Patient Performed Reading of a Phototest – A Reliable Method?

L. Thorslund¹ and M. Falk*¹,²

¹Department of Medical and Health Sciences, Division of Community Medicine, Primary Care Linköping University, 581 85 Linköping, Sweden
²The Research and Development Unit for Local Health Care, County of Östergötland, 581 85 Linköping, Sweden

Abstract: In various situations, in clinical practice or for prevention purposes directed at skin cancer, a broadened use of phototesting to estimate individual skin UV-sensitivity may be warranted. The aim of the present study was to investigate, in a primary health care population, the reliability of patient performed reading of a UVB phototest, when compared to the reading of a trained physician. Thirty-two subjects, all patients recruited in a primary health care population, underwent a UVB phototest, applied on the forearm. Test reading was performed after 24 hours, by the subjects themselves, by counting the number of erythematic reactions (0-6) detectable, and immediately after this, an independent control reading performed by a doctor was also done. The results showed a 72% absolute agreement between the subjects’ readings and the control readings, and with a weighted kappa-value of 0.78 (95 CI: 0.64 – 0.91), i.e. corresponding to “substantial agreement”. In conclusion, patient performed self-reading of a UVB phototest appears to be a fairly reliable method for estimation of individual skin UV-sensitivity, when compared to the reading of a trained observer. The finding opens up for a broadened use of phototesting in clinical practice and for preventive initiatives aiming at identifying at-risk individuals and reducing sun exposure.

Keywords: Phototesting, Ultraviolet radiation, Self-assessment, Test reliability, Skin cancer, Prevention, Risk classification, Skin type, UV-sensitivity, Skin Malignancies.

INTRODUCTION

Phototesting is routinely used in dermatology, primarily in the clinical assessment and diagnosis of patients with suspected photodermatoses, and for dosimetry purposes in phototherapy. Phototest equipments may vary in design and performance, but the basic principle of phototesting is to provoke the skin with different doses of UV-light, and to grade the skin reaction following provocation. The outcome of provocation is classified in terms of minimal erythema dose (MED), which is defined as the lowest UV-dose capable of causing skin erythema. Testing can be performed for different spectra of UV-light, including both UVA and UVB, narrowband or broadband, and the dose can be regulated either by altering illumination time, or by use of different kinds of photo filters [1, 2].

Another field of interest recently investigated in studies, is the possibility of using a phototest in skin cancer prevention [3, 4]. The connection between people’s sun exposure habits in western societies and increasing skin cancer incidence during recent decades, is well documented, and individuals with a sensitive skin type is at extra risk [5-8]. The traditional and most common way of classifying skin reactivity to UV-light is Fitzpatrick’s classification, based on self-reported estimation of tendency to burn and tan [9]. The procedure is, however, highly dependent on subjective parameters and has in several studies been demonstrated to correlate poorly with actual UV sensitivity, objectively measured by phototest [10, 11]. Moreover, people’s self-estimated perception of their own individual UV-sensitivity has been found to affect sun exposure habits to higher extent than the actual UV-sensitivity [12]. The idea of using a phototest in the preventive situation has been to increase awareness of individual UV-sensitivity, but also to identify and target prevention towards individuals who are at specific risk from exposing themselves with excessive UV radiation. Also, in the general management of patients with different dermatological conditions, it might be of interest to estimate the patient’s reactivity to sun light, as well as to other exposures. Thus, both in the clinical situation, for prevention purposes and in research studies, a broadened ability to perform a phototest would be beneficial.

One of the main obstacles for the performance of a phototest is the fact that it is rather time consuming, since in clinical routine both an initial provocation visit and a second, or even repeated visits, for test reading are needed. Furthermore, test reading is most often performed by specialised personnel, such as a doctor or a trained nurse. A possible way to save time, as well as clinical resources, would be if the patients themselves would be able to read the phototest. In a previous study, this has been tried in a student population, in which reading of a UVB phototest and a
sodium lauryl sulphate (SLS) patch test was done by
the subjects, and compared with readings of a trained
observer [13]. In contrast to the “traditional”
assessment, the test reading was based on a
dichotomous outcome (erythema present or not
present), and no detailed scoring of the reactions was
required. For the patch test, results showed
“substantial agreement” according to kappa-analysis
(weighted kappa value 0.76), and the phototest “almost
perfect agreement” (weighted kappa value 0.83),
between the subjects’ and the doctor’s readings.
However, since the study population, consisting of
medical students in preclinical stage of education,
might not be considered to be representative for a
normal population, a similar study performed in
patients, would be valuable.

