Non-therapeutic research with minors: how do chairpersons of German research ethics committees decide?

C Lenk, K Radenbach, M Dahl, C Wiesemann

**Objective:** Clinical trials in humans in Germany—as in many other countries—must be approved by local research ethics committees (RECs). The current study has been designed to document and evaluate decisions of chairpersons of RECs in the problematic field of non-therapeutic research with minors. The authors’ purpose was to examine whether non-therapeutic research was acceptable for chairpersons at all, and whether there was certainty on how to decide in research trials involving more than minimal risk.

**Design:** In a questionnaire, REC chairpersons had to evaluate five different scenarios with (in parts) non-therapeutic research. The scenarios described realistic potential research projects with minors, involving increasing levels of risk for the research participants. The chairpersons had to decide whether the respective projects should be approved.

**Methods:** A total of 49 German REC chairpersons were sent questionnaires; 29 questionnaires were returned. The main measurements were approval or rejection of research scenarios.

**Results:** Chairpersons of German RECs generally tend to accept non-therapeutic research with minors if the apparent risk for the participating children is low. If the risk is clearly higher than “minimal”, the chairpersons’ decisions differ widely.

**Conclusion:** The fact that there seem to be different attitudes of chairpersons to non-therapeutic research with minors is problematic from an ethical point of view. It suggests a general uncertainty about the standards of protection for minor research participants in Germany. Therefore, further ethical and legal regulation of non-therapeutic research with minors in Germany seems necessary.

For the last ten years in Germany, so called non-therapeutic research with study participants not competent to consent has been a target of passionate criticism. For example, in 1997, the “Eisingen case” caused a stir when a trainee of the St Josef’s Home for the Handicapped in Bavaria blew the whistle on human genetic research with mentally handicapped inmates that had taken place without informed or proxy consent. A doctoral candidate from the Institute for Human Genetics of the University of Wuerzburg had taken blood samples from 179 residents without informing the people concerned or their parents or legal guardians. Criticism focussed not only on the lack of consent but also, and predominantly, on the alleged immorality of research without potential direct benefit.

The public reaction was fuelled by a debate in Germany on the European Convention on Human Rights and Biomedicine, in particular article 17 of the convention which deals with the “Protection of persons not able to consent to research”. Human rights activists objected to the permission of minimal risk research with the non-competent. The Church and organisations for the disabled, in particular, warned against a deterioration of existing ethical and legal standards in Germany, should the convention be ratified. Moreover, for the same reason the revision of the Helsinki Declaration on Human Research (Edinburgh 2002), which allowed for this type of research, was received critically by German lawyers as well as the public. It is doubtful, however, whether Germany has reached high standards in the ethical and legal regulation of medical research.

As in most industrialised countries, in Germany clinical trials involving humans must be approved by a local research ethics committee (REC). The relevant law on the approval of prescription drugs (Arzneimittelgesetz/AMG) demands that researchers ensure the safety of participants in clinical testing. The law regulates clinical testing according to basic standards in medical ethics: minimisation of risks for study participants, informed consent, and good clinical practice. In addition, clinical research with minors has to fulfil the following requirements: the drug in question is designed for the diagnosis or treatment of paediatric diseases, clinical trials with adults are not expected to yield adequate results, and parents (and if possible the child too) have to consent to study participation. However, the provisions in the AMG have been criticised for not being precise and detailed enough. It remains unclear, for example, whether the law allows for non-therapeutic research in minors or not (for a critical appraisal of this type of research see Nicholson and Brock).

Further regulations are to be found in a statement of the “Central Ethics Commission” (Zentrale Ethikkommission) at the German Federal Medical Council “On protecting non-competent persons in medical research.” According to these guidelines, minimal risk or burden in certain cases of non-therapeutic research may be acceptable even for vulnerable groups. Only if a vulnerable person shows significant unwillingness to participate in a study, has research to be stopped. However, the status of these recommendations is unclear, as they might be in conflict with existing laws. This situation creates a remarkable amount of uncertainty for the public, as well as for RECs.

