RESEARCH SUBJECT CONSENT FORM

Title:	Ezra Faster Scan Study
Protocol No.:	Ezra_002 WCG IRB Protocol #20222192
Sponsor:	Ezra AI, Inc.
Investigator:	David Girard, BSc Ezra AI, Inc. 600-419 Park Avenue South New York, NY 10016 USA David.girard @ezra.com
Study-Related Phone Number(s):	516.340.1221
Study-Related Email:	research.fasterscan@ezra.com

RESEARCH CONSENT SUMMARY

You are being asked for your consent to take part in a research study. This document provides a concise summary of this research. It describes the key information that we believe most people need to decide whether to take part in this research. Later sections of this document will provide all relevant details. We recommend that discussions occur in a private location to ensure your privacy and confidentiality.

What should I know about this research?

- Someone will explain this research to you.
- Taking part in this research is voluntary. Whether you take part is up to you.
- If you don't take part, it won't be held against you.
- You can take part now and later drop out, and it won't be held against you
- If you don't understand, ask questions.
- Ask all the questions you want before you decide.

How long will I be in this research?

We expect that your taking part in this research will be no longer than 1 month. This includes the time to schedule and provide your paid or commercial Full Body MRI service if applicable. Study activities are anticipated to take less than 2.5 hours of your time.

Why is this research being done?

Magnetic resonance imaging (MRI) is a type of non-invasive imaging technology that can "visually" detect early and late-stage cancer, and other pathologies. MRIs use a magnetic field and radio waves to create a 3-dimensional (3D) image of your body without using radiation.

The purpose of this research is to decrease the amount of time required by an individual to complete a comprehensive magnetic resonance imaging (MRI) scan, such as the Ezra Full Body MRI service, an early cancer screening tool. If the time required to conduct a comprehensive scan is reduced, the cost of the scan may also decrease. By decreasing consumer costs and time, more individuals may be able to afford this early cancer screening technology.

What happens to me if I agree to take part in this research?

If you decide to take part in this research study, and you have paid for an Ezra Full Body MRI appointment, we do not anticipate that the length of your appointment will change. We will add a short series of accelerated imaging or "research imaging" to your MRI appointment. In most cases, research imaging will require 5-10 minutes of time and your overall appointment will not exceed the original 1-hour time slot. All services conducted by Ezra AI Inc. and Ezra Health P.C. to complete your paid Full Body MRI (if applicable) will not be altered, including the return of findings. You will be required to complete a study medical questionnaire. We will provide a honorarium (credit) to you for your participation. There is no cost to you to participate, other than your time. We will use the images and data captured from your Full Body MRI scan as a reference or "control" for our research. We will compare your control images to accelerated MRI imaging to see how the changes we make impact the quality of the research images that are observed. The comparison of the images and data gathered is investigational.

If we contacted you to participate in this research study because you are a prior Ezra member or you became aware of the study through clinicaltrials.gov, we will conduct a reference or "control" MRI scan to compare to an accelerated or "research imaging" MRI scan. We expect this imaging to take less than 30 minutes. You will be required to complete a study medical questionnaire to ensure your safety. There is no cost to you to participate, other than your time. You will not receive a report from Ezra. We will provide an honorarium for your participation. We will compare your control images to accelerated MRI imaging to see how the changes we make impact the quality of the research images that are observed.

We will use these images to develop software that improves the quality of an MRI image. We will use your de-identified MRI images to test mathematical models in a process known as "machine learning". These models may enhance the quality of an image that is acquired through accelerated MRI scans making it near-indistinguishable from the reference or "control" image. The software that is developed from this study may then be applied to other MRI imaging services to decrease the time required to complete an MRI scan.

If an unknown life-threatening pathology or "incidental finding" is observed in de-identified research imaging, and it was not previously disclosed to you, you may elect to be notified. If you opt to be notified, we may re-identify your research images or associate these images to your personal information if we observe an incidental finding. We will refer your MRI research results to a board-certified radiologist for review. A radiologist may make recommendations for further diagnostic imaging. Ezra Health P.C. may review these results and contact you.

Could being in this research hurt me?

The accelerated MRI research imaging scans do not present a greater risk to you than standard MRI imaging. If you decide to participate, you will be required to complete a brief medical questionnaire. We will ask you about any implants or medical devices that you may have that may not be compatible with MRI technology and about your medical history; your answers may determine that you can not participate in this study.

