Brucellosis in pregnant women from Pakistan: an observational study

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Abstract

Background: Brucella species occasionally cause spontaneous human abortion. Brucella can be transmitted commonly through the ingestion of raw milk or milk products. The objective of this study was to determine the sero-prevalence of and to identify potential risk factors for brucellosis in pregnant women from Rawalpindi, Pakistan.

Methods: We conducted a cross-sectional study at the Gynecology Outdoor Patient department of the Benazir Bhutto Hospital, Rawalpindi, Pakistan from March to June 2013. Data related to potential risk factors and clinical history was collected by individual interviews on the blood sampling day. The 429 serum samples collected were initially screened by Rose Bengal Plate Agglutination test for the detection of Brucella antibodies. We applied standard descriptive statistics and logistic regression analyses.

Results: Twenty five (5.8 %; 95 % confidence interval (CI): 3.8 % -8.5 %) serum samples were found to be seropositive. Brucellosis-related clinical symptoms were recorded in various seropositive cases. Animal contact, raw milk consumption, having an abortion history and the experience of an intrauterine fetal death were associated with seropositivity for brucellosis in univariate analyses (all \( p <0.05 \)). In multiple logistic regression models only the contact with animals remained as independent and robust risk factor (odds ratio 5.21; 95 % CI: 1.88-13.75; \( p = 0.001 \)) for seropositivity.

Conclusion: Brucellosis is a serious threat for pregnant women and their unborn children in Pakistan. Pregnant women having brucellosis-related symptoms or previous history of abortions, miscarriages, intrauterine fetal death and other brucellosis-related manifestations should be screened for brucellosis – especially those exposed to animals given the increased risk – and medication should be administered according to state of the art.

Keywords: Brucellosis, Pregnant women, Pakistan, Abortion, Serology

Background

Human brucellosis is a serious, life threatening disease. Four species of the Brucella (B.) genus (B. abortus, B. melitensis, B. suis and B. canis) are causative agents of brucellosis in humans [1]. Brucellosis is transmitted to humans by direct contact with Brucella-infected animals or consumption of their products [2, 3]. Milkers, livestock farmers, abattoir workers, shepherds, veterinarians, meat processing workers and laboratory workers are at high risk of getting infected [4–7].

Brucellosis during pregnancy was first reported in 1908 when Malta fever, i.e. brucellosis, was clinically described [8]. Later on, spontaneous abortions in pregnant women could be associated with the isolation of brucellae from placenta and aborted fetuses [9, 10]. In endemic areas, 1.3 to 12.2 % of pregnant women were found to be seropositive for brucellosis and active brucellosis may be associated with a high abortion rate [11–13]. Complications recorded are abortion, premature delivery, contagious or neonatal brucellosis, intrauterine infection or intrauterine fetal death. Brucella-positive infants may develop aspiration pneumonia and meningitis [14–18]. While brucellosis in farm animals is well controlled in most developed countries, it still exists in the Middle East, South and Central America, Africa, the Caribbean, the Mediterranean Basin and in Asia. In Pakistan, brucellosis is endemic in bovines but only a few

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Studies addressed human brucellosis-related risk factors and clinical outcome so far. Recently, 6.9% of high risk professionals were found to be seropositive. Occupation, locality and consumption of raw milk were identified as associated risk factors. Symptoms were fever, sweating, joint pain, back pain, body pain, abdominal pain, fatigue, lack of appetite and chills [2, 19–21]. Regarding pregnancy-related brucellosis, only a few studies have been conducted world-wide to date [13, 17, 22, 23]. No data are available for Pakistan.

The objective of the present cross-sectional study was to determine if anti-Brucella antibodies can be detected in pregnant women from Pakistan at all. Moreover, we were interested in identifying potential risk factors, clinical outcome and symptoms that are associated with a positive serological result.

