

Review Module J1 IRB
Level Quantum Biofeedback Practitioner

Remember to record your hours and score for your IMUNE Qualification Application.

No.	Lev	Question.....	Ans.
General			
1.	QBP	IRB stands for International Review Board	
2.	QBP	IRB is to report information to a board in order to gather data on conditions and diseases, this information is confidential and the client's name is not listed on it.	
3.	QBP	The IRB is very important for the USA/Canada as many of the facilities are experimental and it is essential that data is accumulated on the effect of these therapies.	
4.	QBP	The IRB participation is mandatory in the USA/Canada if you plan to use the portions of the program which are marked (IRB)	
5.	QBP	The IRB facilities can be used outside the USA/Canada freely	
6.	QBP	QX request firmly that ALL users use the IRB to send QX data to help in assessing the effect of the IRB therapies.	
7.	QBP	To use the IRB programs you <u>must</u> follow the informed consent process.	
8.	QBP	USA users may use any of the therapy programs including those tagged IRB without any particular consideration or client permission.	
9.	QBP	Any practitioner can use the REG tagged programs.	
10.	QBP	If your client refuses to sign an IRB participation consent form you as the technician <u>may not</u> do those sections of the program marked IRB	
11.	QBP	If you do not want to be involved in the IRB you can just ignore it and no one will know or care and then just use the program in whatever way you choose	
12.	QBP	The IRB information is in the Tools and IRB Transfer panel and there is written information regarding it also in there if the technician or client does not understand its reason to be collected.	
13.	QBP	Sending the IRB information to be collected has to be done via the internet.	
IRB Specifics			
14.	QBP	The manufacturer sees some functions of the software as experimental.	
15.	QBP	The USA law allows for studies of such software to be used on people.	
16.	QBP	USA law requires an institutional review board for experimental aspects.	
17.	QBP	The IRB is a group of independent professionals who will review the completed research of the device.	
18.	QBP	The IRB can be located anywhere in the world.	

19.	QBP	The IRB can be located only in America.	
20.	QBP	The manufacturer has an IRB arrangement in the USA	
21.	QBP	The IRB must be properly registered with any government authority where it is located.	
22.	QBP	An investigational device exemption (IRB) is only granted where there is insignificant risk to the client.	
23.	QBP	Any client who participates in the study must be allowed to decide on his participation.	
24.	QBP	A verbal acceptance of the form of consent is acceptable.	
25.	QBP	A written and signed consent form is preferable.	
26.	QBP	A practitioner must be certified or licensed to use the IRB facilities in the program	

If a question or the answer is unclear please discuss with colleagues/your trainer. It will not be possible for IMUNE to respond to individuals for clarification. If after discussing as above clarification is required then please do contact QBP@imune.net.