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Improving superficial hyperthermia treatment

Temperature matters

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CHAPTER 2

TEMPERATURE AND THERMAL DOSE DURING RADIOTHERAPY AND HYPERTHERMIA FOR RECURRENT BREAST CANCER ARE RELATED TO CLINICAL OUTCOME AND THERMAL TOXICITY: A SYSTEMATIC REVIEW

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ABSTRACT

Background: Hyperthermia (HT), heating tumors to 40 - 44 °C, is a known radiotherapy (RT) and chemotherapy sensitizer. The additional benefit of HT to RT for recurrent breast cancer has been proven in multiple randomized trials. However, published outcome after RT-HT varies widely. We performed a systematic review to investigate whether there is a relationship between achieved HT dose and clinical outcome and thermal toxicity for patients with recurrent breast cancer treated with RT-HT.

Methods: Four databases, EMBASE, PubMed, Cochrane library and clinicaltrials.gov, were searched with the terms breast, radiotherapy, hyperthermia and their synonyms. Final search was performed on April 3rd 2019. Twenty-two articles were included in the systematic review, reporting on 2330 patients with breast cancer treated with RT-HT.

Results: Thirty-two HT parameters were tested for a relationship with clinical outcome. In studies reporting a relationship, the relationship was significant for complete response in 10/15 studies, in 10/13 studies for duration of local control, in 2/2 studies for overall survival and in 7/11 studies for thermal toxicity. Patients who received high thermal dose, had on average 34 % (range 27 - 53 %) more complete responses than patients who received low thermal dose. Patients who achieved higher HT parameters had increased odds/ probability on improved clinical outcome and on thermal toxicity.

Conclusion: Temperature and thermal dose during HT had significant influence on complete response, duration of local control, overall survival and thermal toxicity of patients with recurrent breast cancer treated with RT-HT. Higher temperature and thermal dose improved outcome, while higher maximum temperature increased incidence of thermal toxicity.

INTRODUCTION

Hyperthermia (HT), increasing the tumor temperature to 40 - 44 °C, is a known radiotherapy (RT) and chemotherapy sensitizer. The additional benefit of HT to RT and chemotherapy has been proven in randomized trials for melanoma, sarcoma, recurrent breast cancer, cervical cancer and other tumor types [1–5]. A recent systematic review showed the additional value of HT to RT for patients with locoregional recurrent breast cancer [6]. Complete response rate increased from 38.1 % for patients treated with RT to 60.2 % for patients treated with RT-HT, respectively. The complete response of RT-HT varied widely throughout the included studies; range 33.3 - 95.0 % [7, 8]. An important factor contributing to this wide variation was hypothesized to be the variability in HT delivery, i.e. the HT technique, sequencing of RT and HT, duration of HT and temperature and thermal dose achieved during HT treatment [6].

Techniques used in the past and in the present vary in effective field size and penetration depth depending on the design and used frequency of the technique [9], resulting in variations in temperature and thermal dose achieved during HT treatment. Furthermore, the amount of radiosensitization by HT is strongly dependent on the sequence and time-interval between RT and HT, where the effect is largest when HT and RT are given simultaneously [10, 11]. This maximum radiosensitization for simultaneous application is observed for both tumor and normal tissue.

A thermal dose-effect relationship for HT has been reported in the literature, both in pre-clinical and clinical studies. In vitro studies show that there is an increase in cell death when tumor cells are heated longer or to a higher temperature [12–14]. Several clinical studies underline the importance of duration of HT and temperature and thermal dose achieved during HT treatment [15, 16]. The effect of HT increases with a longer heating time, but during and after exposure to heating the onset of thermotolerance is induced, which results in a temporarily reduced sensitivity to heating [17, 18]. This reduced effectiveness of HT is a transient effect and persists for several days [18]. Therefore, clinical HT treatments are limited to 1 - 1.5 h, once or twice weekly. As a result of the observed thermal dose-effect relationship, HT treatments are usually quantified by calculating a thermal isoeffective dose, which captures both temperature and time. The most commonly used thermal isoeffective dose is the number of equivalent minutes at 43 °C (CEM43) [19, 20]. Other parameters representing minimum or median tumor temperature levels achieved during HT, include T90, T50 and TRISE [21–23].

We performed a systematic review to determine whether the large variation in outcome observed in clinical studies for patients with recurrent breast cancer treated with RT-HT can be explained by variation in achieved temperatures and thermal dose during HT treatment.

METHODS

The systematic review was conducted following the PRISMA guidelines [24]. Four databases were searched from inception to April 3rd 2019: EMBASE, PubMed, Cochrane library and clinicaltrials.gov. An initial search was performed in PubMed, and relevant articles were found by citation tracking. The synonyms for breast, hyperthermia and radiotherapy reported in those articles were used as entrance parameters in the final search. For each database an individual search was performed. In PubMed the following search was executed: (“Breast”[Mesh] or breast[tw] or mamma[tw] or mammary[tw]) AND (“Hyperthermia, Induced”[Mesh] or hyperthermia[tw] or heat[tw] or thermotherapy[tw] or thermoradiotherapy[tw]) AND (“Combined Modality Therapy”[Mesh] or “Radiotherapy”[Mesh] or radio[tw] or radiation[tw] or irradiation[tw] or reirradiation[tw] or re-irradiation[tw] or radiotherapy[tw] or thermoradiotherapy[tw]). In clinicaltrials.gov studies for breast cancer and hyperthermia were searched, while a filter for the intervention type, i.e. not drugs, was applied. The search was not limited to any date, both articles and conference proceedings were allowed, a filter for English articles and human studies was applied. When warranted we contacted study authors to identify additional studies.

Inclusion criteria

Single-arm, double-arm, retrospective and prospective studies (randomized and non-randomized) fulfilling the following criteria were included: Patients with recurrent breast cancer treated with external beam RT and local HT.

Study selection

After exclusion of duplicates, articles were screened according to their titles and abstracts (Figure 1). Articles were excluded when:

- Articles were not clinical studies.
- Patients received concurrent chemotherapy or drugs.
- The endpoint of the study was not outcome.
- The relationship between thermal dose parameters and outcome was not analyzed.
- Less than ten patients or lesions with breast cancer treated with RT-HT were reported.
- Articles were updated in a later publication by the same author(s).
- Mixed patient groups were reported and the outcomes of breast cancer patients were not analyzed separately.

Articles from the same authors describing the temperature analysis of the HT treatment in more detail were taken into account.

Data extraction and quality assessment

The primary endpoints of interest were the relationships between temperature and/or thermal dose during HT with clinical outcome: complete response (CR), duration of local control (LC), overall survival (OS) as well as thermal toxicity (i.e. acute blisters). Details of the (pre) treatment patient characteristics and present treatment parameters were investigated, as well as clinical outcome.

The relationships between thermal dose parameters and CR, LC, OS and thermal toxicity were tested in univariate and multivariable analysis. A p -value < 0.05 was considered significant, p between 0.05 and 0.1 a trend and for $p \geq 0.1$ we concluded that there was no relationship. Furthermore, we registered whether a parameter was mentioned by the authors but not reported as related to outcome, which we interpreted as that the analysis was either not performed or not reported. This selective reporting within studies is a possible publication bias. An overview of the relationships of thermal dose parameters to CR, LC, OS and thermal toxicity is presented in a graph. In this graph, the thermal dose parameters are grouped in categories as either high-end temperature (e.g. maximum temperature, T10, T20), mean temperature (e.g. mean and median temperature) and low-end temperature (e.g. T90 and minimum temperature) and as either high-end dose (e.g. maximum CEM43, time > 43 °C), mean dose (e.g. CEM43T50) and low-end dose (e.g. CEM43T90, minimum CEM43, time < 40 °C). The number of HT treatments is considered as a separate category. T10, T20 and T90 are the 10th, 20th, and 90th percentile of all temperature measurements, respectively. For each category, a study is taken into account once. We selected the parameter that had the strongest relationship with the outcome variables.

The odds ratio (OR), risk ratio (RR) and risk difference (RD) were calculated for studies which reported the clinical outcome measures for patients grouped based on the value of their thermal dose parameters during HT treatment. The OR and RR are expressed as the logarithmic average with the 95 % confidence interval (CI). The RD is expressed as the average with the 95 % CI. When one of the events was zero, e.g. patients with a low thermal dose had no CR, we first added 0.5 to all categories and then calculated the OR, RR or RD (Haldane-Anscombe correction) [25]. The statistical analysis was performed in R (version 3.5.1) with package “metafor” (version 2.0-0). No meta-analysis was performed due to the strong variation in definitions of thermal dose parameters.

The study selection and risk of bias assessment (QUIPS) [26] were performed by author AB. Data extraction was done by authors JZ and AB.

RESULTS

A total of 1249 articles were identified through the search and were screened (Figure 1). Twenty-five studies described in twenty-two articles were included in the systematic review, reporting a total of 2330 patients with breast cancer. Two additional articles presenting a detailed temperature analysis of the studies presented by Vernon et al. [4] were taken into account [15, 27].