The REC at Goettingen University, for example, in 2002 received a total of 13 research protocols dealing with minors. Eleven of the 13 protocols had elements of non-therapeutic research. So far, little is known about decisions German RECs actually make in the case of non-therapeutic research with minors: How do chairpersons decide in these cases? Are they willing to accept studies with no direct benefit for the
Using an additional 5 ml blood from a blood sample primarily drawn for diagnostic/therapeutic reasons.

2 Various tests with healthy children: neurological examination, electroencephalogram, hearing test, questionnaires (duration: 7–8 hours over a period of 3 days).

3 Additional myocardial biopsy in the course of a heart operation performed for therapeutic purposes.

4 Additional bone marrow biopsies in leukaemia patients (6 out of 10 exclusively for non-therapeutic reasons).

5 Controlled clinical trial with toddlers, involving a placebo group which would have to undergo several intramuscular injections of sodium chloride solution.

Figure 1 Evaluation of five different studies by chairpersons of German research ethics committees (n = 29).

Results of the respondents are shown in the figure. From the respondents’ point of view, there was a significant difference between the appraisals of studies 1 and 2, and studies 3, 4, and 5. While research scenarios 1 and 2 would have been approved by all REC chairpersons (although sometimes with slight restrictions), trials 3 to 5 were strongly rejected by a number of participants. In our view, this division of opinion is caused by the far more invasive character of the latter research projects. While the study design of research projects 1 and 2 involved only a very small risk for the participants’ health, the medical interventions in studies 3 to 5 were far riskier and could have caused severe side effects.

The design of study 1 had the highest acceptance (28/35; 80%). It included the use of an additional 5 ml blood from a blood sample for diagnostic reasons from children with a non-specific mental retardation. Although the study design as far as risks are concerned strongly resembled the German “Eisingen case” (see above), it seemed quite acceptable to chairpersons. The respondents could add comments to their evaluation of the respective research projects. Most of the comments on the evaluation of study 1 correctly referred to the necessity of obtaining the parents’ or children’s informed consent or assent.

The proposal for study 4 received the highest number of rejections. Children at the age of 6–11 suffering from acute lymphatic leukaemia were supposed to participate in a randomised, two armed study. Its purpose was to compare the outcome of a new chemotherapy with standard therapy. In the course of the study, weekly bone marrow biopsies were to be performed in both study groups. Six out of 10 bone marrow biopsies were intended for purely scientific purposes, without direct benefit for the participant involved. Most of the respondents (17/58.6%) refused approval of the study. In our view, this difference between the appraisals of studies 1 and 2, and studies 3, 4, and 5. While research scenarios 1 and 2 would have been approved by all REC chairpersons (although sometimes with slight restrictions), trials 3 to 5 were strongly rejected by a number of participants. In our view, this division of opinion is caused by the far more invasive character of the latter research projects. While the study design of research projects 1 and 2 involved only a very small risk for the participants’ health, the medical interventions in studies 3 to 5 were far riskier and could have caused severe side effects.

The design of study 1 had the highest acceptance (28/35; 80%). It included the use of an additional 5 ml blood from a blood sample for diagnostic reasons from children with a non-specific mental retardation. Although the study design as far as risks are concerned strongly resembled the German “Eisingen case” (see above), it seemed quite acceptable to chairpersons. The respondents could add comments to their evaluation of the respective research projects. Most of the comments on the evaluation of study 1 correctly referred to the necessity of obtaining the parents’ or children’s informed consent or assent.

The proposal for study 4 received the highest number of rejections. Children at the age of 6–11 suffering from acute lymphatic leukaemia were supposed to participate in a randomised, two armed study. Its purpose was to compare the outcome of a new chemotherapy with standard therapy. In the course of the study, weekly bone marrow biopsies were to be performed in both study groups. Six out of 10 bone marrow biopsies were intended for purely scientific purposes, without direct benefit for the participant involved. Most of the respondents (17/58.6%) refused approval of the study. In our view, this difference between the appraisals of studies 1 and 2, and studies 3, 4, and 5. While research scenarios 1 and 2 would have been approved by all REC chairpersons (although sometimes with slight restrictions), trials 3 to 5 were strongly rejected by a number of participants. In our view, this division of opinion is caused by the far more invasive character of the latter research projects. While the study design of research projects 1 and 2 involved only a very small risk for the participants’ health, the medical interventions in studies 3 to 5 were far riskier and could have caused severe side effects.

