

***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.de](mailto:ubauer@kompetenznetz-ahf.de)  CEO 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 |