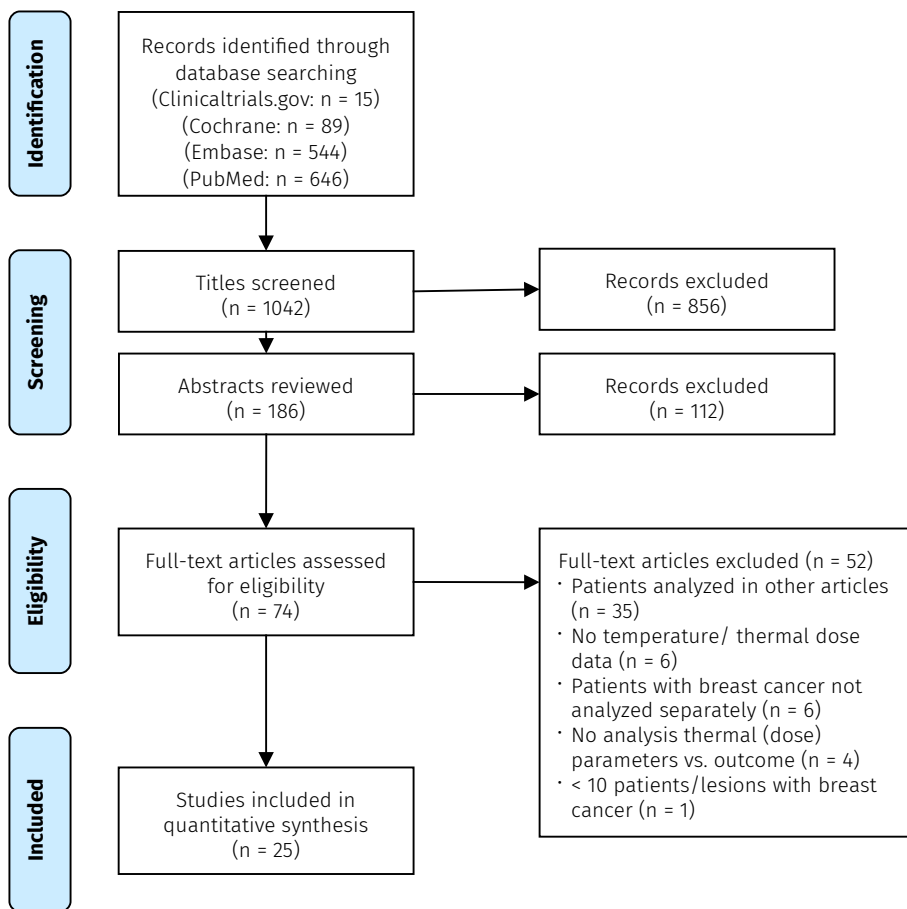


Figure 1. PRISMA flow diagram of the literature search [24].

Patient and treatment characteristics

In total 1749 patients with macroscopic disease (77.0 %) and 563 patients with microscopic disease (23.0 %) were reported. Most patients were treated for recurrent disease (n = 2304, 98.9 %), while 26 patients (1.1 %) were treated for primary disease. Patients received RT-HT for the breast (n = 256), chest wall (n = 1861), and lymph nodes (n = 181) (Table 1).

Most patients had previously received extensive treatment on the areas treated with RT-HT. RT was previously given with a median dose of 50 Gy (range 32 - 65) to 2007 patients (86.1 %). At least 1773 patients (76.1 %) underwent surgery in the currently treated area, 1319 patients (56.6 %) received prior chemotherapy and 971 patients (41.7 %) received prior hormonal therapy.

During RT-HT treatment, patients were treated with a median RT dose of 36 Gy (range 29 - 60) and either microwave HT (n = 2155), radiofrequency HT (n = 135), interstitial HT (n = 31), ultrasound HT (n = 26), capacitive HT (n = 24) or with a mix of HT devices with different frequencies (n = 8) (Table 1). HT treatments had an average duration of 52.8 ± 10.4 minutes (SD). HT was given mostly once or twice per week and in one study every two weeks. Patients received five HT treatments (median, range 2 - 10). In one study RT was given after HT (as soon as possible), while in 19 studies HT was given after RT. The time interval between RT and HT varied from < 30 min (9 studies), < 45 min (3 studies), < 60 min (8 studies), 80 min (1 study) to > 90 min (1 study). Three studies did not report the time interval between RT and HT (Table 1).

Temperature and thermal dose measurement

Temperature during HT was measured invasively in all 25 studies, but not in all patients. The type of thermometry equipment and the number of temperature sensors were poorly reported (Table 1). A total of 32 thermal dose parameters were tested for a relationship with outcome.

Thermal dose parameters were derived from superficial temperature sensors, invasive temperature sensors or from a combination. Thermal dose parameters measured superficially showed no relationship with CR, LC and OS, but did show a relationship with thermal toxicity. Thermal dose parameters derived from invasive measurements correlated with all outcome measures, i.e. CR, LC, OS and thermal toxicity (Table 2, supplementary materials).

Table 1. Salient features and results of the assessment on the risk of bias [26] of the 22 articles, pertaining to 25 individual studies that have been included in the systematic review.

	Arcangeli et al. (1991)	Bakker et al. (2017)	Dragović et al. (1989)	Engin et al. (1993)	Gabriele et al. (2009)	Gonzalez-Gonzalez et al. (1988)	Hehr et al. (2001)	Kapp et al. (1991)	Kapp et al. (1992)	Lee et al. (1998)	Li et al. (2004)
Patients											
Patients (n)	1	262	30	23	23	35	39	89	89	151	73
Lesions (n)	n.r.	n.r.	n.r.	23	44	45	n.r.	241	262	196	85
Women (%)	100	0	n.r.	n.r.	96	n.r.	n.r.	99	99	n.r.	n.r.
Age (years; mean/ median)	75	n.r.	58	58	65	59	57	56	53	59	54
Tumor											
Primary (%)	0	0	0	0	0	0	0	0	0	0	0
Recurrent (%)	100	100	100	100	100	100	100	100	100	100	100
Macroscopic (%)	100	100	100	100	100	100	77	100	0	91	100
Microscopic (%)	0	44	0	0	0	0	23	0	100	9	0
Breast (%)	n.r.	15	n.r.	0	0	0	0	0	0	0	0
Chest wall (%)	n.r.	81	n.r.	100	100	100	100	92	81	77	86
Lymph nodes (%)	n.r.	5	n.r.	0	0	0	0	8	19	23	14
Previous treatment											
Surgery (%)	100	95	n.r.	88	100	100	100	98	100	n.r.	n.r.
Chemotherapy (%)	0	62	n.r.	96	0	50	64	86	71	83	n.r.
Hormonal therapy (%)	100	n.r.	n.r.	100	17	n.r.	33	58	51	51	n.r.
Radiation therapy (%)	0	100	93	88	100	89	62	68	54	51	48
RT dose (Gy; mean)	n.r.	n.r.	50.0	62.0	45-63	32.0	50.0	54.2	50.6	40.0	58.0
Present treatment											
RT dose (Gy; mean/ median)	38.9	32.0	32.0	42.0	31.8	29.4	56.4	39.9	42.4	40.0	50.5
HT technique	MW/C	MW	MW	MW	MW	MW	RF	MW/U	MW	MW	C/MW/U
HT duration (min)	45	60	60	60	30	60	60	45	45	45	30-60
HT sessions/ week	1	1	2	≤ 4	2	1-2	2	1-2	1-2	1-2	1-2
HT sessions/ patient	1 vs. 4	4	8	10	5	5-8	5.6	3.1	5.1	8	4.5
Temperature probes	TCP	TCP	T	FO	TCP	TCP	n.r.	FO/T	FO/ T	FO/ T	TCP
Thermal mapping (y/n)	n	n	y	n.r.	n.r.	n	n.r.	y	y	67%	y
Invasive sensors (y/n)	y	y	y	y	y	y	y	y	y	y	y
# Invasive sensors	3	0	n.r.	1-16	1-2	3-6	1-2	n.r.	n.r.	1-8	n.r.
# Skin sensors	3	42	n.r.	n.r.	n.r.	3-8	6	<30	n.r.	n.r.	n.r.

Vernon et al. (1996) PMH trial	Vernon et al. (1996) MRC BRB trial	Vernon et al. (1996) ESHO trial	Vernon et al. (1996) DHG trial	van der Zee et al. (1999)	Seegenschmiedt et al. (1989)	Sannazzari et al. (1989)	Refaat et al. (2015)	Phromratapanongse et al. (1991)	Oldenberg et al. (2015)	Oldenberg et al. (2010)	Linhorst et al. (2015)	Linhorst et al. (2013)	Lindholm et al. (1995)
17	n.r.	90	100	58	60	63	56	57	57	52	61	60	61
0	n.r.	0	0	0	0	0	20	0	0	0	0	0	0
100	100	100	100	100	100	100	80	100	100	100	100	100	100
100	100	100	100	89	100	100	86	100	100	100	100	0	100
0	0	0	0	11	0	0	14	0	0	0	0	100	0
0	28	4	5	n.r.	6	18	24	0	13	44	7	23	0
94	72	89	79	97	71	73	68	82	78	47	91	73	89
6	0	7	16	n.r.	20	9	8	18	9	9	2	4	11
100	n.r.	n.r.	n.r.	94	100	89	81	n.r.	91	100	100	100	98
47	57	52	37	51	73	n.r.	57	n.r.	54	42	69	42	100
76	77	67	47	52	35	n.r.	n.r.	n.r.	58	46	63	56	27
59	88	100	21	100	78	100	43	100	100	100	100	100	100
n.r.	n.r.	n.r.	n.r.	45.0	56.0	<45.0	n.r.	59.7	50.0	65.0	49.0	48.0	48.0
32-50	28,8-50	32,0	32-50	32,0	36,8	20-40	40-60	29,4	32,7	32,0	32,0	32,0	34,5
MW	MW	MW/RF	MW	MW	MW	MW	MW/l	MW	MW/RF	MW	MW	MW	MW
28	60	60	60	60	45	30-45	45	60	60	60	60	60	45
0.5	1	2	2	2	2	2	2	1-2	1-2	1	1	1-2	1-2
2	3-6	4-8	4-8	8	8.6	8-10	5	2-10	4-6	4	4	4.4	4
n.r.	FO/TCP	n.r.	n.r.	FO/TCP	FO	T	T	FO/T	n.r.	n.r.	FO	FO	T
y	n	n	n	y	y	y	y	y	y	y	y	y	n.r.
y	y	y	y	y	y	y	y	y	y	y	y	y	y
2	10	6	7	19.3	4.2	1-8	n.r.	1-3	n.r.	n.r.	13	11	n.r.
6	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	3-4	n.r.	n.r.	n.r.	n.r.	n.r.

