500P devices of two potential device function problems (1) and a missing maintenance supplement (2) for a limited number of devices. Out of an abundance of caution, Stryker is issuing this notice.

Product description The HeartSine samaritan PAD is a small, lightweight, portable, battery operated Automated External Defibrillator (AED) designed to treat victims of cardiac arrest.

Product issue (1)(PR3977961 – FA318) Stryker informed customers in May 2025 of a manufacturing process **issue related to a circuit board component** that **may** impair the device's ability to function and/or cause failure. This issue was identified during extensive quality testing. We thank you for taking the time to respond to this field action.

(1)(PR4280335 – FA319) Since your response, Stryker has recently identified an **issue with the membrane tail**, an internal component, which could cause an intermittent fault. This failure could occur at any point when the device is holding a charge in preparation to deliver therapy, while delivering a shock, or after shock delivery. The device becomes inoperable after the failure occurs.



(2)(PR4257283 – FA334) Stryker has also identified that a limited number of HeartSine samaritan PAD devices were **distributed without the required maintenance supplement**. This supplement contains periodic maintenance instructions implemented as an additional risk-control measure. While most devices contain the maintenance supplement, this voluntary correction is being initiated to ensure all end users have access to the supplemental maintenance information. This issue was identified through a customer complaint.

Potential risks

(1) (PR3977961 – FA318 & PR4280335 – FA319) Due to internal component issues, the device **may** fail to deliver the intended therapy during use, potentially leading to a delay in treatment or no treatment being delivered during use. **These issues were observed during non-standard rework process and quality testing, not during patient use.**

(2) (PR4257283 - FA334) The omission of this supplement may result in users not performing the recommended quarterly power-cycle check. If the check is not performed, the device may have an undetected audio-prompt failure that could result in reduced device readiness and user confusion. This issue was identified through a customer complaint.

There have been no adverse events reported related to this device recall notification.

If your device experiences any of the above issues during use, please seek an alternative defibrillator and contact **your Authorized Distributor or HeartSine Technologies Technical Support at heartsinesupport@stryker.com.**

Customer actions needed:

1. **Identify impacted devices** by verifying if your device serial numbers are included in the list at the link below. For this field action notice, please select “AED + Pad-Pak”.

<https://heartsinerecall.com/HeartSineSamaritanPAD/LookUp>

A. For devices available in your inventory & Stryker direct customers/distributors:

- a. Instructions for where to locate device serial numbers on the device are found in Appendix A*.
- b. If you have more than 15 devices and would like a list of impacted serial numbers, please email **<Insert Local Email>@stryker.com:**
 - Subject: FA318/FA319/FA334 - Impacted device list - <<your company name>>
 - Be sure to include your name and phone number in case we need to reach out to you.

B. For devices you have transferred to other users (Non-Stryker direct customers/distributors):

- a. Instructions for where to locate device serial numbers on the device are found in Appendix A*.
- b. If you have further distributed any affected product, please provide this notification to the relevant recipients. **Recipients who are not direct Stryker customers should respond to the facility or organization from which they receive this notification and **not directly to Stryker.***
- c. **Email your Stryker representative at xxxxx@stryker.com when you have collected all the responses from your end-users.**

2. **Download** a copy of the missing maintenance supplement when acknowledging the recall and store with your device's user manual until a replacement is received.



3. **Submit your response/acknowledgement** to Stryker electronically for all field actions via this link: <https://heartsinerecall.com/HeartSineSamaritanPAD/LookUp> and/or email xxxxxx@stryker.com the enclosed Business Reply Form by **<Insert Date>**.

- a. We appreciate your previous response to the FA318 field action and your replacement device will be prioritized.

Commented [DA1]: [Name, Name] should this say "previous response"?

4. Until a replacement is available, Stryker recommends keeping your HeartSine samaritan PAD in service if you do not have an alternative public access defibrillator. This recommendation is based on internal testing demonstrating a low probability of failure due to these product manufacturing issues.

5. Maintain awareness of this communication internally and near the affected unit until all required actions have been completed within your facility and the unit has been replaced.

Stryker's planned actions:

Stryker is notifying all customers who have received affected HeartSine samaritan PAD devices to perform the actions outlined above. Once your response is received and a replacement is available, Stryker will be in contact as soon as possible to arrange the next steps for a replacement device.

