

Digital Blood Pressure Monitor



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1. Introduction

1.1. Features of Dr Trust Aone Galaxy

Dr Trust Aone Galaxy (with integrated time/date display) is a fully automatic, digital blood-pressure measuring device for use on the arm, which enables very fast and reliable measurement of the systolic and diastolic blood-pressure as well as the pulse frequency by way of the oscillometric method of measuring. The device offers very high and clinical tested measurement accuracy and has been designed to provide a maximum of user-friendliness.

The device is intended for self-use in home.

Before using, please read through this instruction manual carefully and then keep it in a safe place.

For further questions on the subject of blood-pressure and its measurement, please contact your doctor.



1.2. Important information about self-measurement

- · Substitution of a different component might result in measurement error.
- Cuff is replaceable only by an original.
- Do not use with neonatal patients.
- · Do not intend to use with pregnant or pre-eclamptic patients
- It will cause harmful injury to the patient or affect the blood pressure due to connection tubing kinking.
- · Too frequent measurements can cause injury to the patient due to blood flow interference.
- The application of the cuff over a wound can cause further injury.
- The application of the cuff and its pressurization on any limb where intravascular access or therapy, or

an arteriovenous (A-V) shunt, is present because of temporary interference to blood flow and could result in injury to the patient.

- · Do not let the cuff and its pressurization on the arm on the side of a mastectomy
- Pressurization of the cuff can temporarily cause loss of function of simultaneously used monitoring ME equipment on the same limb.
- The need to check that operation of the automated sphygmomanometer does not result in prolonged impairment of patient blood circulation.
- Not intended to be used together with HF surgical equipment.
- Do not forget: self-measurement means control, not diagnosis or treatment. Unusual values must always be discussed with your doctor. Under no circumstances should you alter the dosages of any drugs prescribed by your doctor.
- The pulse display is not suitable for checking the frequency of heart pacemakers!
- In cases of cardiac irregularity (Arrhythmia), measurements made with this instrument should only be evaluated after consultation with the doctor.

Electromagnetic interference

The device contains sensitive electronic components (Microcomputer). Therefore, avoid strong electrical or electromagnetic fields in the direct vicinity of the device (e.g. mobile telephones, microwave cookers). These can lead to temporary impairment of the measuring accuracy.

2. Important information on the subject of blood-pressure and its measurement

2.1. How does high/low blood-pressure arise?

The level of blood-pressure is determined in a part of the brain, the so-called circulatory centre, and adapted to the respective situation by way of feedback via the nervous system. To adjust the blood-pressure, the strength and frequency of the heart (Pulse), as well as the width of circulatory blood vessels is altered. The latter is effected by way of fine muscles in the blood-vessel walls. The level of arterial blood-pressure changes periodically during the heart activity: During the «blood ejection» (Systole) the value is maximal (systolic blood-pressure value), at the end of the heart's «rest period» (Diastole) minimal (diastolic blood-pressure value). The blood-pressure values must lie within certain normal ranges in order to prevent particular diseases.

2.2. Which values are normal?

Blood pressure is too high if at rest, the diastolic pressure is above 90 mmHg and/or the systolic blood-pressure is over 160 mmHg. In this case, please consult your doctor immediately. Long-term values at this level endanger your health due to the associated advancing damage to the blood vessels in your body.

Should the systolic blood-pressure values lie between 140 mmHg and 160 mmHg and/or the diastolic blood-pressure values lie between 90 mmHg and 100 mmHg, likewise, please consult your doctor. Furthermore, regular self-checks will be necessary.

With blood-pressure values that are too low, i.e. systolic values under 100 mmHg and/or diastolic values under 60 mmHg, likewise, please consult your doctor. Even with normal blood-pressure values, a regular self-check with your blood-pressure monitor is recommended. In this way you can detect possible changes in your values early and react appropriately. If you are undergoing medical treatment to control your blood pressure, please keep a record of the level of your blood pressure by carrying out regular self-measurements at specific times of the day. Show these values to your doctor. Never use the results of your measurements to alter independently the drug doses prescribed by your doctor.