Table 1. Continued.

	Arcangeli et al. (1991)	Bakker et al. (2017)	Dragović et al. (1989)	Engin et al. (1993)	Gabriele et al. (2009)	Gonzalez-Gonzalez et al. (1988)	Hehr et al. (2001)	Kapp et al. (1991)	Kapp et al. (1992)	Lee et al. (1998)	Li et al. (2004)
Treatment outcome											
CR (%)	n.r.	n.r.	57	91	66	60	40	72	n/a	63	52
PR (%)	n.r.	n.r.	36	87	30	18	60	0	n/a	19	40
NR (%)	n.r.	n.r.	7	0	5	22	0	28	n/a	16	8
Local recurrence rate (%)	n.r.	n.r.	19	5	27	15	19	29	15	25	n.r.
DLC (months; median)	n.r.	n.r.	n.r.	n.r.	n.r.	7	n.r.	n.r.	n.r.	n.r.	n.r.
3-year LC (%)	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	68	33	n.r.
Survival (months; median)	n.r.	n.r.	17	7	n.r.	n.r.	28	n.r.	n.r.	11	14
3-year OS (%)	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	55	n.r.	n.r.
FU (months; mean/ median)	18	n.r.	n.r.	n.r.	18	n.r.	28	11	9	11	n.r.
Thermal toxicity (%)	n.r.	26	n.r.	50	n.r.	14	15	4	5	n.r.	n.r.
Relationship thermal dose parameter to clinical outcome											
CR (univariate)	n.r.	n.r.	y	n.r.	y	y	n.r.	n.s.	n.r.	n.r.	n
CR (multivariate)	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n	n.r.	n.r.	n.r.
DLC (univariate)	n.s.	n.r.	n.r.	n.r.	n.r.	n.r.	y	y	y	y	n.r.
DLC (multivariate)	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n	y	y	n.r.
OS (univariate)	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.
OS (multivariate)	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.
Thermal toxicity (univariate)	n.r.	n.r.	n	y	n.r.	n.r.	n.r.	n.r.	y	n.r.	n.r.
Thermal toxicity (multivariate)	n.r.	y	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	y	n.r.	n.r.
Risk of bias assessment (QUIPS tool)											
1. Study participation	mod.	low	mod.	mod.	mod.	mod.	mod.	low	low	low	low
2. Study attrition	low	low	mod.	low	low	low	low	low	low	low	mod.
3. Prognostic factor measurement	low	low	mod.	mod.	mod.	high	low	mod.	mod.	mod.	high
4. Outcome measurement	low	low	low	low	low	low	low	low	low	low	low
5. Study confounding	mod.	low	mod.	low	mod.	mod.	mod.	low	low	low	low
6. Statistical analysis and reporting	low	low	high	low	high	high	high	low	low	low	high

Abbreviations: RT= radiotherapy; HT= hyperthermia; CR= complete response; PR= partial response; NR= no response; DLC= duration of local control; LC= local control; OS= overall survival; FU= follow-up; y=yes; n= no; n.r.= not reported; n.s.= not significant; MW= microwave HT;

	Vernon et al. (1996) PMH trial	Vernon et al. (1996) MRC BRR trial	Vernon et al. (1996) ESHO trial	Vernon et al. (1996) DHG trial	van der Zee et al. (1999)	Seegenschmiedt et al. (1989)	Sannazzari et al. (1989)	Refaat et al. (2015)	Phromratapanpongse et al. (1991)	Oldenborg et al. (2015)	Oldenborg et al. (2010)	Linthorst et al. (2015)	Linthorst et al. (2013)	Lindholm et al. (1995)
	29	n.r.	n.r.	n.r.	74	62	59	53	41	58	n/a	70	n/a	71
	n.r.	n.r.	n.r.	n.r.	0	24	41	27	23	28	n/a	n.r.	n/a	29
	n.r.	n.r.	n.r.	n.r.	26	9	0	21	36	14	n/a	n.r.	n/a	0
	4	6	38	21	36	11	20	43	33	25	27	32	18	69
	n.r.	n.r.	n.r.	n.r.	32	n.r.	n.r.	n.r.	6	5	n.r.	15	16	9
	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	25	78	40	83	n.r.
	n.r.	n.r.	n.r.	n.r.	21	n.r.	n.r.	16	n.r.	17	59	19	82	13
	59	21	68	62	n.r.	n.r.	n.r.	30	n.r.	37	66	32	75	n.r.
	n.r.	n.r.	n.r.	n.r.	21	16	12	13	n.r.	17	64	32	42	n.r.
	24	7	22	16	37	25	n.r.	n.r.	24	13	23	23	24	14
		n.r.			y	y	n.s.	y	y	n	n	n.r.	n.r.	Y
		y			n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.
		n.r.			y	y	n.r.	n.r.	n.r.	n	n	y	y	n.s.
		y			n	n	n.r.	n.r.	n.r.	n.r.	n.r.	n	n	y
		n.r.			n.r.	n.r.	n.r.	y	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.
		y			n.r.	n.r.	n.r.	n.r.	n.r.	n	n	n	y	n.r.
		n.r.			n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n.r.	n	n.r.
	low	low	low	low	low	low	mod.	low	mod.	low	low	low	low	low
	low	low	low	low	low	low	low	low	low	low	low	low	low	low
	mod.	mod.	mod.	mod.	mod.	mod.	mod.	mod.	mod.	high	high	mod.	mod.	mod.
	low	low	low	low	low	low	low	low	low	low	low	low	low	low
	low	low	low	low	low	low	low	low	low	low	low	low	low	low
	low	low	low	low	low	high	high	high	high	low	low	low	low	low

RF= radiofrequent HT; C= capacitive HT; I= interstitial HT; U= ultrasound HT; ASAP= as soon as possible; FO= fiber optic probes; T= thermistors; TCP= thermocouple probes; mod= moderate.

Table 2. The location of temperature sensors (skin and/or invasively) used for calculation of a thermal dose parameter influences the relationship of the parameter with outcome. The result is the number of significant relationships ($p < 0.05$) per thermal dose parameter category divided by the total number of reported relationships (significant $p < 0.05$, trend $0.05 < p < 0.1$, or no relationship $p \geq 0.1$).

	Univariate relationship			Multivariable relationship		
	Skin	Invasive	Skin + Invasive	Skin	Invasive	Skin + Invasive
Complete response	0/0	6/19	2/7	0/0	1/6	0/0
Duration of local control	0/0	9/14	0/8	0/0	3/15	0/4
Overall survival	0/0	0/0	1/1	0/0	1/3	1/1
Thermal toxicity	2/2	2/10	4/8	4/4	1/1	0/4

Clinical outcome

The log OR, log RR and RD are above zero for all clinical outcome measures in every study (Figure 2). Thus, patients achieving a higher thermal dose parameter during HT treatment had an increased odds/ probability of CR, longer duration of LC, longer OS and more thermal toxicity. An overview of all reported thermal dose parameters in the included studies with their respective statistical relationships to the outcome measures and their respective temperature/thermal dose category is available as supplementary materials.

Complete response

Eight studies reported a significant univariate relationship between CR and thermal dose [28, 30–32] or temperature [29, 34, 41, 42] (Figure 3, Table 1). Four studies [30, 33, 36, 41] reported a trend ($p < 0.1$) between temperature and CR, while two studies did not find any relationship between CR and thermal dose parameters [43, 44].

Multiple studies showed a significant relationship between the achieved thermal dose invasively (CEM42.5, CEM43, time > 42.5 °C, time > 43 °C) and CR (Figure 4). Patients who received RT and a high thermal dose had on average 34 % (range 27 - 53 %) more CR than patients who received RT and a low thermal dose [4, 15, 28–34].

Several studies investigated the relationship between temperature and CR. Gabriele et al. [29] found that the CR rate was correlated with the average maximum (skin + invasive) temperature during HT treatment (≥ 42 °C 76.5 % CR vs. < 42 °C 30 % CR), while there was no correlation with average temperature or minimum temperature (Figure 3). Similarly, van der Zee et al. [42] showed that the CR rate increased with a higher maximum temperature

in normal tissue ($p < 0.04$), while neither the maximum temperature in tumor tissue ($p = 0.29$) nor the T90 for normal ($p = 0.62$) or tumor tissue ($p = 0.30$) showed an association with CR. Lindholm et al. [41] found that none of the investigated thermal factors predicted CR. However, for the subgroup treated with 915 MHz ($n = 55/59$), the minimum temperature in the HT session with the highest temperatures, was predictive for CR ($p = 0.02$). Seegenschmiedt et al. [34] found a significant relationship between the mean minimum temperature, the minimum temperature of the first session and of the best session and CR. A minimum invasive temperature above 41 °C resulted in a significantly better CR. Sannazzari et al. [33] also found that higher minimum temperatures increased the CR rate (Figure 2).