If you have any questions or concerns, please contact Stryker Customer Service at **<Insert Info Here>**, your Authorized Distributor, **or HeartSine Technologies Technical Support at heartsinesupport@stryker.com.**

In line with the recommendations of the Medical Device Coordination Group Guidance document Ref MDCCG 2023-3 and EU MDR 2017/745, we can confirm that this FSQA has been notified appropriately to the National Competent Authority for your country.

On behalf of Stryker, we thank you sincerely for your help and support in submitting your additional response by **MMM DD YYYY**. We regret any inconvenience that may be caused and would like to reassure you that we are committed to meeting our high internal quality standards and your expectations.

Appendix A

HeartSine samaritan PAD SAM 350P/SAM 360P/SAM 450P/SAM 500P

Instructions to Identify Impacted Devices

- 1) To find your device serial number and model number, see the labels on the rear of your device as shown below:

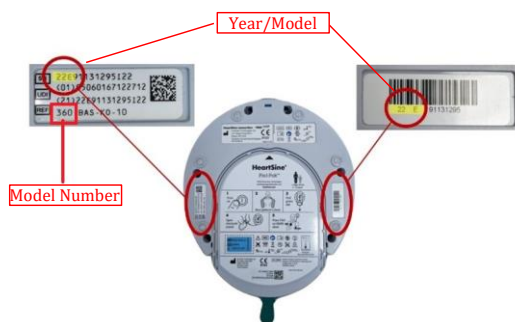


Figure 1 – Serial & Model Number location

Urgent:

Medical device field action

HeartSine® samaritan® PAD

SAM 350P/SAM 360P/SAM 450P/SAM 500P

Customer name: <<merge file>>

Customer #: <<merge file>>

Attn: Safety Manager

Field action number:

PR3977961 – FA318

PR4280335 – FA319

PR4257283 – FA334

<Month DD, YYYY/ DD Month YYYY>



Model	Serial Numbers	UDI-DI
350P	A subset of serial numbers has been affected by this notice. Please check if your AED serial number is affected by visiting the HeartSine PFA Global Landing Page below: https://heartsinerecall.com/HeartSineSamaritanPAD/LookUp If your device is listed, please follow the instructions provided in this letter.	08858251002132S4
360P		
500P		

Stryker has initiated a serial-specific Urgent Recall and a Safety Notice for the devices listed in the table above. This device recall notification is being issued to alert customers with HeartSine® samaritan® PAD 350P/360P/500P devices of two potential device function problems (1) and a missing maintenance supplement (2) for a limited number of devices. Out of an abundance of caution, Stryker is issuing this notice.

Product description The HeartSine samaritan PAD is a small, lightweight, portable, battery operated Automated External Defibrillator (AED) designed to treat victims of cardiac arrest.

Product issue (1)(PR3977961 – FA318) Stryker informed customers in May 2025 of a manufacturing process **issue related to a circuit board component** that **may** impair the device's ability to function and/or cause failure. This issue was identified during extensive quality testing.

(1)(PR4280335 – FA319) In addition, Stryker has recently identified an **issue with the membrane tail**, an internal component, which could cause an intermittent fault. This failure could occur at any point when the device is holding a charge in preparation to deliver therapy, while delivering a shock, or after shock delivery. The device becomes inoperable after the failure occurs.

(2)(PR4257283 – FA334) Stryker has also identified that a limited number of HeartSine samaritan PAD devices were **distributed without the required**

maintenance supplement. This supplement contains periodic maintenance instructions implemented as an additional risk-control measure. While most devices contain the maintenance supplement, this voluntary correction is being initiated to ensure all end users have access to the supplemental maintenance information. This issue was identified through a customer complaint.

Potential risks

(1) (PR3977961 – FA318 & PR4280335 – FA319) Due to internal component issues, the device **may** fail to deliver the intended therapy during use, potentially leading to a delay in treatment or no treatment being delivered during use. **These issues were observed during non-standard rework process and quality testing, not during patient use.**

(2) (PR4257283 - FA334) The omission of this supplement may result in users not performing the recommended quarterly power-cycle check. If the check is not performed, the device may have an undetected audio-prompt failure that could result in reduced device readiness and user confusion. This issue was identified through a customer complaint.

There have been no adverse events reported related to this device recall notification.

If your device experiences any of the above issues during use, please seek an alternative defibrillator and contact **your Authorized Distributor or HeartSine Technologies Technical Support at heartsinesupport@stryker.com.**

Customer actions needed:

1. **Identify impacted devices** by verifying if your device serial numbers are included in the list at the link below. For this field action notice, please select “AED + Pad-Pak”.

