UNIT II: BIOSIGNAL CHARACTERISTICS AND ELECTRODECONFIGURATIONS 9

Bio-signal characteristics – frequency and amplitude ranges. ECG – Einthoven"s triangle, standard 12 lead system. EEG – 10-20 electrode system, unipolar, bipolar and average mode. EMG– unipolar and bipolar mode. Electrodes

ECG - Lead System, Recording Methods and Typical Waveforms

Introduction:

- □ Electrocardiography (or) Electrokardiography deals with the recording and study of electrical activity of the heart muscles.
- The potentials originated in the individual fibers of heart muscles are added to produce the ECG waveform.
- □ The ECG waveform reflects the rhythmic electrical depolarization and repolarization of the heart muscles associated with the contractions and relaxation of the atrium and ventricles.
- □ The electrocardiogram gives details of the state of the heart and any disturbance in the heart rhythmic (ar hythmia) can be diagnosed.
- □ The typical ECG wave consists of P wave, QRS complex and T wave.
- □ The electrical potentials of the heart are measured by placing suitable electrodes.
- \Box Eithersurface electrodes with proper electrode paste orneedle electrode can be used. <u>ECG Lead</u>

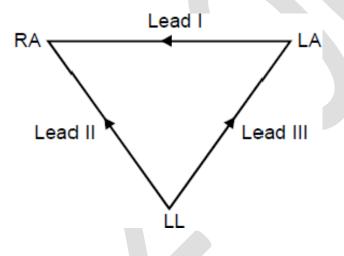
system:

□ There are fourdifferent ECG lead systems used universally, o

Bipolarlimb lead (or) standard lead system

- o Augmented limb lead system
- o Chest lead (or) Pre-cordial system
- *o* Frank lead system (or) co**r** ected orthogonal lead system
- □ <u>Bipolarlimb leads:</u>
- \Box In bipolarlimb leads system ECG is recorded with two electrodes at a time.
- □ The final trace is due to difference in potential of 2 electrodes kept at 2 differentlocations on the body.
- \Box This is also called the standard lead system.
- \Box For this system, the potentials are tapped from 4 locations of the body namely,
 - o Right arm white colourelectrodes

- o Left arm Black colourelectrodes
- o Right leg Green colourelectrodes
- o Left leg Red colourelectrodes
- \Box The right leg electrode is used as reference electrode.
- \Box The three different leads of the system are,
 - o Lead I V1 (Voltage drop from left arm (LA) to right arm (RA)
 - o Lead II V2 (Voltage drop from left leg (LL) to right arm (RA)
 - o Lead III V3 (Voltage drop from left leg (LL) to left arm (LA)
- □ The closed path between RA to LA to LL and back to RA is called the Einthoven triangle.



Einthoven Triangle

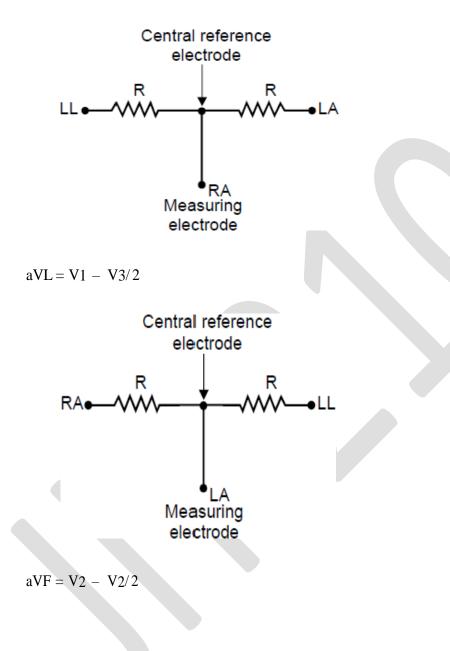
- \square R wave amplitude of lead II is equal to sum of R wave amplitude of leads I and leads III.
- \Box Forexample, if V1 = 0.5 mV, V2 = 0.7 mV, then,
- \Box V3 = 0.2 mV. (i.e) V2 = V1 + V3.
- 2. Augmented Unipolar Limb leads:
- □ This type of lead system is introduced by Wilson.
- □ ECG trace is due to potential between single measuring electrode and central referenceelectrode.
- □ The central reference electrode is built up by tying two electrodes with two equal and large resistors in between them.
- □ A pairof limb electrodes is tied up with 2 large resistors to make the central referenceelectrode and the third limb electrode is the measuring electrode.
- aVR augmented voltage RA aVL
- augmented voltage LA aVF -

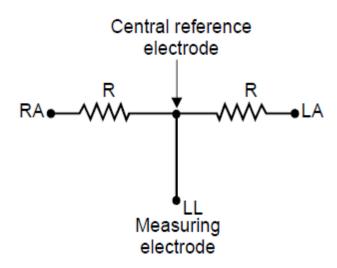
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augmented voltage foot

□ By Kirchoff's law,

aVR = -V1 - V3/2

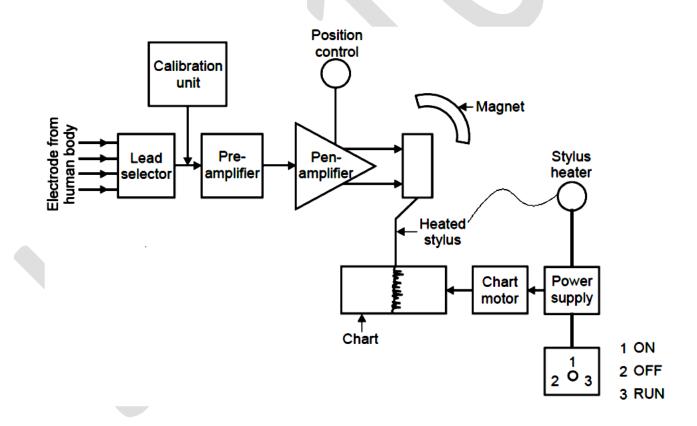




- 3. Unipolarchest lead:
- □ In Unipolarchest leads system, in addition to electrodes present in augmented Unipolarlimb leads there are two more electrodes placed in the chest, close to heart.
- By connecting 3 large equal resistors between LA, RA and LL reference electrode centrepoint obtained, this is the central electrode.
- □ This system includes an integration of 3 unipolarleads, 3 bipolarleads and 6 chestleads.
- □ Location of chest leads (V1 till V6)
 - o V1 4th intercoastal space at right sternal margin
 - $o V2 4^{th}$ intercoastal space at left sternal margin
 - o V3 Midpoint of V2 and V4
 - o V4 5th intercoastal space at mid-clavicularline
 - o V5 Same level as V4 anterior auxiliary line
 - o V6 Same level as V4 Mid auxiliary line
- 4. Frank lead system:
- \Box Same as chest lead system.
- Heart's dipole field is resolved into three mutually perpendicular components andhence state of heart is studied three dimensionally.
- ECG recording set-up (or) ECG Instrumentation:
- □ The connecting wires for the patient electrode originate at the end of a patient cable.
- \Box The wires from the electrodes connect to the lead selectorswitch
- □ A push button allows the insertion of a standardization voltage of 1 mV to standardizeorcalibrate

the recorder.