Both studies that did an in-depth temperature analysis on the trials reported in Vernon et al. [4], found statistically significant multivariable relationships with minimum thermal dose [15, 27], after adjusting for maximum tumor depth and RT regimen (radical or palliative RT) [27] and after adjusting for systemic disease at entry and tumor depth [15].

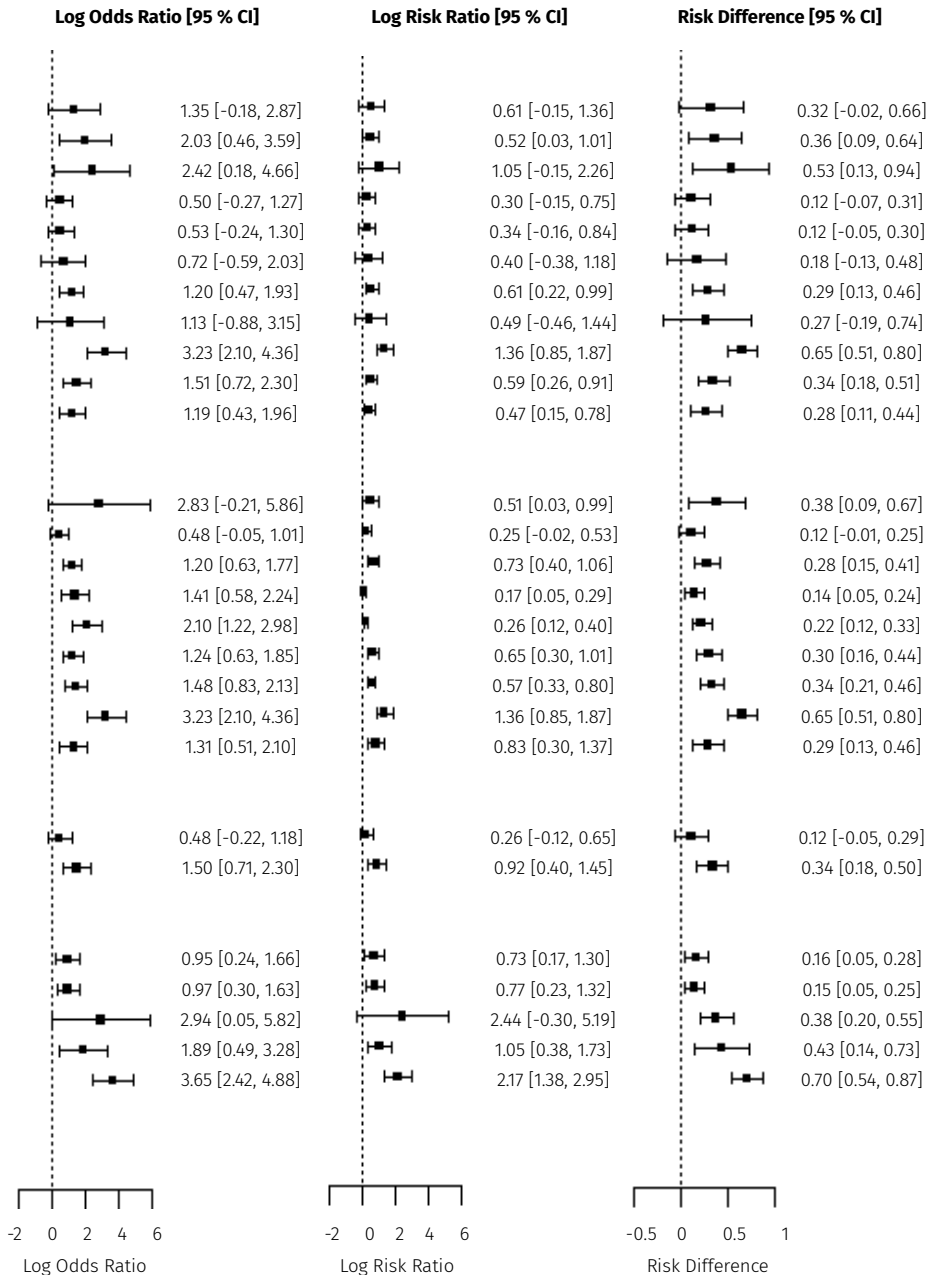
Local control

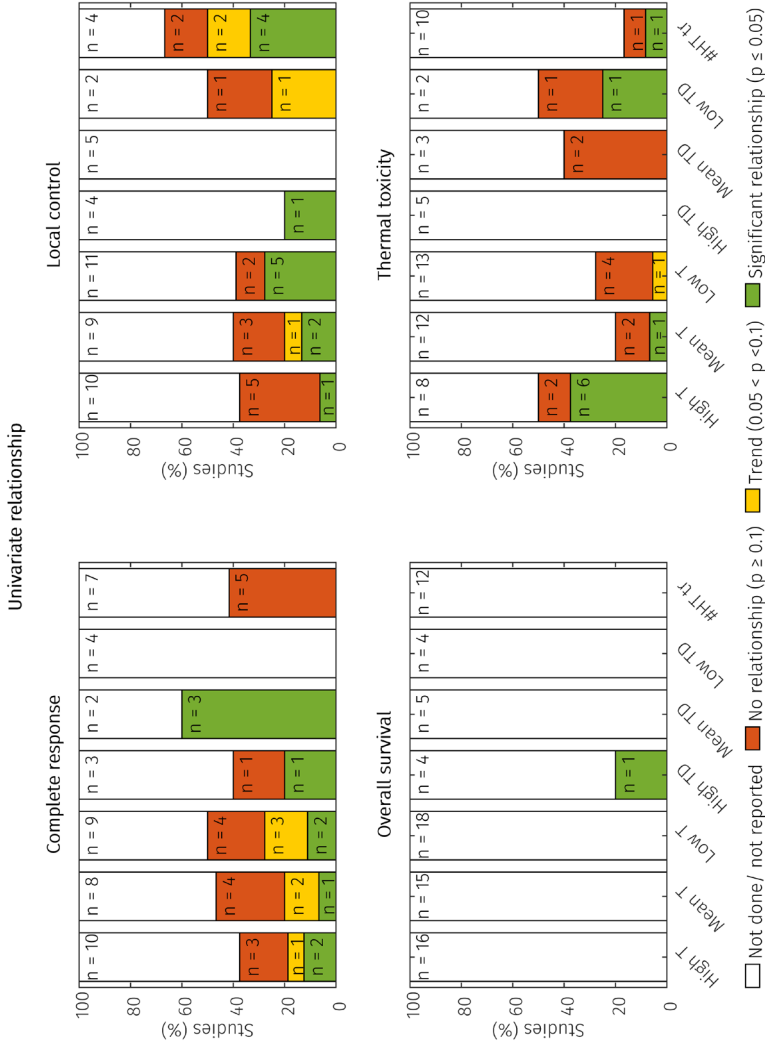
In univariate analysis, duration of LC was significantly correlated with minimum temperature [34, 36, 37], T90 [36, 40, 42] and average temperature [40]. Furthermore, Kapp et al. [45] found that the average percentage of temperatures ≥ 40 °C was significantly correlated with duration of LC ($p < 0.05$) (Figure 2). Van der Zee et al. [42] also found that duration of LC was improved by a higher maximum normal tissue temperature ($p = 0.02$) (Figure 3). Three studies found no relationship (Table 1).

Kapp et al. [36] found that macroscopic tumor lesions that had an average minimum temperature ≥ 40.75 °C or with < 20 % temperatures < 41 °C had longer LC. In patients with microscopic disease, Kapp et al. [37], confirmed that patients achieving an average minimum temperature ≥ 39.7 °C or 96 % temperatures ≥ 40 °C resulted in significantly longer LC. Seegenschmiedt et al., [34] found a significant relationship between LC and the mean minimum temperature, the minimum temperature of the first session and of the best session. Where an invasive minimum temperature above 41 °C resulted in significantly better duration of LC [34]. Lee et al. [38] found that patients who achieved temperatures > 43 °C for a longer duration showed improved duration of LC [38] (Figure 2).