Range & broadcasting	Systolic Blood-pressure	DiastolicBlood-pressure	Measures
Hypotension	lower than 100	lower than 60	Consult your doctor
optimal	between 100 and 120	between 60 and 80	Self-check
normal	between 120 and 130	between 80 and 85	Self-check
high to normal	between 130 and 140	between 85and 90	Consult your doctor
slight hypertension	between 140 and 160	Between90and 100	Seek medical advice
medium hypertension	between 160 and 180	Between100and 110	Seek medical advice
strong hypertension	Higher than 180	Higher than 110	Urgently seek medical
			advice!

Table for classifying blood-pressure values (unit: mmHg) according to World Health Organization:

Further information

- If your values are mostly standard under resting conditions but exceptionally high under conditions of
 physical or psychological stress, it is possible that you are suffering from so-called «labile
 hypertension». Please consult your doctor if you suspect that this might be the case.
- Correctly measured diastolic blood-pressure values above 120mmHg require immediate medical treatment.

3. Various components of the blood-pressure monitor



4. Putting the blood-pressure monitor into operation

4.1 Inserting the batteries

a) Insert the batteries (4 x size AA 1.5V), thereby observing the indicated polarity.

b) If the battery warning icon appears in the display, the batteries remain 20% power to warn user the batteries will be run out.

c) If the battery warning icon appears in the display, the batteries are empty and must be replaced by new ones

Attention! • After the battery warning icon appears, the device is blocked until the batteries have been replaced.

- Please use «AA» Long-Life or Alkaline1.5V Batteries.
- The use of 1.2V Accumulators is not recommended.
- If the blood-pressure monitor is left unused for long periods, please remove the batteries from the device.

4.2 Reading the set date

Please press TIME button and the date will be shown in the display.

4.3 User selection and setting the time / date

User selection: This advanced blood pressure monitor allows you to track blood pressure readings

for 2 individuals independently

a) Before measurement, make sure you set the unit for the intended user. The unit can track results

- for 2 individuals. (User 1, User 2)
- b) Press the TIME button for at least 3 seconds. The display now indicates the set user, during which

the set user blink, to confirm, press ON/OFF button.

- c) Click the MEMORY button to select User
- d) We suggest the first person to take their pressure to be User 1.

Setting the time, date

This blood-pressure monitor incorporates an integrated clock with date display. This has the advantage, that at each measurement procedure, not only the blood-pressure values are stored, but also the exact moment of the measurement. After new batteries have been inserted, the clock begins to run TIME 12:00 and DATE 1-01.You must then re-enter the date and current time. For this, please proceed as follows.

- Press the TIME button for at least 3 seconds firstly, user icon will blink. Then press TIME button again the display now indicates the set year, during which the four characters blink.
- 2. The correct year can be entered by pressing the MEMORY button
- Press the TIME button again. The display now switches to the current date, during which the first character (month) blinks.
- 4. The corresponding month can now be entered by pressing the MEMORY button.
- 5. Press the TIME button again. The last two characters (day) starts blinking.
- 6. The corresponding day can now be entered by pressing the MEMORY button.
- Press the TIME button again. The display now switches to the current time, during which the first character (Hour) blinks.
- 8. The corresponding hour can now be entered by pressing the MEMORY button.
- 9. Press the TIME button again. The last two characters (Minutes) now blink.
- 10. The exact time can now be entered by pressing the MEMORY button.
- 11. Press TIME button (or TIME / DATE or TIME): the unit of measurement will flash.
- 12. Press the "MEMORY to set the unit of measurement (mmHg or kPa)
- Once you have made your settings, press the TIME button (or TIME / DATE or TIME). The setting is confirmed and the clock starts running.
- 14. Now after all settings have been made, press the TIME button once again. The date is briefly displayed and then the time. The input is now confirmed and the clock begins to run.

Further Information

With each press of the button (TIME, MEMORY) one input is made (e.g. switching over from hours to minutes mode, or altering the value by +1). However, if you keep the respective button depressed, you can switch more quickly to find the desired value respectively.

5. Carrying out a measurement

5.1. Before the measurement

- Avoid eating, smoking as well as all forms of exertion directly before the measurement. All these
 factors influence the measurement result. Try and find time to relax by sitting in an armchair in a
 quite atmosphere for about ten minutes before the measurement.
- Measure always on the same arm (normally left).
- Attempt to carry out the measurements regularly at the same time of day, since the blood-pressure changes during the course of the day.

5.2. Common sources of error

Note: Comparable blood-pressure measurements always require the same conditions! These are normally always quiet conditions.