During an MRI imaging session, you may experience the following:

- During your MRI session, while you lay on the MRI table, you may feel claustrophobic due to the small space surrounding you while the scan is occurring.
- If you have an implanted medical device the surrounding tissue may heat during an MRI scan potentially causing burns and discomfort. The magnetic field could cause active medical devices to malfunction or fail.
- If you have an older tattoo(s) that used metallic ink your tattoo(s) may become warm during an MRI.
- You may experience localized heating and discomfort from MRI radiofrequencies.
- If you fail to wear hearing protection during your MRI appointment, or it is not positioned properly, you may experience hearing loss. You may also experience ringing in the ear (tinnitus) after your appointment, even when you wear the provided hearing protection equipment.
- During your MRI scan, you may experience muscle or nerve stimulation, or a "twitching feeling" triggered by changing magnetic fields during your MRI.
- Any metallic objects that you fail to remove or place in the vicinity of the MRI may be rapidly drawn toward the MRI causing physical injury to you or others in the room.
- The MRI table upon which you lay during an MRI scan may suddenly move leading to pinched feet, hands or skin that may be painful. If you lose your balance and/or fall from an MRI table, you may experience injury.

If you experience any of the above, or other issue during your MRI appointment, we would like you to report it to us.

We are collecting and storing personal health information (PHI). It is possible that the privacy and security of your PHI may be compromised by illicit actors or from data breaches. Illicit use of your information from data breaches could result in discrimination which may impact your ability to obtain healthcare or life insurance, or the type of coverage provided, or negatively influence employer relationships such as hiring, promotion and/or other terms of employment.

During this study, if you elect to be informed of life-threatening pathology discovered during research MRI imaging (accelerated imaging), it may be emotionally stressful to you.

Will being in this research benefit me?

Participating in this research will not personally benefit you. In future, accelerated MRI imaging protocols and software developed from this study may reduce the costs of commercial MRI services. This may enable more individuals to use MRI technology to screen and detect early-stage cancer.

What other choices do I have besides taking part in this research?

You may decide not to participate in this study.

If you have paid for an Ezra Full Body MRI scan, you do not need to contact Ezra about your existing appointment even if you decide not to participate in this study.

What else should I know about this research?

Participation in this study is voluntary and you may withdraw your consent at any time.

To prevent fraud, we are required to confirm your identity at the beginning of a video call, and we may request that you show study personnel official government documentation such as a birth certificate, passport, or driver's license with your photo. We may also ask you security questions to confirm your identity when speaking with you over the phone. We may use digital technologies to capture your e-signature in place of a physically recorded (paper) signature.

To participate, you must have access to a computer connected to the internet, or a smart phone with a data plan or access to Wi-Fi. We may contact you via text (SMS), email, video conference or phone for the purposes of this study. We may contact you if you require assistance with study activities including questionnaires, consent form documents, appointments, reporting an incident or how to use your participant portal. You may also contact study personnel or arrange to speak with study personnel about this study.

We request that you report to us if you experience an injury or discomfort from your MRI imaging appointment by calling us, or through completion of an online form through your participant portal within 1 week. The MRI facility may also require information from you if you experience an injury during your appointment, this information will be provided to study personnel. If you experience a serious injury at the MRI facility, an on-site medical professional will direct you to appropriate medical care. If after you leave the MRI facility you become aware of an injury or experience continued discomfort, seek immediate medical attention from your physician, or call 911.

Participating in this study will not impact your medical care, nor prevent you from participating in other clinical studies or seeking other diagnostic or wellness services. This study and any MRI imaging services, or MRI imaging research performed as part of this study do not replace other FDA approved cancer screening or confirmatory diagnostics; research results from this study will not be returned to you.

DETAILED RESEARCH CONSENT

You are being invited to take part in a research study. A person who takes part in a research study is called a research subject, or research participant.

What should I know about this research?

- Someone will explain this research to you.
- This form sums up that explanation.
- Taking part in this research is voluntary. Whether you take part is up to you.
- You can choose not to take part. There will be no penalty or loss of benefits to which you are otherwise entitled.
- You can agree to take part and later change your mind. There will be no penalty or loss of benefits to which you are otherwise entitled.
- If you don't understand, ask questions.
- Ask all the questions you want before you decide.

Why is this research being done?

Magnetic resonance imaging (MRI) is a type of non-invasive imaging technology that can "visually" detect early and late-stage cancer, and other pathologies. MRIs use a magnetic field and radio waves to create a 3-dimensional (3D) image of your body without using radiation.