Methods
Study design, setting, participants and data collection
This study has a cross-sectional study design and no follow-up investigation was planned. Blood samples from pregnant women who visited the Gynecology Outdoor Patient (OPD) department from the Benazir Bhutto Hospital, Rawalpindi, Pakistan for routine follow-up controls and who were willing to participate in the study were collected from March to June 2013. No data for those patients who were not willing to participate was recorded at all to guarantee privacy. Data related to potential risk factors for brucellosis, i.e., age, urbanicity, socio-economic status, contact with animals, consumption of raw milk, abortion in household animals, abortion history, contact with women who had aborted, intrauterine fetal death and number of pregnancies before the study, were collected by individual interviews at the time of blood sampling. It was presumed that these patients could not give the exact exposure time because of common ignorance that zoonotic brucellosis does exist in Pakistan. Thus, self-reported clinical symptoms including headache, fever, sweating, muscle or joint pain, joint swelling, general body malaise or backache, fatigue, lack of appetite and mental inattention or depression were recorded for the time interval between pregnancy start and individual interview date for Rose Bengal plate test (RBPT) [24]. RBPT antigen was purchased from Institute Pourquier, France. Equal amounts of serum and antigen (30 μL) were mixed on a glass plate. The glass plate was agitated for 4 min and the serum was considered positive, if any agglutination occurred. The test was done at the day of admission to the hospital.

Statistical methods and sample size
Standard descriptive statistics were used to summarize the data (continuous variables: quartiles/count data: absolute and relative frequencies). First, we performed univariate analyses using logistic regression to test for associations between presence/absence of human brucellosis seropositivity and quantitative continuous variables as well as count data. Secondly, we performed multiple logistic regression analyses, including all variables with a univariate p-value ≤ 0.2. In case of 5 or less counts in the cross-tables, we report Fisher’s exact test and estimation results only. All reported p-values are two-sided and not corrected for multiplicity. In this explorative study, we applied a significance level α = 0.05 (two-sided).

A sample size of 400 participants has a comparison-wise power of 80% to detect generic exposure-outcome associations for a rate of 5% exposed participants and odds ratios >2.8 (significance level α = 0.05 (two-sided) for a X² test and a 1:1 distribution of the outcome); for higher rates of e.g. 10% exposed participants odds ratios >2.2 are detectable with a similar comparison-wise power (proc power of SAS 9.3; SAS Institute Inc., Cary, NC, USA). Thus, the study was well-powered to detect strong associations with presence/absence of human brucellosis seropositivity.

All statistical analyses were done using the statistical language R 3.0.2.

Results
Four hundred twenty nine pregnant women participated in the study (see Table 1) and 25 (5.8%; 95% CI: 3.8%–8.5%) were found to be RBPT positive.

All pregnant women reported more than one of the symptoms usually associated with clinical brucellosis. The time of pregnancy did not influence the clinical presentation of disease i.e. there was no correlation found between pregnancy status and number of reported symptoms or self-reported severity of these symptoms. The time of infection or onset of disease could not be elucidated by the design of our questionnaire.

The results of the association analyses concerning presence/absence of human brucellosis seropositivity are given in Table 2. Animal contact (p < 0.001), raw milk intake (p < 0.001, Fisher’s exact test), abortion history (p = 0.016) and the experience of an intrauterine fetal death (p = 0.050, Fisher’s exact test) were associated with
seropositivity. For all these univariate associations we estimated strong association effects (see Table 2) but the estimation itself was highly imprecise due to the small number of only 25 seropositive women. Additionally, we modeled the impact of the variables jointly and detected that only the contact with animals was a significant, independent and relatively robust predictor of presence/absence of seropositivity (odds ratio 5.21; 95% CI: 1.88-13.75; \( p = 0.001 \)).

**Discussion**

Brucellosis is an infectious disease which affects both animals and humans. It is endemic in ruminants in Pakistan but little is known on human brucellosis, especially in pregnant women who are considered to be at special risk of abortion.

The various studies on human brucellosis and pregnancy have to be compared with caution due to the different cohorts investigated, techniques used or prevalent epidemiological situation [25]. It is obvious that our study population is not representative for the group of ‘pregnant women in Pakistan’ as it was not random due to restrictions in the acquisition of participants. In a comparable study population 3.5% of pregnant women from rural areas of Saudi Arabia were seropositive [11]. In the present study, 5.8% of pregnant Pakistani women were found to be seropositive and indeed women from rural areas were more often seropositive than those from urban areas. These women are often engaged in animal keeping and thus are in close contact to infected animals or abortion material. Thus, our women may be exposed to a similar risk as high risk professionals in Pakistan - a group in which 6.9% of the persons were recently found to be seropositive [2]. Our findings are corroborated by the data of a survey from Turkey, where 59% of pregnant women with brucellosis were stockbreeders [16]. 25% of Rwanda women suffering from abortions and stillbirth were found to be infected with brucellosis. Again, these women were in close contact to cattle, goats and sheep [26].