- □ From the lead selectorswitch the ECG signal goes to a pre-amplifier, which is a differential amplifier with high common mode rejection ratio, high gain factor, highinput impedance and lowoutput impedance.
- □ It is AC coupled to avoid problems with small DC voltages that may originate from polarization of the electrodes.
- □ The pre-amplifier, is followed by a DC amplifiercalled the pen amplifierorpower amplifier, which provides the power to drive the pen motorthat records the actual ECGtrace.
- A position control in the pen amplifiermakes it possible to centre the pen on therecording paper.
- □ ECG recorders use heat sensitive paper, and the pen is actually an electrically heated stylus, the temperature of which can be adjusted with a stylus heat control foroptimal recording trace.



ECG recording set-up (or) ECG Instrumentation

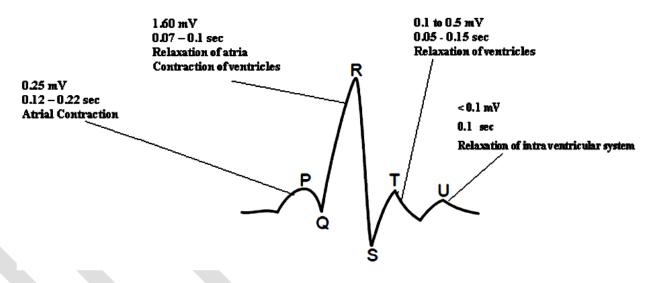
- □ There is a marker stylus that can be actuated by a push button and allows the operator mark a coded indication of the lead being recorded at the margin of the electrocardiogram.
- □ Normally electrocardiogram is recorded at a paperspeed of 25mm/ sec but a fasterspeed of

50mm/sec is provided to allow resolution of the QRS complex at very highheart rates.

- □ The power switch of an ECG record has three positions. In the "ON" position the powerto the amplifieris turned on but the paperdrive is not running. In "RUN" position paper drive starts running. In "OFF" position power is switched off.
- A push button allows the operator to check whether the recorder is connected to the power line with polarity.
- \Box This is done to avoid any shock hazard for the patient.
- □ Isolated orfloating input amplifiers are used for the safety aspect of electrical connections to the patient.

ECG – Typical Waveform:

The complete waveform is called electrocardiogram with labels PQRSTU indicating important diagnostic features.

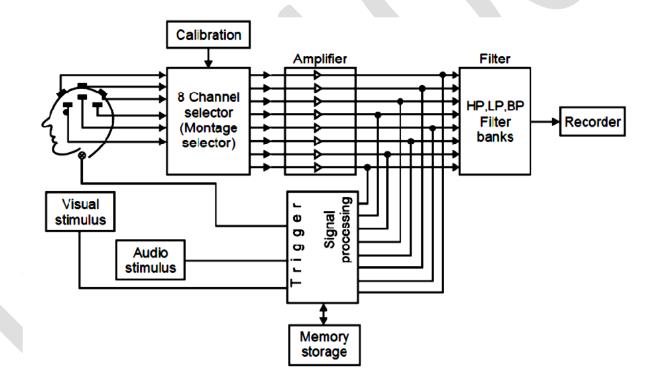


ECG – Typical Waveform

- □ The instrumental set up used for recording is called electrocardiograph.
- □ The process of recording and studying electrical potentials of heart is called electrocardiography.
- □ If the PR interval is more than 0.22 sec, then it is an indication of AV block (first degreeblock)
- \Box When QRS complex duration is more than 0.1 sec, then it is an indication of the bundleblock.

EEG – Lead System, Recording Methods and Typical Waveforms Recording System:

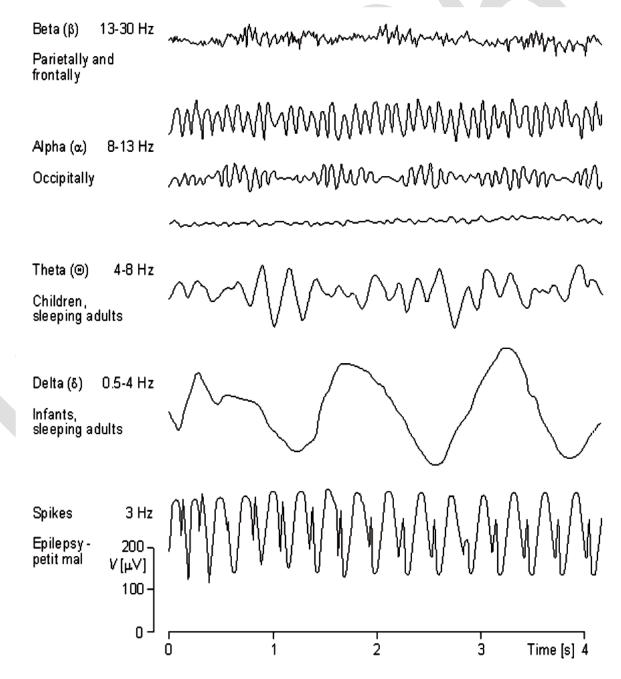
- Electro encephalography (EEG) deals with the recording and study of electrical activity of the brain.
- □ By means of electrodes attached to the skull of a patient, the brain waves can be picked up and recorded.
- □ Electrical potentials of the brain are due to gradient in concentration of dendrite gradedpotentials.
- □ EEG recording set up includes the patient cable consisting of 21 electrodes and isconnected to the eight channel selector.
- □ Every channel of the channel selectorconsists of an individual, multistage, ac coupled, differential, adjustable gain amplifier.
- □ These amplifiers must have high gain and lownoise characteristics since the EEGpotentials are in micro Volt range.



EEG – Recording Set Up

- □ The common mode rejection ratio of the EEG amplifiers should be high to minimizestray interference signals.
- \Box They should have input impedance and lowoutput impedance.
- \Box The amplifiermust be free from drift so as to prevent the slowmovement of recordingpen.
- □ The EEG signal frequency ranges between large values so it becomes necessary to use set of filters including lowpass, high pass and band pass.