The purpose of this research is to decrease the amount of time required by an individual to complete a comprehensive magnetic resonance imaging (MRI) scan, such as the Ezra Full Body MRI service. If the time required to conduct a comprehensive scan is reduced, the cost of the scan may also decrease. By decreasing consumer costs and time, more individuals may be able to afford MRI-based early cancer screening technology.

A minimum of 814 participants will take part in this research. We may later increase or decrease the number of required participants depending on how well our machine learning tools perform.

How long will I be in this research?

We expect that your taking part in this research will last 1 month. This includes the time to schedule and conduct your research and/or commercial Full Body MRI service as conducted by Ezra AI Inc and Ezra Health P.C. or "Ezra". Your participant portal containing your consent form, California Bill of Experimental Rights (residents of California only), study withdrawal documentation, surveys, and incident event (injury/incident) reports submitted by you will remain accessible to you for as long as the study is actively recruiting participants, or until you withdraw from the study. This study and study-related activities may continue for 2-3 years.

Your records, including forms, questionnaires and imaging data may be securely stored by Ezra for a minimum of 2 years from the date this study is terminated or completed, or 2 years after submission and/or approval of study documentation by regulatory agencies. Any document signed by you will be retained for a minimum of 6 years from date of signature.

What happens to me if I agree to take part in this research?

If you decide to take part in this research study, and you have paid for an Ezra Full Body MRI appointment, we do not anticipate that the length of your appointment will change. We will add a short series of accelerated imaging or "research imaging" to your MRI appointment. In most cases, research imaging will require 5-10 minutes of time and your overall appointment will not exceed the original 1-hour time slot. All services conducted by Ezra AI Inc. and Ezra Health P.C. to complete your paid Full Body MRI (if applicable) will not be altered, including the return of findings. You will be required to complete a study medical questionnaire. We will provide an honorarium (credit) to you for your participation. There is no cost to you to participate, other than your time. We will use the images and data captured from your Full Body MRI scan as a reference or "control" for our research. We will compare your control images to accelerated MRI imaging to see how the changes we make impact the quality of the research images that are observed.

If we contacted you to participate in this research study because you are a prior Ezra member or you became aware of the study through clinicaltrials.gov, we will conduct a reference or "control" MRI scan to compare to an accelerated or "research imaging" MRI scan. We expect this imaging to take less than 30 minutes. You will be required to complete a study medical questionnaire to ensure your safety. There is no cost to you to participate, other than your time. You will not receive a report from Ezra. We will provide an honorarium for your participation. We will compare your control images to accelerated MRI imaging to see how the changes we make impact the quality of the research images that are observed.

We may also use these images to develop software that improves the quality of an MRI image. We will use your de-identified MRI images to test mathematical models in a process known as "machine learning". These models may enhance the quality of an image that is acquired through accelerated MRI scans making it near-indistinguishable from the reference or control image.

The software that is developed from this study may then be applied to other MRI imaging services to decrease the time required to complete an MRI scan.

This study will not impact your medical care, nor are you required to consult with study personnel before seeking medical advice or treatment. Medical advice will not be provided to you by study personnel. Your healthcare provider(s) including your primary physician are responsible for your medical care. If you have a medical emergency while participating in this study, call 911 or seek medical attention from your physician or medical care facility.

If an unknown life-threatening pathology or "incidental finding" is observed in de-identified research imaging, and it was not previously disclosed to you, you may elect to be notified. If you opt to be notified, we may re-identify your research images or associate these images to your personal information if we observe an incidental finding. We will refer your MRI research results to a board-certified radiologist for review. A radiologist may make recommendations for further diagnostic imaging. Ezra Health P.C. may review these results and contact you.

If you decide to take part in this study and you have paid for an Ezra Full Body MRI appointment or have had an Ezra MRI service in the past, study personnel may review and retain a copy of your existing medical questionnaire and records. We may "copy" or link information stored at Ezra about you that you provided to Ezra to this study's database.

Regulatory bodies, accreditation agencies, and institutional review boards (ethics review) may require documentation of your electronic health records and study results to evaluate our research findings, confirm that we keep accurate records and verify that we strictly follow our study protocol. If you are a commercial Ezra member, we may access your Ezra electronic health record (EHR) and/or Ezra Personal Health Record (PHR, MyEzra). We will retain documentation from your records related to your answers in medical questionnaires, imaging data, and Ezra reports in a secure, restricted-access database.