- All efforts by the patient to support the arm can increase the blood-pressure. Make sure you are
 in a comfortable, relaxed position and do not activate any of the muscles in the measurement
 arm during the measurement. Use a cushion for support if necessary.
- The performance of the automated sphygmomanometer can be affected by extremes of temperature, humidity and altitude.
- Avoid compression or restriction of the connection tubing.
- A loose cuff causes false measurement values.
- With repeated measurements, blood accumulates in the respective arm, which can lead to false
 results. Correctly executed blood-pressure measurements should therefore first be repeated

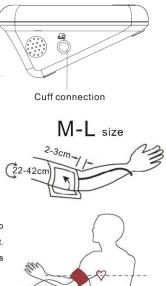
after a 5 minute pause or after the arm has been held up in order to allow the accumulated

blood to flow away (after at least 3 minutes).

5.3. Fitting the cuff

Insert air connector into air outlet shown in left photo and please make sure the fitting of the air connector completely and properly to avoid air leakage.

a) The distance between the edge of cuff and the elbow should be approx. 2~3cm.



b) Secure the cuff with the Velcro fastener, so that it lies comfortably and not too tight, whereby no space should remain between the cuff and the arm.

c) Lay the arm on a table, with the palm upwards. Support the arm a little with a rest (cushion), so that the cuff rests at about the same height as the heart. Take care, that the cuff lies free. Remain so for 2 minutes sitting quietly, before beginning with the measurement.

 d) Let legs uncrossed, feet flat on the floor, back and arm supported.

5.4. Measuring procedure

After the cuff has been appropriately positioned, the measurement can begin:

a) Press the ON/OFF button, the pump begins to inflate the cuff. In the display, the

increasing cuff-pressure is continually displayed.

b) After reaching the inflation pressure, the pump stops and the pressure slowly falls away. The cuff-pressure (large characters) is displayed during

the measurement. When the device has detected the pulse, the heart symbol in the display begins to blink for every pulse beat. c) When the measurement has been concluded. The measured systolic and diastolic blood-pressure values as well as the pulse frequency are now displayed.

Example (Fig.): Systole 118, Diastole 73, Pulse 75

The measurement results are displayed, until you switch the device off. If no button is pressed for 3 minutes, the device switches automatically off, to save the batteries.

5.5. Discontinuing a measurement

If it is necessary to interrupt a blood pressure measurement for any reason (e.g. the patient feels unwell), the "ON/OFF" power button can be pressed at any time. The device then immediately lowers the cuff-pressure automatically.

5.6. Memory - storage and recall of the measurements

The blood-pressure monitor automatically stores each of the last 120 measurement values. By pressing the MEMORY button, an average value of the last 3 measurements as well as the last

ON-DEE





measurementand the further last 120 measurements (MR119,MR118,...,MR1)can be displayed one after the other



(MR1: Values of the last measurement) (MR2-MR120: Values of the measurement before MR1)

5.7. Memory full

Pay attention that the maximum memory capacity is not exceeded. When the memory is full, the old

values are automatically over written with new ones. When memory is full , the display shown 1 second as follows to remind you " memory full "

5.8. Memory- cancellation of all measurements Attention!

Before you delete all readings stored in the memory, make sure you willnot need \ refer to the readings at a later date. Keeping a written record is prudent and may provide additional information for your doctor's visit. In order to delete all stored readings, depress the MEMORY button for at least 5 seconds, the display will show the symbol «CL» and then release the button.Topermanently clear the memory, Press the MEMORY button while «CL» is flashing. To indicate deletion of stored readings.





6. Appearance of the pulse Arrhythmia Indicator for early Detection

This symbol \checkmark indicates that certain pulse irregularities were detected during the measurement. In this case, the result may deviate from your normal blood pressure – repeat the measurement. In most cases, this is no cause for concern. However, if the symbol appears on a regular basis (e.g. several times a week with measurements taken daily) we advise you to tell your doctor. Please show your doctor the following explanation

Information for the doctor on frequent appearance of the Arrhythmia indicator

This instrument is an oscillometric blood pressure monitor that also analyses pulse frequency during measurement. The instrument is clinically tested. The arrhythmia symbol is displayed after the measurement, if pulse irregularities occur during measurement. If the symbol appears more frequently (e.g. several times per week on measurements performed daily) we recommend the patient to seek medical advice.