- □ The amplified EEG signals are passed through this filter bank.
- □ Typical cut off frequencies for the lowpass filters are 5.3, 1.6, 0.53 and 0.16Hz.
- □ Band pass filters tuned for 60 Hz are used to eliminate mains frequency interference.
- □ The high pass filters have typical cut off frequencies 15, 30, 70, 300 Hz.
- □ The output voltage from the amplifiermay eitherbe applied directly to the eight channeldisplay through the filter bank orit may be stored as data on a tape recorder or computermemory forfurther processing.



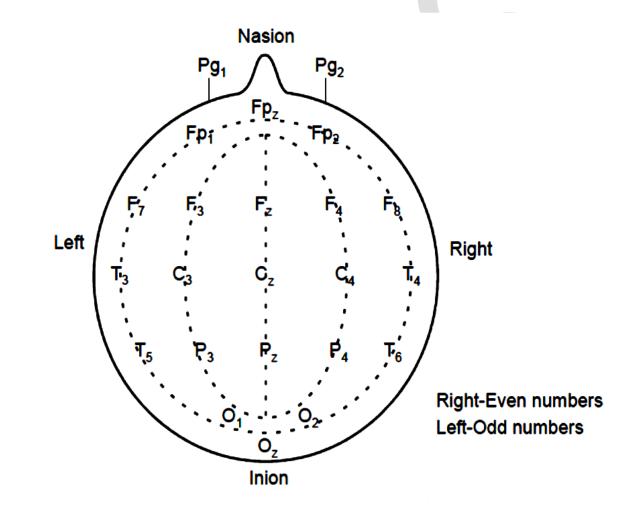
EEG – Typical Waveform

- □ There are otherfacilities available to record evoked potentials from the brain due to external stimuli like visual stimulus, audio stimulus and tactile stimulus.
- □ The time delay between the stimulus and response can also be measured in the signal processing unit and can be accounted forevoked potentials.
- □ EEG helps in diagnosing disorders orabnormalities related to brain such as, tumors, Epilepsy, sclerosis, sleep disorders, mis-functioning of neurons etc.

EEG - Electrode System:

- □ The commonly used electrode system for recording EEG signals is termed as 10-20electrode system or Montage electrode System.
- □ The electrodes are placed at a distance of 10% and 20% appropriately from the total distance between the extreme end points of the skull namely appropriately from the total distance between the extreme end points of the skull namely appropriately from the total distance between the extreme end points of the skull namely namely appropriately from the total distance between the extreme end points of the skull namely namely appropriately from the total distance between the extreme end points of the skull namely namely appropriately from the total distance between the extreme end points of the skull namely namel
- □ This type of a system is otherwise called montage electrode system.
- □ In montage system the electrodes are placed symmetrically on both sides of the skull.
- Both bipolarand unipolarsystem can be used. In bipolarthe differences in potential betweentwo neighboring electrodes are considered.
- □ For Unipolartype the reference electrodes are placed on the non-active part of the headlike forehead oron the earlobes.
- □ Firstly the distance between nasion and inion is determined and the distance between the right and left earlobe is also determined.
- □ The central electrode is placed at the intersection of the imaginary lines joining nasionand inion and right earand the left ear.
- Electrode placement along imaginary line between inion and nasion, o Fpz –
 10% of distance between nasion-inion
 - o Fz 20% of distance between nasion-inion
 - o Cz Centre point of nasion-inion and right earlobe-left earlobe
 - o Pz 20% of distance between nasion-inion
 - o Oz 10% of distance between nasion-inion
- □ Electrode placement along imaginary line between right earlobe and left earlobe,
 - o T3 10% of distance between right and left earlobe
 - o C3 20% of distance between right and left earlobe
 - o Cz Already placed [centre point of nasion-inion and right and left earlobe]
 - o C4 20% of distance between right and left earlobe
 - o T4 10% of distance between right and left earlobe
- □ Electrode placement between Fpz and Oz passing through T3

- o Fp1 10% of distance between Fpz and Oz
- o F7 20% of distance between Fpz and Oz
- o T3 already placed (10% distance between right and left lobe at the left)
- o T5 20% of distance between Fpz and Oz
- o O1 10% of distance between Fpz and Oz



10-20 EEG Electrode System

- \Box Electrode placement between Fpz and Oz through T4,
 - o Fp2 10% of distance between Fpz and Oz
 - o F8 20% of distance between F¬pz and Oz
 - o T4 already there (10% distance between right and left earlobe at the right)
 - o T6 20% distance between Fpz and Oz

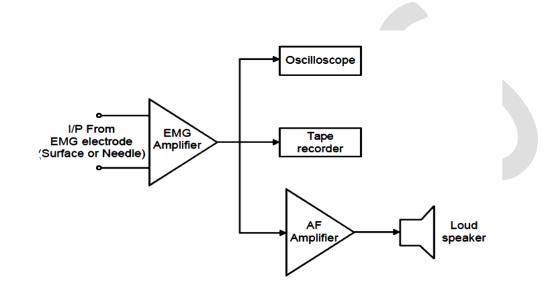
- o O2 10% distance between Fpz and Oz
- \Box Electrode placement between Fp2 and O2 through C4,
 - o F4 25% of distance of Fp2 and O2
 - o C4 20% of distance between right and left earlobe (already present) (right side)
 - o P4 25% of distance between Fp2 and O2
- \Box Electrode placement between Fp1 and O1 through C3,
 - o F3 25% of distance between Fp1 and O1
 - o C3 20% of distance between right and left earlobe (already present) (left side).
 - o P3 25% of distance between Fp1 and O1

EMG – Lead System, Recording Methods and Typical Waveforms:

- Electromyograph is the instrument for recording and interpreting the electrical activity of muscles action potential.
- □ The electrical activity of the underlying muscle can be measured by placing surfaceelectrodes on the skin.
- □ To record the action potential of individual motorunit the needle electrode is inserted into the muscle.
- □ EMG indicates the amount of activity of a given muscle or a group of muscles.
- □ EMG waveform appears, very much like a random noise signal.
- □ Contraction of a muscle produces action potential. When a muscle is relaxed there isno action potential.
- \Box The surface of the skin is cleaned and electrode paste is applied.
- □ The electrodes are kept in place by means of elastic bands.
- □ The amplitude of the EMG signals depends upon the type and placement of electrodesused and degree of muscularexertions.
- □ The surface electrode picks up many overlapping spikes and produces an average voltage from various muscles and motorunits.
- \Box The EMG signal ranges from 0.1 mV to 0.5 mV.
- □ The frequency components of the EMG signal vary from 20 KHz to 10 KHz and they are restricted to the frequency range of 20Hz to 200Hz forclinical purposes by using a low

pass filter.

- \Box The normal frequency is 60 Hz.
- □ Slowspeed strip chart recorders and the signals are displayed or CRO and photographic recordings are also made.