Study	Thermal dose parameter	High		Low	
		CR	PR+NR	CR	PR+NR
Complete response					
Dragovic et al., 1989	≥ 2 sessions ≥ 30 min CEM43	12	5	5	8
Gabriele et al., 2009 (group A)	Tmax ≥ 42 °C	26	3	8	7
Gonzalez-Gonzalez et al., 1988	CEM43 > 20 min	9	2	2	5
Hand et al., 1997 (Vernon MRC BrB)	CEM43 session 1 ≥ 10 min	17	20	33	64
	Σ1-3 CEM43 ≥ 10 min	30	40	15	34
Phromratanapongse et al., 1991	CEM42.5 > 50 min	16	14	5	9
Refaat et al., 2015	Time > 42.5 °C ≥ 200 min	41	23	22	41
Sannazzari et al., 1989	Tmin > 43 °C	7	3	3	4
Seegenschmiedt et al., 1989	Tmin > 41 °C	37	5	12	41
Sherar et al., 1997 (Vernon)	ΣCEM43 > 10 min	48	14	25	33
	Max CEM43 > 6 min	46	16	27	31
Local control at 12 months					
Arcangeli et al., 1991	4 vs. 1 HT sessions	11	0	7	5
Kapp et al., 1991	Tmin ≥ 40.75 °C	45	40	64	92
	% T <41 °C ≥ 20%	41	35	43	122
Kapp et al., 1992	Tmin ≥ 39.7 °C	177	11	59	15
	% T ≥ 40 °C ≥ 96%	182	8	53	19
Lee et al., 1998	Time > 43 °C ≥ 10 min	76	46	24	50
	> 8 HT sessions	58	17	53	68
Seegenschmiedt et al., 1989 *	Tmin > 41 °C	37	5	12	41
Sherar et al., 1997 (Vernon)	Max CEM43 > 6 min	32	30	13	45
Overall survival at 18 months					
Refaat et al., 2015	Time > 42.5 °C ≥ 200 min	33	31	25	38
Sherar et al., 1997 (Vernon)	ΣCEM43 > 10 min	35	27	13	45
Thermal toxicity					
Linthorst et al., 2013	Tmax ≥ 43 °C	31	67	14	78
Linthorst et al., 2015	Tmax ≥ 43 °C	37	93	15	99
Lindholm et al., 1995 (915 MHz)	Tmax > 43 °C	18	26	0	13
	Tmax > 44 °C	8	4	10	33
Seegenschmiedt et al., 1989	Tmax > 45 °C	23	6	6	60

Figure 2. Forest plots of the odds ratio (OR), risk ratio (RR) and risk difference (RD) for complete response (CR) [4, 15, 28–34], local control (LC) at 12 months [15, 35–38] and * 6 months [34], overall survival (OS) at 18 months [15, 32], and thermal toxicity [34, 39–41]. The values were calculated from subgroups of patients, which were grouped based on their variation in thermal dose parameters during hyperthermia treatment; thermal dose parameters above a certain cut-off value (high) vs. thermal dose parameters below the cut-off value (low). The OR and RR are presented as the logarithmic mean with 95 % confidence interval (CI). The RD is presented as the mean with 95 % CI. *Abbreviations:* CEM43= cumulative equivalent minutes at 43 °C [19, 20], Tmax= maximum temperature, Tmin= minimum temperature, CR= complete response, PR= partial response, NR= no response, LC= local control, Tox= thermal toxicity.





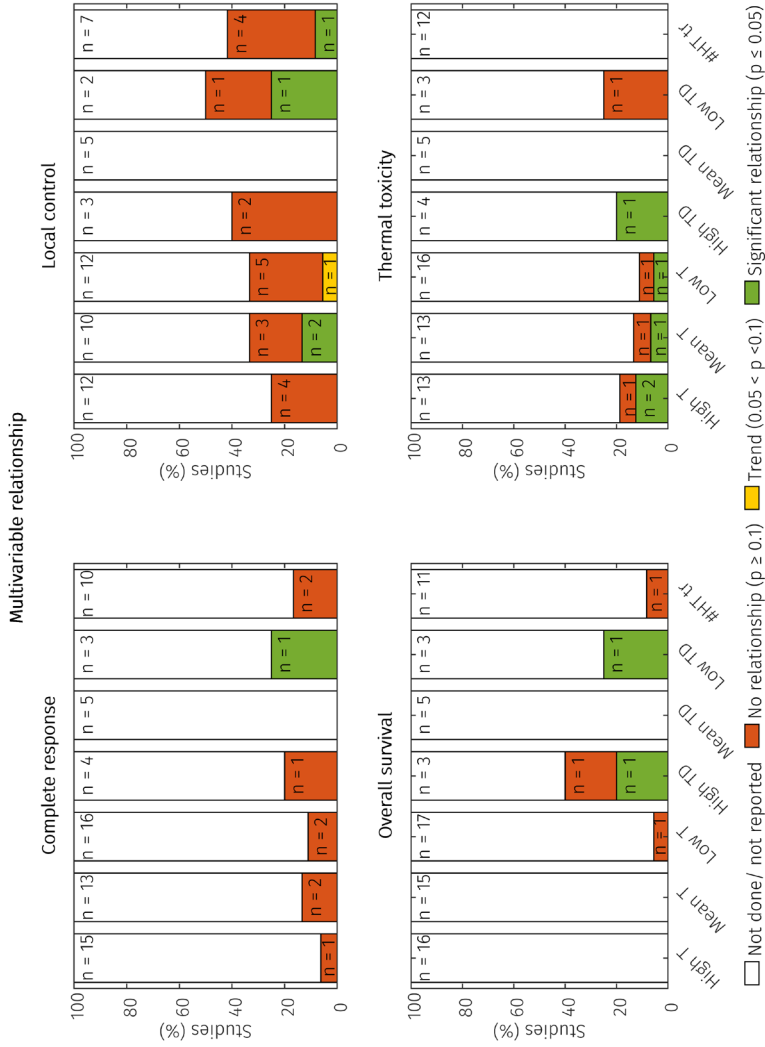


Figure 3. The number of studies where a high-end, mean or low-end thermal parameter (T), thermal dose parameter (TD) or number of hyperthermia treatments (#HT tr), was reported in relation to complete response, duration of local control, overall survival and thermal toxicity. The univariate or multivariable relationship of the thermal dose parameters with one of the outcome measures is labeled as either a significant relationship ($p < 0.05$) a trend (p -value between 0.05 and 0.1), no relationship ($p \geq 0.1$) or as not done or not reported. The labels inside the bars represent the number of studies.

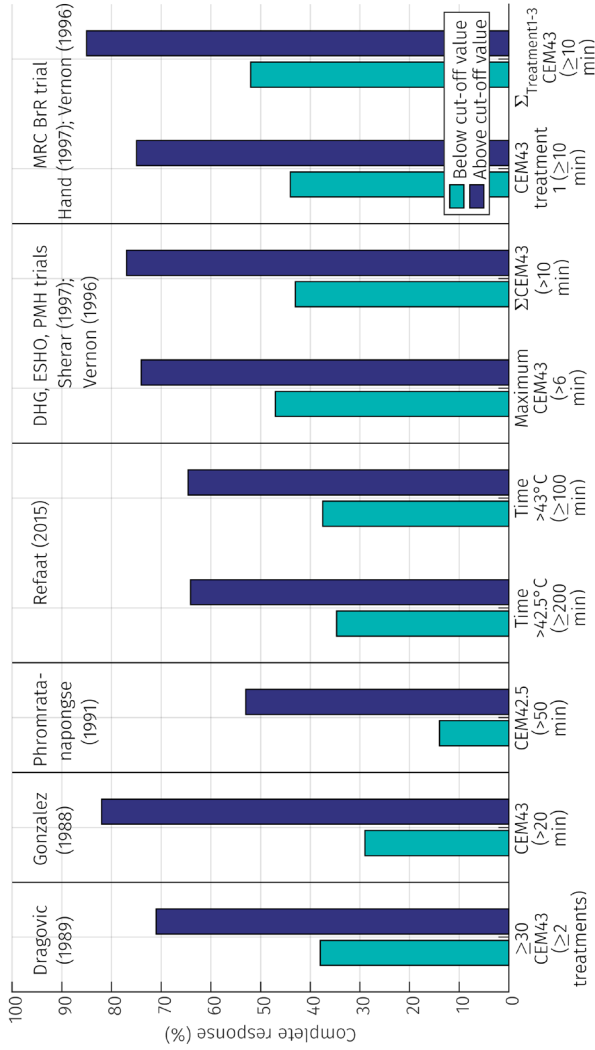


Figure 4. The relationship between tumor response and thermal dose below/above certain cut-off values (noted between the brackets) [4, 15, 28–34].

Eight studies investigated the relationship between the number of HT treatments and LC. Four found a significant relationship [37–39, 46] and two a trend [35, 41]. Kapp et al., [37] found that \geq two HT treatments improved duration of LC compared to one treatment in total. Hehr et al. [46] found that $>$ seven HT sessions improved duration of LC. Lee et al. [38] treated patients with a median of eight treatments, where more treatments were better, which remained significant in multivariable analysis. Arcangeli et al. [35] treated one patient with 40 lesions, where a lesion was either treated with one or four HT treatments. When a lesion received four HT treatments there was less tumor regrowth and longer freedom from local progression (Figure 2).

In multivariable analysis, several thermal dose parameters remained significant for a relationship with duration of LC:

- The minimum thermal dose after adjusting for systemic disease at entry [4, 15].
- The average temperature after adjusting for lesion size and time interval between diagnosis primary tumor and present treatment [41].
- The percentage temperature ≥ 40 °C after adjusting for estrogen receptor status, initial T-stage, time interval between initial breast cancer to first failure, age at HT and concurrent RT [37].
- The number of HT treatments and ≥ 10 minutes > 43 °C after adjusting for initial SAR, current RT dose, tumor volume and tumor thickness [38].

Overall survival

Both studies investigating the relationship between thermal dose parameters and OS found a positive relationship. Most relevant thermal dose parameters correlating with survival are:

- The time > 42.5 °C. Refaat et al. [32] reported that when patients were treated with more than 200 minutes above 42.5 °C, patients had better OS. This remained significant in multivariable analysis after adjusting for microscopic disease, RT dose and mastectomy.
- CEM43T100. The DHG, ESHO and PMH trials [4, 15] found in multivariable analyses that OS at 18 months after treatment was significantly better for patients who received more than 6 minutes CEM43T100 compared to patients who received less than 6 minutes CEM43T100, 57 % versus 22 %, respectively, after adjusting for systemic disease at entry, age, and tumor area (Figure 2).

