DATE: May 11, 2007 Description of the standard state in the standard state involving research by the IRB provides the subjects or other unanticipated or adverse events involving risks or harms to human research subjects or other unanticipated or adverse events involving risks or harms to human research subjects or other unanticipated or adverse events involving risks or harms to human research subjects or other unanticipated or adverse events involving risks or harms to human research subjects or other unanticipated or adverse events involving risks or harms to human research subjects or other unanticipated or adverse events involving risks or harms to human subjects according review and research is eligible for expective or or adverse events involving risks or harms to human research subject research. 8. Report promptly proposed changes in previously approved human subjects research involving risks or harms to human research subjects, your responsibilities include the following: 1. Report promptly proposed changes in previously approved human subject research adverse events involving risks or harms to human research subjects or others. 2. Report promptly to the IRB fine tudy is closing (i.e., data collecting and data analysis complete at Virginia Tech.) If the study is closing or experiment and state involving risks or harms to human research subjects adverse events involving risks or harms to human research subjects adverse events involving the other intinuing review with a request for continuing review and approval for the IRB before the study sector other involving human subjects adverse events involving the preval is not obtained (nules the subjects.) 2. Report promptly to be the IRB the sector the IRB before the study sector othere involving human subje	872	Virgini	aTech	Office of Research Compliance Institutional Review Board 1880 Pratt Drive (0497) Blacksburg, Virginia 24061 540/231-4991 Fax: 540/231-0959 E-mail: moored@vt.edu www.irb.vt.edu	
 TC: Marilyn D. Casto Marilyn Whitney FROM: David M. Moor Continuing Continuing Review Due Date: 5/25/2008 FROM: David M. Moor Continuation 2: "Licensing for Interior Designers: Three Case Bubles", IRB Expedited Continuation 2: "Licensing for Interior Designers: Three Case Bubles", IRB # 05-379 This memo is regarding the above referenced protocol which was previously granted expedited approval by the IRB. The proposed research is eligible for expedited review according to the specifications authorized by 45 CFR 46.110 and 21 CFR 56.110. Pursuant to your request, as Chair of the Virginia Tech Institutional Review Board, I have granted approval for extension of the study for a period of 12 months, effective as of June 9, 2007. Approval of your research by the IRB provides the appropriate review as required by federal and state laws regarding human subject research. As an investigator of human subjects, your responsibilities include the following: Report promptity proposed changes in previously approved human subject research activities to the IRB, including changes to your study forms, procedures and investigators, regardless of how minor. The proposed changes must not be initiated without IRB review and approval, except where necessary to eliminate apparent involving risks or harms to human research subjects or others. Report promptity to the IRB on the study's closing (i.e., data collecting and data analysis complete at Virginia Tech). If the study is be continue past the expiration date (listed above), investigators must submit a request for continuing review prior to the continuing review due date (listed above). It is the researcher's responsibility to obtained (uness the study has been reported to the IRB as closed) prior to the expiration date, all activities involving human subjects and data analysis must be able estis. The responsibility to obtained (uness the study has been reported to the IRB as closed) prior to the expiration date,		DATE:	May 11, 2007		
 Marilyn Whitney Approval date: 6/9/2007 Continuing Review Due Date:5/25/2008 FROM: David M. Moore Continuation 2: "Licensing for Interior Designers: Three Case Studies", IRB Expedited Continuation 2: "Licensing for Interior Designers: Three Case Studies", IRB # 05-379 This memo is regarding the above referenced protocol which was previously granted expedited approval by the IRB. The proposed research is eligible for expedited review according to the specifications authorized by 45 CFR 46.110 and 21 CFR 56.110. Pursuant to your request, as Chair of the Virginia Tech Institutional Review Board, I have granted approval for extension of the study for a period of 12 months, effective as of June 9, 2007. Approval of your research by the IRB provides the appropriate review as required by federal and state laws regarding human subject research. As an investigator of human subject, your responsibilities include the following: 1. Report promptly proposed changes in previously approved human subject research activities to the IRB. including changes to your study forms, procedures and investigators, regardless of how minor. The proposed changes must not be initiated without IRB review and approval, except where necessary to eliminate apparent immediate hazards to the subjects. 