EMG – Recording Set Up

- □ Usually two cathode ray tubes one forviewing and otherfor recording is used.
- □ A light sensitive papermoves over the recording cathode ray tube and the image is produced on that paper.
- □ Forcontinuous recording, the paperspeed is about 5 to 25 cm/sec.
- The amplifiershould have uniform frequency response in the frequency range from 10Hz to 1 KHz with high CMRR (100 dB) and input impedance greater than 10 M Ω with lowoutput impedance.
- \Box The signal is also recorded in a tape recorder forfuture reference.

Contraction of muscle ·····M···

Typical EMG Waveform

□ EMG waveforms are very useful forstudying the neuromuscular function, neuromuscular condition, reflex responses, extent of nerve lesion and diagnosing themusculardiseases.

Nerve conduction Velocity:

- □ Nerve conduction velocity is used to identify the location and extent of nerve lesion.
- To find out the nerve conduction velocity two electrodes are used namely, 1)
 Stimulating electrode and 2) Measuring electrode.
- □ When a short brief electric pulse of 0.1 ms or 0.2 ms is given to the muscle, the muscleundergoes contraction ortwitch giving rise to action potential.
- □ The stimulating electrode gives the stimulation pulse.
- ☐ The measuring electrode records the EMG and hence the action potential due to contraction.
- □ The time between stimulation and contraction helps in determining velocity, if the distance between them is known.
- □ Firstly, the stimulating and measuring electrode are separated by a known distance 11.
- □ The time between stimulation and contraction is noted t1. This time is also called latency.
- □ Then the stimulating and measuring electrode are removed and replaced at differentlocations separated by a distance 12, which is lesserthan 11.
- \Box Nowlatency cor esponding to 12, which is t2 is noted.
- \Box The conduction velocity, V is calculated by, $11 \frac{12}{t1-t2}$.
- \Box Normal nerve conduction velocity is 50 m/s.
- \Box If it is below 40 m/s, then a disorder orlesion is detected at the location.

 \square B – Due to later receptor potentials produced by syruptic ending of the photo receptors.C to D – Wave recorded at the offset of the light stimulus.

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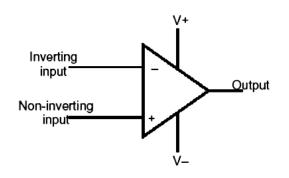
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UNIT III SIGNAL CONDITIONING CIRCUITS

Need for bio-amplifier - differential bio-amplifier, Impedance matching circuit, isolation amplifiers, Power line interference, Right leg driven ECG amplifier, Band pass filtering

Amplifiers

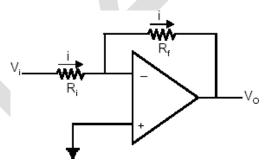
- \Box Amplifieris a device which amplifies or increases an input of cur ent or voltage.
- □ Most bio-electric signals are of very lowamplitude thus requiring amplification.
- □ Amplifiers are also used forinterfacing sensors/transducers that senses body motions,temperature and chemical concentrations to signal conditioners.
- □ Thus it can be said that an amplifierused to process bio-potentials are called bio-electric amplifiers.
- □ Some bio-electric amplifiers are ac-coupled and some are dc-coupled.
- Linearintegrated circuits are being used in a number of electronic applications including medical electronic instruments.
- □ An important linearintegrated circuit is operational amplifier.
- □ The operational amplifier(OP-AMP) is a multi-terminal, high gain differential amplifier.
- □ It is normally used in circuits that have characteristics determined by external negative feedback networks.
- □ The circuit symbol foran OP-AMP is shown in figure below.
- □ The inverting input produces an output signal that is 180 degrees out of phase with the input signal, which is termed as inversion of the signal.
- □ The non-inverting input produces an output signal that is in phase with the input signal.



Operational Amplifier Circuit Diagram

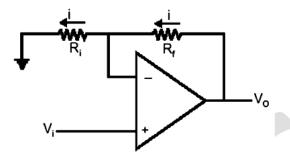
- \Box The properties of an ideal op-amp include,
 - *i*. Infinite open loop voltage gain.

- ii. Zero output impedance
- *iii*. Infinite input impedance
- iv. Infinite frequency response
- v. Zero noise contribution
- vi. Both input followeach otherin feed back circuits.
- $\hfill\square$ There are three basic classes of voltage amplifiers namely,
- □ Inverting follower
- \Box Non-inverting follower with gain
- \Box Unity gain non inverting follower.
- □ There are two basic rules orinput terminal restrictions that are very helpful in designing op-amp circuits.
- \Box They are,
- □ When the op-amp output is in its linear range, the two input terminals are at the same voltage.
- \Box No cur ent flows into eitherinput terminal of the op-amp.
- □ In the basic inverting amplifiercircuits, a portion of the output voltage Vo is fed back via feed back resistor Rf to negative input terminal.
- \Box The positive input of the op-amp is at 0V. This condition is called as virtual ground.
- \Box Since the input voltage is Vi the cur ent through Ri is i = Vi/Ri



Inverting Amplifier

- \Box By rule (2) no cur ent can enterthe op-amp and thus I flows through Rf.
- \Box Hence there is a voltage drop i Rf across Rf.
- \Box Therefore, Vo = -i.Rf
- $V_0 = -V_i R_f / R_i$
- (or) $V_0/V_i = -R_f/R_i$
- □ In the basic non inverting amplifiercircuit Vi is connected to positive input terminal.



Non-inverting AmplifierBy

rule (1), Vi also exists at negative terminal.

This causes cur ent $i = V_i/R_i$ to flow to ground through Rf.

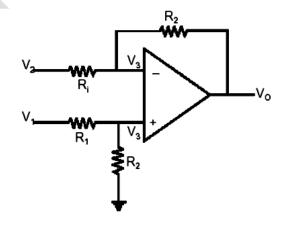
$$Vo = i(Rf + Ri)$$

Vo/ Vi = i(Rf + Ri) / i .RiVo/

$$V_i = (R_f + R_i) / R_i$$

Differential Amplifier

- In medical electronics, the differential amplifieris employed when it is necessary to measure the voltage difference between two points, both of them varying in amplitudeat different rates and at different patterns.
- Example: Differential amplification of heart generated voltages picked up by means ofelectrodes on the arms and legs.
- □ A differential amplifierproduces an output voltage that is proportional to the difference between the voltages applied to the two input terminals.



Differential Amplifier

Since an operational amplifierhas a pairof differential input terminals, it is easily connected foruse in a

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differential amplifierconfiguration.

The differential voltage at the input terminals of the op-amp is zero, that is nodes a and bare at the same potential designated as V3.