Thermal toxicity

Overall, seven studies report a relationship between toxicity and maximum temperature [34, 37, 39–41, 47, 48], of which two studies also report a relationship with thermal dose [39, 47]. Furthermore, Lindholm et al. [41] found that two HT treatments per week gave more severe skin reactions than one treatment per week. Linthorst et al. [39] found that the average temperature on the skin had a relationship with thermal toxicity. While four studies report no relationship [28, 31, 42, 44] (Figure 3, Table 1).

Several studies showed a relationship between the recorded maximum temperature on the skin and thermal toxicity (Figure 2). Engin et al. [48] found that patients without a skin reaction had a maximum temperature on the skin of 42.1 ± 0.5 °C (mean \pm SD), while patients with thermal blisters had a maximum temperature of 43.7 ± 0.2 °C ($p < 0.01$). Seegenschmied et al. [34] found that thermal adverse events (superficial blisters and deep burns) were significantly correlated with maximum temperatures of more than 45 °C ($p < 0.01$). Lindholm et al. [41] found that one of the prognostic factors for blisters was the highest average maximum skin temperature during a given HT treatment ($p = 0.03$). The correlation between normal tissue damage and maximum skin temperature was even stronger ($p = 0.01$) when only tumors treated with 915 MHz were considered ($n = 55/59$). Linthorst et al. [39] found that invasively measured thermal parameters had no influence on thermal toxicity, whereas the incidence of grade 2 and 3 burns clearly increased with a higher maximum skin temperature. Patients with a maximum skin temperature < 43 °C had a grade 2 or 3 burn incidence of 15 %, while incidence was 32 % in patients with a maximum skin temperature > 43 °C ($p = 0.006$). In 2015, Linthorst et al. [40] again showed that the incidence of grade 2 and 3 skin burns increased with a higher maximum skin temperature; 13 % incidence when the maximum skin temperature was < 43 °C and 28 % when ≥ 43 °C ($p = 0.003$), see Figure 5. Likewise, Bakker et al. [47] found in multivariable analysis a significant relationship between the maximum temperature on the skin and the occurrence of thermal skin damage adjusted for patient and tissue type, they derived probability models for skin toxicity for scar and skin tissue (Figure 5).

Multivariable analysis by Kapp et al. [37] revealed a higher maximum invasive temperature in fields developing adverse events (cutaneous and/or subcutaneous induration/fibrosis, normal tissue burns/ ulceration, edema, pain) compared with fields free of adverse events (45.8 °C vs. 45.2 °C, respectively; $p = 0.005$), after adjusting for HT device, tumor stage and site. Furthermore they found a correlation for the percentage of invasive temperature ≥ 45 °C, after adjusting for HT device, tumor stage and site.

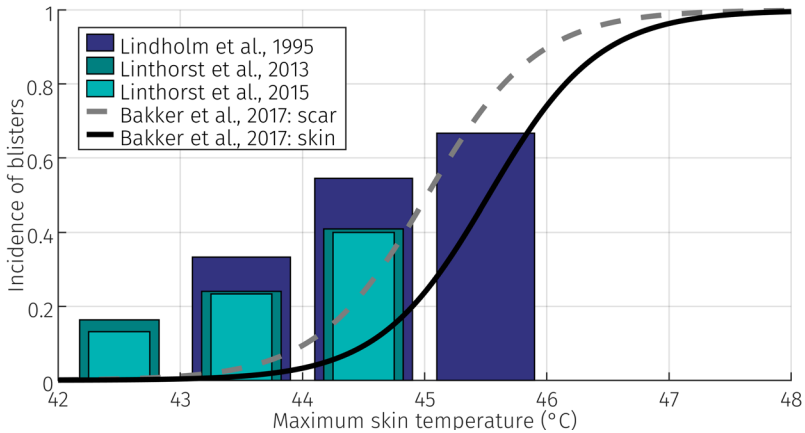


Figure 5. The incidence of thermal toxicity (blisters) increases with a higher maximum skin temperature, data adapted from Lindholm et al. [41], Linthorst et al. [39] and Linthorst et al. [40] combined with probability models for thermal toxicity for scar and skin tissue [47].

Three studies [31, 42, 44] found no correlation between thermal parameters and adverse events. Although, van der Zee et al. [42] did find a correlation between the technique used to deliver HT, e.g. 434 MHz treatments caused much less acute damage than 2450 MHz treatments ($p = 0.001$). They reported that the acute damage for 2450 MHz systems was likely due to having no water bolus cooling. Indeed, when Lindholm et al. [49] added water bolus cooling to a 2450 MHz system, toxicity decreased from 55 % to 17 %.

DISCUSSION

This systematic review shows that for patients with locoregional recurrent breast cancer treated with RT-HT, higher temperature and/or thermal dose during HT improves CR, LC and OS [4, 15, 27–32, 34, 36–38, 40–42], while increasing the incidence of thermal toxicity [34, 37, 39–41, 47, 48]. It is intriguing to see that the CR of patients who received RT alone vs. patients who received RT with low dose HT were comparable; 38.1 % [6] vs. 36.2 % (Figure 4), respectively. Comparable benefit in CR was found when either RT-HT or RT with high dose HT was received; 60.2 % [6] vs. 66.6 % (Figure 4), respectively. The achieved temperature and thermal dose during HT thus at least partly explains the reported variation in CR after RT-HT in patients with recurrent breast cancer [6].

Optimizing the temperature and thermal dose of HT treatment in patients with locoregional recurrent breast cancer in re-irradiated area is therefore of utmost importance. The included studies aimed for minimum temperatures between 41 °C and 43 °C, which were rarely achieved. Whereas the achieved maximum temperature was frequently higher than the beforehand specified allowed maximum temperature, which varied between 43 °C and 50 °C. This implies a very heterogeneous temperature distribution which allowed maximum temperatures to vary widely between institutes. The recently published ESHO guidelines for superficial HT quality assurance might help to improve overall quality assurance practice and reduce variation between institutes. These guidelines advise to aim for a T90 > 40 °C and T50 > 41 °C to achieve local tumor control, with maximum temperatures of 43 - 45 °C [50]. Furthermore, hyperthermia techniques have improved over the years and more uniform and adequate heating is presently feasible.

Temperature and thermal dose measurement

Although achieved temperatures and thermal dose during HT treatment determined treatment quality, these treatment parameters were often not reported. Six studies concerning HT and RT in patients with recurrent breast cancer were excluded from this review because no temperature data were reported. In the studies that were included, the reported thermal dose parameters were often not tested for a relationship with outcome (four studies) or the result of the tests were not reported (Figure 3).

In total, the values of 27 different thermal dose parameters were presented and 32 parameters were tested for relationships. These parameters ranged from simple temperature statistics (minimum, average and maximum temperature, etc.) to more complex thermal dose parameters (CEM43, minutes > 42.5 °C, % temperature > 45 °C, etc.). Besides that, there is also an institutional variation in the calculation of thermal dose parameters. In the rare studies where thermal dose parameters were reported, the reported parameters varied

widely, e.g. low-end versus high-end temperatures. Although some correlation is expected because thermal dose parameters from the high-end, mean and low-end temperatures are correlated [27, 51]; comparison of these parameters was nevertheless difficult.

The measured temperature during HT is vastly dependent on the type, extent and location of thermometry used; e.g. the number of sensors, type of thermometry equipment [52], location of sensors (e.g. invasive, skin, macroscopic tumor, scar, healthy tissue) [47], but also on the use of a (circulating) water bolus during HT treatment [49]. For example, it is essential to have a sufficient number of sensors, because sampling the true temperature distribution during HT treatment with a small number of temperature sensors may introduce significant errors in the measured and calculated thermal dose variables compared to the true underlying temperature distribution [53–57]. Several studies attribute their inability to find a dose-effect relationship to the low number of temperature sensors [43, 48, 58]. Thus, adequate thermometry feedback during treatment and registration of thermal dose parameters are essential to ensure treatment quality. To allow comparison of the achieved temperature and thermal dose between different institutes, a uniform registration, calculation and reporting of thermal dose parameters is required.

Abovementioned arguments were a leading factor in scoring a moderate to high risk of bias in one of the domains of the QUIPS tool; i.e. the prognostic factor measurement. Only three out of the 22 included studies had a low risk of bias in this respective domain (Table 1).

