



Dear Sir/Madam,

We are writing regarding the Swiss Personalised Health Network. In order to ensure global privacy of sensitive genetic information, we are seeking clarification on your data sharing policies.

We note that the project is a public-private partnership with industrial partners, and thus there will be access to data by third parties. We thus seek clarification on who may register for data access, including private companies, insurance companies, law enforcement or immigration services, and exactly what data is shared, and if it is identifiable information.

We further seek clarification what privacy protections are in place to prevent access to identifiable genetic (including re-identifiable) information in the hands of the project and any third parties by law enforcement and immigration services. This includes what policies are in place to prevent data breaches by your organisation or third party collaborators. Further, we seek clarification on whether such data is sold to third party companies.

We also request access to the consent forms that do not appear to be publicly available on the website. Moreover, we could not find any information on what occurs to data following withdrawal of participants and whether data is retained or deleted, and request clarification on this.

The information that is provided, or not provided to us, will be shared with the public in order to promote transparency and to inform on the privacy implications of growing genetic databases. We thus thank you for any information that you can share with us to facilitate this process.

Yours sincerely,

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