The nodal equation at 'a' is

((V3 - V2)/R1) + ((V3 - V0)/R2) = 0.....(1)

The nodal equation at 'b' is

((V3 - V1)/R1) + (V3/R2) = 0(2)

Rear anging we get,

((1/R1) + (1/R2))V3 - V2/R1 = Vo/R2(3)

((1/R1) + (1/R2))V3 - V1/R1 = 0(4)

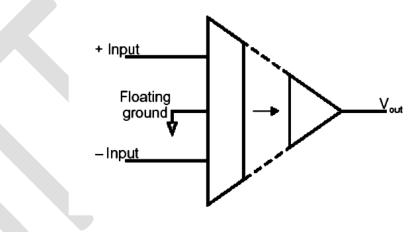
Subtracting (4) from (3) we get, 1/

 $R1.(V\neg 1 - V2) = V0/R2$

$$V0 = R2/R1.(V1 - V2)$$

Isolation Amplifier

- □ Isolation amplifiers are used for providing protection against leakage cur ents.
- □ Isolation amplifiers are used mainly in ECG recording, to prevent accidental internal cardiac shock.



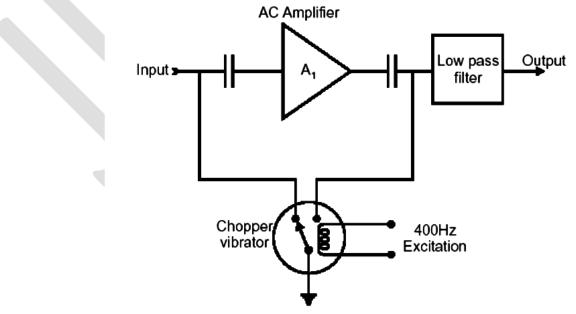
Isolation Amplifier

- □ Isolation amplifiers provide insulation between the patient connectorand the ac powermains line cord.
- □ Isolation amplifiers are mainly used to protect hospital patients, susceptible toelectrical shock hazards.
- □ They break the ohmic continuity of electric signals between the input and output of the amplifier.

- $\hfill\square$ These type of bio-electric amplifiers provide as much as $10^{12}\,\Omega$ of insulation.
- □ The insulation amplifier composed of an input amplifier, modulator, an isolation bar ier, a demodulator and an output amplifier.
- $\hfill\square$ The isolation amplifier is really an energy converter.
- □ There is an input common and an output common that are electrically isolated from one another.
- □ Isolation bar iermay be optical, magnetic transformer, capacitive oreven heat transfer.
- □ In the isolation amplifierelectrical energy on the modulatorside is converted to some non electrically conductive energy in the ba**r** ierand then converted back to electrical energy on the demodulatorside.
- □ Isolation amplifiers actually operate on the principle of attenuation.
- □ A high bar ier impedance acts in series between input and output.
- □ Isolation amplifierbreak ground loops to permit incompatible circuits to be interfaced together while reducing noise.
- They amplify signals while passing only lowleakage cur ent to prevent shock to peopleordamage to equipment.
- Depending on the design the isolation amplifiers are classified as battery powered, car ier, optically coupled and cur ent loading.

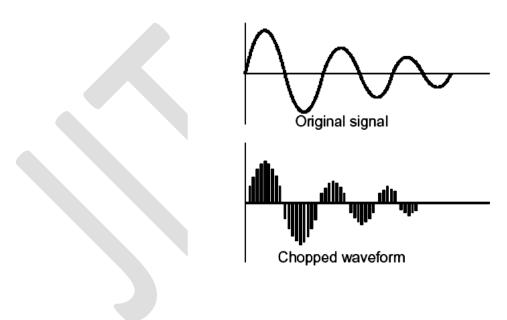
Chopperamplifiers

Chopperamplifieris used in medical electronic devices as it achieves adequate low frequency response while avoiding drift present in direct coupled amplifier. Drift is thechange in gain caused by thermal effects on the amplifiercomponents.



Chopper Amplifier

- □ Chopperamplifiers use a chopping device, which converts a slowly varying direct cur ent to an alternating form with amplitude proportional to the input direct cur ent.
- □ Phase dependent on polarity of the original signal.
- □ This alternating voltage is then amplified by a conventional amplifier whose output isrectified back to get an amplified direct cur ent.
- □ The differential input chopperamplifieris shown in figure below,
- □ The chopperis a vibrator driven single pole, double throw switch that grounds the amplifierinput and output terminals on alternate swings of the switch.
- \Box The choppervibrator coil is excited by a 400 Hz ac car iersignal.
- \Box The chopperis on the input circuit only.
- □ The centre tap of the input transformer is connected to the input connector, while the two winding extremities are connected to the chopper.
- \Box The pole of the chopperswitch becomes the other terminal of the input connector.
- □ Gain is provided by the ac-coupled amplifier.



Chopped Waveforms

□ In medical field, chopperamplifiers are used in the amplification of small DC signals of a few micro volts. Chopperamplifiers are used with transducers such as strain gauge, thermistors,

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thermocouples etc.

Electrical safety in medical environment: shock hazards – leakage cur ent-

Instruments forchecking safety parameters of biomedical equipments

Introduction:

- Medical Technology has substantially improved health care in all medical specialties and has reduced mortality forcritically ill patients.
- □ The increased complexity of medical devices and theirutilization in more procedures result in about 10,000 device related patient injuries according to a survey.
- Most of these injuries are attributable to improper use of a device as a result of inadequate training and lack of experience.
- Medical procedures usually expose the patient to more hazards than the typical home or workplace, because in medical environments the skin and mucous membranes are frequently penetrated oraltered and because there are many sources of potentially hazardous substances and energy forms that could injure the patient.
- □ Sources of potential hazards in medical environment include fire, air, earth, water, chemicals, drugs, microorganisms, vermin, waste, sound, electricity, natural and unnatural disasters, sur oundings, gravity, mechanical stress and people responsible foracts of omission and radiation from x-rays, ultrasound, magnets, ultraviolet light, microwaves and lasers.

□ Electrical safety is important to be considered in the medical instrumentation field. <u>Physiological Effects of electricity:</u>

- □ For a physiological effect to occur, the body must become part of an electric circuit.
- □ Cur ent must enterthe body at one point and leave at some otherpoint.
- □ The magnitude of the cur ent is equal to the applied voltage divided by the sum of theseries impedances of the body tissues and the two interfaces at the entry points.
- □ The largest impedance is often the skin resistance at the contact surface.
- \Box Three phenomenons can occur when electric cur ent flows through biological tissue:
 - 1) Electric stimulation of excitable tissue (nerve and muscle)

- 2) Resistive heating of tissues
- 3) Electrochemical burns and tissue damage fordirect cur ent and very high voltages.
- □ Psychophysical and physiological effects that occurin human as the magnitude of applied electric cur ent progressively increases.
- □ The approximate range of cur ents needed to produce each effect when 60 Hz cur ent is applied for 1 to 3 sec via copper wires that a 70 Kg human hold in each hand.

