Nature	Dissemination level	Leading institution
R	**CO**	**AP-HP**

D6.9 - Ethical Monitoring V1

WP6 - Ethics and Social Challenges

Nature	Dissemination level	Leading institution
R	**CO**	**AP-HP**

DELIVERABLE REFERENCE NUMBER	**D6.9**
TITLE	Ethical Monitoring V1
Version	V05
LEADING BENEFICIARY	**AP-HP**
NATURE	**R - Document, report**

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DISSEMINATION LEVEL

CO	Confidential, only for members of the consortium and the Commission Services

REVISION HISTORY

PAGE 2 – TITLE 2	DATE	AUTHOR	ORGANISATION	DESCRIPTION
V.1	10.01.2025	D. Bouton	AP-HP	*First Draft*
V.2	15.01.2025	M. Camus	AP-HP	*Corrections and comments*
V .3	20.01.2025	M. Beauchesne	HUGO	*Layout*
V.4	27.01.2025	C. Demoor	HUGO	*Final Draft*
V.5	27.01.2025	M. Beauchesne	HUGO	*Final Layout*

Nature	Dissemination level	Leading institution
R	**CO**	**AP-HP**

Project number: 101136549
Duration: January 2024- December 2027

LIST OF PARTNERS

BEN – Beneficiaries / **AFF** – Affiliated partners / **ASS** – Associated partners

1.	BEN	GCS	HUGO Groupement de Coopération Sanitaire des Hôpitaux Universitaires Grand Ouest
1a.	AFF	CHUA	Centre Hospitalier Universitaire d'Angers
1b.	AFF	ICO	ICO Institut de Cancérologie de l'Ouest
2.	BEN	HUPON	Magyar Gyermekonkologiai Halozat – Magyar Gyermekonkologusok es Gyermekhematologusok Tarsasaga
3.	BEN	UNIOSL	Universitetet I Oslo
4.	BEN	CHUSE	Centre Hospitalier Universitaire Saint Étienne
5.	BEN	IGG	Istituto Giannina Gaslini
6.	BEN	RESILIENCE	Resilience
7.	BEN	OUH	Oslo Universitetssykehus HF
8.	BEN	AUH	Aarhus Universitetshospital
9.	BEN	AU	Aarhus Universitet
10.	BEN	TUCH	Varsinais-Suomen Hyvinvointialue
11.	BEN	HUS	HUS-YHTYMA
12.	BEN	ISGLOBAL	ISGlobal
13.	BEN	KDBZ	Klinika za Dječje Bolesti Zagreb
14.	BEN	CLB	Centre de Lutte Contre le Cancer Léon Bérard
15.	BEN	CINECA	Cineca Consorzio Interuniversitario
16.	BEN	UKESSEN	Universitaetsklinikum Essen
17.	BEN	CCBL	Javna Zdravstvena Ustanova Univerzitetski Klinicki Centar Republike Srpske
18.	BEN	ONKO-I	Onkoloski Institut Ljubljana
19.	BEN	EPICONCEPT	Epiconcept
20.	BEN	ALP	Asociatia Little People Romania
21.	BEN	CUSL	Cliniques Universitaires Saint-Luc ASBL
22.	BEN	AP-HP	Assistance Publique des Hôpitaux de Paris
23.	BEN	UMC-Mainz	Universitaestsmedizin der Johannes Guterberg-Universitaet Mainz

24.	BEN	PANCARE	PanCare
25.	ASS	UNILU	Universität Luzern
26.	ASS	IGR	Institut Gustave Roussy
27.	ASS	ERINTETTEK	Érintettek Egyesület
28.	ASS	AGUERRIS	Association Les Aguerris
29.	ASS	YCE	Fundatia Youth Cancer Europe
30.	ASS	RMH	The Royal Marsden National Health Service Trust