### Table 1

Characteristics of the 429 Rose Bengal Plate Agglutination test (RBPT) tested pregnant women from Pakistan

| Characteristic [unit] | RBPT tested (%) | RBPT positive (%) |
|-----------------------|-----------------|-------------------|
| **Age [Years]**       |                 |                   |
| Median (Q1;Q3)\(^b\)  | 26 (22;32)      | 27 (25;36)        |
| 18-25                 | 211 (49.18)     | 11 (44.00)        |
| 26-33                 | 118 (27.51)     | 7 (28.00)         |
| 34-41                 | 72 (16.78)      | 2 (8.00)          |
| >41                   | 28 (6.53)       | 5 (20.00)         |
| **City\(^c\)**        |                 |                   |
| Urban                 | 255 (59.44)     | 11 (44.00)        |
| Rural                 | 174 (40.56)     | 14 (56.00)        |
| **Socio-economic status\(^d\)** |              |                   |
| Low                   | 260 (60.61)     | 17 (68.00)        |
| Medium                | 169 (39.39)     | 8 (32.00)         |
| **Contact with animal** |                |                   |
| No                    | 389 (90.68)     | 16 (64.00)        |
| Yes                   | 40 (13.99)      | 9 (36.00)         |
| **Raw milk intake**   |                 |                   |
| No                    | 412 (96.04)     | 12 (48.00)        |
| Yes                   | 17 (3.96)       | 13 (52.00)        |
| **Contact with abortion in animals** |           |                   |
| No                    | 421 (98.14)     | 24 (96.00)        |
| Yes                   | 8 (1.86)        | 1 (4.00)          |
| **Contact with person with abortion** |         |                   |
| No                    | 410 (95.57)     | 22 (88.00)        |
| Yes                   | 19 (4.43)       | 3 (12.00)         |
| **Number of pregnancies** |           |                   |
| Median (Q1;Q3)\(^b\)  | 2 (1;3)         | 2 (2;4)           |
| 1                     | 150 (35.05)     | 5 (20.00)         |
| 2                     | 106 (24.77)     | 9 (36.00)         |
| >2                    | 172 (40.19)     | 11 (44.00)        |
| **Abortion history**  |                 |                   |
| No                    | 388 (90.44)     | 19 (76.00)        |
| Yes                   | 41 (9.56)       | 6 (24.00)         |
| **Intrauterine fetal death** |           |                   |
| No                    | 414 (96.50)     | 22 (88.00)        |
| Yes                   | 15 (3.50)       | 3 (12.00)         |
| **Symptoms\(^e\)**   |                 |                   |
| Headache              | ND              | 13 (52.00)        |
| Fever                 | ND              | 12 (48.00)        |
| Sweating              | ND              | 21 (84.00)        |
| Muscle or joint pain  | ND              | 14 (56.00)        |
| Joint swelling        | ND              | 2 (8.00)          |

\(^a\)for “Age” and “Number of pregnancies”, median and interquartile ranges are also reported

\(^b\)(Q1;Q3) is the interquartile range

\(^c\)defined as self-reported location of residence; “rural” means that participants report to live in villages and “urban” means that they report to live in cities

\(^d\)defined as self-reported economic and social position in relation to others

\(^e\)symptoms were assessed in the 25 RBPT positive women only; multiple symptoms can be reported for each woman.
results of the logistic regression analyses, Odds ratios (ORs) with 95 % confidence intervals (CIs) and two-sided p-values are provided for the univariate and multiple models (Details see text). Note that the left column lists both units (for quantitative variables) or reference values for categorical variables in parenthesis.