Threshold of Perception:

When the local cu**r** ent density is large enough to excite nerve endings in the skin, the subject feels a tingling sensation.

Cur ent at the threshold of perception is the minimal cur ent that an individual can detect. When someone with moistened hands grasps small copper wires, the lowest thresholds are about 0.5 mA at 60 Hz, thresholds fordc cur ent range from 2 to 10 mA.

Let – Go Cur ent:

The let - go cur ent s defined as the maximal cur ent at which the subject can withdrawvoluntarily. The minimal threshold for let - go cur ent is 6 mA. Still highercur ents cause involuntary contraction of respiratory muscles severe enough to bring about asphyxiation if the cur ent is not inter upted.

Micro-shock & Macro-shock:

Many devices have a metal chassis and cabinet that can be touched by the medical attendants and patients. If they are not grounded, then an insulation failure orshort circuitresults and leads to macro-shock ormicro-shock.

Micro-shock:

A physiological response to a cur ent applied to the <u>surface of the heart</u> that results in unwanted stimulation like muscle contractions ortissue injury is called micro-shock. Micro

-shock is caused when cur ents in excess of 10 micro Amperes flowthrough an insulated catheterto the heart. The cathetermay be an insulated, conductive fluid filled tube, ora solid wire pacemakercable.

Micro-shock Hazards:

1) Leakage Cur ents:

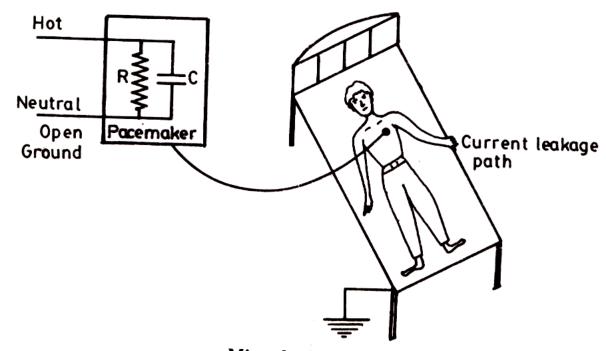
Most of the accidents occurdue to improper grounding and leakage cur ents. <u>The leakagecur ent is an</u> <u>extraneous cur ent flowing along a path other than those which is intended to flow.</u> Leakage cur ent may be caused due to resistive, inductive orcapacitive couplings with the mains orsome electronic equipment.

As shown in figure below patient with a pacemakerlying in an electrically operated bed with a bipolarcathetergoing to the right ventricle of the heart via the right jugularvein. Thepacemaker's case is connected to the ground of the power cord. The three wire 2.5 metre power cord is nowconnected to a two wire 3 metre extension cord which is plugged into the three wire power outlet.

The bed frame is properly grounded to the power system. The patient's left hand is restingon the bed frame. Since the ground of the pacemakeris floating, by capacitive coupling a total of 180 micro Amperes of leakage cur ent is present in the pacemaker. In this case

leakage cur ent is passed from the pacemakerthrough the catheterinto the heart, through

the body core of the left hand and then to the ground via the bed frame. The heart undergoes ventricular fibrillation. This dangerous accident arises because of the open ground of the pacemakerby using a two wire extension cord. The leakage cur ent flowis due to 1) Ungrounded Equipment, 2) Broken ground wire and 3) Unequal ground potentials

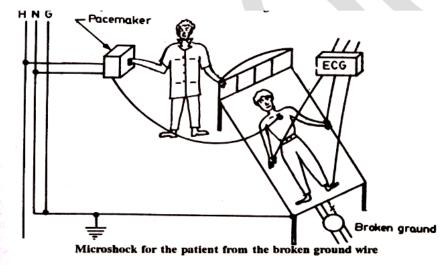


Microshock due to leakage current

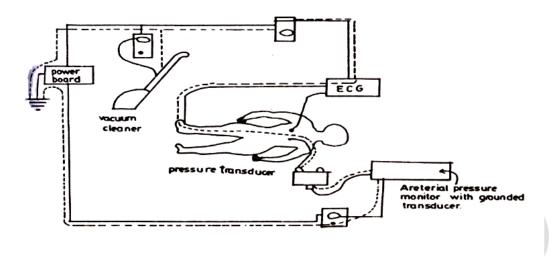
As

illustrated in figure belowthere is a possibility that a doctorholding the pacemaker's wireby his one hand and touching the electrical bed frame by his otherhand can increase the risk of leakage cur ent. Broken ground connection on the electric bed allows a voltage to exist on the bed frame due to capacitive coupling between the bed frame and power line. The pacemaker wire is going into the heart of the patient.

The heart activity is monitored by an ECG recorder. Hence a leakage cur ent flows from themotorof the bed frame to the medical attendants hand and to the patient's heart through the catheterorpacemaker wire and then to the ground of the ECG unit. In the figure below H stands forhot, N forneutral and G stands for Ground.



2) Static Electricity:



Mircoshock from static electricity

Static Electricity may be dangerous to people and sensitive equipment having integrated circuits. Sparks from static electricity could ignite flammable gases, causing an explosion. Shocks from static electricity could cause cardiac ar est if applied to a pacemakercatheter. Carpets used on the floorare a source of static electricity build up.

As illustrated in figure above an accident due to static electricity may be due to a simple vacuum cleaner.

A vacuum cleaneris plugged into a wall power outlet as the same circuitas the ECG monitorand the arterial pressure monitor. The frame of the vacuum cleaneris

connected to ground. The motorin the cleanerdue to dust collection and moisture may provide a leakage path from line to the outercasing. If a fault cur ent of 1 A flows from the vacuum cleanercasing to ground of the power board and the resistance to power board ground is 0.08 ohms, then a voltage of 80 mV is developed. For acardiac cathetersaline solution the resistance may be 500 ohms. 80 mV across 500 ohms drives 160 mA cur ent

which is very much greater than 10micro Ampere, the maximum safe cur ent through the

heart in the case of micro-shock. The micro-shock arises due to static electricity due to dust collection in the motorof the vacuum cleanerand by the saline catheteroffering a lowresistance path thereby raising the cur ent density of the heart. Even a potential difference of 5 mV across the electronic instrument probes can be potentially lethal problem in the case of micro-shock.