Table of contents

1. Ethical Monitoring role of AP-HP within the e-QuoL project

1.1 Ethic Structure of the project

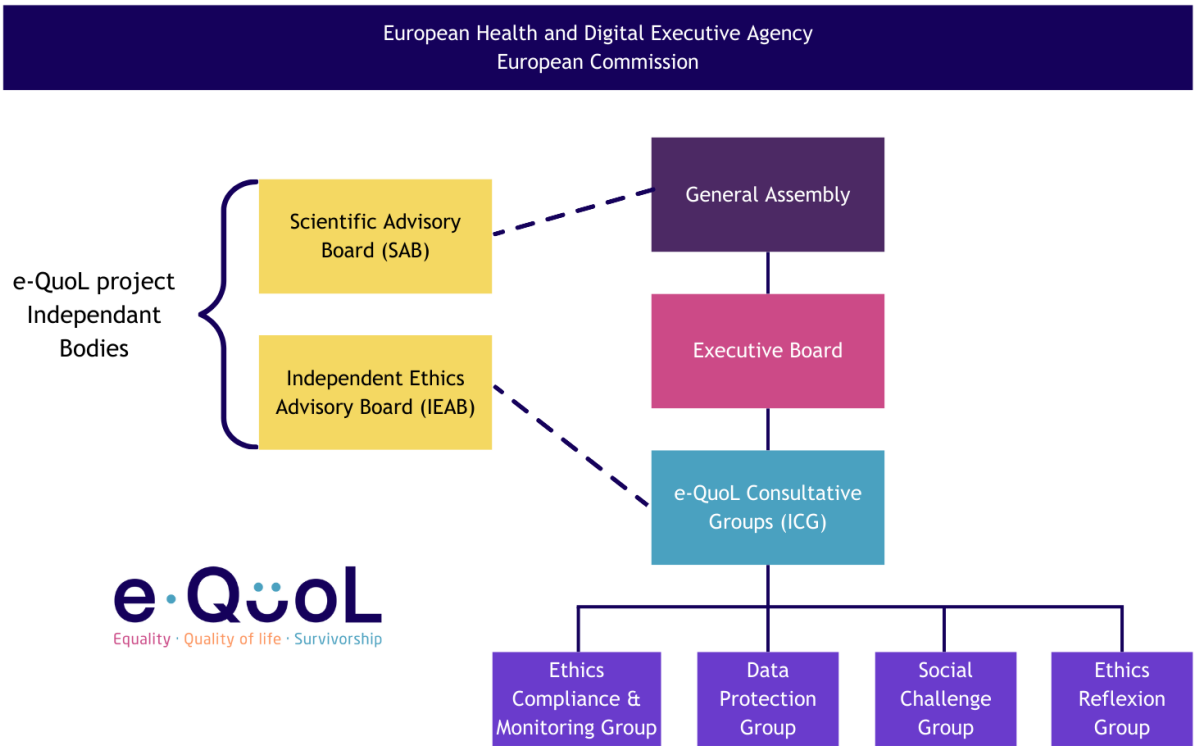

Figure 1 - Organisation scheme of the e-QuoL project, including ethics structures

1.2 Organisation of the Ethical Monitoring

AP-HP is providing ethical support on protocol, informed consent, privacy, data protection, etc. by conducting evaluation of study related documents provided to AP-HP by partners responsible for study implementation within their WPs and tasks in order to ensure compliance with ethics requirements, GDPR and national regulations. Any partner within the e-QuoL project who is wishing to have some documents evaluated or experience iossues with ethical submission are free to contact AP-HP to get help or recommendations.

2 Ethical Monitoring - report at month 12

2.1 Surveys / activities carried out during the 1st year

- February 2024, WP2, HUGO Partner: ethical assessment of the Questionnaire Specification e-QuoL "Digital tools after a childhood, adolescent or young adulthood cancer" (WP2)
    - Description of the study: this survey collected opinions from HCPs and association members regarding digital tools for CAYACS survivorship. It did not involve personal or medical data collection.
    - Ethical and Regulatory concern: none identified as the survey was concerned opinion research.
    - Conclusion: ethical submission was deemed unnecessary.
- March 2024: Assessment of Psychosocil Support Needs CAYACS - e-QuoL NS (WP2)
    - Study description: evaluation of psychosocial and health condition needs of survivors and their families using an anonymous questionnaire.

