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### **Medically Necessary Item Allowance Request**

(Reference MNL-352286)

Instructions: First save this PDF file to desktop or folder to enable form features. Complete sections 1, 2, 3 & 4 and sign "User Agreement." Select the "Submit to OHS" button to email to <a href="PantexMedicalProviders@pxy12.doe.gov">PantexMedicalProviders@pxy12.doe.gov</a>. Contact the Prohibited and Controlled Articles hotline (806) 477-4444 for any questions.

Note: Any individual attempting access into a security area with a medically necessary item to include Medical Support Aid (see Section 2) that cannot reasonably be removed to facilitate x-ray search procedures at Protected Area (PA) and Material Access Area (MAA) security stations, equipped with Personnel Positive Identity and Verification (PPIV) Booth, must present a Safeguard and Security (S&S) Allowance Card for access.

### 1). Information on Individual

Full Name (First, Middle, Last)	Employee	Badge #	Phone #	Site Host/	Badge #	Contact phone #	Contact Email
	Contractor			Supervisor Name			
	Site visitor						
	New Hire						
	Other						

#### 2). Information on Item/Device

Type Make Model						Medical Support Aid						
	Hearing aid					Med	Medical Support Any device, instrument, apparatus or applian			ance to be used		
	Glucose monitor					Aid	Required?	after a medical procedure and/or in tandem with a metal				
	Insulin pump						implant or prosthetic for medical aid – either			er temporary or		
	Heart monitor						Yes	perman	permanent. Examples include, crutches, wheelchairs		neelchairs, braces,	
	Metal							slings, ca	slings, casts, etc. If yes, indicate duration of use in Section 5			
	Implant/Prosthetic					□ No		Occupational Health Section (OHS)				
	Other											
					Duration of use		Start		Estimated			
							Date:	Date: End Date:				
Item Location (Check all that apply):						Peripheral Device						
	Left Center Right			Peri	Peripheral Device							
	Left Arm		Head/Neck		Right Arm	Necessa	Any Medical Personal Electronic Device (MEDPED) the requires the use of a peripheral device (companion d				•	
	Left Elbow	□ Chest □ Right Elbow		Right Elbow		Yes		•				
	Left Waist/Hip		Back (Upper)		Right Waist/Hip		No	both wire and wireless) necessary for proper function of the MEDPED require a separate PX-6390. Approval of a MEDPED does not constitute approval of any peripheral device.				
	Left Glutei		Back (Lower)		Right Glutei							
	Left Shin/Calf □ Abdomen □ Right Shin/Calf							does not constitute approval of any peripheral device.			ciai uevice.	
Com	ments/other description:						•			•		

OFFICIAL USE ONLY

May be exempt from public release under the Freedom of Information Act (5 U.S.C. 552). Department of Energy review required before public release.

Exemption: 7, Law Enforcement Name: Clint Olson Org: CNS SS&ES Date: 08/17/2022

Guidance: CG-SS-5, 7/16, DOE OC CNS eDC/RO ID: 482454



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### 3). Area of Use (check all that apply):

Security Areas	Explosive Safety Areas	Comments:
General Access Area (GAA)	General Purpose Area	
Property Protection Area (PPA)	Explosive Area	
Limited Area (LA)	Nuclear Explosive Area	
Protected Area (PA)		
Material Access Area (MAA)		

### 4). User Agreement - Electronic Devices Only

#### Consolidated Nuclear Security, LLC Personal Medical Device User Agreement

(Read this document carefully before signing)

**General**: Consolidated Nuclear Security, LLC (CNS) utilizes the Technical Review Request process to evaluate and approve all Portable Electronic Devices (PEDs) considered Controlled Articles in accordance with Department of Energy (DOE) Order (O) 473.1a, *Protection Program Operations*, for introduction and use at Pantex Plant (Pantex). Upon submission and approval of this PX-6390, the user agrees to abide by device/equipment use restrictions. Violations of this user agreement and identified use restrictions may constitute an Incident of Security Concern (IOSC).

**Employees/Contractors/Vendors:** Users of approved controlled articles understand that it is their responsibility to maintain control of their approved device/equipment and ensure it is not used in an unauthorized manner. The user of the device/equipment agrees to follow these restrictions when introducing or using this device at any onsite or off site Pantex facility:

- The device will not enter any Nuclear Explosive Areas or Hazardous Locations identified in MNL-00055, Pantex Plant Non-Nuclear Facilities Safety Systems Manual, unless otherwise expressly approved in PX-6390, Issued MEDPED Card, and List-0185, Approved Personal Medical Device List.
- The device contains no audio or video recording capability.
- The device will not connect to peripheral device capable of transmitting audio or video information while in a security area.
- The device will not connect to site network or telephone equipment.
- If equipped with wireless capabilities, the device will transmit at power levels less than or equal to 100 miliwatts.
- The device is authorized by Site Medical as a necessary MEDPED in accordance with WI 02.01.01.01.20, Reporting and Processing Off-the-Job Injury or Illness.

I agree to all terms and conditions outlined in the above	Signature:
user agreement.	

Instructions: Select the "Submit to OHS" button to email to PantexMedicalProviders@pxy12.doe.gov.

Submit to Occupational Health Safety



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Site	Use	On	ly
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# 5). Occupational Health Section (OHS)

MEDPED IS on current LIST-0185.	OHS Comments:
MEDPED is NOT on current evaluated LIST-0185.	
N/A – device is a metal implant/prosthetic/medical aid.	
Further medical examination was needed – Appointment was conducted on:	
Date: Time:	
OHS has determined the device to be medically justifiable.	OHS Signature:
OHS has determined the device is NOT medically justifiable.	
The User is a visitor, no medical records are available	

Submit to S&S Operations



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NOTE: NO ACCESS WILL BE GRANTED WHERE ELECTRONICALLY POWERED MEDICAL DEVICES (EMPDs) ARE PROHIBITED

o). I active Engineering Section							
□ Device is not on List-0185 – please review for Master Technical Review Req	Facility	Facility Engineering Comments:					
For MEDPED Exceptions, the device is NOT AUTHORIZED in the following (Check N/A (Not-Applicable) or all the areas the device is NOT authorized to the control of the contro							
□ Nuclear Explosives Area (e.g., Zone 12, Zone 4) □ Material Access A	reas (MAAs)	Facility	y Engineering Signatu	re:			
☐ Explosives Areas (e.g., Zone 11 explosives ☐ Hazardous locatio processing) ☐	ns						
□ N/A (Approved in all areas)							
□   Approved − Electrical Equipment Review (EER) # (Send t Security)	o Technical		ER needed (Send to S	&S Ops)			
7). Technical Security Section							
For MEDPED Exceptions, the device is NOT AUTHORIZED in the following							
areas	_						
Check N/A or all areas where the device is NOT authorized	N/A	Tier 0	Tier 1	Tier 2	Tier 3	Tier 4	
Technical Security Signature:	Technica	Security Comments:					
Submit to S&S Operations							
8). S&S Operations Section							
Approved – Permanent Allowance Card Issue date Allowance Card ID # Exp		S&S Signature:					
□ Not Approved – Requestor has been notified that device requires a TRR/EE							