<https://heartsinerecall.com/HeartSineSamaritanPAD/LookUp>

A. For devices available in your inventory & Stryker direct customers/distributors:

- a. Instructions for where to locate device serial numbers on the device are found in Appendix A*.
- b. If you have more than 15 devices and would like a list of impacted serial numbers, please email **<Insert Local Email>@stryker.com:**
 - Subject: FA318/FA319/FA334 - Impacted device list - <<your company name>>
 - Be sure to include your name and phone number in case we need to reach out to you.

B. For devices you have transferred to other users (Non-Stryker direct customers/distributors):

- a. Instructions for where to locate device serial numbers on the device are found in Appendix A*.
- b. If you have further distributed any affected product, please provide this notification to the relevant recipients. **Recipients who are not direct Stryker customers should respond to the facility or organization from which they receive this notification and **not directly to Stryker.***
- c. **Email your Stryker representative at xxxxx@stryker.com when you have collected all the responses from your end-users.**

2. **Download** a copy of the missing maintenance supplement when acknowledging the recall and store with your device’s user manual until a replacement is received.
3. **Submit your response/acknowledgement** to Stryker electronically for all field actions via this link: <https://heartsinerecall.com/HeartSineSamaritanPAD/LookUp> and/or email xxxxxx@stryker.com with the enclosed Business Reply Form by **<Insert Date>.**

4. Until a replacement is available, Stryker recommends keeping your HeartSine samaritan PAD in service if you do not have an alternative public access defibrillator. This recommendation is based on internal testing demonstrating a low probability of failure due to these product manufacturing issues.
5. Maintain awareness of this communication internally and near the affected unit until all required actions have been completed within your facility and the unit has been replaced.

Stryker's planned actions:

Stryker is notifying all customers who have received affected HeartSine samaritan PAD devices to perform the actions outlined above. Once your response is received and a replacement is available, Stryker will be in contact as soon as possible to arrange the next steps for a replacement device.

In line with the recommendations of the Medical Device Coordination Group Guidance document Ref MDCG 2023-3 and EU MDR 2017/745, we can confirm that this FSCA has been notified appropriately to the National Competent Authority for your country.

If you have any questions or concerns, please contact Stryker Customer Service at <Insert Info Here>, your Authorized Distributor, or HeartSine Technologies Technical Support at heartsinesupport@stryker.com.

On behalf of Stryker, we thank you sincerely for your help and support in submitting your response by MMM DD YYYY. We regret any inconvenience that may be caused and would like to reassure you that we are committed to meeting our high internal quality standards and your expectations.

Appendix A

HeartSine samaritan PAD SAM 350P/SAM 360P/SAM 450P/SAM 500P

Instructions to Identify Impacted Devices

- 1) To find your device serial number and model number, see the labels on the rear of your device as shown below:

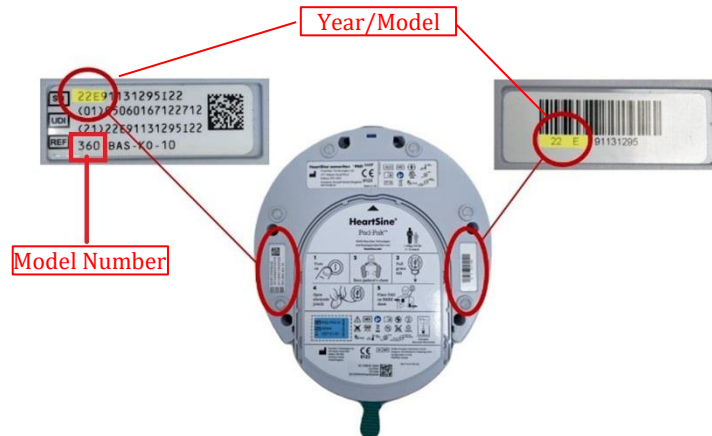


Figure 1 – Serial & Model Number location

stryker

Urgent:

Medical device field action

HeartSine® samaritan® PAD
SAM 350P/SAM 360P/SAM 450P/SAM 500P

Customer name: <<merge file>>

Customer #: <<merge file>>

Attn: Safety Manager

Field action number:

PR3977961 – FA318

PR4257283 – FA334

<Month DD, YYYY/ DD Month YYYY>



Model	Serial Numbers	UDI-DI
350P	A subset of serial numbers has been affected by this notice. Please check if your AED serial number is affected by visiting the HeartSine PFA Global Landing Page below: https://heartsinerecall.com/HeartSineSamaritanPAD/LookUp If your device is listed, please follow the instructions provided in this letter.	08858251002132S4
360P		
500P		

Stryker has initiated a serial-specific Urgent Recall and a Safety Notice for the devices listed in the table above. This device recall notification is being issued to alert customers with HeartSine® samaritan® PAD 350P/360P/500P devices of a potential device malfunction (1) and a missing maintenance supplement (2) for a limited number of devices. Out of an abundance of caution, Stryker is issuing this notice.

