CONSENT WAIVER CRITERIA

To request a waiver or alteration of the required elements of informed consent, address the following criteria in Section 7 of Initial Application Part 2:

(1) The research involves no more than minimal risk to the subjects

(2) The waiver or alteration will not adversely affect the rights and welfare of the subjects

(3) The research could not practicably be carried out without the waiver or alteration

(4) Whenever appropriate, the subjects will be provided with additional pertinent information after participation.

You may request a waiver of informed consent by including responses, in your IRB application, to the above listed waiver criteria.