The instrument does not replace a cardiac examination, but serves to detect pulse irregularities at an early stage

7. Error messages /malfunctions

If an error occurs during a measurement, the measurement is discontinued and a corresponding error code is displayed.

Error No.	Possible cause(s)		
ERR 1	No pulse has been detected.		
ERR 2	Unnatural pressure impulses influence the measurement result. Reason: The arm		
	was moved during the Measurement (Artefact).		
ERR 3	The inflation of the cuff takes too long. The cuff is not correctly seated.		
ERR 5	The measured readings indicated an unacceptable difference between systolic		
	and diastolic pressures. Take other reading following directions carefully. Contact		

	you doctor if you continue to get unusual readings.		
Err8	The pressure in the cuff is too high (over 290 mmHg)		

Further Information - The level of blood-pressure is subject to fluctuations even with healthy people. Important thereby is, that comparable measurements always require the same conditions (Quiet conditions)! If, in spite of observing all these factors, the fluctuations are larger than 15mmHg, and/or you hear irregular pulse tones on several occasions, please consult your doctor. For licensing, the device has been subjected to strict clinical tests, by which the computer program used to measure the blood-pressure values was tested by experienced specialist doctors in Germany. The same computer program is used in every individual device, and has thus also been clinically tested. The manufacture of the devices takes place according to the terms of the European standard for blood-pressure measuring devices (see technical data) you must consult your specialist dealer or chemist if there are technical problems with the blood-pressure instrument. Never attempt to repair the instrument yourself! Any unauthorized opening of the instrument invalidates all warrantee claims!

Other possible malfunctions and their elimination

If problems occur when using the device, the following points should be checked and if necessary,

the corresponding measures are to be taken:

Malfunction	Remedy
The display remains empty when the	1. Check batteries for correct polarity and if necessary insert
instrument is switched on although the	correctly.
batteries are in place.	2. If the display is unusual, re-insert batteries or exchange them.
The device frequently fails to measure	1. Check the positioning of the cuff.
the blood pressure values, or the values	2. Measure the blood-pressure again in peace and quiet under
measured are too low (too high).	observance of the details made under point 5.

Every measurement produces a different value although the instrument functions	1. Please read the following information and the points listed under «Common sources of error». Repeat the measurement.
normally and the values displayed are	Please note: Blood pressure fluctuates continually so
normal	successive measurements will show some variability.
Blood pressure measured differs from	1. Record the daily development of the values and consult your
those values measured by the doctor.	doctor. Please note: Individuals visiting their doctor frequently
	experience anxiety which can result in a higher reading at the doctor
	than obtained at home under resting conditions.

8. Care and Maintenance, Recalibration

a) Do not expose the device to extreme temperatures, humidity, dust or direct sunlight.

b) The cuff contains a sensitive air-tight bubble. Handle this carefully and avoid all types of straining through twisting or buckling.

c) Clean the device with a soft, dry cloth. Do not use petrol, thinners or similar solvent. Spots on the

cuff can be removed carefully with a damp cloth and soapsuds. The cuff must not bewashed!

d) Do not drop the instrument or treat it roughly in any way. Avoid strong vibrations.

e) Never open the device! Otherwise the manufacturer calibration becomes invalid!

9. Safey, care and disposal

A Safety and protection

- This instrument may be used only for the purpose described in this booklet. The manufacturer cannot be held liable for the damage caused by incorrect application.
- This instrument comprises sensitive components and must be treated with caution. Observe the storage and operating conditions described in the "Technical specifications" section!

- Protect it from water and moisture, extreme temperatures, impact and dropping, contamination and dust, direct sunlight, heat and cold.
- The cuffs are sensitive and must be handled with care.
- Only pump up the cuff once fitted.
- Do not use the instrument close to strong electromagnetic fields such as mobile telephones or radio installations.
- · Do not use the instrument if you think it is damaged or notice anything unusual.
- If the instrument is not going to be used for a prolonged period the batteries should be removed.



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Read the additional safety instructions in the individual sections of this booklet. Ensure that children do not use the instrument unsupervised: some parts are small enough to be swallowed.