Product description The HeartSine samaritan PAD is a small, lightweight, portable, battery operated Automated External Defibrillator (AED) designed to treat victims of cardiac arrest.

Product issue (1)(PR3977961 – FA318) Stryker informed customers in May 2025 of a manufacturing process **issue related to a circuit board component** that may impair the device's ability to function and/or cause failure. **This issue was identified during extensive quality testing.**
We thank you for taking the time to respond to this field action.

(2)(PR4257283 – FA334) Since your response, Stryker has identified that a limited number of HeartSine samaritan PAD devices were **distributed without the required maintenance supplement**. This supplement contains periodic maintenance instructions implemented as an additional risk-control measure. While most devices contain the maintenance supplement, this voluntary correction is being initiated to ensure all end users have access to the



supplemental maintenance information. This issue was identified through a customer complaint.

Potential risks

(1)(PR3977961 – FA318) Due to internal component issues, the device **may** fail to deliver the intended therapy during use, potentially leading to a delay in treatment or no treatment being delivered during use. These issues were observed during non-standard rework process and quality testing, not during patient use.

(2)(PR4257283 – FA334) The omission of this supplement may result in users not performing the recommended quarterly power-cycle check. If the check is not performed, the device may have an undetected audio-prompt failure that could result in reduced device readiness and user confusion. This issue was identified through a customer complaint.

There have been no adverse events reported related to this device recall notification.

If your device experiences any of the above issues during use, please seek an alternative defibrillator and contact your **Authorized Distributor or HeartSine Technologies Technical Support at heartsinesupport@stryker.com**.

Customer actions needed:

1. **Identify impacted devices** by verifying if your device serial numbers are included in the list at the link below. For this field action notice, please select “AED + Pad-Pak”.

<https://heartsinerecall.com/HeartSineSamaritanPAD/LookUp>

A. For devices available in your inventory & Stryker direct customers/distributors:

- a. Instructions for where to locate device serial numbers on the device are found in Appendix A*.
- b. If you have more than 15 devices and would like a list of impacted serial numbers, please email **<Insert Local Email>@stryker.com**:
 - Subject: FA318/FA334 - Impacted device list - <<your company name>>
 - Be sure to include your name and phone number in case we need to reach out to you.

B. For devices you have transferred to other users (Non-Stryker direct customers/distributors):

- a. Instructions for where to locate device serial numbers on the device are found in Appendix A*.
- b. If you have further distributed any affected product, please provide this notification to the relevant recipients. **Recipients who are not direct Stryker customers should respond to the facility or organization from which they receive this notification and **not directly to Stryker**.*

- c. **Email your Stryker representative at xxxxx@stryker.com when you have collected all the responses from your end-users.**

2. **Download** a copy of the missing maintenance supplement when acknowledging the recall and store with your device’s user manual until a replacement is received.

3. **Submit your response/acknowledgement** to Stryker electronically for the **additional** field actions via this link: <https://heartsinerecall.com/HeartSineSamaritanPAD/LookUp> **and/or** email **xxxxxx@Stryker.com** the enclosed Business Reply Form by **<Insert Date>**.

- a. We appreciate your previous **response** to the FA318 field action and your replacement device will be prioritized.



4. Until a replacement is available, Stryker recommends keeping your HeartSine samaritan PAD in service if you do not have an alternative public access defibrillator. This recommendation is based on internal testing demonstrating a low probability of failure due to these product manufacturing issues.
5. Maintain awareness of this communication internally and near the affected unit until all required actions have been completed within your facility and the unit has been replaced.

Stryker's planned actions:

Stryker is notifying all customers who have received affected HeartSine samaritan PAD devices to perform the actions outlined above. Once your response is received and a replacement is available, Stryker will be in contact as soon as possible to arrange the next steps for a replacement device.

If you have any questions or concerns, please contact Stryker Customer Service at <Insert Info Here>, your Authorized Distributor, or HeartSine Technologies Technical Support at heartsinesupport@stryker.com.