Occasionally, we may request that you return for an additional MRI imaging session for research purposes. You may decide to decline at that time. If you agree to an additional MRI appointment, you will be provided an additional honorarium based on the time required.

Participation in this study will not impact your medical care by your healthcare team or primary physician. If you choose to, you may inform your healthcare professional(s) about your participation in this study, including this consent document. Participating in this study does not prevent you from participating in other clinical studies. If you do participate in other clinical studies, we ask that you disclose this information (where legally permitted) to study personnel. You may disclose your participation in this study to other investigators from other clinical studies that you may currently participate in.

We may use digital health technologies, software, or services from third parties in the conduct of this study to securely store, archive or analyze data, to retrieve and/or record your medical information, to provide your participant portal, or to communicate with you. These tools may

require that you agree to end user license agreements or terms of service as a condition of use that is specific to these platforms. We may use digital technologies to capture your electronic signature and the signature of the individual conducting this consent in place of a physically recorded (pen and paper) signature.

We may process your data on restricted-access secure cloud computing servers located within the United States and Canada. We may share your de-identified data with commercial (for-profit) partners and/or academic research partners and/or study staff located in the United States or Canada for research purposes.

When you enroll, we will provide you with access to your participant portal. Your participant portal contains a copy of your signed consent form, a study withdrawal form, and an incident report form. You may submit your desire to withdraw from the study, or to file an incident report with us. Do not share your participant portal password with anyone else. If you believe your account has been compromised, please notify study personnel immediately.

During this study, or after this study ends, Ezra may conduct other research studies to develop new assays or develop other investigational devices or software. You may, if you choose (optional), provide consent to Ezra to use the information collected from this study for a future study. If you choose not to provide consent for future research, it will not impact your participation in this study, nor your choice to participate in any future study conducted by Ezra.

De-identified (anonymized) summary findings from this study may be used for publication in scientific journals, abstracts, or oral/video presentations. Your participation in this study will not be disclosed. We may deposit your de-identified data in open (publicly available) or closed access (authorized user only) databases as required for scholarly publication, or to facilitate research collaborations.

Any software or devices developed from this research study are for investigational purposes only and are not approved by the Food and Drug Administration (FDA). Results from this study may result in the development of software and/or a marketed commercial (for-profit) product (device) and may be used to support a pre-market approval submission to the FDA. If the FDA approves software or devices developed from this research study, they may become available to you through paid commercial services. You will not be financially compensated for any software or device that may be developed; except for the honorariums you receive for study participation.

We encourage you to take your time to review this study and your decision to participate before providing consent. If you choose to, you may discuss this study with your family members, friends, elders, healthcare professionals or others. You may email, schedule a phone call, or request a video call to answer any additional questions that you may have before providing consent, or at any time throughout your participation in the study. You will be sent a signed copy of this consent form. If you are a resident of California, you must receive a copy of the California Experimental Subject's Bill of Rights before signing this form.

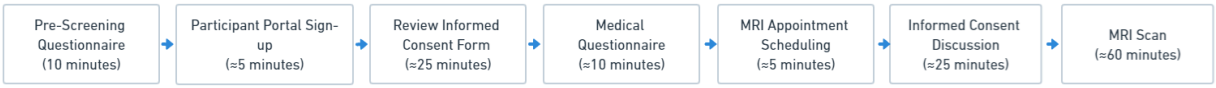
What are my responsibilities if I take part in this research?

If you decide to participate in this study, you will be required to provide, complete, or permit the following:

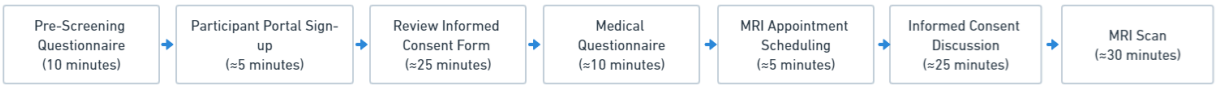
- Complete a brief questionnaire (10 minutes) about language preferences, hearing, or cognitive impairments, issues that may impact your ability to concentrate or communicate with us, or if someone is helping you complete the questionnaire.
- Permit study personnel to review your electronic health records from Ezra Health P.C. and Ezra AI Incorporated, including those contained within your MyEzra portal, and medical questionnaires completed as part of your paid Ezra services.
- Agree to sign up and use a secure online participant portal to complete study questionnaires, review consent documentation, file incident reports or withdraw from the study. You may also contact us to perform these activities.
- Agree to report any MRI-related incidents you experience during, or after your appointment by email, phone, or via your participant portal.
- Permit study staff to contact you via text (SMS), email, phone, or secure video conference at a mutually agreeable time, as required to complete study questionnaires, assist with obtaining documentation or records, assist with your participant portal, or to provide an honorarium to you for your participation.
- If you choose to, you may evaluate the person conducting this informed consent discussion, or later evaluate your MRI service through your participant portal.