Besides phototesting and patch testing, the self-
reading concept has been tried in a few other
situations, but the literature is sparse, and results vary.
In studies on tuberculin tests, results range from
inadequate agreement levels [14, 15] to consistently
reliable [16-19], in part probably explained by subject
material- or instruction-related differences. The
dichotomous character of the reading procedure of the
tuberculin test, however, is similar to that of the
described photostudy [13]. In a study on self-reading
of skin reactions following chickenpox vaccination, a
90% agreement was found, and the procedure
was stated to be usable for routine vaccination assessment
[20]. Other examples where self-reporting by patients
have been found to be reliable are reading of a patch
test for nickel allergy [21], scoring of hand eczema [22],
and self-assessment of vaginal pH measurement [23,
24]. In some medical fields, a more advanced progress
in terms of self-reading has developed, for example
self-reporting of blood pressure in patients with
hypertension, for which implementation of the
procedure has been reported to lead to increased use
and reinforced preventive benefits [25, 26]. This may
be viewed as an example of how patient performed
self-reading can increase quality of care and at the
same time save clinical resources.

The aim of the present study was to investigate, in a
primary health care population, the reliability of patient
performed reading of a UVB phototest, when compared
to the reading of a trained physician.

MATERIALS AND METHODS

Study Population

The study was performed at a primary health care
centre in south east Sweden. Participants for the study
were recruited by a poster located in the waiting room
of the health care centre. Age ≥18 years were inclusion
criteria, and known history of photodermatosis or
present intake of any UV-sensitizing drugs were
exclusion criteria. All participants gave informed written
consent on their voluntary participation in the study.

Testing Procedure

The phototest used for the study (Dermalight 80
MED-tester, A.L.T Lichtterapietechnik, Germany) with a
broadband fluorescent UV lamp (Philips PL 9W/12)
consisted of six 12×12 mm provocation fields emitting
separate increasing doses of UV light, and was applied
on the palmar side of the lower arm of the patient. The
illumination time was 25 s, which resulted in the
following UV doses: 18, 36, 51, 63, 87 and 105
mJ/cm². Test provocation was done at the primary
health care centre, and after 24 h, the subjects
revisited the centre to read the test. This was
performed by the subjects themselves, by counting the
number of erythematous reactions detectable and then
reporting the results on a specific test protocol. They
were instructed that each visible reaction, even a
barely perceptible erythema, was to be classified as a
reaction, a procedure based on the findings of Lock-
Anderson et al. [2] that this criterion for minimal
erythema dose (MED) has been shown to be more
consistent, in terms of inter-observer variability, than
that relating to sharp-bordered reactions. For blinding
purpose, the test protocols were placed in closed
envelopes, and immediately after the subjects’ own test
readings, a doctor (one of the authors, M.F.) performed
an independent test reading, following the same
reading criteria. Furthermore, an additional subscoring
of each reaction was also conducted, as follows:

- = negative
(+ ) = barely perceptible erythema
1+ = erythema with a clear border
2+ = erythema and oedema
3+ = erythema, oedema and papules
4+ = erythema, oedema and vesicles.

Statistical Analysis

The agreement between patient-performed self-
reading and the doctor’s control reading, according to
the number of positive reactions on each subject, was

4+ = erythema, oedema and vesicles.
estimated in percent, and by calculation of weighted Kappa. The statistical benefit of Kappa is that it takes into account the possible agreement that could happen by chance, thus being a valuable complement to percentage presentation solely. The smaller sample, the better agreement is needed to gain a high kappa value, since the probability that the observed agreement might be an expression of chance increases. The following guidelines for interpretation of kappa value are usually used: 0-0.20 = poor agreement, 0.21-0.40 = fair agreement, 0.41-0.60 = moderate agreement, 0.61-0.80 = substantial agreement, 0.80-1.00 = almost perfect agreement [27].

Also, Pearson’s $\chi^2$-test was performed to detect possible age and gender related differences in the cases of disagreement between the subjects’ and doctor’s test readings. In the calculations age was dichotomized to a higher or lower value then the median in the sample.

**Ethics Approval**

The study was approved by the Regional Ethics Review Board in Linköping (Dnr M192-09).