3. Report promptly to the IRB any injuries or other unanticipated or adverse events involving risks or harms to human research subjects or others. 3. Report promptly to be IRB any injuries or other unanticipated above), it is the researcher's responsibility to obtained unless the study has been reported to the IRB as a closed) prior to the expiration date (listed above), it is the researcher's responsibility to obtained unless the study has been reported to the IRB as closed) prior to the expiration date, all activities involving human subjects and data analysis complete at Virginia Tech). If the study has been reported to the IRB as closed) prior to the expiration date,		MEMORANDU	JM		
 Approval date: 6/9/2007 Continuing Review Due Date: 5/25/2008 Expiration Date: 6/8/2008 SUBJECT: IRB Expedited Continuation 2: "Licensing for Interior Designers: Three Case Studies", IRB # 05-379 This memo is regarding the above referenced protocol which was previously granted expedited approval by the IRB. The proposed research is eligible for expedited review according to the specifications authorized by 45 CFR 46.110 and 21 CFR 56.110. Pursuant to your request, as Chair of the Virginia Tech Institutional Review Board, I have granted approval for extension of the study for a period of 12 months, effective as of June 9, 2007. Approval of your research by the IRB provides the appropriate review as required by federal and state laws regarding human subject research. As an investigator of human subjects, your responsibilities include the following: Report promptly proposed changes in previously approved human subject research activities to the IRB, including changes to your study forms, procedures and investigators, regardless of how minor. The proposed changes must not be initiated without IRB review and approval, except where necessary to eliminate apparent immediate hazards to the subjects. Report promptly to the IRB of the study's closing (i.e., data collecting and data analysis complete at Virginia Tech). If the study is to continue past the expiration date (listed above), it is the researcher for responsibility to obtained re-approval is not obtained (unless the study has been reported to the IRB as a closed) prior to the continuing review due date (listed above). It is the research accided the analysis must cease immediately, except where necessary to eliminate apparent immediate hazards to the subjects. If re-approval is not obtained (unless the study has been reported to the IRB as closed) prior to the continuing review prior to the continuing review prior to the continuing review prior to the expiration date, all activities		TO:			
 Studies¹, IRB # 05-379 This memo is regarding the above referenced protocol which was previously granted expedited approval by the IRB. The proposed research is eligible for expedited review according to the specifications authorized by 45 CFR 46.110 and 21 CFR 56.110. Pursuant to your request, as Chair of the Virginia Tech Institutional Review Board, I have granted approval for extension of the study for a period of 12 months, effective as of June 9, 2007. Approval of your research by the IRB provides the appropriate review as required by federal and state laws regarding human subject research. As an investigator of human subjects, your responsibilities include the following: Report promptly proposed changes in previously approved human subject research activities to the IRB, including changes to your study forms, procedures and investigators, regardless of how minor. The proposed changes must not be initiated without IRB review and approval, except where necessary to eliminate apparent immediate hazards to the subjects. Report promptly to the IRB any injuries or other unanticipated or adverse events involving risks or harms to human research subjects for continuing review prior to the continuing review due date (listed above). It is the researcher's responsibility to obtained re-approval from the IRB before the study's expiration date. If re-approval is not obtained (unless the study has been reported to the IRB as closed) prior to the expiration date, all activities involving human subjects and data analysis must cease immediately, except where necessary to eliminate apparent immediate hazards to the subjects. 		FROM:		Continuing Review Due Date:5/25/2008	