Relationship of thermal dose parameters with outcome

An association between treatment outcome and thermal dose parameters was not found in all studies. This is not surprising, since most of the included studies were not designed to investigate the presence of a HT dose-response effect. Studies aimed at giving a uniform thermal dose which, if successful, will result in little variation in HT parameters, and thus a relationship of a HT parameter with outcome is more difficult to prove. Nevertheless, most included studies (22/25) showed enough variation in several of the thermal dose parameters to find significant positive relationships with outcome in both univariate and multivariable analysis. Three studies [43, 44, 59] did not find any relationship between thermal dose parameters and CR, LC and OS. Both studies of Oldenburg et al. [44, 59] only investigated parameters derived from the skin, while Li et al. [43] used a low number of invasive temperature sensors. Furthermore, univariate analysis does not take into account other patient-, tumor- and treatment related parameters, which also contribute to the final outcome of RT-HT. In this systematic review, only ten studies have performed multivariable analysis to adjust for patient- tumor- and treatment related parameters [15, 27, 47, 32, 36–42]. Of these studies, in seven of the 37 multivariable analyses performed

for CR, LC and OS, the addition of a thermal dose parameter improved the model. To clarify the need for multivariable analysis, for example Hehr et al. [46] found that more HT treatments yielded better duration of LC, but those patients probably received more RT treatments and thus a higher RT dose (range 30 - 68 Gy). The RT dose is known to influence treatment outcome in cancer patients. Patients with primary breast cancer treated with a high RT dose benefit less from the addition of HT than patients treated with a lower dose palliative RT scheme [4, 16]. Performing a multivariable analysis to investigate the relationship of outcome with thermal dose parameters is thus essential to establish a reliable correlation. Not performing a multivariable analysis yields a high risk of bias in this situation.

Several studies find a relationship between the number of HT treatments and duration of LC. However, two randomized studies investigating the number of HT treatments found no difference between groups. In a prospective randomized study, comparing once versus twice weekly HT applications in 127 patients with chest wall recurrences of breast cancer, no difference was found in CR and duration of LC [60] (patients also reported in [34]). Similarly, Kapp et al. [61] compared two versus six HT treatments in 70 patients with superficially located tumors and found no difference in CR and duration of LC. For both studies, the RT dose was similar between groups.

In general, the addition of HT to RT is well tolerated and adds no significant late toxicity in patients treated with RT-HT, although more blistering occurs during RT-HT compared to RT alone [4]. This acute thermal toxicity is caused by higher maximum temperatures [34, 37, 39-41, 47, 48] and thermal dose [39, 47] on the skin during HT treatment. In untreated healthy skin, a higher thermal dose on the skin also increases blistering [62, 63].

As shown in Table 2, thermal dose variables derived superficially were predictive for thermal toxicity, while thermal dose variables derived from invasive temperature sensors were predictive for both CR, LC, OS and thermal toxicity. Surface temperature measurements during HT reflect an average of the water bolus and the true surface temperature. Nevertheless, the true surface temperature might indeed have a significant relationship with outcome. Therefore, it is important to measure both invasive and superficial temperatures during treatment and to report them separately.

Other HT parameters that are related to outcome are the specific absorption rate (SAR) coverage [38], type of HT device [38, 42] and frequency [37, 42]. These HT parameters influence the achieved temperature and thermal dose during HT.

Relationship between thermal dose parameters and outcome in other tumor types

Relationships between thermal dose parameters and clinical outcome have not only been found in recurrent breast cancer, but also in cervical carcinomas [21, 22, 64], sarcomas [56, 65], rectal cancer [66], brain tumors [67] and melanomas [2, 68]. Thermal dose variables (maximum CEM43, CEM43T50, CEM43T90, CEM43T100, TRISE) [2, 21, 22, 56, 64, 65], minimum temperature variables (minimum temperature, T90) [65, 67, 69] and maximum temperature variables [2] were found to relate to outcome.

The first randomized controlled study showing a relationship between thermal dose achieved during HT and outcome was published in 1984 [70]. Dewhurst and Sim randomized 236 dogs and cats with a variety of cancers to either receive RT alone or RT-HT. In the RT-HT group (n = 116) a relationship between CEM43T100 and both CR and duration of LC was found, while the maximum CEM43 was related to thermal injury. Animals achieving an average CEM43T100 ≤ 1 during all HT treatments had a CR comparable to the CR in the group treated with RT alone (n = 120). Furthermore, extensive temperature data from animal studies indicated that the coolest part of the tumor determined the response to RT-HT [70–72]. Interestingly, in animals with tumors that showed very heterogeneous heating more skin toxicity was found while the response rate decreased [72, 73]. It is likely that the maximum temperature limited the HT operator to increase the temperature in the coolest part of the tumor. These pioneering studies paved the way for subsequent human trials.

CEM43

The suitability of CEM43 to represent thermal dose is a widely debated subject, as excellently summarized by van Rhoon in 2016 [17]. CEM43 is based on the direct cytotoxic effect of heat. The amount of cell death depends on the temperature and duration of heating and has been shown to be suitable for predicting the risk of adverse events of RT-HT in normal tissue [47, 63]. CEM43 does not include all underlying synergistic mechanisms of HT to RT when used to predict treatment efficacy of combined RT-HT; e.g. inhibiting DNA damage repair [74], selective killing of radioresistant hypoxic tumor cells [75] and increased radiosensitivity by enhanced tissue perfusion and reoxygenation [76]. Each of these mechanisms display a different dose-effect relationship [10]. Furthermore, instead of representing the direct effect of HT quality, the measured CEM43T90 might instead actually represent tumor characteristics (i.e. perfusion level), which are predictive of sensitivity to RT. The ability to realize a uniform and high temperature is likely to be associated with a homogeneously perfused and thus well oxygenated tumor. Such a tumor would generally respond well to treatment, even with RT alone. Therefore, the development of a new dose parameter incorporating the combined RT-HT effect and/or employing thermoradiotherapy planning [77] might more accurately represent the synergistic effect of RT and HT in the whole area treated with RT-HT.

Nevertheless, most underlying HT mechanisms display increasing HT effectiveness with higher temperatures. This might explain why multiple clinical studies included in this review did find a relationship between CR and CEM43 or other thermal dose parameters (Figure 4). Consequently, CEM43T90 has been deployed in a prospective setting to investigate whether patients with superficial tumors treated with $CEM43T90 \leq 1$ or $CEM43T90 > 10$ had different outcome. Indeed, superficial tumors treated with a thermal dose $CEM43T90 > 10$ had considerably longer duration of LC than patients with $CEM43T90 \leq 1$ [16]. So although the CEM43 concept has shortcomings, at present this parameter does show a relationship with outcome in many studies.

CONCLUSION

This systematic review shows that higher temperature and thermal dose during hyperthermia therapy significantly improve clinical outcome, e.g. complete response, local control and overall survival; and increase thermal toxicity for patients with recurrent breast cancer treated with radiotherapy and hyperthermia. A sufficiently high dose hyperthermia is required to achieve the radiosensitizing effect of hyperthermia. Thermal dose parameters derived from the surface have a relationship with thermal toxicity, whereas invasive thermal dose parameters have a relationship with response, local control and overall survival as well as with thermal toxicity. Achieving a clinically relevant effect of hyperthermia when added to radiotherapy for patients with locoregional recurrent breast cancer in previously irradiated area, requires ensuring high dose hyperthermia.

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SUPPLEMENTARY MATERIALS

Supplementary Table 1. The categorization of the reported thermal dose parameters measured with sensors positioned on the skin/ and or invasively and their statistical relationships (1= significant $p < 0.05$; trend= $0.05 < p < 0.01$, 0= not significant $p > 0.1$) with outcome and thermal toxicity as reported in the included studies. When multiple variables per study are within one temperature/ thermal dose category, only one is included in the analysis of the systematic review, the other variables are displayed between brackets.

Study	Thermal dose parameter	Position		Temperature		
		inv.	skin	high	mean	low
Arcangeli et al. (1991)	Tmax (°C)	x		x		
	Tmean (°C)	x			x	
	Tmin (°C)	x				x
	Number HT treatments					
Bakker et al. (2017)	Tmax (°C)		x	x		
	Tmax (sensor) (°C)		x	(x)		
	T10 (°C)		x	(x)		
	T10 (sensor) (°C)		x	(x)		
	T50 (°C)		x		(x)	
	T50 (sensor) (°C)		x		x	
	T90 (°C)		x			(x)
	T90 (sensor) (°C)		x			x
Dragovic et al. (1989)	CEM43 (min)	x				
	CEM43 ≥ 30 min (≥ 2 sessions; ≤ 1)	x				
	CEM43, best session (min)	x				
	CEM43, worst session (min)	x				
Engin et al. (1993)	Tmax, max (°C)		x	x		
	Tmax (°C)		x	(x)		
	Tmean (°C)		x		x	
	Tmin (°C)		x			x
	Tmin, first session (°C)		x			(x)
	CEM43 (min)		x			

Supplementary Table 1. Continued.

Study	Thermal dose parameter	Position		Temperature		
		inv.	skin	high	mean	low
Gabriele et al. (2009)	Tmax (°C)	x	x	x		
	Tmax (°C) (≥42; <42)	x	x	(x)		
	Tmean (°C)	x	x		(x)	
	Tmin (°C)	x	x			x
Gonzalez-Gonzalez et al. (1988)	Tmax, max (°C)	x		x		
	Tmean (°C)	x			x	
	Tmin, worst session (°C)	x				x
	CEM43 (min)		x			
	CEM43 (min) (>20; ≤20)	x				
Hehr et al. (2001)	Tmax (°C)		x	x		
	Tmax (°C)	x		(x)		
	T20 (°C)	x		(x)		
	T50 (°C)	x			x	
	Tmean (°C)		x		(x)	
	Tmean (°C)	x			(x)	
	T90 (°C)	x				x
	Number HT treatments (≥7; <7)					
Kapp et al. (1991)	Tmax (°C)	x		x		
	Tmean (°C)	x			x	
	T90 (°C)	x				x
	T90 (°C) (≥41; <41)	x				(x)
	Tmin (°C)	x				x
	% T > 43.5°C (%)	x		(x)		
	% T < 41°C (%)	x				(x)
	Number HT treatments					
Kapp et al. (1992)	Tmax (°C)	x		x		
	Tmean (°C)	x			(x)	
	Tmin (°C) (≥39.7; <39.7)	x				x
	% T > 43.5°C (%)	x		(x)		
	% T ≥ 45°C (%) (>3; <3)	x		(x)		
	% T ≥ 40°C (%)	x			x	

Thermal dose			#tr	CR		DLC		OS		Tox	
high	mean	low		uni	multi	uni	multi	uni	multi	uni	multi
				1							
				1							
				0							
				0							
				trend							
				trend							
				trend							
	(x)										
	x			1							
			x			1					
				0	0	0	0				
				0	0	trend	0				
				trend	0	trend	0				
						1	0				
				0	0	1	0				
				0	0		0				
				0	0	1	0				
			x	0	0	0	0				
						0	0		1	1	
						0	0				
						1	trend		0		
						0	0				
						0	0		1	1	
						1	1				

Supplementary Table 1. Continued.