In line with the recommendations of the Medical Device Coordination Group Guidance document Ref MDCG 2023-3 and EU MDR 2017/745, we can confirm that this FSCA has been notified appropriately to the National Competent Authority for your country.

On behalf of Stryker, we thank you sincerely for your help and support in submitting your additional response by MMM DD YYYY. We regret any inconvenience that may be caused and would like to reassure you that we are committed to meeting our high internal quality standards and your expectations.

Appendix A

HeartSine samaritan PAD SAM 350P/SAM 360P/SAM 450P/SAM 500P

Instructions to Identify Impacted Devices

- 1) To find your device serial number and model number, see the labels on the rear of your device as shown below:

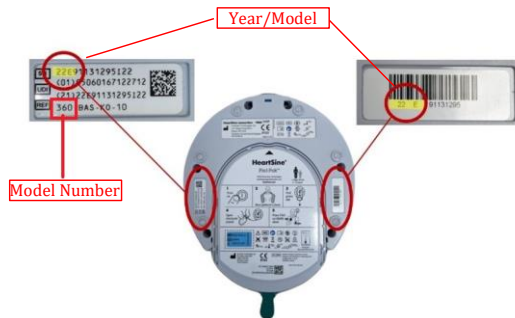


Figure 1 – Serial & Model Number location

Urgent:

Medical device field action

HeartSine® samaritan® PAD

SAM 350P/SAM 360P/SAM 450P/SAM 500P

Customer name: <<merge file>>

Customer #: <<merge file>>

Attn: Safety Manager

Field action number:

PR3977961 – FA318

PR4257283 – FA334

<Month DD, YYYY/ DD Month YYYY>



Model	Serial Numbers	UDI-DI
350P	A subset of serial numbers has been affected by this notice. Please check if your AED serial number is affected by visiting the HeartSine PFA Global Landing Page below: https://heartsinerecall.com/HeartSineSamaritanPAD/LookUp If your device is listed, please follow the instructions provided in this letter.	08858251002132S4
360P		
500P		

Stryker has initiated a serial-specific Urgent Recall and a Safety Notice for the devices listed in the table above. This device recall notification is being issued to alert customers with HeartSine® samaritan® PAD 350P/360P/500P devices of a potential device malfunction (1) and a missing maintenance supplement (2) for a limited number of devices. Out of an abundance of caution, Stryker is issuing this notice.

Product description The HeartSine samaritan PAD is a small, lightweight, portable, battery operated Automated External Defibrillator (AED) designed to treat victims of cardiac arrest.

Product issue (1)(PR3977961 – FA318) Stryker informed customers in May 2025 of a manufacturing process **issue related to a circuit board component** that **may** impair the device’s ability to function and/or cause failure. This issue was identified during extensive quality testing.

(2)(PR4257283 – FA334) Stryker has also identified that a limited number of HeartSine samaritan PAD devices were **distributed without the required maintenance supplement**. This supplement contains periodic maintenance instructions implemented as an additional risk-control measure. While most devices contain the maintenance supplement, this voluntary correction is being

initiated to ensure all end users have access to the supplemental maintenance information. This issue was identified through a customer complaint.

Potential risks

(1)(PR3977961 – FA318) Due to internal component issues, the device **may** fail to deliver the intended therapy during use, potentially leading to a delay in treatment or no treatment being delivered during use. **These issues were observed during non-standard rework process and quality testing, not during patient use.**

(2)(PR4257283 – FA334) The omission of this supplement may result in users not performing the recommended quarterly power-cycle check. If the check is not performed, the device may have an undetected audio-prompt failure that could result in reduced device readiness and user confusion. This issue was identified through a customer complaint.

There have been no adverse events reported related to this device recall notification.

If your device experiences any of the above issues during use, please seek an alternative defibrillator and contact your **Authorized Distributor or HeartSine Technologies Technical Support at heartsinesupport@stryker.com.**

Customer actions needed:

1. **Identify impacted devices** by verifying if your device serial numbers are included in the list at the link below. For this field action notice, please select “AED + Pad-Pak”.

<https://heartsinerecall.com/HeartSineSamaritanPAD/LookUp>

A. For devices available in your inventory & Stryker direct customers/distributors:

- a. Instructions for where to locate device serial numbers on the device are found in Appendix A*.
- b. If you have more than 15 devices and would like a list of impacted serial numbers, please email **<Insert Local Email>@stryker.com:**
 - Subject: FA318/FA334 - Impacted device list - <<your company name>>
 - Be sure to include your name and phone number in case we need to reach out to you.