We anticipate ≈2.5 hours of your time will be required to complete all study activities, including your MRI appointment.

If you paid for an Ezra Full Body Scan, your time commitment is as follows:



If you are a previous Ezra member or are joining the study from information posted on clinicaltrials.gov, your time commitment is as follows:



Could being in this research hurt me?

The accelerated MRI imaging scans added to your Full Body MRI appointment or conducted as part of a research imaging series do not present a greater risk to you than standard MRI imaging. When receiving an MRI scan, you will hear loud clicking and banging noises, sometimes

frequently, other times occasionally. Incidents requiring medical intervention related to MRI imaging occur in less than 0.29% of appointments (less than 3 people per 1000 appointments). When you schedule your paid Full Body MRI or a research MRI appointment, a medical questionnaire must be completed. These questions will ask you about any implants or medical devices that you may have that may not be compatible with MRI technology.

Less than 1 in 1000 individuals may experience the following, unless otherwise noted:

- During your MRI session, you may feel claustrophobic due to the small space surrounding you on the table that you lie upon while the scan is occurring. This could cause you anxiety (common).
- If you have an implanted medical device the surrounding tissue may heat during an MRI scan potentially causing burns and discomfort. The magnetic field could cause active medical devices to malfunction or fail.
- If you have an older tattoo(s) that used metallic ink, your tattoo(s) may become warm during an MRI.
- You may experience localized heating and discomfort from MRI radiofrequencies that may lead to skin reddening, blisters, or burns (less than 2 people per 1000).
- If you fail to wear hearing protection during your MRI appointment, or it is not positioned properly, you may experience hearing loss. You may also experience ringing in the ear (tinnitus) after your appointment, even when you wear the provided hearing protection equipment.
- During your MRI scan, you may experience muscle or nerve stimulation, or a “twitching feeling” triggered by changing magnetic fields during your MRI.
- Any metallic objects that you fail to remove or place in the vicinity of the MRI may be rapidly drawn toward the MRI causing physical injury to you or others in the room.
- The MRI table upon which you lay during an MRI scan may suddenly move leading to pinched feet, hands or skin that may be painful. If you lose your balance and/or fall from an MRI table, you may experience injury.

If you experience any of the above, trained staff at the MRI facility will assess if medical intervention is necessary and recommend you visit a healthcare facility or arrange for medical transport to a medical facility. If following your appointment, you experience continued discomfort because of your MRI appointment and it was not previously reported to the MRI facility, seek treatment from your physician, or call 911. When your medical issue has been safely addressed, inform study personnel by phone, email or through your participant portal by completing an incident report form. Ezra and study personnel will also receive an incident report from the MRI facility, and it will be added to your study record.

We are collecting and storing personal health information (PHI) related to your health. It is possible that the privacy and security of your PHI may be compromised by illicit actors or from data breaches. Illicit use of your information from data breaches could result in discrimination which may impact your ability to obtain healthcare or life insurance, or the type of coverage provided, or negatively influence employer relationships such as hiring, promotion and/or other terms of employment. If we become aware of a data breach, we will contact you. If you believe

your participant portal account has been compromised, please inform study personnel immediately.

During this study, if you elect to be informed of life-threatening pathology discovered during research MRI imaging (accelerated imaging), it may be emotionally stressful to you.

Will it cost me money to take part in this research?

Participation in this study is voluntary. There are no fees or expenses charged to you for research imaging or participating in this study. Any text message (SMS), cellular or internet fees incurred by you to participate will not be reimbursed. You will not be reimbursed for parking expenses associated with attending your MRI appointment. If you provide consent for future research use of your information, no fees or expenses will be charged to you.

