

***INQUIRY***

**Regarding the use of data/biomaterial and/or the infrastructure of the Competence Network and the National Register for Congenital Heart Defects**

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| *Please e-mail to*ubauer@kompetenznetz-ahf.deCEO of the National Register for Congenital Heart Defects (**NRCHD**) and the Competence Network for Congenital Heart Defects (**CNCHD**)Dr. Ulrike Bauer |

***(Details of this inquiry will be treated as confidential)***

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| **1 Research topic/title**      |
| **2 Principal investigator(s)**First name/surname      Organisation/institution      Address:      Tel./Fax:      Email:      Co-investigator(s):       |
| **3 Short description of research question and aims**      |
| **4 Description of patient population**Diagnoses, phenotype, sample size, inclusion/exclusion criteria      |
| **5 Detailed description of data that are required for evaluation/calculation****Data that we generally provide*:*** *sex, age, main and secondary diagnoses, hereditary, fetal and neonatal diagnoses (linked to IPCCC and/or ICD10). Additional data in the case of biomaterial: age at sampling***All further anamnestic and clinical data upon request (please add a table, if appropriate)**      |
| **6 Required biomaterial** (if applicable)Type/amount of samples      |