### Table 1: Outcome of Patient Performed Self-Readings of their Phototests, and the Corresponding Control Readings by Doctor, also Showing the Results of the Detailed Scoring of Each Reaction (+)=Barely Perceptible Erythema, 1+ = Erythema with a Clear Border

<table>
<thead>
<tr>
<th>Number of subjects</th>
<th>Subject reading (number of reactions)</th>
<th>Control reading by doctor (number of reactions)</th>
<th>Detailed scoring by doctor, UV-dose (mJ/cm²):</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>18</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>0</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>0</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>0</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>0</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>0</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>1</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>1</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>1</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>1</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>1</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>1</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>0</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>0</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>2</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>2</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>2</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>2</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>n = 32</td>
<td>2</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>2</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>3</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>3</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>3</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>3</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>3</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>3</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>3</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>3</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>3</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>3</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>3</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>4</td>
<td>4</td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>3</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>4</td>
<td>4</td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>4</td>
<td>4</td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>4</td>
<td>4</td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>4</td>
<td>4</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>4</td>
<td>4</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>4</td>
<td>4</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>4</td>
<td>4</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>4</td>
<td>4</td>
<td></td>
</tr>
</tbody>
</table>
RESULTS

32 voluntary subjects fulfilling inclusion criteria, 11 male and 21 female, between 18 – 81 years of age, with a mean age of 49 years, and the median value 47.5 years, joined the study. No subjects needed to be excluded due to the exclusion criteria. Of the 32 subject, 25 subjects reacted to the highest UV dose (105 mJ/cm²), 22 subjects to the second highest (87 mJ/cm²), 16 subjects to the third highest (63 mJ/cm²), and 6 subjects to the fourth highest dose (51 mJ/cm²), while the remaining 7 subjects did not react to any of the provoked doses. For none of the subjects were the two lowest doses able to cause any reaction.

The results of the subjects’ self-readings and doctor’s control readings are displayed in Table 1, also showing the outcome of the detailed subscoreing performed by the doctor. Absolute agreement between subjects’ and doctor’s test readings were found in 72% of cases, and the weighted kappa-value was 0.78 (95 CI: 0.64 – 0.91), i.e. corresponding to “substantial agreement”. Six subjects overestimated, and three subjects underestimated, the number of reactions when compared to the doctor’s reading. From the detailed scoring of the reaction, it could be stated that in the cases of underestimation by the subjects, all missed reactions were scored as “barely perceptible erythema”.

In Table 2, age and gender distribution according to the level of agreement between subject’s and the doctor’s control reading is presented. No statistically significant differences in distribution according to these parameters could be detected (Pearson’s χ²-test).

DISCUSSION

In the light of increasing skin cancer incidence and excessive sun exposure habits in western societies during the past decades, objective methods and strategies to identify at-risk individuals and to communicate sun exposure risks need to be developed and promoted. The environment for this may be in primary health care, at dermatology clinics or in public health situations, but the tools used can likely be similar. In the present study it has been demonstrated that a UVB phototest with a patient performed self-reading procedure, is a fairly reliable method to grade individual skin UV-sensitivity, without the need for excessive clinical resources, which opens up for a broadened use in various situations. The method per se has the advantage of being non-invasive, easy to administer and pedagogical in nature, and – without the need for revisit – also quickly performed.

The traditional Fitzpatrick’s classification of skin type has since long been the predominant method of grading skin UV-sensitivity in the broader perspective, i.e. not in the clinical situations where a more accurate estimation, such as in the diagnosis of photodermatoses, is warranted. Several studies, however, have shown the method to be quite unreliable in relation to actual UV-sensitivity [10, 11, 28], and it has also been demonstrated that its self-estimated outcome may be affected of age and gender, young people (14-18 years of age), and especially young males, more often tend to underestimate their skin-sensitivity [10]. Since young people constitute one of the most important target group for preventive measures directed at skin cancer, the performance of a phototest, revealing the actual UV-sensitivity, would thus be preferable compared to self-estimation by means of Fitzpatrick’s classification, with high risk of underestimation, in order to communicate individual risk. In a previous study, self-estimated Fitzpatrick’s skin type was stronger associated with the level of sun protection than was the actual UV-sensitivity, when measured by phototest [12], which indicates that there