Study	Thermal dose parameter	Position		Temperature		
		inv.	skin	high	mean	low
Kapp et al. (1992) continued.	% T < 41°C (%)	x				(x)
	Number HT treatments					
Lee et al. (1998)	Time > 43 ≥ 10 min (yes; no)	x				
	Time > 43 ≥ 15 min (yes; no)	x				
	Time > 43 ≥ 20 min (yes; no)	x				
	Time > 43 ≥ 45 min (yes; no)	x				
	Number HT treatments (>8; 5-8; 2-4; <2)					
Li et al. (2004)	Tmax, at end treatment (°C)	x		x		
	Tmean (°C)	x			x	
	Tmin, at end treatment (°C)	x				x
	% T > 42°C (%)	x				
	Number HT treatments					
Lindholm et al. (1995)	Tmax (°C)		x	(x)		
	Tmax (°C)	x		(x)		
	Tmax, max (°C)		x	x		
	Tmean (°C)	x			x	
	Tmean (°C)		x		(x)	
	Tmin (°C)	x				(x)
	Tmin, best session (°C)	x				(x)
	Tmin, first session (°C)	x				(x)
Number HT treatments (1; 2 per week)						
Lindholm et al. (1995) subgroup 915 MHz	Tmax (°C)	x		(x)		
	Tmean (°C)	x			(x)	
	Tmin, best session (°C)	x				x
Linthorst et al. (2013)	Tmax (°C)	x		x		
	Tmax (°C)		x	x		
	Tmean (°C)	x			x	
	Tmean (°C)		x		x	
	T90 (°C)	x				x
	T90 (°C)		x			x

Thermal dose			#tr	CR		DLC		OS		Tox	
high	mean	low		uni	multi	uni	multi	uni	multi	uni	multi
						0	0				
			x			1	0				
x						1	0				
(x)						1	0				
(x)						1	0				
(x)						1	0				
			x			1	1				
				0							
				0							
				0							
x				0							
			x	0							
				0						0	
				0						1	
				0		0	1			0	
				0							
				0							
				0							
			x	0		trend				1	
				trend							
				trend							
				1							
						0	0			0	0
										1	
						0	0			0	0
										1	
						0	0			0	0
										trend	

Supplementary Table 1. Continued.

Study	Thermal dose parameter	Position		Temperature		
		inv.	skin	high	mean	low
Linthorst et al. (2013) continued.	CEM43 T90 (min)	x				
	CEM43 T90 (min)		x			
	Number HT treatments (1-4; 5-8)					
Linthorst et al. (2015)	Tmax (°C)	x		x		
	Tmax (°C)		x	x		
	Tmean (°C)	x			x	
	T90 (°C)	x				x
	CEM43 T90 (min)	x				
Oldenberg et al. (2010)	T10 (°C)	x		x		
	T10 (°C)		x	(x)		
	T50 (°C)	x			x	
	T50 (°C)		x		(x)	
	T90 (°C)	x				x
	T90 (°C)		x			(x)
	CEM43 T50 (min)	x				
	CEM43 T50 (min)		x			
	CEM43 T90 (min)	x				
CEM43 T90 (min)		x				
Oldenberg et al. (2015)	Tmax (°C)	x	x	x		
	T10 (°C)	x	x	(x)		
	T50 (°C)	x	x		x	
	T90 (°C)	x	x			x
	Number HT treatments (1; 2 per week)					
Phomratanapongse et al. (1991)	CEM42.5 (min) (>50; ≤50)	x				
	Number HT treatments (>4; ≤4)					
Refaat et al. (2015)	Time > 42.5 (min) (≥200; <200)	x	x			
	Time > 43°C (min) (≥100; <100)	x	x			
Sannazzari et al. (1989)	Tmin (°C) (>42; <42)	x				x

Supplementary Table 1. Continued.

Study	Thermal dose parameter	Position		Temperature		
		inv.	skin	high	mean	low
Seegenschmiedt et al. (1989)	Tmax (°C) (>45; <45)	x		x		
	Tmax, best session (°C) (>45; <45)	x		(x)		
	Tmax, first session (°C) (>45; <45)	x		(x)		
	Tmin (°C) (>41; <41)	x				x
	Tmin, best session (°C) (>41; <41)	x				(x)
	Tmin, first session (°C) (>41; <41)	x				(x)
	Number HT treatments					
van der Zee et al. (1999)	Tmax (°C)	x		(x)		
	Tmax, normal tissue (°C)	x		x		
	T90, best session (°C)	x				x
	T90, best session (°C)		x			(x)
Vernon et al. (1996) all studies data retrieved from Sherar et al. (1997)	T90, best session (°C)	x				x
	Time > 42°C, max (min)	x				
	Time > 42°C, total (min)	x				
	CEM43 T100, max (min)	x				
	CEM43 T100, total (min)	x				
	Number HT treatments					
Vernon et al. (1996) MRC BrR trial data retrieved from Hand et al. (1997)	Tmax, max (°C)	x		(x)		
	Tmax, min (°C)	x		(x)		
	Tmax (°C)	x		(x)		
	T50 (°C)	x			x	
	T90 (°C)	x				(x)
	Tmin (°C)	x				(x)
	Tmax - Tmin (°C)	x			(x)	
	% Tmax > 42°C (%)	x		(x)		
	% Tmax > 43°C (%)	x		(x)		
	% Tmax > 44°C (%)	x		(x)		
	% Tmean > 42°C (%)	x		(x)		
	% Tmean > 43°C (%)	x		(x)		
	% Tmean > 44°C (%)	x		(x)		

Thermal dose			#tr	CR		DLC		OS		Tox	
high	mean	low		uni	multi	uni	multi	uni	multi	uni	multi
											1
											1
											1
				1		1					
				1		1					
				1		1					
			x								0
				0		0					0
				1		1	0				0
				0		1	0				0
				0		0					0
					0	0		0			
(x)					0	0		0			
x					0	0		0			
		(x)			1	1		0			
		x			1	1		1			
			x		0	0		0			
				0							
				0							
				0							
				1	0						
				trend	0						
				1	0						
				1	0						
				1	0						
				1	1						
				1	0						
				1	0						
				0							
				0							

Supplementary Table 1. Continued.

Study	Thermal dose parameter	Position		Temperature		
		inv.	skin	high	mean	low
Vernon et al. (1996)	Time > 41°C, max (min)	x				
MRC BrR trial	Time > 42°C, max (min)	x				
data retrieved from	Time > 43°C, max (min)	x				
Hand et al. (1997)	Time > 44°C, max (min)	x				
continued.	Time > 42°C, max total (min)	x				
	Time > 43°C, max total (min)	x				
	Time > 44°C, max total (min)	x				
	Time > 40°C, min (min)	x				
	Time > 41°C, min (min)	x				
	Time > 42°C, min (min)	x				
	Time > 43°C, min (min)	x				
	Time > 44°C, min (min)	x				
	Time > 40°C, min total (min)	x				
	Time > 41°C, min total (min)	x				
	Time > 42°C, min total (min)	x				
	Time > 43°C, min total (min)	x				
	Time > 44°C, min total (min)	x				
	CEM43 max (min)	x				
	CEM43 max, total (min)	x				
	CEM43 T100 (min)	x				
	CEM43 T100, total (min)	x				

Abbreviations: inv= invasive; #tr= Number of HT treatments; CR= complete response; DLC= duration of local control; OS= overall survival; Tox= thermal toxicity; uni= univariate analysis; multi= multivariable analysis; Tmax= maximum temperature; Tmax, max= highest maximum temperature;

Thermal dose			#tr	CR		DLC		OS		Tox	
high	mean	low		uni	multi	uni	multi	uni	multi	uni	multi
(x)			1	0							
(x)			0								
(x)			0								
(x)			0								
(x)			0								
(x)			0								
(x)			0								
		(x)	1	0							
		(x)	1	0							
		(x)	1	1							
		(x)	0								
		(x)	0								
		(x)	1	0							
		(x)	1	0							
		(x)	1	0							
		(x)	0								
		(x)	0								
(x)			0								
(x)			0								
		(x)	1	trend							
		(x)	1	1							

Tmean= average temperature; Tmin = minimum temperature; T10, T20, T50, T90, T100= the 10th, 20th, 50th, 90th and 100th percentile of all temperature measurements, respectively; CEM42.5, CEM43= the cumulative equivalent minutes at 42.5°C or 43°C, respectively.