      Using a quantitative survey, the project aims to: 1) assess the information and psychosocial support needs in CAYACSs and their families across Europe; 2) analyse association with: a) cancer-related and socio-demographic characteristics, b) knowledge, skills and confidence in managing their own wellbeing, c) quality of life.
    - Method:
        - Inclusion criteria: a) CAYACSs diagnosed with cancer at age 0-25 and loved ones of CAYACSs, b) age study: 16 years old and older, c) living in Europe.
        - Recruitment strategy (depending on countries): a) direct contract by physician or in a hospital / clinic, b) patients / parents / survivors associations and foundations, c) work of mouth (participants are invited to share questionnaire in their networks), d) social media, newsletters and internet.
        - Measurements: a) information and psychosocial needs: CCSS-NAQ, b) Quality of Life: PROMIS Global Health 10 item / PROMIS Global Health Pediatric 7+2 item, c) patient activation: PAM® 13 item (patient) / CG-PAM® 13 item

(informal caregiver), d) cancer-related and sociodemographic characteristics: self-developed questions.

- Ethical concern: the protocol, in addition to psychosocial related questions, contain also health condition questions. The promotor received the following ethic approval:
  - From the Ethics Committeee Northwest and Central Switzerland (EKNZ), the Clarification of responsibility that the study does not fall under the Human Research Act (Project ID: Reg-2024-00750) (June 14th, 2024).
  - From the Institutional Review Board of the University of Lucerne, the Ehtics Board Review Clearance for the research project (IRB #2024-008) (June 19th, 2024).
- Data protection concern: the questionnaires have been answered online through web connection? IP addresses collection or any tracking method have not been used to conform with GDPR and local Data Protection authorities and avoid the necessity to get an authorisation from them. Data are centralised on a database at UNILU in Switzerland.
- Actions undertaken: 1) out of 14 participating countries, 8 required an ethical submission, 2) no IP address or tracking methods were used.
- Conclusions: the questionnaire was not spread in the countries where additional ethics approval was needed until it had been received positively.

2.2 Preparatory works for the future clinical study and qualitative study

- September 2024: preliminary review of the e-QuoL effectiveness protocol (WP5)
  - Study description: this protocol involves evaluating e-PSCT, a mobile app sharing personal and health data.
  - Ethical concerns: medical device investigation under EU regulations, potential GDPR implications.
  - Conclusion: submission to data protection authorities will be necessary In the different countries.
  - Action undertaken: timeline taken Into account for ethics approvals before the clinical study.
- November 2024: focus group on neurocognitive impairments (WP2, 3)
  - Study description: qualitative study on usability and acceptability of e-health tools for CAYACSs with neurocognitive impairments.

- Ethical concerns: it is a psychosocial investigation which does not include the collection of health data.
  - For participant unable to give informed consent and for which legal guardians or representatives informed consent is obtained, patient approval should also be seeked. In case of negative answer, they won't be able to participate.
  - To invite participants at the different meetings: email address collection for feedback should not be centralised and remain at the site where meetings are taking place. Email should be sent in a way that recipients do not see other recipients' email addresses.
- Conclusion: as it is a psychosocial investigation which does not includee the collection of health data, it does not require ethical submission.
- Action undertaken: no specific action needs to be undertaken. Nevertheless, a lot of the countries have decided to obtain an ethical approval.

2.3 Workshops and activities in Living-Labs

These activities do not collect any personal information, nor collect any health data. In this context, no ethical approval was necessary.

The participants involved in the testimonials, whether patients or professionals, have participated willingly and provided their consent by signing an image right release form. It ensures their understanding and agreement regarding the purpose and use of the materials.

Wen sending information to the user panel, anyone who is not a member of the consortium has their email addresses hidden to other recipients.

3 Annex 1 - image use authorisation form for Resilience video recordings

AUTHORIZATION FOR THE USE OF IMAGE

BETWEEN:

RESILIENCE, a simplified joint-stock company with a capital of 18,838.56 euros, registered in the Paris Trade and Companies Register under number 893 834 713, whose registered office is located at 6, rue d'Armaillé - 75017 Paris, represented Mrs. **Marion ZUNZ** as VP Patient Experience,

Hereinafter "**RESILIENCE**",

On the one hand,

AND

Mrs/Mr **NAME,** born on DATE OF BIRTH, living at ADDRESS;

Hereinafter "**You**",

On the other hand.