3) Inter uption of power:

Inter uption of electrical power to life support equipment can be hazardous. If a delay

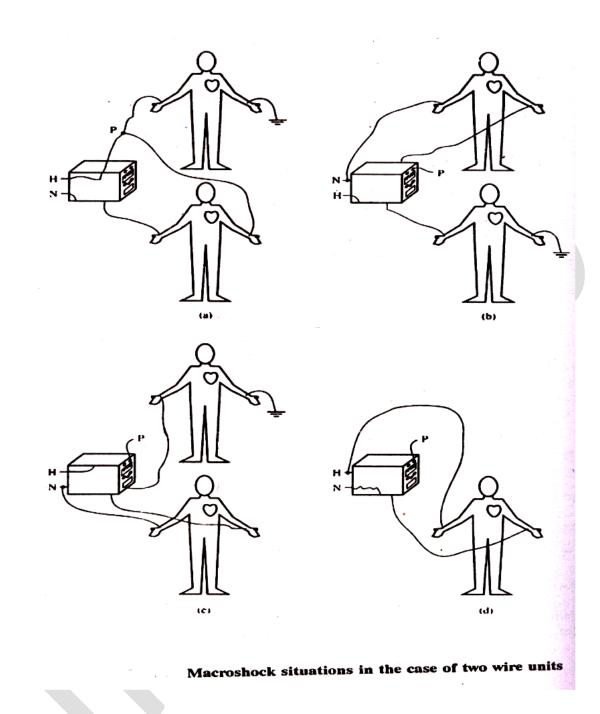
occurs before emergency power is brought into operation, the failure of a respirator, monitor, defibrillator, pacemakerorotherlife support equipment can be fatal. The possibility of a power failure must be considered in the planning of a power distribution system. Electrical service to life support system should be un-inter uptible.

Macro-shock:

A physiological response to a cur ent applied to the <u>surface of the body</u> that produces unwanted orunnecessary stimulation like muscle contractions ortissue injury is called macro-shock. All hospital patients and medical attendants are exposed to macro-shocksfrom defective electric devices and biomedical equipment.

Macro-shock Hazards:

Macro-shock occurs with two wire systems. With two wire equipment it is always dangerous to get between the hot H and neutral N wires. If the patient touches H and N wires simultaneously with two limbs, then the cur ents are flowing directly through vitalorgans of circulation and



respiration. Because N wires are internally grounded, touching H and G wires can produce macro-shock. Figure above illustrates additional hazardous situations that result from faults that may occurin the equipment. In part (a), the H lead shorts to the patient lead P. Thus a macro- shock results if the patient touches ground orthe chassis. In part (b), the hot wire H and neutral N are reserved because the two wire plug has been reserved. A grounded patient is therefore shocked upon touching the chassis. In part (c) the H wire shorted to the chassis, causing the shock configuration shown as the patient touches eitherneutral orground and the chassis. In part (d) the neutral wire accidentally shorts to the equipment case, leading to a shock situation of H to chassis or H to ground. If the H line faults to N, no shock occurs unless the patient touches H or N and ground.

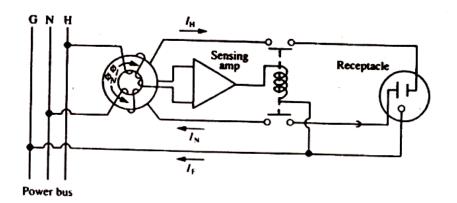
To protect from hazards of a two wire plug, a third ground wire is added as in a 3-pin plug. This wire is usually connected to the chassis of the equipment and ensures that it will not rise to a high voltage. Anothermethod is to double-insulate that chassis that is to place alayerof insulation between the circuit board chassis and the equipment case which is exposed to the user.

Devices to protect against electrical Hazards:

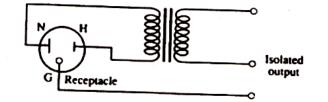
Several devices are available to protect patients from health care coworkers from hazardous electrical cur ents. These range from devices to protect against high voltage macro-shock hazards to procedures that minimize the probability that a micro-shock willoccur. Some of the devices include,

1. Ground Fault Inter upter:

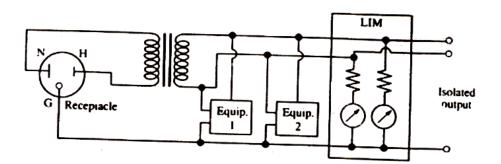
A ground fault inter upter(GFI) protects against a shock that occurs if a person touches the hot lead with one hand and the ground with the other. The GFI opens the power lead if the hot lead cur ent differs by more than approximately 2 mA from the neutral lead cur entfora duration of longerthan 0.2 second. The GFI as shown in figure belowconsists of a magnetic coil on which the hot lead and the neutral lead are mound with the same number of turns, but in opposite directions. When the system is normal, IN is equal to IH, and the magnet flux, Φ , in the coil due to these cur ents cancels. Underthis condition the sensing coil does not have a voltage induced in it. However, when the hot lead faults, oris touched by a person, the fault cur ent IF is shunted to ground. Then IN = IH – IF and IH is not equal to IN. underthis fault condition the cor esponding fluxes in the coil are unequal, and a net flux exists in the coil. This induces a voltage into the



Ground fault interrupter



(a) An isolation transformer



(b) An isolation transformer connected to a line isolation monitor and other equipment

sensing amplifier. If the cur ent IF exceeds 2 mA for 0.2 second, the relay opens the line and prevents a macro-shock from injuring the person, as well as preventing further damage to the equipment. The GFI can be conveniently mounted in the power receptacle. It is required in wet areas.

2. Isolation Transformer:

The isolation transformer provides a second means of protecting against an H-lead to G- lead macroshock. It also prevents sparks when the H lead touches ground, a particularlyimportant protection I an explosive orflammable environment, such as when flammable anesthetics orexcessive oxygen is present. Figure belowclearly shows that a fault such as a short circuit from eithersecondary lead of the transformer to ground will car y no cur ent. Therefore, a secondary lead to ground spark, orshock, is prevented. However, when the isolation transformer is in use, and equipment is plugged into the secondary, thestray capacitance and the input impedance of the hardware tend to make a conductive path to ground. This reduces the isolation by completing the circuit from eithersecondary lead to ground and then to the othersecondary lead. If a fault occurs in the secondary, a hazardous cur ent will flow.

3. Line Isolation Monitor:

A Line Isolation monitor(LIM) puts a relatively large impedance from eithersecondary leadthrough an ammeterto ground of the isolation transformer. If there is a conductive path through the equipment shown in figure below, the meterin the LIM will read a cur ent. The meteron the LIM is calibrated to read what cur ent would flowthrough a short circuit fault if it should occurfrom eithersecondary to ground. An alarm in the LIM is usually set off when it is calculated that a short circuit fault between a secondary lead and ground would draw 2 to 5 mA of cur ent. This alarm merely indicates that the back up system failed and the equipments is no longerisolated. It does not mean that the dangerous cur ents are already flowing. Therefore, if the equipment is critically needed, the LIM alarm may, sometimes, justifiably be over idden.