B. For devices you have transferred to other users (Non-Stryker direct customers/distributors):

- a. Instructions for where to locate device serial numbers on the device are found in Appendix A*.
- b. If you have further distributed any affected product, please provide this notification to the relevant recipients. **Recipients who are not direct Stryker customers should respond to the facility or organization from which they receive this notification and **not directly to Stryker.***
- c. **Email your Stryker representative at xxxxx@stryker.com when you have collected all the responses from your end-users.**

2. **Download** a copy of the missing maintenance supplement when acknowledging the recall and store with your device’s user manual until a replacement is received.
3. **Submit your response/acknowledgement** to Stryker electronically for all field actions via this link: <https://heartsinerecall.com/HeartSineSamaritanPAD/LookUp> and/or email xxxxxx@Stryker.com the enclosed Business Reply Form by **<Insert Date>.**

4. Until a replacement is available, Stryker recommends keeping your HeartSine samaritan PAD in service if you do not have an alternative public access defibrillator. This recommendation is based on internal testing demonstrating a low probability of failure due to these product manufacturing issues.
5. Maintain awareness of this communication internally and near the affected unit until all required actions have been completed within your facility and the unit has been replaced.

Stryker's planned actions:

Stryker is notifying all customers who have received affected HeartSine samaritan PAD devices to perform the actions outlined above. Once your response is received and a replacement is available, Stryker will be in contact as soon as possible to arrange the next steps for a replacement device.

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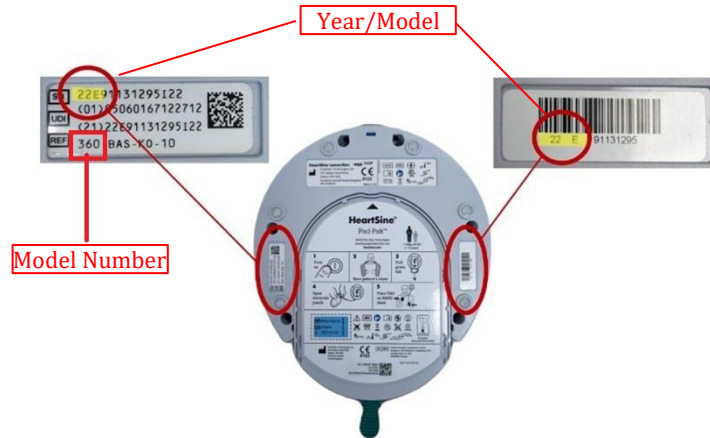


Figure 1 – Serial & Model Number location

Urgent:

Medical device field action

HeartSine® samaritan® PAD

SAM 350P/SAM 360P/SAM 450P/SAM 500P

Customer name: <<merge file>>

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Model	Serial Numbers	UDI-DI
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Product description The HeartSine samaritan PAD is a small, lightweight, portable, battery operated Automated External Defibrillator (AED) designed to treat victims of cardiac arrest.

Product issue (1)(PR4280335 – FA319) Stryker has recently identified an **issue with the membrane tail**, an internal component, which could cause an intermittent fault. This failure could occur at any point when the device is holding a charge in preparation to deliver therapy, while delivering a shock, or after shock delivery. The device becomes inoperable after the failure occurs.

(2)(PR4257283 – FA334) Stryker has also identified that a limited number of HeartSine samaritan PAD devices were **distributed without the required maintenance supplement**. This supplement contains periodic maintenance instructions implemented as an additional risk-control measure. While most devices contain the maintenance supplement, this voluntary correction is being initiated to ensure all end users have access to supplemental maintenance information. This issue was identified through a customer complaint.

Potential risks

(1)(PR4280335 – FA319) Due to the internal component issue, the device **may** fail to deliver the intended therapy during use, potentially leading to a delay in treatment or no treatment being delivered during use. This issue was observed during non-standard rework process and quality testing, not during patient use.

(2)(PR4257283 – FA334) The omission of this supplement may result in users not performing the recommended quarterly power-cycle check. If the check is not performed, the device may have an undetected audio-prompt failure that could result in reduced device readiness and user confusion. This issue was observed through a customer complaint.

There have been no adverse events reported related to this device recall notification.