Article 1. Use of your Image

By the present Agreement, you accept that RESILIENCE can use your personality attributes, including your face, your image, your voice (on photos and/or videos), your name and your first name (hereinafter your "**Image**"), within the framework of interviews and video shootings, creations of RESILIENCE, intended to collect your testimony and your expertise on X [OR ALTERNATIVELY during the period from X to X].

You agree to the use of your Image by RESILIENCE worldwide and in all its forms, in relation to RESILIENCE's activities. In this context, you agree that your Image may be used by RESILIENCE on its website, the website of its Affiliates and/or its partners, its social networks or those of its Affiliates, in television programs, in publications promoting the activities of RESILIENCE or its Affiliates, and/or in mobile applications and/or technological tools developed by RESILIENCE or its Affiliates.

An Affiliate is defined as any company under the control of RESILIENCE or any company under the control of which RESILIENCE is placed or placed under the same control as RESILIENCE, the term "control" being understood here as defined in Article L233-3 of the French Commercial Code.

This use may be made by any process, any technique and on any medium, including (a) television (in particular by radio, internet, cable or satellite), (b) magnetic or digital recording media (in particular videocassettes, CD- Rom, DVD-Rom, Blu-Ray Disc, HD-DVD, hard disk, USB storage media, memory card), (c) intranet or Internet (in particular via websites, mobile applications, software or any content

distribution platform and/or on (d) paper (in particular by means of posters, flyers, posters or press), for the purposes of illustration, information, promotion and/or publicity in connection with RESILIENCE's activities.

RESILIENCE will use my Image only for the purposes and on the communication media defined above, without harming my dignity, my reputation or causing me any prejudice.

RESILIENCE will exercise all the exploitation rights attached to the recordings and/or photographs made, which will remain its exclusive property.

You also acknowledge that you are not bound by any exclusive contract on the use of your Image.

Article 2. Entry into force – Duration

This Agreement shall take effect on the date of the signature of the Agreement *OR ALTERNATIVELY* take effect retroactively from the DATE OF FILMING and for a period of twenty (20) years.

This agreement is granted worldwide.

Article 3. Counterparty

You acknowledge and expressly accept that the present transfer of your rights on your Image is granted free of charge. However, RESILIENCE undertakes to reimburse you for all reasonable travel and accommodation expenses, after prior agreement from RESILIENCE on the amounts to be incurred and upon presentation of invoices.

Article 4. Data protection

RESILIENCE collects and processes your personal data for the following purposes:

- Collection, recording and preservation of your Image;
- Photography, sound and/or video recording;
- Keeping a record so that we can contact you again.

We invite you to consult our website (section "**Privacy Policy**") for more information on the management of your personal data and your rights.

Article 5. Assignment or transfer of the Agreement

RESILIENCE will be free to transfer this authorization and all or part of the rights and obligations to any Affiliate, or by way of merger, acquisition, demerger, partial contribution of assets or transfer of business.

Article 6. Electronic signature

The Parties acknowledge and agree to use the electronic signature (e.g. via the DocuSign service) as a means of signing any contractual document (this Agreement, any amendment, etc.) in accordance with the requirements of articles 1366 et seq. of the French Civil Code. Consequently, the Parties acknowledge that this electronic signature has the same value as their handwritten signature (in particular last signature.concerning the content of the document, the identity of the signatory and his

consent to the obligations and consequences arising from the document) and to confer a date certain to that attributed to the

Article 7. Applicable law - Dispute

The Agreement shall be governed by and construed in accordance with the laws of France. Any dispute relating to its conclusion, validity, interpretation, execution and/or resolution shall fall within the exclusive jurisdiction of the Paris Court of Appeal.

Done in duplicate,

For **RESILIENCE**
Mrs Marion ZUNZ,
VP Patient Experience

For **NAME**
Mrs / M. NAME,

4 Annex 2 - image use authorisation form for MOOC production

ADVITA Productions

• FILMS • DOCUMENTARIES •

Tel: +33 (0)4 76 21 40 35

Fax: +33 (0)4 76 21 72 86

e-mail: info@advita.com

6, rue Commandant Gilot

38000 GRENOBLE

FRANCE

Image Usage Authorization

I, the undersigned, ...,

Residing at: .....