The design of study 1 had the highest acceptance (28/35; 80%). It included the use of an additional 5 ml blood from a blood sample for diagnostic reasons from children with a non-specific mental retardation. Although the study design as far as risks are concerned strongly resembled the German “Eisingen case” (see above), it seemed quite acceptable to chairpersons. The respondents could add comments to their evaluation of the respective research projects. Most of the comments on the evaluation of study 1 correctly referred to the necessity of obtaining the parents’ or children’s informed consent or assent.

The proposal for study 4 received the highest number of rejections. Children at the age of 6–11 suffering from acute lymphatic leukaemia were supposed to participate in a randomised, two armed study. Its purpose was to compare the outcome of a new chemotherapy with standard therapy. In the course of the study, weekly bone marrow biopsies were to be performed in both study groups. Six out of 10 bone marrow biopsies were intended for purely scientific purposes, without direct benefit for the participant involved. Most of the respondents (17/58.6%) refused approval of the study. In our view, this difference between the appraisals of studies 1 and 2, and studies 3, 4, and 5. While research scenarios 1 and 2 would have been approved by all REC chairpersons (although sometimes with slight restrictions), trials 3 to 5 were strongly rejected by a number of participants. In our view, this division of opinion is caused by the far more invasive character of the latter research projects. While the study design of research projects 1 and 2 involved only a very small risk for the participants’ health, the medical interventions in studies 3 to 5 were far riskier and could have caused severe side effects.

The design of study 1 had the highest acceptance (28/35; 80%). It included the use of an additional 5 ml blood from a blood sample for diagnostic reasons from children with a non-specific mental retardation. Although the study design as far as risks are concerned strongly resembled the German “Eisingen case” (see above), it seemed quite acceptable to chairpersons. The respondents could add comments to their evaluation of the respective research projects. Most of the comments on the evaluation of study 1 correctly referred to the necessity of obtaining the parents’ or children’s informed consent or assent.

The proposal for study 4 received the highest number of rejections. Children at the age of 6–11 suffering from acute lymphatic leukaemia were supposed to participate in a randomised, two armed study. Its purpose was to compare the outcome of a new chemotherapy with standard therapy. In the course of the study, weekly bone marrow biopsies were to be performed in both study groups. Six out of 10 bone marrow biopsies were intended for purely scientific purposes, without direct benefit for the participant involved. Most of the respondents (17/58.6%) refused approval of the study. In our view, this difference between the appraisals of studies 1 and 2, and studies 3, 4, and 5. While research scenarios 1 and 2 would have been approved by all REC chairpersons (although sometimes with slight restrictions), trials 3 to 5 were strongly rejected by a number of participants. In our view, this division of opinion is caused by the far more invasive character of the latter research projects. While the study design of research projects 1 and 2 involved only a very small risk for the participants’ health, the medical interventions in studies 3 to 5 were far riskier and could have caused severe side effects.

The design of study 1 had the highest acceptance (28/35; 80%). It included the use of an additional 5 ml blood from a blood sample for diagnostic reasons from children with a non-specific mental retardation. Although the study design as far as risks are concerned strongly resembled the German “Eisingen case” (see above), it seemed quite acceptable to chairpersons. The respondents could add comments to their evaluation of the respective research projects. Most of the comments on the evaluation of study 1 correctly referred to the necessity of obtaining the parents’ or children’s informed consent or assent.

The proposal for study 4 received the highest number of rejections. Children at the age of 6–11 suffering from acute lymphatic leukaemia were supposed to participate in a randomised, two armed study. Its purpose was to compare the outcome of a new chemotherapy with standard therapy. In the course of the study, weekly bone marrow biopsies were to be performed in both study groups. Six out of 10 bone marrow biopsies were intended for purely scientific purposes, without direct benefit for the participant involved. Most of the respondents (17/58.6%) refused approval of the study. In our view, this difference between the appraisals of studies 1 and 2, and studies 3, 4, and 5. While research scenarios 1 and 2 would have been approved by all REC chairpersons (although sometimes with slight restrictions), trials 3 to 5 were strongly rejected by a number of participants. In our view, this division of opinion is caused by the far more invasive character of the latter research projects. While the study design of research projects 1 and 2 involved only a very small risk for the participants’ health, the medical interventions in studies 3 to 5 were far riskier and could have caused severe side effects.