<table>
<thead>
<tr>
<th>Gender</th>
<th>Patient readings &lt; control readings (n)</th>
<th>Patient readings = control readings (n)</th>
<th>Patient readings &gt; control readings (n)</th>
<th>Sign. (χ²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male</td>
<td>0</td>
<td>8</td>
<td>2</td>
<td>p = 0.47</td>
</tr>
<tr>
<td>Female</td>
<td>3</td>
<td>15</td>
<td>4</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Age</th>
<th>Patient readings &lt; control readings (n)</th>
<th>Patient readings = control readings (n)</th>
<th>Patient readings &gt; control readings (n)</th>
<th>Sign. (χ²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt; 47.5 years</td>
<td>1</td>
<td>10</td>
<td>5</td>
<td>p = 0.18</td>
</tr>
<tr>
<td>&gt; 47.5 years</td>
<td>2</td>
<td>13</td>
<td>1</td>
<td></td>
</tr>
</tbody>
</table>
might be a need to enlighten at-risk individuals to protect themselves appropriately in regard to how UV-sensitive they in fact are.

In the previous study based on self-reading performed by medical students, the results showed a somewhat better agreement with the trained observer reading (85%) [13], than what was observed in the present study on patients. Additionally, in contrary to the results of the previous study, there was a tendency to overestimate reactions among subjects. There are several possible explanations for this. First, the study populations differed markedly, both with regard to age and educational level, and it is likely to believe that the patient population used in the present study is more representative of a “normal” population, and certainly an ordinary primary health care population, than the student sample. Another circumstance that might have affected the outcome, is the fact that the test reactions in the student-based study was round in shape, instead of square shaped as in the present study, although the significance of this difference appears to be more doubtful. The fact that six subjects overestimated compared to the doctor’s reading is interesting, and raises the question whether the development and progress of the erythemal reactions may have played a role. Usually UV-induced erythema is as most pronounced between 10-15 hours after provocation [1, 29, 30], and then declines slowly. It can therefore not be excluded that the subjects may have noticed weak erythemal reactions that had then gone to regress within 24 hours, when the final reading was performed. This could possibly have been avoided technically by covering up the test area after provocation, a procedure that in practice, however, complies poorly with the common test procedure. On the whole, the sparse difference in test reading outcome possibly related to this is likely to be of minor importance in clinical practice.

The location for the application of the phototest may be discussed, since the forearm is perhaps not the most common location used in clinical situations and in studies. Commonly phototests are applied on the upper back or on the buttocks, partly because these areas are quite flat, and partly because they are in general less exposed to sunlight [31-33]. However, application on the inner forearms, as in the present study, is also stated to be appropriate [34], and was a prerequisite for the self-reading procedure. Previous studies indicate that UV-sensitivity tend to be somewhat higher on the back and buttocks than on the arms, but that the difference was slight [31, 35]. For dosimetry purpose prior to phototherapy, the back and buttocks are likely to be preferable, in order to avoid unwanted excessive sunburn, but e.g. for the use in preventive situations, directed at skin cancer, the difference has reasonably little significance. It is also important to point out that patient performed self-reading is not intended to replace phototest reading by experts, but to be a valuable complement when appropriate.

A weakness of the study is the relatively limited sample size. However, the kappa-analysis takes sample size into account, so that a small sample size demands higher level of percentage agreement to generate a high kappa value. The agreement level found in the present study must thus be considered to be reliable. Another limitation of the study is the fact that control reading was performed by one single doctor. This makes it difficult to relate to the possible effects of inter-observer variability that could have affected the results if there were more than one physician performing the control readings, which has in a previous studies shown to be substantial [2, 36]. Subsequently, the ultimate setup would be a study with both a somewhat larger subject sample size, and multiple trained observers performing the control readings. There was also a skewness in distribution between gender, females being overrepresented in the study population. This probably reflects that females are in general found to be more interested in health and lifestyle matters [37], and perhaps also that they have been found to look more seriously on sunburn damage than males [38-40].

In conclusion, patient performed self-reading of a UVB phototest appears to be a fairly reliable method for estimation of individual skin UV-sensitivity, when compared to the reading of a trained observer. The finding opens up for a broadened use of phototesting in clinical practice and for preventive initiatives aiming at identifying at-risk individuals and reducing sun exposure.

REFERENCES


Received on 07-04-2012 Accepted on 10-05-2012 Published on 25-06-2012

http://dx.doi.org/10.6000/1927-7229.2012.01.01.12

© 2012 Thorslund and Falk; Licensee Lifescience Global. This is an open access article licensed under the terms of the Creative Commons Attribution Non-Commercial License (http://creativecommons.org/licenses/by-nc/3.0/) which permits unrestricted, non-commercial use, distribution and reproduction in any medium, provided the work is properly cited.