

# GAPS allowable uses and register linkage

If study subgroups or cases and controls have different information, please use one form for each.

Study name: \_\_\_\_\_

Study subgroup (if applicable): \_\_\_\_\_

ICD codes and phenotypes allowed for study by GAPS: \_\_\_\_\_  
\_\_\_\_\_

Can these samples be used to calculate and report aggregated Sweden allele frequencies? \_\_\_\_\_

Can these samples be used for UNICORN (used in aggregate as controls in association studies)? \_\_\_\_\_

Can these samples be uploaded to Sanger HRC for imputation? \_\_\_\_\_

Is it permitted for the Honest Broker to link the genetic data to the national registers? \_\_\_\_\_

Is re-contact possible? \_\_\_\_\_ By genotype, phenotype or both? \_\_\_\_\_

## Register linkage

<u>Register</u>	<u>Linkage allowed</u>		<u>Linkage done</u>	
Patient register	yes	no	yes	no
Cause of death register	yes	no	yes	no
Prescribed drug register	yes	no	yes	no
Multi-generation register	yes	no	yes	no
Conscription	yes	no	yes	no
Grades, testing	yes	no	yes	no
Medical birth register	yes	no	yes	no
SES (family status, income, education)	yes	no	yes	no
Population register (birthplace, migration)	yes	no	yes	no
Other _____	yes	no	yes	no
Other _____	yes	no	yes	no

What month/year was the last register linkage done? \_\_\_\_\_

Additional comments \_\_\_\_\_  
\_\_\_\_\_

Signature \_\_\_\_\_