



Additional patient information for participation in central data collection for national quality assurance in the treatment of congenital heart defects

Dear patient, dear parents,

the European Union's General Data Protection Regulation (EU-GDPR) applies from the 25th of May 2018. This means some changes for you. Among other things, you have **more information rights** than before.

We would like to inform you that you have the **right to complain** to the responsible data protection supervisory body if you believe that data protection provisions are being violated. This is as follows for the upholding of data protection regulations in the National Register:

Berliner Beauftragte für Datenschutz und Informationsfreiheit
Friedrichstr. 219, 10969 Berlin, Tel: 030 13889-0, Fax: 030 2155050
E-mail: mailbox@datenschutz-berlin.de,
www.datenschutz-berlin.de

The following is responsible for any possible violations of data protection regulations **in the clinic responsible for your treatment:**

(please enter here)

You can request information about the data saved about you at any time. You can also request its deletion. Furthermore, you also have the **right to receive information and free copies of documents** concerning the data collected about you or your child during the course of our research. The 'Nationale Register für angeborene Herzfehler e. V.' is responsible for data processing

Contact:
Nationales Register für angeborene Herzfehler e. V.
Managing Director: Dr Ulrike Bauer
Augustenburger Platz 1, 13353 Berlin
Tel 030 4593 7276
ubauer@kompetenznetz-ahf.de
The National Register's data protection agent is
Mr Stefan Beudt
beudt@kompetenznetz-ahf.de



Patient's personal information

Surname: _____

Forename: _____

Date of birth: _____

I hereby declare my willingness to continue my participation in the project outlined above.

Location, date

Patient/guardian signature

Location, date

2nd guardian* signature

* Both guardians' signatures are required if the patient is a minor