 approval by the ĪRB. The proposed research is eligible for expedited review according to the specifications authorized by 45 CFR 46.110 and 21 CFR 56.110. Pursuant to your request, as Chair of the Virginia Tech Institutional Review Board, I have granted approval for extension of the study for a period of 12 months, effective as of June 9, 2007. Approval of your research by the IRB provides the appropriate review as required by federal and state laws regarding human subject research. As an investigator of human subjects, your responsibilities include the following: Report promptly proposed changes in previously approved human subject research activities to the IRB, including changes to your study forms, procedures and investigators, regardless of how minor. The proposed changes must not be initiated without IRB review and approval, except where necessary to eliminate apparent immediate hazards to the subjects. Report promptly to the IRB any injuries or other unanticipated or adverse events involving risks or harms to human research subjects or continue past the expiration date (listed above), investigators must submit a request for continuing review prior to the continuing review due date (listed above). It is the researcher's responsibility to obtained (re-approval from the IRB before the study's expiration date. If re-approval is not obtained (unless the study has been reported to the IRB as closed) prior to the expiration date, analysis must cease immediately, except where necessary to eliminate apparent immediate hazards to the subjects. 		SUBJECT:		"Licensing for Interior Designers: Three Case	
 Iaws regarding human subject research. As an investigator of human subjects, your responsibilities include the following: Report promptly proposed changes in previously approved human subject research activities to the IRB, including changes to your study forms, procedures and investigators, regardless of how minor. The proposed changes must not be initiated without IRB review and approval, except where necessary to eliminate apparent immediate hazards to the subjects. Report promptly to the IRB any injuries or other unanticipated or adverse events involving risks or harms to human research subjects or others. Report promptly to the IRB of the study's closing (i.e., data collecting and data analysis complete at Virginia Tech). If the study is to continue past the expiration date (listed above), investigators must submit a request for continuing meview prior to the continuing review due date (listed above). It is the researcher's responsibility to obtained re-approval from the IRB before the study's expiration date. If re-approval is not obtained (unless the study has been reported to the IRB as closed) prior to the expiration date, all activities involving human subjects and data analysis must cease immediately, except where necessary to eliminate apparent immediate hazards to the subjects. 		approval by the IRB. The proposed research is eligible for expedited review according to the specifications authorized by 45 CFR 46.110 and 21 CFR 56.110. Pursuant to your request, as Chair of the Virginia Tech Institutional Review Board, I have granted approval for extension of the study for			
 activities to the IRB, including changes to your study forms, procedures and investigators, regardless of how minor. The proposed changes must not be initiated without IRB review and approval, except where necessary to eliminate apparent immediate hazards to the subjects. Report promptly to the IRB any injuries or other unanticipated or adverse events involving risks or harms to human research subjects or others. Report promptly to the IRB of the study's closing (i.e., data collecting and data analysis complete at Virginia Tech). If the study is to continue past the expiration date (listed above), investigators must submit a request for continuing review prior to the continuing review due date (listed above). It is the researcher's responsibility to obtained re-approval from the IRB before the study's expiration date. If re-approval is not obtained (unless the study has been reported to the IRB as closed) prior to the expiration date, all activities involving human subjects and data analysis must cease immediately, except where necessary to eliminate apparent immediate hazards to the subjects. 					
		 activities to the IRB, including changes to your study forms, procedures and investigators, regardless of how minor. The proposed changes must not be initiated without IRB review and approval, except where necessary to eliminate apparent immediate hazards to the subjects. 2. Report promptly to the IRB any injuries or other unanticipated or adverse events involving risks or harms to human research subjects or others. 3. Report promptly to the IRB of the study's closing (i.e., data collecting and data analysis complete at Virginia Tech). If the study is to continue past the expiration date (listed above), investigators must submit a request for continuing review prior to the continuing review due date (listed above). It is the researcher's responsibility to obtained re-approval from the IRB before the study's expiration date. 4. If re-approval is not obtained (unless the study has been reported to the IRB as closed) prior to the expiration date, all activities involving human subjects and data analysis must cease immediately, except where necessary to eliminate 			
		cc: File		——— Invent the Future	