



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

Deidentified to
protect patient
confidentiality

AFFIX PATIENT LABEL HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given

protect patient

Name of Participant (please print)

Signature

Date

11/3/16

Name of Witness* to
Participant's Signature

protect patient

Signature

Date

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

Deidentified to
protect patient
confidentiality

AFFIX PATIENT LABEL HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print)

protect patient

Signature

Date 11/3/16

Name of Witness* to
Participant's Signature (please print)

protect patient

Signature

Date 11/3/16

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.

Participant Information and Consent Form Version 2, 27th Nov 2015



Participant Information and Consent Form
Version: 1.0 Dated: 19th Sept 2015

Deidentified to
protect patient
confidentiality

AFFIX PATIENT LABEL HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 1, dated 19th Sept 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print)

protect patient

Signature

Date

11/3/16

Name of Witness* to
Participant's Signature (please print)

protect patient

Signature

Date

11/03/16

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

Deidentified to
protect patient
confidentiality

AFFIX PATIENT LABEL HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print) [redacted]

Signature [redacted]

Date 9/3/16

Name of Witness* to Participant's Signature (please print) [redacted]

Signature [redacted]

Date 9-3-16

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.



Participant Information and Consent Form
Version: 3.0 Dated: 3rd Dec 2015

Deidentified to
protect patient
confidentiality

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 3, dated 12th Dec 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print) [redacted]

Signature [redacted]

Date 3-4-2016

Name of Witness* to Participant's Signature (please print) [redacted]

Signature [redacted]

Date 5-4-16

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

Deidentified to
protect patient
confidentiality

AFFIX PATIENT LABEL HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print) [redacted]

Signature [redacted]

Date 08/03/2016

Name of Witness* to Participant's Signature (please print) [redacted]

Signature [redacted]

Date 08/03/2016

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

Deidentified to
protect patient
confidentiality

AFFIX PATIENT LABEL HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print)

Signature

Date

Name of Witness* to
Participant's Signature (please print)

Signature

Date

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.

Participant Information and Consent Form Version 2, 27th Nov 2015



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

Deidentified to
protect patient
confidentiality

AFFIX PATIENT LABEL HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print)

Signature

protect patient

Date

Name of Witness* to
Participant's Signature (please print)

Signature

Deidentified to protect
patient confidentiality

Date

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.

Participant Information and Consent Form Version 2, 27th Nov 2015



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

Deidentified to
protect patient
confidentiality

AFFIX PATIENT LABEL HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print)

Signature

protect patient

Date

Name of Witness* to
Participant's Signature (please print)

Signature

protect patient

Date

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.

Participant Information and Consent Form Version 2, 27th Nov 2015



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

Deidentified to
protect patient
confidentiality

AFFIX PATIENT LABEL HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print)

Signature

Date

Name of Witness* to

Participant's Signature (please print)

Signature

Date

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.

Participant Information and Consent Form Version 2, 27th Nov 2015



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

Deidentified to
protect patient
confidentiality

AFFIX PATIENT LABEL HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print)

Signature

Date

Name of Witness* to

Participant's Signature (please print)

Signature

Date

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.

Participant Information and Consent Form Version 2, 27th Nov 2015



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

U.R.N
Surname
Given
Date

Deidentified to
protect patient
confidentiality

AFFIX PATIENT LABEL HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Signature

Name of Participant (please print)

Signature

protect patient

protect patient

Date

Name of Witness* to

Participant's Signature (please print)

Signature

protect patient

Date

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.

Participant Information and Consent Form Version 2, 27th Nov 2015



Participant Information and Consent Form
Version: 1.0 Dated: 19th Sept 2015

U.R.
Sur
Giv
Da

Deidentified to
protect patient
confidentiality

AFFIX PATIENT LABEL HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 1, dated 19th Sept 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print)

protect patient

Signature

Date 19/9/15

Name of Witness* to
Participant's Signature (please print)

protect patient

Signature

Date 2/12/16

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

Deidentified to
protect patient
confidentiality

AFFIX PATIENT LABEL HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print)

protect patient

Signature

Date 29/12/2016

Name of Witness* to
Participant's Signature (please print)

protect patient

Signature

Date 29/12/16

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

Deidentified to
protect patient
confidentiality

AFFIX PATIENT LABEL HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print)

Deidentified to protect patient confidentiality

Signature

Date 26-2-2016

Name of Witness* to
Participant's Signature (please print)

protect patient

Signature

Date 26-2-16

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

U R Nu
Sumarr
Given N
Date of .