- Must use the recognized accessories, detachable parts and materials, if the use of other parts or materials can degrade minimum safety.
- · A warning to remove primary batteries if the instruments is not likely to be used for sometime

Instrument care

Clean the instrument only with a soft, dry cloth

Disposal



Batteries and electronic instruments must be disposed of in accordance with the locally applicable regulations, not with domestics waste.

10. Reference to Standards

Device standard: Device corresponds to the requirements of the European standard for

Standard (for arm BPM with adapter)

IEC60601-1-6:2010+A1:2013/ EN60601-1-6:2010+A1:2015

IEC60601-1:2005+A1:2012/EN60601-1:2006+A11:2011+A1:2013+A12:2014

IEC60601-1-2:2014/ EN60601-1-2:2015

IEC/EN60601-1-11:2015

IEC80601-2-30:2009+A1:2013/EN80601-2-30:2010+A1:2015

The stipulations of the EU-Guidelines 93/42/EEC for Medical Products Class IIa have been fulfilled.

11. Remark:

X	Some electrical and electrical equipments forbid to abandon and disposal at will	C € ⁰¹⁹⁷	TUV NO.
	Manufacturer's name and address	\$	Reading Instruction Book before use
0-3	Inapplicable baby	★	Type B equipment
Ø	Cuff Connector	Ĵ	Keep Dry





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12. Technical specifications

documents

Measurement Procedure:	Oscillometric , corresponding to Korotkoff method: Phase I : systolic , Phase V : diastolic		
Display:	Digital display		
Measuring range:	Pressure: 30 to 280 mmHg (in 1 mmHg increment) Pulse: 40 to 199 beat/minute		
Static accuracy:	Pressure: ±3mmHg / Pulse: ±5% of reading		
Measuring resolution :	1mmHg		
Inflation:	Automatic inflation by internal pump		
Memory function:	2 x 120 memories for 2 users (SYS, DIA, Pulse)		
Decompression:	Constant exhaust valve system		
Power source:	4- size "AA" alkaline Batteries (adapter : micro USB DC5.0V / 1.0A) optional		
Operation temperature:	5~40°C/41~104°F		
Operation humidity:	15%~85%RH maximum		
Storage temperature:	-10~55°C/14~131°F		
Storage humidity:	10%~95%RH maximum		

Dimensions :	135×112×71±1.0 mm			
Weight :	510 g±5g (including batteries and cuff)			
Cuff pressure display range:	0~290mmHg/0~38.7kPa			
Electrical shock protection:	Internal power unit			
Safety classifications:	Type B equipment			
Mode of operation:	Continuous operation			
Protection against ingress of water:	IP22			
Accessories:	M-size Cuff , 4 "AA" batteries, instruction manual			

Please be noticed the power adapter (micro USB DC5.0V / 1.0A) is not supplied from the origin ,users can buy

the adapter in the market which must comply to EN60601-1,EN60601-1-2

13. Manufacturer's Declaration

Dr Trust Aone Galaxy is intended for use in the electromagnetic environment specified below. The customer or the user of Dr Trust Aone Galaxy should assure that it is used in such an environment.

Emission Test	Compliance	Electromagnetic Environment	
RF emission CISPR 11	Group 1	Dr Trust Aone Galaxy uses RF energy only for	
		internal functions. Therefore, this RF emission is	
		extremely weak and there is little chance of it	
		creating any kind of interference whatsoever with	
		nearby electronic equipment.	
RF emissions CISPR 11	Class B	Dr Trust Aone Galaxy is suitable for use in all	
Harmonic emissions IEC	Not applicable	establishments, including domestic establishments	
61000-3-2		and those directly connected to the public low	

Electromagnetic Emissions: (IEC60601-1-2)

Voltage fluctuations/flicker IEC	
61000-3-3	

Not applicable voltage power supply network that supplies buildings used for domestic purposes.