| Variable [unit or reference (REF)] | Univariate logistic models | Multiple logistic model |
|-----------------------------------|---------------------------|-----------------------|
| Age [per year]                    | 1.05 (0.99, 1.10)         | 0.093                 |
| Rural city [REF urban]*           | 1.94 (0.86, 4.48)         | 0.110                 |
| Medium socio-economic status [REF low]* | 0.71 (0.28, 1.64) | 0.437 |
| Contact with animal [REF no contact] | 6.77 (2.67, 16.34) | <0.001 |
| Raw milk intake [REF no raw milk intake]* | 102.86 (27.35, 390.43) | <0.001 |
| Contact with abortion in animals [REF no contact]* | 2.36 (0.10, 18.69) | 0.384 |
| Contact with person with abortion [REF no contact]* | 3.29 (0.76, 12.40) | 0.091 |
| Number of pregnancies [per one]   | 1.16 (0.89, 1.49)         | 0.265                 |
| Presence of abortion history [REF absence] | 3.33 (1.15, 8.47) | 0.016 |
| Experience of intrauterine fetal death [REF no experience]* | 4.43 (1.00, 16.24) | 0.050 |

*defined as rural mean patients live in village and urban mean patients live in city
*defined as patient belong to middle class versus patients of poor families
*result of Fisher’s exact
*not calculated due to too few observations

Pakistani women from rural, low income families were more often seropositive (6.5 %) than those from better situated families of urban areas (4.7 %). Pregnant women with an abortion history, with contact to animals that aborted or with contacts to women who had an abortion were more often seropositive (14.6 %, 12.5 % and 15.8 %, respectively), although the latter two factors were not significantly associated with seropositivity in our study. These findings may jointly indicate that brucellosis has already become chronic in our women and may have caused multiple abortions. Alternatively, reinfections might have occurred frequently given that their living conditions were not altered. It may be speculated that these women simply had no adequate access to medical care due to poor economic situation as all these women reported symptoms of active brucellosis. Thus, brucellosis proved again to be a disease sustained by poverty. Childlessness caused by (chronic) brucellosis may have also negative impact on the social standing of a family in the society or for security in old age. Especially the mental pressure on the childless woman may be high. Future studies should reflect on these issues, too.

Pregnant women consuming raw milk were more often seropositive (76.5 %) compared to those never consuming raw milk (2.9 %). However, the overall rate of reported raw milk consumption was relatively low (~4 %). Nevertheless, raw milk consumption-related high-frequent Brucella seropositivity was also observed in pregnant women from Rwanda and Turkey [16, 25, 26]. The lack of knowledge about the presence of brucellosis in livestock, its transmissibility and adequate countermeasures e.g. heating of milk, or indifference fosters human brucellosis. Programs to educate illiterate agricultural population are needed in Pakistan to reduce disease incidence. Efficient regulations on food hygiene should be placed into force.

However, brucellosis in pregnant women poses also severe risks for newborns. Infection was shown to be a leading cause of neonatal mortality among infants delivered at a tertiary hospital in Karachi, Pakistan [27]. Congenital transmission of brucellosis was reported from Argentina [18, 28] in an infant whose parents were livestock farmers. Transmission by breast milk has already been described [29–31]. Now, future studies are needed to determine the share of brucellosis on neonatal mortality in Pakistan.

Our study has several limitations. First, while the Rose Bengal plate test (RBPT) is a feasible and reliable test especially as screening test in our study setting, confirmation should be done using other tests (e.g. Complement Fixation Test or Enzyme Linked Immunosorbent Assay). Secondly, the number of 25 RBPT positive women results in highly imprecise odds ratio estimators. Thirdly, a major drawback of our study is that the prevalent Brucella species remains unknown and molecular epidemiology cannot be used to identify chains of infection. Use of genus or species specific quantitative real-time polymerase chain reaction (qRT-PCR) and molecular typing directly from the sample will help to overcome these weaknesses. For the moment we cannot exclude that a relevant number of cases might have been caused by veterinary vaccine which still have a considerable pathogenicity for humans. Handling these vaccines or freshly vaccinated animals still excreting the vaccine strain will pose a special risk for pregnant women in rural areas. Both,
vaccination and awareness for the pathogenicity of vaccines have to be objects of future questionnaires.

We admit that the data obtained have only restricted significance. However, they might convince public health policy makers, physicians and families having religious reservations to contribute to future studies investigating the risks of brucellosis in pregnancy. Thus, counterprograms for veterinary and human public health can be tailored and implemented in Pakistan.