The design of study 1 had the highest acceptance (28/35; 80%). It included the use of an additional 5 ml blood from a blood sample for diagnostic reasons from children with a non-specific mental retardation. Although the study design as far as risks are concerned strongly resembled the German “Eisingen case” (see above), it seemed quite acceptable to chairpersons. The respondents could add comments to their evaluation of the respective research projects. Most of the comments on the evaluation of study 1 correctly referred to the necessity of obtaining the parents’ or children’s informed consent or assent.
In the course of the study, cases of myocarditis in each group would be counted. Study participants with myocardial inflammation were to receive standard treatment. As the chart in the figure shows, the placebo controlled study (5) was more easily accepted by chairpersons than study 4. Seven of the respondents (24.1%) would have approved the study without further restrictions, while 12 (41.4%) would have rejected it.

CONCLUSION
The results of our study show that in the case of non-therapeutic research with minors involving a higher than minimal risk, decisions of chairpersons of German RECs vary to a disturbingly high degree. In contrast to widespread public criticism, chairpersons do not object on principle to non-therapeutic research with only minimal risk or burden (for a critical discussion of the notion “minimal risk” see Kopelman and Maio). There seems to be a kind of consensus among chairpersons with regard to the evaluation of less invasive interventions for non-therapeutic research, as the undisputedly positive evaluation of studies 1 and 2 has shown. More invasive interventions like those involved in studies 3 to 5, however, are seen as more controversial. This reflects the ethical discussion about non-therapeutic research with minors in Germany. From our point of view, it is disturbing that RECs in Germany seem to arrive at widely differing decisions where more invasive interventions are concerned. Whereas four out of 29 committees, according to the vote of their chairpersons, would probably have refused the approval for at least three of the five studies, two committees might have approved all studies without any restrictions.

We believe this situation is not acceptable. We are aware of the fact that the opinion of chairpersons cannot simply be equated with the decision of the committees as such. However, chairpersons are usually experienced and well informed members of ethics committees as far as ethical and legal regulations are concerned. Their interpretation of what is legally and ethically acceptable on the basis of current German regulations should, where more than minor risks are at stake, at least not differ to such a wide extent. The European Convention on Human Rights and Biomedicine, for example, strictly rules out this type of research. That the opinions of German chairpersons on these issues vary to such a high degree reveals significant uncertainty as to the ethical standards in non-therapeutic research. This creates problems for study participants, researchers, and the public, who are left uncertain about the standards of protection of research participants and in risk benefit analyses. This problem is particularly prominent in multicentre research projects where several local RECs are involved. Our conclusion is that there is an urgent need for a more detailed, comprehensive, and unambiguous regulation of research with the non-competent in Germany that does not permit such a wide range of interpretation. Meanwhile, the ratification of the European Convention on Human Rights and Biomedicine seems to be a good way to guarantee at least minimal ethical standards in these cases in Germany.

REFERENCES
Non-therapeutic research with minors: how do chairpersons of German research ethics committees decide?
C Lenk, K Radenbach, M Dahl and C Wiesemann

*J Med Ethics* 2004 30: 85-87
doi: 10.1136/jme.2003.005900

Updated information and services can be found at:
[http://jme.bmj.com/content/30/1/85](http://jme.bmj.com/content/30/1/85)

**References**
This article cites 2 articles, 0 of which you can access for free at:
[http://jme.bmj.com/content/30/1/85#BIBL](http://jme.bmj.com/content/30/1/85#BIBL)

**Email alerting service**
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

**Topic Collections**
Articles on similar topics can be found in the following collections

- Research and publication ethics (490)

**Notes**

To request permissions go to:
[http://group.bmj.com/group/rights-licensing/permissions](http://group.bmj.com/group/rights-licensing/permissions)

To order reprints go to:
[http://journals.bmj.com/cgi/reprintform](http://journals.bmj.com/cgi/reprintform)

To subscribe to BMJ go to:
[http://group.bmj.com/subscribe/](http://group.bmj.com/subscribe/)