Postal Code: ..... City: ..... Country: .....

Telephone: .....

E-mail: .....

authorize ADVITA Productions to use my image as part of the MOOC "Cancer in Childhood, Living Well Afterward" produced by Michèle and Bernard Dal Molin, in collaboration with Dr. Claire Berger and Charlotte Demoor Goldschmidt, project manager Léonie Casagranda, and educational engineer Virginie Lanlo.

- Distribution on the internet or TV: sites of the project's partners and/or content distribution platforms.
- Use of all or part in the form of DVDs, animated presentations, books, CD-ROMs, or any other known or unknown medium to date, in any format, and worldwide.

I also assure that I will not request any rights or any remuneration in exchange for the use of these images in these films for any type of distribution.

Done at ....., on the .....

Signature: .....

5 Annex 3 - user panel consent form

DO YOU WANT TO PARTICIPATE IN THE RESEARCH PROJECT

E-QUOL: E-HEALTH TOOLS TO PROMOTE EQUITY IN QUALITY OF LIFE FOR CHILDHOOD TO YOUNG ADULTHOOD CANCER PATIENTS AFTER TREATMENT/SURVIVORS AND THEIR FAMILIES – DEVELOPMENT OF THE E-PSCT TOOL?

PURPOSE OF THE PROJECT AND WHY YOU ARE BEING ASKED TO PARTICIPATE

Would you like to participate in a research project aimed at developing content for a digital application for survivors of cancer that occurred in childhood, adolescence, or young adulthood and their families?

The purpose of this study is to co-create content for a phone app, with childhood, adolescent, and young adult cancer survivors (CAYACSs), their families, and health care professionals (HCPs). The phone app will provide support and information for the psychological and social (psychosocial) needs of these groups.

This study is part of a larger project, e-QuoL (e-Health tools to promote Equity in Quality of Life for childhood to young adulthood cancer patients after treatment/survivors and their families), which is a multi-country initiative. You have received this invitation to participate because you either have had cancer, you have been nominated by someone who has had cancer as a child or adolescent as a member of their “local support network”, or you are a health care professional with experience in the follow up of childhood cancer survivors. In e-QuoL, we invite people like yourself from every participating country to form our “User panel”. We do this because it is essential that the app we are developing is meeting the needs of survivors and their families across Europe. In this project, the app will then be tested out in follow up care after childhood cancer in five European countries, and will be available for every childhood cancer survivor and their families in Europe. The aim is to see if such an app can help meet the information and support needs of survivors and their families by providing tailored information and advice on how to manage possible psychosocial consequences of the cancer and its treatment.

WHAT DOES THE PROJECT ENTAIL?

If you agree to participate, you will be invited to attend co-creation workshops, which will be held online in large- and small-groups (break-out rooms). Each workshop will last approximately 2 hours and there will be a total of 6 workshops. You will be asked to provide your input and opinions on various discussion topics at each workshop. The specific activities and material presented will be determined iteratively, but some examples include brainstorming important psychosocial needs/topics and providing input on a needs screening tool that will be included in the final phone app. In some instances, you may also be asked to provide feedback on content for the app or its lay-out via emails in between workshops.

Data from these co-creation workshops and online collaborative groups will include:

- Recording of the online meetings
- Transcripts of audio recordings
- Field notes (notes written by the researchers during the discussions based on their observations of the discussions)

- Written input from the workshop participants (e.g. notes/post-it notes/digital whiteboard tools in workshops or comments in the collaborative groups)
- Basic sociodemographic information: For all: age, gender, role in workshop, country of residence, experience using health-related apps, and for survivors: Age at diagnosis, diagnosis, and whether you are in follow-up care or not.

No new health data will be collected.

POSSIBLE ADVANTAGES AND DISADVANTAGES

Possible Advantages:

- You will meet other survivors and their family members or support persons during co-creation workshops
- You will have a hands-on role in the creation of a phone application tailored to child, adolescent, and young adult cancer survivors and their families
- You will have the opportunity to test out the application which will be able to provide you with tailored information and advice regarding survivorship issues that are relevant for you.