This study is funded by Ezra AI Inc. The study investigator(s) and study personnel are employees and receive salary from Ezra AI Inc; they may also receive options for shares or shares of Ezra AI Inc as compensation. MRI facility employees and contractors that conduct your imaging may receive payments for their services from Ezra AI Incorporated. Physician partners that refer you to commercial services of Ezra AI Inc may receive referral incentive payments. Study consultants or investigators participating in this study may also receive payments from Ezra AI Incorporated.

Will being in this research benefit me?

Participating in this research will not personally benefit you. In future, accelerated MRI imaging protocols and software developed from this study may reduce the costs of commercial services. If you consent to the use of your information for future research (optional), new techniques or software may be developed that later become commercially available. This may enable more individuals to use MRI technology to screen and detect early-stage cancer.

What other choices do I have besides taking part in this research?

You may decide not to participate in this study. You can have your planned MRI without participating in this study. Study personnel will keep a record of your name and contact information in a study database so that we do not contact you again to participate in this study. Taking part in this research is not required to utilize other commercial (for-profit) services provided by Ezra or other future services that may become available, nor will participation in this study influence the performance of these commercial services by Ezra.

What happens to the information collected for this research?

Your private information, Ezra electronic health record(s), and information collected from this study may be shared with individuals and organizations that conduct or watch over this research, including:

- The research sponsor
- People who work with the research sponsor, including monitors and auditors
- Medical record software platform(s)
- MRI service facilities (information relevant to your appointment and safety)
- The Research and Advisory Panel of California (California residents)
- Other regulatory authorities
- Government agencies, such as the Food and Drug Administration (FDA)
- WCG IRB, the Institutional Review Board (IRB) that reviewed this research

We may publish the results of this research. However, we will keep your name and other identifying information confidential. We protect your information from disclosure to others to the extent required by law. We cannot promise complete secrecy.

A description of this clinical trial will be available on http://www.ClinicalTrials.gov, as required by U.S. Law. This Web site will not include information that can identify you. At most, the Web site will include a summary of the results. You can search this Web site at any time.

If you consent to participate in this study, we will collect information from your self-reported questionnaires. Your medical history or other reported factors may determine if you are ineligible to participate in this study. We will keep a copy of your initial consent form and any completed questionnaires in our study database so that we do not contact you again to participate in this study.

Data or information collected for this study may be de-identified and used for future research or distributed to another investigator for future research if you consent to future research (optional) without additional informed consent. De-identified data or information from this study may also be shared with other commercial (for-profit) or academic partners for research purposes during this study.

In the future, analysis of this study's results may lead to new research questions that cannot be answered by this study. These new research questions could be based on existing health information about you. If this occurs, we may create a new study that may require additional information or participation from you. You may elect to be contacted to participate in a future study.

You will receive a copy of this signed informed consent form. If you are a resident of California, you will receive a copy of the California Experimental Subject's Bill of Rights.

Who can answer my questions about this research?

If you have questions, concerns, or complaints, or think this research has hurt you or made you sick, contact the research team at 516.340.1221 or at research.fasterscan@ezra.com.

This research is being overseen by WCG IRB. An IRB is a group of people who perform independent review of research studies. You may talk to them at 855-818-2289 or researchquestions@wcgirb.com if:

- You have questions, concerns, or complaints that are not being answered by this study's research team.
- You are not getting answers from the research team.
- You cannot reach the research team.
- You want to talk to someone else about the research.
- You have questions about your rights as a research subject.

What if I am injured because of taking part in this research?

If you have an MRI-related or other injury at your scheduled appointment, trained staff at the MRI facility will assess if medical intervention is necessary and recommend you visit a healthcare facility or arrange for medical transport to a medical facility. Ezra and study personnel will receive an incident report from the MRI facility, and it will be added to your study record. If following your appointment, you experience continued discomfort because of your MRI appointment and it was not previously reported to the MRI facility, seek treatment from your primary physician, or call 911. When your medical issue has been safely addressed, inform study personnel. You may also inform study personnel through your participant portal by completing an incident report form.

Your insurance may be billed for treatment if you are injured. The sponsor will pay any charges that are not covered by insurance policy or the government, provided the injury was not due to your underlying illness or condition and was not caused by you or some other third party and was the direct result of participating in this study. No other payment is routinely available from the sponsor.

As a participant in this study, you have the right of suit to recover compensation for damages directly caused by research procedures.

Can I be removed from this research without my approval?