Power Line Interference

Electrical safety in medical environment: shock hazards – leakage cur ent-

Instruments forchecking safety parameters of biomedical equipments

Introduction:

- Medical Technology has substantially improved health care in all medical specialties and has reduced mortality forcritically ill patients.
- □ The increased complexity of medical devices and theirutilization in more procedures result in about 10,000 device related patient injuries according to a survey.
- Most of these injuries are attributable to improper use of a device as a result of inadequate training and lack of experience.
- Medical procedures usually expose the patient to more hazards than the typical home or workplace, because in medical environments the skin and mucous membranes are frequently penetrated oraltered and because there are many sources of potentially hazardous substances and energy forms that could injure the patient.
- Sources of potential hazards in medical environment include fire, air, earth, water, chemicals, drugs, microorganisms, vermin, waste, sound, electricity, natural and unnatural disasters, sur oundings, gravity, mechanical stress and people responsible foracts of omission and radiation from x-rays, ultrasound, magnets, ultraviolet light, microwaves and lasers.

□ Electrical safety is important to be considered in the medical instrumentation field. <u>Physiological Effects of electricity:</u>

- □ For a physiological effect to occur, the body must become part of an electric circuit.
- □ Cur ent must enterthe body at one point and leave at some otherpoint.
- □ The magnitude of the cur ent is equal to the applied voltage divided by the sum of theseries impedances of the body tissues and the two interfaces at the entry points.
- □ The largest impedance is often the skin resistance at the contact surface.
- □ Three phenomenons can occur when electric cur ent flows through biological tissue:
 - 4) Electric stimulation of excitable tissue (nerve and muscle)

- *5*) Resistive heating of tissues
- 6) Electrochemical burns and tissue damage fordirect cur ent and very high voltages.
- □ Psychophysical and physiological effects that occurin human as the magnitude of applied electric cur ent progressively increases.
- □ The approximate range of cur ents needed to produce each effect when 60 Hz cur ent is applied for 1 to 3 sec via copper wires that a 70 Kg human hold in each hand.

Threshold of Perception:

When the local cu**r** ent density is large enough to excite nerve endings in the skin, the subject feels a tingling sensation.

Cur ent at the threshold of perception is the minimal cur ent that an individual can detect. When someone with moistened hands grasps small copper wires, the lowest thresholds are about 0.5 mA at 60 Hz, thresholds fordc cur ent range from 2 to 10 mA.

Let – Go Cur ent:

The let - go cur ent s defined as the maximal cur ent at which the subject can withdrawvoluntarily. The minimal threshold for let - go cur ent is 6 mA. Still highercur ents cause involuntary contraction of respiratory muscles severe enough to bring about asphyxiation if the cur ent is not inter upted.

Micro-shock & Macro-shock:

Many devices have a metal chassis and cabinet that can be touched by the medical attendants and patients. If they are not grounded, then an insulation failure orshort circuitresults and leads to macro-shock ormicro-shock.

Micro-shock:

A physiological response to a cur ent applied to the <u>surface of the heart</u> that results in unwanted stimulation like muscle contractions ortissue injury is called micro-shock. Micro

-shock is caused when cur ents in excess of 10 micro Amperes flowthrough an insulated catheterto the heart. The cathetermay be an insulated, conductive fluid filled tube, ora solid wire pacemakercable.

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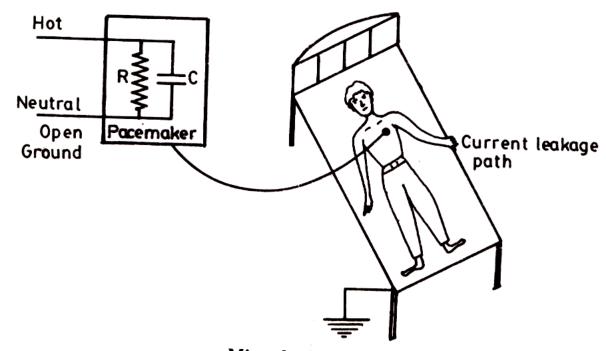
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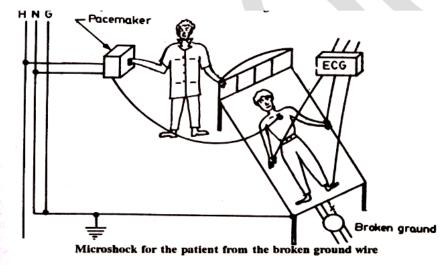


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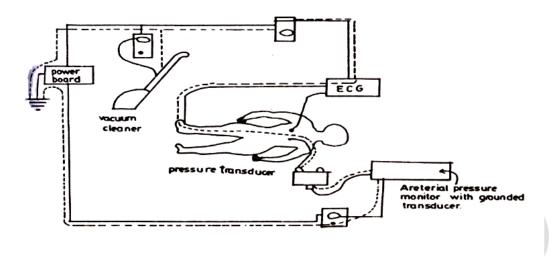
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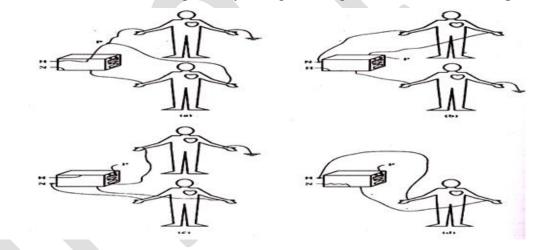
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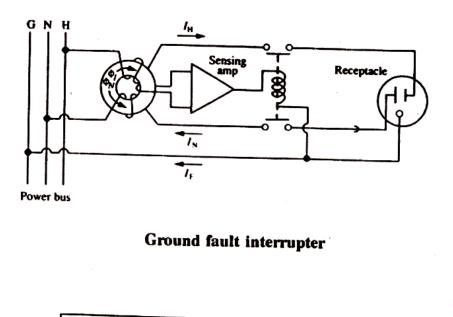
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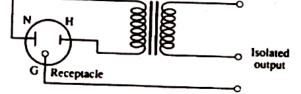
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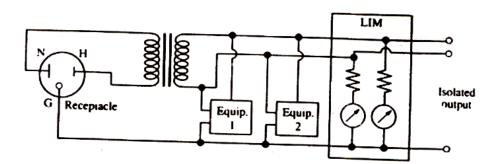
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(a) An isolation transformer



(b) An isolation transformer connected to a line isolation monitor and other equipment

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