

AED MONTHLY CHECKLIST

School/Location: _____	Month/Date	Aug	Sept	Oct	Nov	Dec	Jan	Feb	Mar	Apr	May	June
Inspector's Initials												

Instruction and Recommended Corrective Action												
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<i>Check defibrillator, battery, electrode, packs and cables for damage or foreign substances. If it appears that unit was tampered with, immediately notify Risk Management</i>												
<i>Adult electrode pads plugged into connector. If loose, reconnect electrode pads to AED connector. If pads expired contact Risk Management. Pads expire: _____</i>												
<i>OK or green light is viable on readiness display on handle. If OK or green light is not visible refer to Troubleshooting Manual</i>												
<i>Note USE by date on battery. If date passed, notify Risk Management. Replace battery when exp. date is within 90 days of inspection. Battery expires: _____</i>												
<i>Infant/child electrode pads stored with AED. If missing or expired notify Risk Management. Pads expire: _____</i>												
<i>Other resuscitation equipment stored with AED: Black/Red bag attached to the AED. If missing, notify Risk Management.</i>												

Troubleshooting Readiness Display												
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<p><i>NOTE: Reminder: Complete AED Reporting. USE form after use. DISCARD Pads after each use</i></p> <p><i>At the end of the school year forward completed checklist to Risk Management</i></p>	<p>If battery is not present, notify Risk Management. If battery life is present in unit, press ON and verify that OK appears. If OK is visible, unite is now ready for use. If OK is still not visible, remove the battery to verify expiration date and check pins, then reinsert until "click" is heard. Press ON and verify OK is present. Notify Risk Management 416-5515 for replacement parts or any problems with AED.</p>
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Initials	If AED is inoperable, retrieve another AED, if possible, and continue CPR until EMS arrives
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	Name (Printed)	Signature	Title

AED (Defibrillator) Medical Authorization

The Food & Drug Administration considers defibrillators to be prescription devices pursuant #21 CFR 801.109, and medical authorization is required. Most states provide immunity from civil liability to the physician prescribing on AED. State legislation can be accessed through your state's website, medical board, or <http://aedhelp.com>

This serves as Medical Authorization for External Defibrillators and Automated External Defibrillator(s) (AEDs) as indicated below:

1. Recipient of the AED Medical Authorization (check appropriate box or boxes):

- Individual/Patient
- Business
- Single location
- Multiple locations

1. Name of recipient of AED(s): _____

2. Address of each location at which an AED will be located:

Location name: _____

Street: _____

City/state/zip: _____

Phone number: _____

Contact/title: _____

Location name: _____

Street: _____

City/state/zip: _____

Phone number: _____

Contact/title: _____

If more locations are provided for under this Medical Authorization, please attach a separate piece of paper listing the required contact information for each location.

List any restrictions to this Medical Authorization, if applicable:

Authorization physician (please print):

Name: _____

Street: _____

City/state/zip: _____

Phone: _____

