

International Journal of Hyperthermia



ISSN: 0265-6736 (Print) 1464-5157 (Online) Journal homepage: informahealthcare.com/journals/ihyt20

Letter to the editor: Intracorporeal whole body hyperthermia: toxicity assessment

G. J. Wiedemann, D. M. Katschinski, M. Mentzel, D. Mulkerin & H. I. Robins

To cite this article: G. J. Wiedemann, D. M. Katschinski, M. Mentzel, D. Mulkerin & H. I. Robins (1997) Letter to the editor: Intracorporeal whole body hyperthermia: toxicity assessment, International Journal of Hyperthermia, 13:2, 249-250, DOI: 10.3109/02656739709012388

To link to this article: https://doi.org/10.3109/02656739709012388

	Published online: 09 Jul 2009.
	Submit your article to this journal $oldsymbol{\mathbb{Z}}$
ılıl	Article views: 64
Q ^N	View related articles 🗹

Letter to the editor Intracorporeal whole body hyperthermia: toxicity assessment

We have reviewed with interest the recently published letter by Shecterle et al. (International Journal of Hyperthermia, 12, 569-571, 1996) regarding a 'new invasive device' for whole body hyperthermia (WBH). The authors pointed out that their new methodology avoids the problem of serial cannulation, which can cause complications. In our view, this is not the major problem associated with extracorporeal WBH. In this regard, the most recent peer-reviewed experience (utilizing extracorporeal hyperthermia and chemotherapy) using a dialysis device (Wiedemann et al. 1994), as well as a subsequent experience using a heat exchanger (Wiedemann et al. 1996b), demonstrated grade 4 (WHO) renal toxicity. This was related in part to hypotension and associated decreased renal perfusion. This was observed in spite of the fact that Wiedemann et al. used high dose dopamine and noradrenaline to minimize morbidity and prevent mortality. Interestingly when the same chemotherapy was utilized with exactly the same patient population (i.e. sarcoma patients) all of the significant aforementioned toxicity was eliminated (Wiedemann et al. 1996a) utilizing the Aquatherm radiant heat WBH system (Robins 1994, Robins et al. 1996). We noted that Shecterle et al. provided no data regarding either blood pressure, blood flow rates, fluid rates or follow up creatinines in the animals tested. It would be assumed, based upon the collective experience with extracorporeal hyperthermia over the years, that these values would be abnormal in comparison to non-invasive radiant-heat devices (Robins et al. 1985, Robins 1994). Other issues not addressed include: the potential risk of heparin, and other drugs, e.g. diuretics and steroids (used to treat the common complication of extracorporeal hyperthermia, i.e. pulmonary oedema) as well as antibiotics (which is a requirement with canine and clinical extracorporeal WBH). Beyond this, the authors elude to a significant depression of platelet counts, which is a major clinical toxicity and is unique to extracorporeal WBH. Finally, the authors do not comment on the excessive cost and labour intensive aspects of performing clinical extracorporeal hyperthermia which includes general anaesthesia, Swan-Ganz catheterization, arterial lines, and intensive care unit observations.

In summary, based upon our own clinical experience with the extracorporeal and non-invasive radiant-heat WBH, as well as the information reported by Shecterle et al., we reject the concept that the mere survival of animals following extracorporeal hyperthermia suggests safety or efficiency for treating patients with neoplastic diseases.

Acknowledgment

Studies made with the support of Cancer Res Inc., NY; GJW & DMK Deutsche Forschungsgemeinschaft.

250 Letter

References

- ROBINS, H., KUTZ, M., TUTSCH, K., TIGELAAR, C., PAUL, D., LONGO, W., and D'OLEIRE, F., 1996, Phase I clinical trial of melphalan (L-PAM) and 41.8C whole body hyperthermia (WBH) in cancer patients. *Journal of Clinical Oncology* (in press).
- ROBINS, H. I., SCHMITT-TIGGELAAR, C. L., COHEN, J. D., WOODS, J. P., HEISS, C., GILLIS, W., and D'OLEIRE, F., 1994, A new technological approach to heat whole body Hyperthermia. *Cancer Letters*, **79**, 137–145.
- ROBINS, H. I., FENNIS, W. H., NEVILLE, A. J., SHECTERLE, L., MARTIN, P.A., GROSSMAN, J., DAVIS, T. E., NEVILLE, S., GILLIS, W., and RUSEY, B. F., 1985, A non-toxic system for 41-8°C whole body hyperthermia: Results of a phase I study using a radiant heat device. Cancer Research, 45, 3937-3944.
- ROBINS, H. I., NEVILLE, A. J., and COHEN, J. D., 1992, Whole body hyperthermia: biological, and clinical aspects (Berlin Heidelberg: Springer-Verlag).
- Schecterle, L. M., Kelly, T., and St. Cyr, J. A., 1996, Letter: Safe and efficient extracorporeal method of inducing whole body hyperthermia. *International Journal of Hyperthermia*, 12(4), 569-571.
- Wiedemann, G. J., d'Oleire,, F., Knop, E., Eleftheriadis, S., Bucksy, P., Feddersen, S., Klouche, M., Geisler, J., Mentzel, M., Schmucker, P., Feyerabend, T., Weiss, C., and Wagner, T., 1994, Ifosfamide and Carboplatin combined with 41.8°C whole-body Hyperthermia in patients with refractory sarcoma and malignant teratoma. *Cancer Research*, 54, 5346–5350.
- WIEDEMANN, G. J., KATSCHINSKI, D. M., MENTZEL, M., WAGNER, T., and ROBINS, H. I., 1996a, Systematic hyperthermia and ICE chemotherapy for sarcoma. ASCO: Late breaking abstracts. *Journal of Clinical Oncology*, 14(5), 1751.
- WIEDEMANN, G. J., ROBINS, H. I., GUTSCHE, S., MENTZEL, M., DEEKEN, M., KATSCHINSKI, D. M., ELEFTHERIADIS, S., CRAHE, R., WEISS, C., STORER, B., and WAGNER, T., 1996b, Ifosfamide, carboplatin and etoposide (ICE) combined with 41-8°C whole-body hyperthermia in patients with refractory sarcoma. European Journal of Cancer, 32A(5), 888-892.

G. J. WIEDEMANN†
D. M. KATSCHINSKI‡
M. MENTZEL†
D. MULKERIN‡
and H. I. ROBINS‡

† Medical University of Lubeck, Germany ‡ University of Wisconsin Comprehensive Cancer, Madison, WI USA