Deidentified to
protect patient
confidentiality

AFFIX PATIENT LABEL HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print) Deidentified to protect patient confidentiality

Signature M 23/02/16

Name of Witness* to Participant's Signature (please print) protect patient

Signature [Signature] Date 23/02/16

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.

Participant Information and Consent Form Version 2, 27th Nov 2015



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

Deidentified to
protect patient
confidentiality

AFFIX PATIENT LABEL HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print) Deidentified to protect patient confidentiality

Signature [Signature] Date 15/2/16

Name of Witness* to Participant's Signature (please print) protect patient

Signature [Signature] Date 15/2/16

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.

Participant Information and Consent Form Version 2, 27th Nov 2015



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

Deidentified to
protect patient
confidentiality

AFFIX PATIENT LABEL HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print) Deidentified to protect patient confidentiality

Signature [Signature] Date 15/2/2016

Name of Witness* to Participant's Signature (please print) protect patient

Signature [Signature] Date 15/2/2016

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.

Participant Information and Consent Form Version 2, 27th Nov 2015



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

U.R Number [REDACTED]
 Surname [REDACTED]
 Given Name [REDACTED]
 Date of Birth [REDACTED]
 Affix Patient Label HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print) [REDACTED]

Signature [REDACTED]

Date 15-02-16

Name of Witness* to
Participant's Signature (please print) S. T. O. S. J.

Signature [REDACTED]

Date 15/2/16

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

U.R Number [REDACTED]
 Surname [REDACTED]
 Given Name [REDACTED]
 Date of Birth [REDACTED]
 Affix Patient Label HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print) [REDACTED]

Signature [REDACTED]

Date 12 Feb 16

Name of Witness* to
Participant's Signature (please print) [REDACTED]

Signature [REDACTED]

Date 12 Feb 16

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.



Participant Information and Consent Form
Version: 2.0 Dated: 27th Nov 2015

U.R Number [REDACTED]
 Surname [REDACTED]
 Given Name [REDACTED]
 Date of Birth [REDACTED]
 Affix Patient Label HERE

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print) [REDACTED]

Signature [REDACTED]

Date 9/2/16

Name of Witness* to

Participant's Signature (please print) [REDACTED]

Signature [REDACTED]

Date 9/2/16

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.

Deidentified to protect patient confidentiality

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print)

Deidentified to protect patient confidentiality

Signature  Date 9-2-16

Name of Witness* to Participant's Signature (please print)

protect patient

Signature  Date 9/2/16

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.

Deidentified to protect patient confidentiality

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr P. Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print)

Deidentified to protect patient confidentiality

Signature  Date 13/1/16

Name of Witness* to Participant's Signature (please print)

protect patient

Signature  Date 1-3-16

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.

Deidentified to protect patient confidentiality

Consent Form - Adult providing own consent

Title A prospective single centre randomised controlled clinical trial comparing the physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Short Title Physiology of apnoeic oxygenation vs. low tidal volume ventilation in anaesthetised cardiac surgical patients.

Protocol Number Version 2, dated 27th Nov 2015

Project Sponsor Department of Anaesthesia, Austin Health

Principal Investigators A/Prof Laurence Weinberg, Dr Raymond Hu

Co-Investigators Ms Laura Machan, A/Prof Philip Peyton, Dr Chong Tan, Dr Param Pillai, Dr Louise Ellard, Dr Peter McCall, Dr Ian Harley

Location Austin Health, Heidelberg, Victoria, Australia

Declaration by Participant

I have read the Participant Information Sheet or someone has read it to me in a language that I understand.

I understand the purposes, procedures and risks of the research described in the project.

I give permission for my doctors, other health professionals, hospitals or laboratories outside this hospital to release information to Austin Health concerning my disease and treatment for the purposes of this project. I understand that such information will remain confidential.

I have had an opportunity to ask questions and I am satisfied with the answers I have received.

I freely agree to participate in this research project as described and understand that I am free to withdraw at any time during the study without affecting my future health care.

I understand that I will be given a signed copy of this document to keep.

Name of Participant (please print)

protect patient

Signature  Date 8/3/16

Name of Witness* to Participant's

protect patient

Signature  Date 8/3/16

* Witness is not to be the investigator, a member of the study team or their delegate. In the event that an interpreter is used, the interpreter may not act as a witness to the consent process. Witness must be 18 years or older.