

# INTERNAL ONLY

## ISLHD POLICY COVER SHEET



**Health**  
Illawarra Shoalhaven  
Local Health District

<b>NAME OF DOCUMENT</b>	Patient-supplied Equipment
<b>TYPE OF DOCUMENT</b>	Policy
<b>DOCUMENT NUMBER</b>	ISLHD CORP PD 02
<b>DATE OF PUBLICATION</b>	April 2021
<b>RISK RATING</b>	Medium
<b>REVIEW DATE</b>	April 2024
<b>FORMER REFERENCE(S)</b>	ISLHD CORP PD 02 - Patients Use of Personal Electrical Devices
<b>EXECUTIVE SPONSOR or EXECUTIVE CLINICAL SPONSOR</b>	Chief Information Officer
<b>AUTHOR</b>	Director Clinical Engineering
<b>KEY TERMS</b>	Electrical, Biomedical, equipment, CPAP
<b>FUNCTIONAL GROUP OR HUB</b>	ISLHD, Clinical Engineering
<b>NSQHS STANDARD</b>	Standard 1
<b>SUMMARY</b>	The purpose of this policy is to regulate and control the use of personal electrical devices of convenience within ISLHD sites. It applies to patients, visitors and staff.

### COMPLIANCE WITH THIS DOCUMENT IS MANDATORY

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**1. POLICY STATEMENT**

The use of personal electrical devices within ISLHD sites requires controls to minimise risks and ensure the safety of patient locations.

**2. AIMS**

The purpose of this policy is to regulate and control the use of personal electrical devices of convenience within the ISLHD. It applies to patients, visitors and staff.

The policy is designed to reduce risks by removing all unnecessary power cables and plug packs from patient locations.

Additional precautions specific to the use of communication devices such as mobile phones and walkie talkies are outlined in MoH guideline, [GL2005-045 27/01/2005 Mobile Phones and Wireless Communication Devices – Interference with medical Equipment – use of](#)

**3. TARGET AUDIENCE**

Hub General Managers, Facility Managers, Department Heads, Nurse Unit Managers, Clinical and Nursing staff

**4. RESPONSIBILITIES**

**Specific responsibilities**

In order to ensure that patient-supplied equipment complies with and is maintained to the required standards, it would be preferred for owners of such equipment to obtain permission for use and supply copies of recent electrical testing, up to date maintenance check records and have arrangements in place for its inspection thereafter, as specified by the manufacturer of the device or by a qualified external contractor/service agent of their choice.

**Either the Department Head, Nurse Unit Manager, clinician or nursing staff** (or their delegate) is to visually inspect patient-supplied equipment and associated power cables/plug packs, using attached Appendix 1: Patient-supplied Equipment - Flow Chart, to ensure that there are no signs of damage to the device or cables. This includes battery operated devices.

Note: if patients have been using the device regularly in their home environment, then to the best of our knowledge it can be assumed that the equipment is working correctly. This however does not negate the need to perform the above inspection.

**Admission staff** should be aware of this policy and advise patients that their patient-supplied equipment is required to undergo the aforementioned inspection before being used.

Patients who are admitted for short durations such as Oncology day care patients and dialysis patients should, as far as practical, use battery powered devices only. This will eliminate unnecessary power cables and risks associated with use of mains power in potentially wet environments. Additional batteries should be brought in by the patient if their device is required to run the full length of their stay. Patients with obstructive sleep apnoea may be requested to bring their CPAP machine and ancillary equipment with them at the time of admission.

**Patients** are also to be made aware that their personal electrical devices, cables and battery chargers/plug packs are to be visually inspected for damage prior to use and that battery only operated devices are preferred. **Any damage to the devices, cables or battery chargers/plug packs revealed by visual inspection will result in the item not being allowed to be used on the premises.**

Patients should also be made aware that the ISLHD does not bear liability for the loss of, theft, or damage to personal electrical equipment.

If the NUM or delegated person has any doubts about the risk involved by allowing the device to be used in the ward, then the Clinical Engineering department should be contacted as soon as possible, to have the device inspected or tested.

#### General Principles

Patients and visitors are not to use their personal electrical devices whilst in ISLHD facilities where the use of the device may:

- Constitute an electrical safety risk
- Constitute an EMI risk with the facilities life support medical equipment
- Pose a risk to staff and others (cables on the floor, heavy TVs on cabinets not designed for the weight, cables severed when dropping side rails of beds, cables in vicinity of fluids, etc.)

#### Caution

- In general, permission should be refused for patient-supplied mains-powered equipment to be used in a healthcare facility. (AS/NZS 2500:2020 Clause 5.8.5).
- Use of power boards and extension cords should be discouraged as they may compromise the environmental protection provided in patient locations. Double adaptors and piggy back plugs should not be used. (AS/NZS 2500:2020 Clause 6.7)
- Use of mains powered devices should be discouraged in bathrooms and wet areas.

#### During Use

- Mains operated devices – Where use is permitted by the Department Head or their delegate, ensure any cables are clear of bed rails, are off the floor and remain safe.
- Ensure the mains switch on the power point is turned off, and the device is turned off before plugging the mains cable/plug pack for the device in, turn the power point switch on, turn the device on.

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**After Use**

- Ensure all devices are turned off, the power is turned off at the wall and the power cable is unplugged from the wall and the complete device is safely stored.

**5. DEFINITIONS**

**E.M.I.** – Electromagnetic Interference – interference in a circuit caused by the radiation of an electric or magnetic field.

**Personal electrical device** – refers to any electrical device that is designed for portable personal use and is brought into the ISLHD facilities for the personal use of patients and visitors. This may include but is not limited to, CPAP machines, portable televisions, laptop computers and mobile phones and their chargers.

**Cardiac-type procedure** - is considered to be undertaken when an indwelling electrical conductor in contact with the heart is accessible outside the patient's body, and there is a risk of microshock.

**Patient location** – any intended location of the bed, table or seating arrangements for a patient, whether or not occupied by the patient. Of particular importance in this policy are the power points near a patient's bed or dialysis patient's chair.

**6. DOCUMENTATION**

Appendix 1 – Patient-supplied Equipment – Flow Chart

**7. AUDIT**

Not Required

**8. REFERENCES**

- [Work Health & Safety Regulation 2017 - Part 4.7 General electrical safety in workplace and energised electrical work.](#)
- [Safework NSW Code of Practice – Managing electrical risks in the workplace - August 2019.](#)
- AS/NZS 3760 – In service safety inspection and testing of electrical equipment
- AS/NZS 2500 – Safe use of medical electrical equipment in health care
- AS/NZS 3551 – Technical management programs for medical devices
- [SafeWork NSW website](#)
- Policy ISLHD CORP PD 26 - Biomedical Equipment – Testing, tagging and labelling

**9. REVISION & APPROVAL HISTORY**

<b>Date</b>	<b>Revision No.</b>	<b>Author and Approval / Date</b>
May 2007	0	<b>Author:</b> Manager Biomedical Services
		<b>Approval / Date:</b> SESIH Area Biomedical/Clinical Engineering Committee, Director Operations and Area Executive Committee / May 2007
Jul 2017	1	<b>Author:</b> Manager Clinical Engineering
		<b>Approval / Date:</b> Executive Director Finance and Corporate Services / Mar 2018
Feb 2021	2	<b>Author:</b> Director Clinical Engineering  <b>Approval/Date:</b> Corporate Policy Recommendation committee/ March 2021 <b>Approval/Date:</b> Chief Information Officer / April 2021

**10. APPENDIX 1 – Patient-supplied Equipment – Flow Chart**