Conclusions

Brucellosis is considered to be the most common zoonotic infection worldwide and a serious hazard for pregnant women. Humans with brucellosis-related symptoms or manifestations (including a history of abortions) should be investigated for brucellosis; subsequent care should be according to the state of the art. Rose Bengal plate test (RBPT) should be considered as a screening test that must be confirmed by other reliable tests. This study confirmed that brucellosis is a serious threat and public health concern for pregnant women and their unborn children and more generally for humans in contact with animals or raw milk consumption in Pakistan. The findings reinforce the need for adequate preventive and control measures to be taken against zoonotic brucellosis especially in one of the most susceptible and vulnerable individuals of the human population.

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Availability of data and materials

All the data supporting study findings is contained within the manuscript and will be shared upon request.

Authors’ contributions

SAli designed and coordinated the study, carried out the laboratory work and data analyses, wrote the manuscript and helped to draft and review the manuscript. SAKhter, IK, AA, SQ and QA were involved in the study design, sample collection, sample examination, analyses and writing of the manuscript. HN and FM supervised the study, critically revised the paper. HE was responsible for all correspondence to the journal. AS (supervisor) and MK (supervisor) were involved in the study design, data analyses and interpretation, wrote parts of the paper and revised and approved the final version. All authors read and approved the final version of the manuscript.

Competing interest

The authors declare that they have no competing interests.

Consent for publication

Not applicable.

Ethics approval and consent to participate

This study was approved by the ethical committee of Pir Mehr Ali Shah Arid Agriculture University Rawalpindi, Pakistan. All women provided written informed consent to participate in this study; the study was conducted in accordance with the Declaration of Helsinki [32].

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References

1. Lopes LB, Nicolino R, Haddad JPA. Brucellosis - risk factors and prevalence: a review. Open Vet Sci J. 2010;4:72–84.
2. Ali S, Ali Q, Neubauer H, Mezler F, Eichner M, Khan I, Abatiab EN, Ullah N, Irfan M, Akhter M. Seroprevalence and risk factors associated with brucellosis as a professional hazard in Pakistan. Foodborne Pathog Dis. 2013;10:500–5.
3. Chen S, Zhang H, Liu X, Wang W, Hou S, Li T, Zhao S, Yang Z, Li C. Increasing threat of brucellosis to low-risk persons in urban settings. China Emerg Infect Dis. 2014;20(1):126–30.
4. Osbode HOK, Adah BMJ, Nafarnda WD, Aworh MKF, Bello RH. Sero-prevalence of Brucella abortus antibodies in slaughtered cattle and meat by-product handlers in Ilorin abattoir, Kwara state – Nigeria. Prime J Microbiol Res. 2012;2(01):109–13.
5. Trayler RM, Lehman MW, Bosserman EA, Guerra MA, Smith TL. A literature review of laboratory-acquired brucellosis. J Clin Microbiol. 2013;51(9):3025–62.
6. Kutlu M, Ergonul O, Sayin-Kutlu S, Guven T, Ustun C, Alp-Cavus S, Ozturk SB, Acicbe O, Akalin S, Tekin R, Tekin-Koruk S, Demiroglu YZ, Keskiner R, Gonen I, Sapmaz-Karabag S, Bosnak V, Kazak E. Risk factors for occupational brucellosis among veterinary personnel in Turkey. Prev Vet Med. 2014;117(1):52–8.
7. Pathak AD, Dubal ZB, Dojlad S, Roarane A, Rodrigues S, Naik R, Naik-Gaonkar S, Kalorey DR, Kurkure NV, Naik R, Barbudhese SB. Human brucellosis among pyrexis of unknown origin cases and occupationally exposed individuals in Goa Region, India. Emerg Health Threats J. 2014;22(7):23846. doi:10.3402/ ehtj.v7.23846.
8. Eyre JWH. Melitensis septicemia. Lancet. 1908;1747–52.
9. Young EJ. Human brucellosis. Rev Infect Dis. 1983;5(s8):821–42.
10. Seoud M, Saade G, Awar G, Uwaydah M. Brucellosis in pregnancy. J Reprod Med. 1991;36(6):441–5.
11. Shafir A, Reyes Z, Thomassen P. Screening for brucellosis in pregnant women. J Trop Med Hyg. 1990;93:42–3.
12. Khan M, Mah M, Memish Z. Brucellosis in pregnant women. Clin Infect Dis. 2001;32:1172–7.
13. Eshamy M, Ahmed A. The effects of maternal brucellosis on pregnancy outcome. J Infect Dev Ctries. 2008;2:230–4.
14. deCarle DW. Premature labor in a case of undulant fever. Proc Mayo Clin. 1931;6:469–71.
15. Oran O, Karslioglu A, Secmeer G, Berkman E, Kanra G. A premature infant with Brucella abortus infection. Turk J Pediatr. 1983;25:139–42.
16. Kordoglu M, Adali E, Kordoglu Z, Karahocagil MK, Kolusari A, Yildizhan R, et al. Brucellosis in pregnancy: a 6-year clinical analysis. Arch Gynecol Obstet. 2010;281:201–6.
17. Abo-Shehada MN, Abu-Halaweh M. Seroprevalence of Brucella species among women with miscarriage in Jordan. East Mediterr Health J. 2011;17(11):871–4.
18. Cacace ML, Claroas EA, Erazu KA, Escobar GI, Lucero NE. Congenital brucellosis in an infant. Vector Borne Zoonotic Dis. 2013;13:7513–5.
19. Dean AS, Crump L, Greter H, Hattendorf J, Schelleng E, Zinsstag J. Clinical manifestations of human brucellosis: a systematic review and meta-analysis. PLoS Negl Trop Dis. 2012;6(2), e1929. doi:10.1371/journal.pntd.0001929.
20. Ning P, Guo M, Guo K, Xu L, Ren M, Cheng Y, Zhang Y. Identification and effect decomposition of risk factors for Brucella contamination of raw whole
21. Ali S, Ali Q, Mebeer F, Khan I, Akhter S, Neubauer H, Jamal SM. Isolation and identification of bovine Brucella isolates from Pakistan by biochemical tests and PCR. Trop Anim Health Prod. 2014;46:73–8.