The person in charge of this research can remove you from this research without your approval. Possible reasons for removal include:

- You do not meet study eligibility requirements
- You are unable or fail to complete the online medical questionnaire
- You are unable to keep your scheduled MRI appointment, or can't complete your MRI session

- You decide to participate in another clinical trial or study that may negatively impact either study
- If you discover that you are pregnant before your MRI appointment
- If you refuse to follow COVID-19 safety policies that may exist at an MRI service provider
- It is in your best interest
- The research is canceled by the FDA, IRB or the sponsor

We will tell you about new information that may affect your health, welfare, or choice to stay in this research. This may include a revised consent form that you must later review and provide consent to continue participating, or to collect additional information. You will be provided an opportunity to ask questions and decide if you are willing to continue participating.

What happens if I agree to be in this research, but I change my mind later?

You can decide not to participate in this study. If you agree to take part in this research, but later change your mind, you may withdraw from the study at any time. Whatever your decision there will be no penalty or loss of benefits to which you are otherwise entitled. You may contact the study sponsor by phone, email or register your withdrawal through the participant portal. If you are comfortable providing a reason for your withdrawal, please let us know (optional). If you decide to withdraw from this study, any existing MRI image series and completed questionnaires will be retained in the study database. Information collected prior to your withdrawal may continue to be used by study personnel to ensure the integrity of our analysis, but no new information will be collected about you from any future MRI services you obtain from Ezra for this study. If you have provided consent for future research use (optional), your de-identified information and data may continue to be used for these purposes. We will keep a copy of your consent form for regulatory purposes and retain your contact information so that we do not ask you to participate in this study again. If you elected to be notified of future studies, we may contact you if you are eligible to participate.

Your participant portal may become dormant, and you may no longer be able to access documents within the study portal. It is recommended that you retain a copy of your consent form, withdrawal form, and if a California resident, a copy of the California Experimental Subject's Bill of Rights. If you are an Ezra member and have a MyEzra account, the information contained within it will remain active.

Withdrawing from this study will not prevent you from seeking MRI imaging services from Ezra in the future, nor impact existing services that you have obtained from Ezra. If you have a scheduled Ezra MRI appointment that is not part of this study, your appointment will not be impacted. If you withdraw from this study within 48 hours of your scheduled commercial MRI appointment, you may need to reschedule your commercial appointment with an Ezra care advisor. The information contained within your MyEzra personal health record from paid MRI

services will not be affected by your withdrawal from this study. Talk to an Ezra care advisor if you would like to make changes to your paid MRI imaging services from Ezra.

Your study records may be securely stored by Ezra for a minimum of 2 years from the date this study is terminated or completed, or 2 years after submission and/or approval of study documentation by regulatory agencies. Any document signed by you will be retained for a minimum of 6 years from date of signature. Image files from your MRI appointment may be stored for up to 10 years by an MRI facility. De-identified data or summary findings may be retained indefinitely.

Will I be paid for taking part in this research?

Participation in this study is voluntary. If you are receiving a commercial Full Body MRI and research imaging, your appointment is estimated to take approximately 1 hour. The addition of research imaging to your commercial MRI appointment is not anticipated to change your appointment time and you will receive an honorarium for participating. After you complete all study activities, your honorarium will be applied directly to your commercial fees which are charged to your method of payment at the conclusion of your imaging appointment.

If you are receiving research imaging only, your appointment is anticipated to take 30 minutes. If any appointment exceeds these estimated times, your honorarium (Amazon Gift Card) will increase as detailed below. You must complete your research imaging appointment to receive your honorarium. If we request an additional MRI appointment, and you accept the appointment, you will be compensated again, following research imaging guidelines below. We will send you an email after you complete your research imaging appointment that contains details on how to access your honorarium (Amazon Gift Card).

MRI Appointment Schedule	**Appointment Time**	**Honorarium**
Ezra Full Body + Research Imaging	less than 1 hour	$300 Ezra commercial service refund
Ezra Full Body + Research Imaging	less than 1.5 hours	$350 Ezra commercial service refund
Ezra Full Body + Research Imaging	less than 1.75 hours	$375 Ezra commercial service refund
Research Imaging Only	less than 30 minutes	$200 Amazon Gift Card
Research Imaging Only	30-60 minutes	$300 Amazon Gift Card

If you are a resident of California.

Any consent form and documents related to this study must be provided in the language in which you are fluent. You are also required to receive a copy of the California Experimental Subject's Bill of Rights before signing this consent form.