Electromagnetic Immunity: (IEC60601-1-2)

Immunity test	IEC60601-1-2 test	Compliance level	Electromagnetic environment
	level		-guidance
Electrostatic	±6 kV contact	±6 kV contact	Floors should be wood, concrete or
discharge (ESD) IEC	±8 kV air	±8 kV air	ceramic tile. If floors are covered
61000-4-2			with synthetic material, the relative
			humidity should be at least 30 %.
Electric fast	±2 kV for power	Not applicable	Mains power quality should be that
transient/ burst IEC	supply lines		of a typical commercial or hospital
61000-4-4	±1 kV for		environment.
	input/output lines		
Surge IEC 61000-4-5	±1 kV differential	Not applicable	Mains power quality should be that
	mode		of a typical commercial or hospital
	±2 kV common		environment.
	mode		
Voltage dips, short	<5 % U⊤ (95% dip	Not applicable	Mains power quality should be that
interruptions and	inU_T.) for 0.5 cycle		of a typical commercial or hospital
voltage variations on	40 % U _T (60% dip in		environment. If the user of the
power supply input	$U_{\rm T}$) for 5 cycles		upper arm stlye requires continued
lines IEC 61000-4-11	70 % U _T (30% dip		operation during power mains
	inU⊤) for 25 cycles		interruptions, it is recommended
	<5 % U⊤ (95% dip		that Dr Trust Aone Galaxy be
	inU $_{\rm T}$) for 5 sec.		powered from an uninterruptible
			power supply or a battery.
Power frequency	3 A/m	Not applicable	Not applicable
(50/ 60 Hz) magnetic			
field IEC 61000-4-8			

Immunity test	IEC60601-1-2	IEC60601-1-2	2 Electromagnetic environment - guidance	
	test level	test level		
			Portable and mobile RF communications	
			equipment should be used no closer to any partof	
			Dr Trust Aone Galaxy , including cables, than	
			therecommended separation distance	
			calculatedfrom the equation applicable to the	
			frequency of the transmitter.	
			Recommend separation distance	
			3V	
			d = 1.2×p ^{1/2} 80Mhz to 800 MHz	
			d = 2.3×p ^{1/2} MHz to 2.5 GHz	
			Where P is the maximum output power rating of	
	3 Vrms 150 kHz		the transmitter in watts (W) according to he	
Conducted RF	to 80 MHz 80%		transmitter manufacturer and d is the recom-	
IEC 61000-4-6	AM (2Hz)	3 Vrms	mended separation distance in meters (m).	
			Field strengths from fixed RF transmitters as	
			determined by an electromagnetic site survey ^a ,	
Radiated RF	3 Vrms 80 MHz		should be less than the compliance level in each	
IEC 61000-4-3	to 2.5 GHz 80%	3 V/m	frequency range ^b . Interference may occur in the	
	AM (2Hz)		vicinity of equipment marked with the following	
			symbol:	
			((w))	
Note1: At 80 MH	z and 800 MHz, the h	igher frequency ra	nge applies.	
Note2: These gui	idelines may not appl	y in all situations. E	Electromagnetic propagation is affected by absorption	
and reflection from structures, objects, and people.				

^aField strengths from fixed transmitters, such as base stations for radio (cellular/cordless)telephones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcastcannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measuredfield strength in the location in which the Dr Trust Aone Galaxy is used exceeds the applicable RF compliance level above, Dr Trust Aone Galaxy should be observed to verify normal operation. Ifabnormal performance is observed, additional measures may be necessary, such as reorienting or relocating Dr Trust Aone Galaxy.

^bOver the frequency range 150 kHz to 80MHz, field strengths should be less than 3 V/m.

Recommended Separation Distances:

Recommended separation distance between portable and mobile RF communications equipment and Dr Trust Aone Galaxy.

Dr Trust Aone Galaxy is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of Dr Trust Aone Galaxy can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and Dr Trust Aone Galaxy as recommended below, according to the maximum output power of the communications equipment.

Rated maximum output	Separation distance according to frequency of transmitter m			
power of transmitter (W)	150 kHz to 80 MHz	80 MHz to 800 MHz	800 MHz to 2.5 GHz	
power of transmitter (w)	d = 1.2×p ^{1/2}	d = 1.2×p ^{1/2}	d = 2.3×p ^{1/2}	
0.01	0.12	0.12	0.23	
0.1	0.38	0.38	0.73	
1	1.2	1.2	2.3	
10	3.8	3.8	7.3	
100	12	12	23	

For transmitters rated at a maximum output power not listed above, the recommended separation distance d in meters (m) can be determined using the equation applicable to the frequency of the transmitter, where *P* is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer.

Note1:At 80MHz and 800MHz, the separation distance for the higher frequency range applies

Note2: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

For any query contact:- dr@drtrustusa.com