22. Baud D, Peter Q, Langel C, Regan L, Greub G. Seroprevalence of Coxiella burnetii and Brucella abortus among pregnant women. Clin Microbiol Infect. 2009;15:499–501.

23. Roushan MR, Bajani M, Asnafi N, Saedi F. Outcomes of 19 pregnant women with brucellosis in Babol, northern Iran. Trans R Soc Trop Med Hyg. 2011;105:540–2.

24. Alton GG, Jones LM, Angus RO, Verger JM. Bacteriological methods. In: Alton GG, Jones LM, Angus RO, Verger JM, editors. Techniques for the Brucellosis Laboratory. Paris: Institut National De La Recherche Agronomique; 1988. p. 13–61.

25. Karcaaltincaba D, Sencan I, Kandemir O, Guvendag-Guven ES, Yalvac S. Does brucellosis in human pregnancy increase abortion risk? Presentation of two cases and review of literature. J Obstet Gynaecol Res. 2010;36(2):418–23.

26. Rujeni N, Mbanzamihigo L. Prevalence of Brucellosis among Women Presenting with Abortion/Stillbirth in Huye, Rwanda. J Trop Med. 2014;2014:740479. doi:10.1155/2014/740479.

27. Mustufa MA, Korejo R, Shahid A, Nasim S. Infection remains a leading cause of neonatal mortality among infants delivered at a tertiary hospital in Karachi, Pakistan. Infect Dev Ctries. 2014;8(11):1470–5.

28. Giannacopoulos I, Eliopoulos MI, Ziambaras T, Papanastasiou DA. Transplacentally transmitted congenital brucellosis due to Brucella abortus. J Infect. 2002;45:209–10.

29. Lubani MM, Dudin KI, Sharda DC, Abu Sinna NM, Al-Shab T, AA A-R a, Labani SM, Nasrallah A. Neonatal brucellosis. Eur J Pediatr. 1988;147:520–2.

30. Palanduz A, Palanduz S, Güler K, Güler N. Brucellosis in a mother and her young infant: Probable transmission by breast milk. Int J Infect Dis. 2000;4:55–6.

31. Ceylan A, Köstü M, Tuncer O, Peker E, Kırımı E. Neonatal brucellosis and breast milk. Indian J Pediatr. 2012;79:389–91.

32. World Medical A. Ethical principles for medical research involving human subjects. Bull World Health Organ. 2001;79:373–4.