Possible Disadvantages:

- All of the workshops will be help virtually (Zoom) and after normal working hours which may be cumbersome

Possible Risks:

- There is a possible risk of loss of anonymity. To mitigate this risk, all data will be de-identified (coded) and your name will be stored in a separate document from your data.

VOLUNTARY PARTICIPATION AND RIGHT TO WITHDRAW CONSENT

Your participation in the project is completely voluntary. If you would like to participate, please sign the consent form at the end of this document. You can withdraw your consent at any time without giving a reason. There will be no negative consequences for you if you do not want to participate or if you choose to withdraw at a later stage. If you withdraw your consent, your health data will not be used in any further research. You can request access to the data held on you, and this will be provided within 30 days.

If you want to withdraw at a later stage or have questions about the project, you can contact the project manager (see the contact details at the end of this document).

WHAT HAPPENS TO THE DATA HELD ON YOU?

The data registered about you will only be used as described under the purpose of the project, and is planned for use in for the duration of the project (December 2027) and for five years thereafter for control purposes (December 2032), after which it will be deleted. Use and storage time can only be extended after approval from the Regional Committee for Medical and Health Research Ethics and other relevant authorities. You have the right to access the information that is registered about you and to have any errors in this information corrected. You also have the right to information about the data security measures that apply to the processing of the data. You can lodge a complaint about the processing of your data to the Norwegian Data Protection Authority and the institution's Data Protection Officer.

All data will be processed and stored without names and personal identification numbers or other

directly identifiable information (i.e., coded data). A code links you to your data through a list of names. Only the project members, Hanne C. Lie and Kristen Thornton, will have access to this list. All data will be stored at the secure server for sensitive data at the University of Oslo (TSD). Everything that is discussed during the workshops will be treated as confidential and everyone in the project group with access to the data has declared confidentiality.

Publishing results is a necessary part of the research process. Data in published research will be de-identified to preserve the privacy of individual participants, but we have a duty to inform you that we cannot rule out individuals being identified.

WHAT GIVES US THE RIGHT TO PROCESS PERSONAL DATA ABOUT YOU?

We are processing information about you because the research project is considered to be in the public interest, but you have the right to object if you do not wish to be included in the project.

On behalf of *The University of Oslo,* Sikt – Norwegian Agency for Shared Services in Education and Research, has assessed that the processing of personal data in this project is in accordance with the privacy regulations.

Your rights:

As long as you can be identified in the collected data, you have the right to:

- object
- access the personal data registered about you
- have personal data about you corrected/rectified,
- have personal data about you deleted, and
- file a complaint with the Norwegian Data Protection Authority regarding the processing of your personal data.

SHARING DATA AND TRANSFERRAL ABROAD

As part of the implementation of the project, it may be relevant to transfer de-identified data held on you to our project partners at Resilience, the creator of the phone application platform that will adapt the application based off outcomes from these co-creation workshops. The Resilience tool - co-founded in 2021 in partnership with global experts and the Gustave Roussy Institute (France) - is a class IIa medical device which currently follows more than 5000 patients in France and Belgium. If this will be necessary, this will concern summaries and conclusions from the workshops, not the opinions of individual participants.

The University of Oslo (UiO) is responsible for ensuring that data are transferred in accordance with Norwegian law and the EU General Data Protection Regulation (GDPR). The code linking you to your personally identifiable information will not be disclosed.

FINANCES

Survivors of childhood cancer and their local network representatives (the non-professional user members), will receive remuneration, 600EUR for an estimated 30 hours of project participation, in recognition of their time spent in the project as user experts. Local health care professionals will not receive any such remuneration.

APPROVALS

The Regional Committee for Medical and Health Research Ethics has considered the research ethics in the project and given its approval. [Committee's case number will be entered here]

The University of Oslo and the project manager, Hanne C. Lie, are responsible for privacy and data protection in this project.
According to the new Personal Data Act (LOV-2021-06-18-124), the controller [Universitetssykehuset i Oslo and the University of Oslo] and project manager [Hanne C. Lie] have an independent responsibility to ensure that the processing of your information has a legal basis. This project has a legal basis in the EU's privacy regulation article 6 no. 1a and article 9 no. 2a and your consent.

You have the right to complain about the processing of your data to the Norwegian Data Protection Authority: postkasse@datatilsynet.no.