If your device experiences any of the above issues during use, please seek an alternative defibrillator and contact **your Authorized Distributor or HeartSine Technologies Technical Support at heartsinesupport@stryker.com.**

Customer actions needed:

1. **Identify impacted devices** by verifying if your device serial numbers are included in the list at the link below. For this field action notice, please select “AED + Pad-Pak”.

<https://heartsinerecall.com/HeartSineSamaritanPAD/LookUp>

A. For devices available in your inventory & Stryker direct customers/distributors:

- a. Instructions for where to locate device serial numbers on the device are found in Appendix A*.
- b. If you have more than 15 devices and would like a list of impacted serial numbers, please email **<Insert Local Email>@stryker.com:**
 - Subject: FA319/FA334 - Impacted device list - <<your company name>>
 - Be sure to include your name and phone number in case we need to reach out to you.

B. For devices you have transferred to other users (Non-Stryker direct customers/distributors):

- a. Instructions for where to locate device serial numbers on the device are found in Appendix A*.
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2. **Download** a copy of the missing maintenance supplement when acknowledging the recall and store with your device’s user manual until a replacement is received.
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4. Until a replacement is available, Stryker recommends keeping your HeartSine samaritan PAD in service if you do not have an alternative public access defibrillator. This recommendation is based on internal testing demonstrating a low probability of failure due to these product manufacturing issues.
5. Maintain awareness of this communication internally and near the affected unit until all required actions have been completed within your facility and the unit has been replaced.

Stryker's planned actions:

Stryker is notifying all customers who have received affected HeartSine samaritan PAD devices to perform the actions outlined above. Once your response is received and a replacement is available, Stryker will be in contact as soon as possible to arrange the next steps for a replacement device.

If you have any questions or concerns, please contact Stryker Customer Service at <Insert Info Here>, your Authorized Distributor, *or HeartSine Technologies Technical Support at heartsinesupport@stryker.com.*

In line with the recommendations of the Medical Device Coordination Group Guidance document Ref MDCG 2023-3 and EU MDR 2017/745, we can confirm that this FSCA has been notified appropriately to the National Competent Authority for your country.

On behalf of Stryker, we thank you sincerely for your help and support in submitting your response by MMM DD YYYY. We regret any inconvenience that may be caused and would like to reassure you that we are committed to meeting our high internal quality standards and your expectations.

Appendix A

HeartSine samaritan PAD SAM 350P/SAM 360P/SAM 450P/SAM 500P

Instructions to Identify Impacted Devices

- 1) To find your device serial number and model number, see the labels on the rear of your device as shown below:

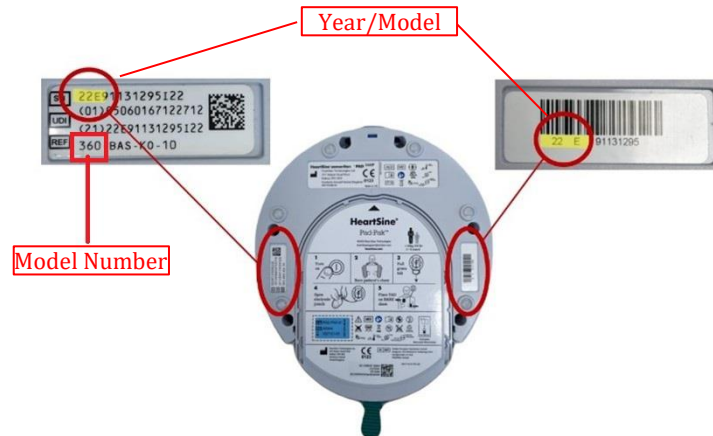


Figure 1 – Serial & Model Number location

Urgent:**Medical device field action****HeartSine® samaritan® PAD****SAM 350P/SAM 360P/SAM 450P/SAM 500P****Customer name:** <<merge file>>**Customer #:** <<merge file>>**Attn: Safety Manager****Field action number:** **PR4257283 – FA334****<Month DD, YYYY/ DD Month YYYY>**

Model	Serial Numbers	UDI-DI
350P 360P 500P	A subset of serial numbers has been affected by this notice. Please check if your AED serial number is affected by visiting the HeartSine PFA Global Landing Page below: https://heartsinerecall.com/HeartSineSamaritanPAD/LookUp If your device is listed, please follow the instructions provided in this letter.	08858251002132S4

Stryker has initiated a serial-specific Safety Notice for the devices listed in the table above. This device notification is being issued to alert customers with HeartSine samaritan PAD SAM 350P/SAM 360P/SAM 450P/SAM 500P devices of a missing maintenance supplement. Out of an abundance of caution, Stryker is completing a voluntary **correction** of these devices.