CONTACT DETAILS

If you have questions about the project, experience any adverse events, or want to withdraw your participation, you can contact Hanne C. Lie at h.c.lie@medisin.uio.no or [+47 22 85 14 66].
If you have questions about data protection in the project, you can contact the Data Protection Officer at the institution: The data protection officer at the University of Oslo: personvernombud@uio.no or Norwegian center for research data AS: personverntjenester@nsd.no or phone: 55582117.

I AGREE TO PARTICIPATE IN THE PROJECT AND CONSENT TO MY PERSONAL INFORMATION BEING USED AS DESCRIBED

Place and date

Participant's signature

Participant's name in block capital letters

6 Annex 4 - Ethics Board review clearance for the study "e-QuoL-NS" (UNILU)

IF, BrLe, 6002 Luzern

Dr. Anica Ilic University of
Lucerne
Faculty of Health Sciences and Medicine Alpenquai 4
6005 Lucerne

Date: June 19, 2024 Page: 1 / 1

Ethics Board Review Clearance for the research project “e-QuoL-NS: Assessment of Psychosocial Support Needs in Childhood, Adolescence, and Young Adulthood Cancer Survivors (CAYACSs) and their Loved Ones” (IRB #2024-008)

To whom it may concern

The Ethics Board at the University of Lucerne authorizes the research described in Anica Ilic’s research proposal “e-QuoL-NS: Assessment of Psychosocial Support Needs in Childhood, Adolescence, and Young Adulthood Cancer Survivors (CAYACSs) and their Loved Ones” (IRB # 2024-008).

Specifically, we have reviewed the information regarding the research plan, survey, the data management plan, and informed consent. We confirm that they comply with all applicable regulations.

We therefore approve the planned research outlined in the submitted documents.

Yours sincerely,

Prof. Dr. Leif Brandes
President of the Ethics Board

FROHBURGSTRASSE 3
P.O. BOX 6002 LUZERN

7 Annex 5 - Ethics Committee clarification for the study e-QuoL-NS (EKNZ)

EKNZ

Ethikkommission
Nordwest- und
Zentralschweiz

Präsident
Prof. Christoph Beglinger
Vizepräsidenten
Prof. Thomas Kühne
Dr. Marco Schörer

Dr. Gisela Michel
Faculty of Health Sciences and Medicine
University of Lucerne
Alpenquai 4
6005 Lucerne

Basel, 14.06.2024 / uw

Clarification of responsibility of the Ethics Committee Northwest and Central Switzerland (EKNZ)

Project-ID	Req-2024-00750
Project title	e-QuoL-NS: Assessment of psychosocial support needs in childhood, adolescence, and young adulthood cancer survivors (CAYACSs) and their loved ones
Submission Date	31.05.2024
Applicant	Dr. Gisela Michel

Decision

The research project does not fall under the scope of the Human Research Act, because your project is using only anonymised data. An authorisation from the ethics committee is therefore not required and the EKNZ is not responsible for its review.

Fees

Tariff code: 6.0 **Amount:** CHF 200.--
In accordance with the current swissethics fee schedule.

Yours sincerely,

Prof. Christoph Beglinger
President of the Ethics Committee
Northwest and Central Switzerland / EKNZ

Nature	Dissemination level	Leading institution
R	**CO**	**AP-HP**

The e-QuoL project, number 101136549, is funded by the European Union. *Views and opinions expressed are however those of the author(s) only and do not necessarily reflect those of the European Union or Horizon Europe, granting authority. Neither the European Union nor the granting authority can be held responsible for them.*

Project funded by

Schweizerische Eidgenossenschaft
Confédération suisse
Confederazione Svizzera
Confederaziun svizra

Swiss Confederation

Federal Department of Economic Affairs,
Education and Research EAER
State Secretariat for Education,
Research and Innovation SERI

This work has received funding from the Swiss State Secretariat for Education, Research and Innovation (SERI)

The UK participant (The Royal Marsden NHS Foundation Trust) in Horizon Europe Project [e-QuoL] is supported by United Kingdom Research and Innovation (UKRI), grant nº10098114.

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e·QuoL
Equality · Quality of Life · Survivorship

Funded by
the European Union