Product description The HeartSine samaritan PAD is a small, lightweight, portable, battery operated Automated External Defibrillator (AED) designed to treat victims of cardiac arrest. Affected devices were received by customers within March 11, 2024 – February 18, 2025.

Product issue Stryker has identified a limited number of HeartSine samaritan PAD devices that were **distributed without the required maintenance supplement**. This supplement contains periodic maintenance instructions implemented as an additional risk-control measure. While most devices contain the maintenance supplement, this voluntary correction is being initiated to ensure all end users have access to supplemental maintenance information. This issue was identified through a customer complaint.

Potential risks Omission of this supplement may result in users not performing the recommended quarterly power-cycle check. If the check is not performed, the device may have an undetected audio-prompt failure that could result in reduced device readiness and user confusion. This issue was identified through a customer complaint.

There have been no adverse events reported related to this device recall notification.

Customer actions needed:

1. **Identify impacted devices** by verifying if your device serial numbers are included in the list at the link below. For this field action notice, please select “AED + Pad-Pak”.

<https://heartsinerecall.com/HeartSineSamaritanPAD/LookUp>

A. For devices available in your inventory & Stryker direct customers/distributors:

- a. Instructions for where to locate device serial numbers on the device are found in Appendix A*.
- b. If you have more than 15 devices and would like a list of impacted serial numbers, please email <Insert Local Email>@stryker.com:
 - Subject: FA334 - Impacted device list - <<your company name>>
 - Be sure to include your name and phone number in case we need to reach out to you.

B. For devices you have transferred to other users (Non-Stryker direct customers/distributors):

- a. Instructions for where to locate device serial numbers on the device are found in Appendix A*.
- b. If you have further distributed any affected product, please provide this notification to the relevant recipients. **Recipients who are not direct Stryker customers should respond to the facility or organization from which they receive this notification and **not directly to Stryker.***
- c. Email your Stryker representative at xxxxx@stryker.com when you have collected all the responses from your end-users.

2. **Download** a copy of the missing maintenance supplement when acknowledging the recall and store with your device’s user manual until a replacement is received.
3. **Submit your response/acknowledgement** to Stryker electronically via this link: <https://heartsinerecall.com/HeartSineSamaritanPAD/LookUp> and/or email xxxxxx@Stryker.com the enclosed Business Reply Form by [<Insert Date>](#).
4. Maintain awareness of this communication internally and near the affected unit until all required actions have been completed within your facility.

Stryker’s planned actions:

Stryker is notifying all customers who have received affected HeartSine samaritan PAD devices to perform the actions outlined above.

If you have any questions or concerns, please contact [Stryker Customer Service at <Insert Info Here>](#), your [Authorized Distributor](#), or [HeartSine Technologies Technical Support at \[heartsinesupport@stryker.com\]\(mailto:heartsinesupport@stryker.com\)](#).

In line with the recommendations of the Medical Device Coordination Group Guidance document Ref MDCG 2023-3 and EU MDR 2017/745, we can confirm that this FSCA has been notified appropriately to the National Competent Authority for your country.

On behalf of Stryker, we thank you sincerely for your help and support in submitting your response by [MMM DD YYYY](#). We regret any inconvenience that may be caused and would like to reassure you that we are committed to meeting our high internal quality standards and your expectations.

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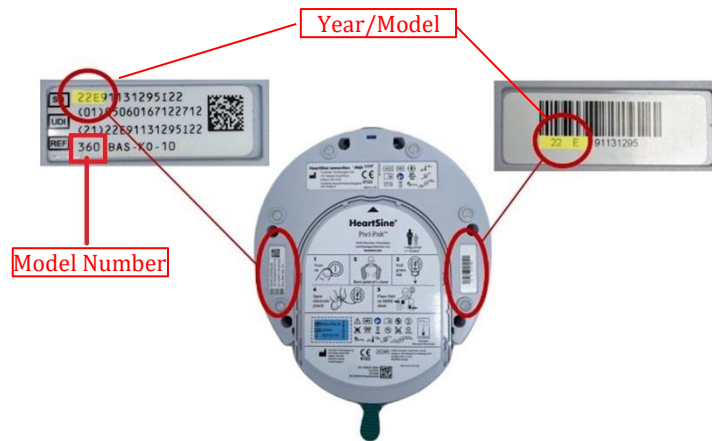


Figure 1 – Serial & Model Number location